# Importation of live garden snails (*Cornu aspersum*) for heliciculture

Final Biosecurity Import Risk Review

Animal Biosecurity Branch | Biosecurity Animal Division

October 2025

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**Cataloguing data**

This publication (and any material sourced from it) should be attributed as**: DAFF (2025). Importation of live garden snails (*Cornu aspersum*) for heliciculture: Final Biosecurity Import Risk Review. Department of Agriculture, Fisheries and Forestry, October 2025**

This publication is available at [agriculture.gov.au/publications](https://www.agriculture.gov.au/publications).

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**Acknowledgement of Country**

We acknowledge the Traditional Custodians of Australia and their continuing connection to land and sea, waters, environment and community. We pay our respects to the Traditional Custodians of the lands we live and work on, their culture, and their Elders past and present.

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## Summary

The Department of Agriculture, Fisheries and Forestry (the department) has prepared this biosecurity import risk review to analyse the risks associated with the proposed importation into Australia of live snails of the terrestrial species, *Cornu aspersum*, as breed-stock to produce edible snails for human consumption. The review considered relevant scientific information, industry practices and operational practicalities. The review was also informed by peer-reviewed publications and a range of other available evidence, and used external specialist expertise and, as appropriate, drew on information provided by commercial parties.

The review encompassed the invasive potential of *C. aspersum*, as well as the diseases and pests considered relevant to the proposed importation of *C. aspersum*. Australia currently permits the importation of (killed) terrestrial snails for human consumption if they have been retorted in accordance with the import conditions (DAFF, 2024), including:

* The goods must have been hermetically sealed in a container before being heat treated to a minimum core temperature of 100°C, obtaining an F0value of at least 2.8.
* The final product must be imported in the hermetically sealed (airtight) container in which it was retorted.
* The goods must be retorted, commercially sterile and shelf stable.

There are also conditions for importation of freshwater snails for laboratory use.

This review focussed on live *C. aspersum*, with risk management measures proposed for the importation of both foundation adult breed-stock and hatchling snails. In accordance with the *Biosecurity Act 2015*, the measures for each will reduce biosecurity risk to a very low level, but not to zero. The measures will also seek to ensure that the correct species of snail is imported, thus mitigating any unassessed invasive potential.

The measures for the importation of either adult or hatchling *C. aspersum* include a prerequisite for the approval of both the exporting country, as a source for this commodity, and the individual snail farming establishments from which the adult or hatchling snails will be sourced. In particular, snail farms seeking to export *C. aspersum* to Australia must be commercial operations and government certified for basic quality standards for food safety and general hygienic operations. The current Quality Standard applicable to the commercial operation which is exporting *C. aspersum* to Australia must be linked to a specific import application and will be reviewed as part of the application assessment process. Some modification or amendment of this Quality Standard may be required.

There are **separate protocols for adult snails and for hatchling snails**, with each addressing both species verification and biosecurity.

Species verification and biosecurity measures for adult snails

* Adult snails exported to Australia must be certified by the exporter and exporting country as *C. aspersum* only. Certification of species may be based on morphological identification or on an approved molecular method. Approval will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity.
* Adult snails will be directed to an Approved Arrangement on arrival at the Australian border. This will be a biosecure facility and may be a government-supervised quarantine facility or may be a snail farming facility that is managed privately. The privately managed Approved Arrangement and site will include (*inter alia*) site design and security, as well as biosecurity protocols for site maintenance, people and the disposal of waste. The privately managed Approved Arrangement and site will also be subject to the department’s requirements for audit and compliance.
* A subset of the snails within each imported consignment will be subject to species verification by morphological or molecular identification. Morphological Identification involves sacrifice of the selected snails as dissection is required. Dissection and identification must be undertaken by a suitably qualified inspector who has been approved by the department to undertake this task. Identification must be based on a diagnostic key that has been approved by the department for use in this context. Molecular identification will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity. The identification procedure will be described in the Approved Arrangement and the protocol for its application will be based on the anticipated size of consignment.
* Imported adult snails will remain under the Approved Arrangement and within the biosecure facility where they will be bred to the next generation. Throughout this period, the imported snails will be isolated from Australian snails and from the definitive host of both *Angiostrongylus* spp. and *Crenosoma vulpis*. This will ensure that the following generation of C. aspersum will be free of *Angiostrongylus* spp. and *Crenosoma vulpis*. Isolation within the biosecure facility will also break the life cycle of vectored plant pathogens. Successive generation(s) of snails will then be eligible for release from the Approved Arrangement and site.

The imported adult snails will not be eligible for release from the Approved Arrangement site. They will either remain within the site for their natural life or will be euthanised. The remains of imported adult snails will be managed as biosecurity waste.

Species verification and biosecurity measures for hatchling snails

* Hatchling snails are snails less than 10 days old that have not been released from the hatching environment and have not been exposed to plant substrate.
* Hatchling snails must be sourced from establishments that employ either an internal or mixed (internal and external) farming system and must only have received commercially processed livestock feed.
* Hatchling snails must be certified by the exporter and exporting country as *C. aspersum* only. Certification of species may be based on morphological identification of breed stock or on an approved molecular method. Approval will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity.
* Hatchling snails will be directed to an Approved Arrangement on arrival at the Australian border. This will be a government-supervised biosecure facility where the snails can be held while species is verified.
* A subset of the hatchling snails within each imported consignment will be subject to species verification by molecular identification. This will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity. The protocol for the application of this procedure will be described in the Approved Arrangement and will be based on the anticipated size of a consignment.

Having verified the species of hatchling snails within a consignment (as above), the consignment may be released from biosecurity control. It will be a condition of the import permit that imported hatchling snails are for breeding purposes only and must not be used for human consumption.

## Introduction

Australia’s biosecurity policies aim to guard against the entry, establishment and spread of exotic diseases and pests which might otherwise threaten Australia’s agricultural industries, natural environments, or people. Biosecurity import risk review enables the Department of Agriculture, Fisheries and Forestry (the department) to assess the level of biosecurity risk that may be associated with proposals to import goods into Australia. If the biosecurity risk exceeds Australia’s appropriate level of protection (ALOP), risk management measures are proposed. If the biosecurity risk cannot be reduced to a very low level, but not to zero, as required under the Biosecurity Actand through means available to the department, the goods will not be imported into Australia.

Biosecurity import risk is assessed by the department using technical and scientific experts from relevant fields and involve consultation with stakeholders at various stages during the process. The assessment may take the form of a biosecurity import risk analysis (BIRA) or a non-regulated risk analysis (such as scientific review of existing policy and import conditions, or scientific advice). This biosecurity import risk review is a non-regulated analysis.

Further information about Australia’s biosecurity framework is provided in the [Biosecurity import risk analysis guidelines 2016](https://www.agriculture.gov.au/biosecurity-trade/policy/risk-analysis/guidelines).

The department recognises that new scientific information and technologies, or other combinations of measures, may provide an equivalent level of biosecurity protection for the disease agents identified as requiring risk management. The department will consider technical submissions that objectively demonstrate suitable efficacy of alternative biosecurity measures.

### Background

In response to members of the public (the applicants) with specific commercial interests in developing the Australian heliciculture industry, the relevant departments of the Australian Government independently undertook analyses for the importation of live *C. aspersum,* as required under Australian law.

Two separate pieces of legislation underpin the required risk review for this import proposal.

The first evaluation (completed June 2020) was undertaken by the (then) Department of the Environment and Energy in accordance with the *Environment Protection and Biodiversity Conservation Act 1999* (EPBC Act)*,* and was the project undertaken at the time of the first contact by the applicants. This evaluation recommended listing *C. aspersum* (Common Garden Snail) in Part 1 of the [List of Specimens taken to be Suitable for Live Import](https://www.legislation.gov.au/F2006B01053/latest/versions) (specimens that can be imported without a permit under the EPBC Act). Importation of C. aspersum would be subject to an importation approval under the Biosecurity Act.

The current biosecurity import risk review is being undertaken by the department with additional technical expertise provided by an external consultant, in accordance with the requirements of the Biosecurity Act.

Completion of both evaluations is a prerequisite to any live animal imports.

The applicants advised that there is significant interest in the restaurant industry in Australia, and internationally, in obtaining high quality *escargots* (edible snails) for human consumption. To service this need, they wished to develop Australian heliciculture businesses by farming the common garden snail (*C. aspersum*). The snails bearing the desirable characteristics would be sourced from eligible overseas producers of edible snails.

The applicants have asserted that the *C. aspersum* currently available in Australia are smaller than the more commercially attractive snails available in Europe. Snails of less than 8 to 10 grams, and of small shell size, are reportedly unsuitable for sale. The applicants wish to use imported snails as a base to breed from, rather than ‘spending years developing snails of a suitable size from domestic stock’.

At the beginning of the 20th century, two subspecies of this snail were defined based on shell characteristics: *Helix aspersa aspersa* (now classified as *Cornu aspersum aspersum)* and *Helix aspersa maxima* (now classified as *Cornu aspersum maximum)*. These subspecies are currently recognised as genetically divergent, although there are differing views on whether they constitute separate subspecies (Guiller and Madec, 2010; Guiller *et al*., 2001; Rygalo-Galewska *et al*., 2022). *C. aspersum* is highly variable morphologically and several distinct morphotypes have also been described. Differences exist with respect to size, shape, thickness and colour of the shell. Some authors assert that the form *maximum* is a distinct subspecies *C. aspersa maximum* ([Taylor, 1914](https://www.cabidigitallibrary.org/doi/10.1079/cabicompendium.26821#core-ref-235)), known in French as 'gros-gris', with *C. aspersum sensu stricto* known as the 'petit-gris'. The status of *C. aspersum maximum* as a valid subspecies is not consistently supported in the literature ([Guiller](https://www.cabidigitallibrary.org/doi/10.1079/cabicompendium.26821" \l "core-ref-130) *[et al](https://www.cabidigitallibrary.org/doi/10.1079/cabicompendium.26821" \l "core-ref-130)*[., 2001](https://www.cabidigitallibrary.org/doi/10.1079/cabicompendium.26821" \l "core-ref-130)).

For the purpose of this biosecurity import risk review, the classification of separate subspecies has been accepted.

*C. aspersum* is farmed or collected in the wild in several countries for human consumption. Both subspecies are farmed. The subspecies *C. aspersum maximum* is larger and has a lifespan of more than 8 years, whereas the subspecies *C. aspersum aspersum* is smaller, more widely distributed and has a lifespan of 3 to 4 years (Russo and Madec, 2011). These subspecies also differ in their shell chemical composition (Ligaszewski, Surowka *et al*., 2009) and metabolism (Czarnoleski, Labecka *et al*., 2016) and potentially in other attributes yet to be defined. *Cornu aspersum maximum* does not occur in Australia.

According to the applicants, noting the information has not been verified by the department:

Heliculture farms in Europe have bred C. aspersum for eating for over 2000 years and have selectively bred snails for a consistent size and growth rate. Importing disease-free snails of the required age and size from a commercial snail farm would ensure that the breeding of snails for restaurants could commence quickly and safely. A single 60 kg shipment of adult snails (approximately 6,000 individuals) would be imported to begin breeding for future generations.

This biosecurity import risk review considers that certain controls over commercial establishments supplying *C. aspersum snails* to Australian importers (generally termed snail farms) must be in place to allow for effective biosecurity risk management. Standards governing the operation of such establishments differ widely and, when present, tend to focus purely on food safety risks. In addition, snail farms often fatten large numbers of snails in an outside environment immediately before sale (Rygalo-Galewska *et al*., 2022). This increases the likelihood that wild snails will enter the snail farm in the country of origin and be included in exported consignments.

Controls over commercial establishments is also considered in relationship to the likelihood that a biosecurity hazard can establish and spread within Australia. There is a significant likelihood that snails imported into Australia and released to commercial snail farms in Australia will enter the broader Australian environment, so hazards of concern must be effectively managed prior to that point.

### Scope

This biosecurity import risk review focussed on the importation of live *C. aspersum* from any country to mainland Australia and Tasmania. The Australian External Territories are not considered. The intent is for imported snails to serve as breed-stock – that is, the imported snails will not be used for human consumption.

### Next steps

This is the final report for the import risk review. The department will work with the applicants to establish the most practical approach to the implementation of the risk management measures.

## Method

This biosecurity import risk review included three subsidiary assessments:

* Quarantine pest potential of *C. aspersum*
* Risk assessments for animal and human pathogens
* Risk assessment for *C. aspersum* as a vector for plant pathogens.

The (then) Department of the Environment and Energy undertook an evaluation of the invasive potential *C. aspersum*. This evaluation recommended listing *C. aspersum* (Common Garden Snail) in Part 1 of the **List of Specimens taken to be Suitable for Live Import** (specimens that can be imported without a permit under the EPBC Act). In the current biosecurity import risk review, a further evaluation was undertaken to determine whether *C. aspersum* fitted the International Plant Protection Convention’s (IPPC) definition of a quarantine pest. Pest categorisation is documented in Section 4.

The biosecurity analyses for animal and human pathogens, documented in Section 5, followed the Department of Agriculture, Fisheries and Forestry’s (the department’s) Biosecurity Import Risk Analysis Guidelines (2016). Biosecurity risk is defined in the Biosecurity Act and refers to the likelihood of a pathogen entering, establishing or spreading in Australian territory, and the potential for the pathogen causing harm to human, animal or plant health, the environment, economic or community activities.

The analyses for animal and human pathogens commenced with a categorisation step that sought to filter a potentially large number of pathogens and focus on the assessment on those that are relevant to the commodity. This step is termed ‘hazard identification’ by the World Organisation for Animal Health (WOAH, previously the OIE), and is summarised in [Appendix A](#_Appendix_A:_hazard).

The evaluation for each pathogen of concern included assessment of the likelihood that the pathogen will enter Australia with the importation of live snails (*C. aspersum*) and that susceptible species within Australia will be exposed to the pathogen.

* Entry assessment describes the pathway for importation as it relates to the introduction of pathogens and estimates the likelihood of this complete pathway occurring. It considers: (a) biological factors; (b) country factors, such as the prevalence of the pathogen; and (c) commodity factors, such as the production system, the quantity to be imported, and any testing, treatment or processing that is part of baseline production.
* Exposure assessment describes the biological pathways necessary for exposure of susceptible animals, plants and the environment to the pathogen and estimates the likelihood of the exposure occurring. It again considers: (a) biological factors; (b) country factors, such as the presence of competent vectors, and relevant geographical and environmental characteristics; and (c) commodity factors, such as the quantity to be imported, its end use and disposal practices.

The likelihood that each pathogen will become established and spread within Australia was examined as part of the assessment of consequences, which focussed on impact at a national level. The likelihood of entry and exposure, and the likely consequences, were then combined using the department’s risk matrix (Table 3) to provide an estimate of unrestricted biosecurity risk.

An overview of this risk assessment process is given in Figure 1. All likelihoods were evaluated qualitatively, using the terms shown in Table 1. Likely consequences were also evaluated qualitatively, and the terms for this are given in Table 2.

The final part of the biosecurity analyses (Section 5.9) delt with the potential for *C. aspersum* to vector key plant pathogens. This part of the review was necessarily less structured than other sections, as it considered a wide range of pathogens and included substantial uncertainty.

Figure 1 Components of the unrestricted risk estimate

Beginning at country of export, it flows to the Australian border, then to exposure of susceptible animals, to establishment in susceptible populations, to spread among susceptible populations, to overall effect of establishment and/or spread.
Country of export, the Australian border, and exposure of susceptible animals are classified under entry and exposure scenarios and assessment.
Establishment in susceptible populations, spread among susceptible populations, and overall effect of establishment and/or spread are classified as outbreak scenarios and are under consequence assessment.

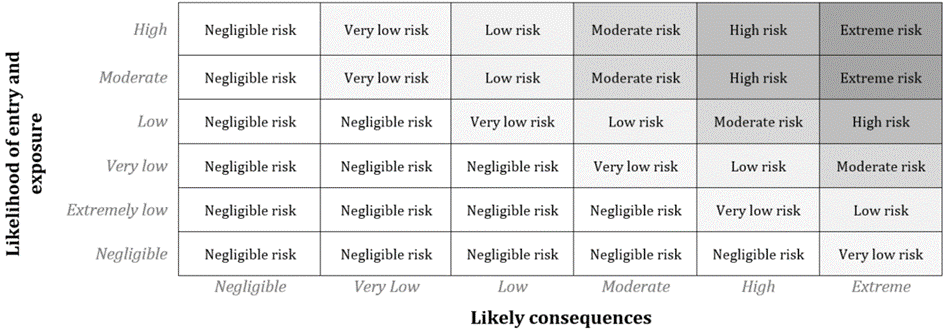
Table 1 Nomenclature for qualitative likelihoods

| Likelihood | Descriptive definition |
| --- | --- |
| High | The event would be very likely to occur |
| Moderate | The event is equally likely to occur or not occur |
| Low | The event would be unlikely to occur |
| Very low | The event would be very unlikely to occur |
| Extremely low | The event would be extremely unlikely to occur |
| Negligible | The event would almost certainly not occur |

Table 2 Rules for determining the likely consequences using effect categories

| Overall effect | Description |
| --- | --- |
| Extreme | The effect is likely to be highly significant at the national level. Implies that economic stability, societal values or social well-being would be seriously affected. |
| High | The effect is likely to be significant at the national level and highly significant within affected zones. Implies that the effect would be of national concern. However, serious effects on economic stability, societal values or social well-being would be limited to a given zone. |
| Moderate | The effect is likely to be recognised on a national level and significant within affected zones. The effect is likely to be highly significant to directly affected parties. |
| Low | The effect is likely to be recognised within affected zones and significant to directly affected  parties. It is not likely that the effect will be recognised at the national level. |
| Very low | The effect is likely to be minor to directly affected parties. The effect is unlikely to be  recognised at any other level. |
| Negligible | The effect is unlikely to be recognised at any level within Australia. |

Table 3 Risk estimation matrix – likelihood of entry and exposure and likely consequences



## *Cornu aspersum*: pest categorisation

### Identification of species and subspecies

The species *C. aspersum*, previous name *Helix aspersa*, exhibits a wide variety of shell patterns and colouration (Chevallier 1977; Kougiagka *et al*., 2022; Madec *et al*., 1998). These morphological characteristics were the basis for the description of many forms, varieties and species of *Helix*, creating a complex and convoluted taxonomy history. The subspecies *C. aspersum maximum* *(*or *C. aspersum maxima,* as referenced elsewhere) (*Helix*(*Pomatia*) *aspersa*var.*maxima*) was initially described by Taylor (1883) based on specimens collected from Algeria. Chevallier (1977) describes a range of ‘morphs’ including *C. aspersum maximum* noting its large size, up to 45 mm, and distribution limited to the Mediterranean.

The native range of *C. aspersum maximum* is believed to be North Africa, although wild populations of the subspecies are unknown (Dupont-Nivet, Guiller and Bonnet 1997; Guiller *et al*., 2001). The situation is further complicated by the selective breeding of *C. aspersum maximum* for commercial purposes.

Madec et al., (1998) conducted studies to compare characteristics of *C. aspersum maximum* to previous studies on *C. aspersum aspersum*, reporting that differences observed between the species – larger clutch size and egg weight – were likely explained by size-effect since *C. aspersum maximum* are physically larger than *C. aspersum aspersum*. Similarly, Çelıket al., (2018) demonstrated that physically larger *C. aspersum aspersum* laid larger eggs and subsequently produced larger juveniles.

Dupont-Nivet, Guiller and Bonnet (1997) suggested that the size of snails is a heritable trait based on breeding studies involving *C. aspersum aspersum* and *C. aspersum maximum*, and Freitaset al., (2023) demonstrated heritably of weight in *C. aspersum maximum*.

However, environmental factors are known to impact the size and development rate of snails (Çelıket al., 2019; Kougiagka *et al*., 2022). For example, stocking density and temperature may affect the growth rate of *C. aspersum maximum* (Freitas *et al*., 2023; Jess and Marks 1998). Inbreeding may also affect fitness and inherited traits in snails (Kougiagka et al., 2022). Dupont-Nivet et al., (2001) observed significant loss of genetic variability in a population of *C. aspersum* artificially selected for breeding.

Czarnoleski, Labecka and Kozłoeski (2016) compared the size of *C. aspersum aspersum* and *C. aspersum maximum* grown at different temperatures; both subspecies produced larger individuals when grown at 20°C than 15°C. However, *C. aspersum aspersum* grown at 20°C, produced larger adults than *C. aspersum maximum* grown at 15°C.

Using partial sequences of the 16S gene, Guiller *et al*., (2001) identified two haplotypes in *C. aspersum aspersum* designating them East and West based on geographic distribution in North Africa. Analysis of *C. aspersum maximum,* obtained from farmed snails in France, formed a separate third haplotype, which was more closely related to the East haplotype of *C. aspersum aspersum*. However, snails collected from Morrocco exhibiting shell measurements consistent with *C. aspersum maximum*, did not align with the farmed French *C. aspersum maximum*, but were related to the Western haplotype of *C. aspersum aspersum*. Guiller *et al*., (2001) advised that their results of genetic analysis did not resolve the taxonomic relationships within the species.

Later phylogenetic studies seeking to resolve the evolutionary history of *C. aspersum aspersum* have used *C. aspersum maximum* as an outgroup (Guiller and Madec 2010; Guiller *et al*., 2012; Sherpa *et al*., 2018).

In investigating the closely related genus *Erctella*, Colomba *et al*., (2015) suggest that *C. aspersum* may be a group of [cryptic] species, but specified further studies are necessary to confirm their hypothesis. It is noted that Colomba *et al*., (2015) only included a small number of *C. aspersum* in their molecular analysis, and that all samples originated from Italy.

In 2022, Kougiagka *et al*., (2022) compared morphological and genetic features of farmed *C. aspersum aspersum* and *C. aspersum maximum* in Greece. Both subspecies were divided into three haplotypes, with genetic variation observed within populations rather that between them.

Gomot-de Vaufleury and Borgo (2001) crossed the two subspecies, *C. aspersum aspersum* and *C. aspersum maximum,* achieving fertile offspring. The resulting hybrids had a high level of juvenile mortality and low fecundity, with poor fitness compared to the parents. The authors noted that although hybrids were obtained, the snails showed a tendency to reproductive isolation, both behavioural and anatomical, and support the validity of the two subspecies (Gomot-de Vaufleury and Borgo, 2001).

Finally, no differences in genitalia structure are recorded between the subspecies *C. aspersum aspersum* and *C. aspersum maximum*, although size differences within and between the subspecies are noted and discussed (Guiller and Madec, 2010; Madec, Bellido and Guiller, 2003; Sherpa *et al*., 2018). Ziȩtek, Ziomek and Wilczyńska (2019) developed a dissection protocol appropriate for use on both subspecies.

In summary, the available genetic information is not sufficient to resolve the taxonomic status of the subspecies *C. aspersum maximum*. Based on morphological diagnosis and the other information presented here, the department supports *C. aspersum maximum* is a valid subspecies of *C. aspersum* and unlikely to be a cryptic species.

Given the absence of wild populations, it appears that the subspecies persists only due to commercial farming efforts in a number of countries. Evidence from published literature supports that *C. aspersum* adult size and weight are inherited; it is likely these traits are maintained in *C. aspersum maximum* through selective breeding and growth under optimal conditions.

### Pest potential and status

IPPC ISPM 11 (Pest Risk Analysis for Quarantine Pests) states that the taxonomic level usually considered in a Pest Risk Analysis is ‘species’ – the use of ‘subspecies’ may be justified if there is scientific evidence demonstrating that differences in characteristics are stable and significantly affect phytosanitary status.

From a plant health perspective, the subspecies *C. aspersum maximum* does not meet the IPPC definition of a quarantine pest for Australia. The available evidence suggests that *C. aspersum maximum* is very similar to *C. aspersum aspersum*, which is already present and widespread in Australia, and any impacts on plant health and the environment are likely to be commensurate with, or less than, those of *C. aspersum aspersum*.

The species *C. aspersum* was likely introduced to Australia shortly after European settlement and is present in all states and territories (Blacket *et al*., 2016). The direct impact of *C. aspersum* on plant health in Australia is limited to agricultural and ornamental plants in human-disturbed environments.

Subspecies of *C. aspersum* are expected to have a similar biology and behaviours. *C. aspersum maximum* is reported to be physically larger, show variation in some shell and have a longer lifespan (Russo and Madec, 2011) and lay more eggs than *C. aspersum aspersum* (Madec *et al*., 1998; Rygało-Galewska, Zglińska and Niemiec, 2022). These traits are not necessarily indicators of increased potential for economic damage as they may be linked to trade-offs in fitness, such as the longer time taken to reach reproductive maturity (Février, Russo and Madec, 2009; Madec *et al*., 1998; Russo and Madec, 2011).

Both *C. aspersum aspersum* and *C. aspersum maximum* are commercially farmed throughout Europe (Kougiagka *et al*., 2022; Rygało-Galewska, Zglińska and Niemiec, 2022). There are reports of *C. aspersum maximum* being commercially farmed in France (Chevallier, 1980; Guiller and Madec, 2010), Greece (Kougiagka *et al*., 2022), Italy (Zucaro *et al*., 2016), Poland (Drozd *et al*., 2017), Spain (Segade *et al*., 2009) and Ukraine (Danilova, 2022), as well as Brazil (Soares, Hayashi and Cocito, 2002). Although both subspecies have been intentionally introduced to multiple countries for heliciculture, only *C. aspersum aspersum* has established outside its native range and become a pest.

There is a recent report of *C. aspersum* from the Lviv region of the Ukraine (Gural-Sverlova and Gural, 2021). Here the authors suggest one pathway of introduction into the environment is escape from snail farms, noting an illegal dumping event as well as an abandoned farm. Another potential introduction pathway discussed is infested imported plants. The authors comment that some of the snails recovered had a dark mantle, characteristic of the large form of *C. aspersum,* specifically *C. aspersum maximum*; no further taxonomic diagnosis is attempted. The authors specifically refer to these reports as ’colonies’, highlighting that their findings indicate small, localised populations that have yet to establish.

The species *C. aspersum* is recognised as highly polyphagous, with a wide plant host range including but not limited to apple, apricot, citrus, peach, barley, oats, wheat, cabbage, carrot, cauliflower, celery, bean, beet, lettuce, onion, peas, tomato, as well as an array of ornamental plants (Dekle and Fasulo, 2021; Iglesias and Castillejo, 1999). Although *C. aspersum* is widely accepted as a serious pest of agricultural and ornamental plants, there is limited information on the direct impact on crop losses or yield (Sanchez, 2010), or the costs associated with management (Jiang *et al*., 2022). Reports of crop damage found include losses in citrus fruits in California (Sakovich, 2002) and grapes in South Africa (Sanderson and Sirgel, 2002); in Australia, Sanderson and Sirgel (2002) highlight the issue of contamination with introduced snails *C. aspersum* and *Theba pisana*, in the dried fruit (sultana) industry, which was ultimately linked to changes in soil-management practices.

*C. aspersum* has not been found in native Australian bushland, despite being widespread in urban and agricultural settings (DAWE, 2020). Holland, McDonnell and Williams (2007) found small numbers of *C. aspersum* (reported as *Helix aspersa*) in remnant grasslands in Victoria, but excluded these from their study of introduced molluscs in native grasslands, due to low numbers and the fact that this species was not found in the study traps. Daniell (1994) commented that the only invasive snails thought to have caused a significant impact on Australian native vegetation are ‘white snails’ of the Helicidae family, specifically *Theba pisana*.

There are no reports in the literature that the host range of the subspecies *C. aspersum maximum* differs from that of *C. aspersum aspersum*. Despite being recorded being farmed in several European countries, there are no reports any crop damage attributed to the subspecies *C. aspersum maximum*.

### Similar farmed species

The most common species of snails farmed in Europe are: *C. aspersum aspersum* (peti gris), *C. aspersum maximum* (gros gris), *Helix lucorum* (Turkish snail), *H. pomatia* (Roman snail/ Burgundy snail) and *Eobania vermiculata* (chocolate banded snail) (Conte 2015; Rygało-Galewska, Zglińska and Niemiec 2022).

Of these farmed species, *Eobania vermiculata* and *Helix pomatia* morphologically resemble *C. aspersum*, especially at the juvenile stage, and are considered (under IPPC nomenclature) to be quarantine pests for Australia.

#### *Eobania vermiculata* (synonym *Massylaea vermiculata*)

*Eobania vermiculata* is native to the Mediterranean region (Schultes, 2014; Welter-Schultes, 2012). This species is reported from the European countries Albania, Bulgaria, Croatia, Cypress, France, Greece, Hungry, Italy, Macedonia, Malta, Montenegro, Spain and Ukraine (Bank and Neubert, 2017; Williams and Bunkley-Williams, 2023). *Eobania vermiculata* is also native to northern Africa occurring in Algeria, Lebanon, Libya, Morocco, the Republic of Türkiye and Tunisia (Bößneck, 2011; Dedov *et al*., 2022; Schultes, 2014).

*Eobania vermiculata* is spreading globally and has been reported from Belgium (Ronsmans and Van den Neucker 2016), Bosnia and Hezegovina (Karaman 2006), Egypt (Desouky and Busais, 2012), Georgia (Mumladze and Paposhvili, 2016), Iraq (Bashê and Al-Qassab, 2024), Israel (Mienis, Rittner and Vaisman, 2016), Japan (Ueshima, Okamoto and Saito, 2004), Jordan (Amr, Baker and Katbeh-Bader, 2019), the Netherlands (Soes, 2014), Qatar (Al-Khayat, 2010), Romania (Grigore, 2021), the Russian Federation (Egorov, 2008), Saudi Arabia (Desouky and Busais, 2012; El-Wakil, Banaja and Amer, 2011), Slovakia (Páll-Gergely, Fehér and Čejka, 2020), South Africa (Herbert, 2010) and the United States (Puizina *et al*., 2013). A specimen has been identified from the United Kingdom (Notton, 2006).

*Massylaea* spp. (which includes *E. vermiculata*) are listed as a National Priority Plant Pest for Australia. There are historic reports of isolated/local populations of *E. vermiculata* in NSW, SA and Tasmania, however these are considered now extinct (Blacket *et al*., 2016; Smith, 1992). The Victorian and South Australian state governments are currently undertaking isolated, local eradication programs for *E.* *vermiculata*, following separate near border detections in 2017 and 2022 respectively.

*Eobania vermiculata* is intercepted by the department at the border on non-commodity goods (e.g. new and used vehicles, machinery and break-bulk cargo) and shipping containers. It is considered a hitchhiker pest due to its frequent association with non-plant pathways, and ability to survive the extended journey to Australia.

Internationally, between 1993–1998, *E. vermiculata* made up approximately 5% of gastropods intercepted by the United States Department of Agriculture (Robinson, 1999). Live *E. vermiculata* have been intercepted on commercial shipments by Israel (Mienis, Rittner and Vaisman, 2016), and on vehicles from Italy in China (GACC, 2019).

As a popular species for human consumption, *E. vermiculata* has been deliberately imported into several countries for production (Notton, 2006; Texas Invasive Species Institute, 2014). Now illegal in the United States, authorities have intercepted smuggled consignments of breeding stock (AJOT, 2017) as well as live snails (consignments up to 10 kg) found routinely in passenger baggage (Robinson, 1999).

*Eobania vermiculata* mainly inhabits coastal regions. Habitats include shrubland, wooded hinterland, fields, gardens, vineyards and agricultural crops. *Eobania vermiculata* has been observed sheltering in crevices in stone walls, behind shrubs and amongst low-growing vegetation (Herbert 2010; Puizina *et al*., 2013), on stems of trees and shrubs, and under rubbish (e.g. cardboard and wood) (Mienis, Rittner and Vaisman 2016). Snails scale fences, trees, palms and bushes to escape heat (Mienis, Rittner and Vaisman, 2016; Yildirim, Kebapҫi and Gümüş, 2004); juveniles are reported to hibernate under stones or leaves (Ronsmans and Van den Neucker, 2016).

Like many snails, *E. vermiculata* is polyphagous; hosts include fruit, vegetables and ornamental plants (Eshra, 2013; Mohammed, 2015). *Eobania vermiculata* has been a pest in Egypt since the mid-1960s (Abo Bakr, Eshra and Hussein, 2007), reported from a range of crops and ornamental plants (Abdel Kader *et al*., 2016; Eshra 2013), including citrus (Hashem and El-Halawany, 1996). In the Republic of Türkiye, *E. vermiculata* is reported to cause damage to nectarines by gnawing fruits (Hazir and Ulusoy, 2012) and was recorded as a pest on avocado for the first time by Kahraman, Kirişik and Kahraman (2020).

##### Morphological similarities

Although adult *E. vermiculata* may have distinct shell patterning, variation within the species is widespread. For example, Welter-Schultes (2012) reports shell colour of *E. vermiculata* to be highly variable, from whitish to greenish yellow, often with colour bands or spots, and Desouky and Busais (2012) reported variation in morphological features between *E. vermiculata* populations from Egypt and Saudi Arabia.

The morphological identification of snail species that exhibit wide variation in shell complicates identification, especially in juvenile specimens (Blacket *et al*., 2016). The geographic range of *C. aspersum* and *E. vermiculata* overlap. In addition, both species are commercially farmed in Europe.

In summary, *E. vermiculata* is a quarantine pest for Australia. It has the ability to establish and spread, demonstrated by its expansion to many countries outside its native range, with climates similar to areas of Australia. *Eobania vermiculata* has consequences for plant health, feeding on a range of crops, fruit and ornamental hosts that are present in Australia.

#### *Helix pomatia*

*Helix pomatia* is not present in Australia. Smith (1992) postulates that illegal breeding colonies may exist in southern Australia, however no further information is provided to support this claim. Blacket *et al*., (2016) suggests that records of *H. pomatia* in Australia refer to intercepted specimens held in collections. No further records of *H. pomatia* in Australia were found.

*Helix pomatia* is native to Central and Western Europe (Egorov 2015; Welter-Schultes 2012). This species is reported to be present in Andora, Austria, Belarus, Belgium, Bosnia and Herzegovina, Bulgaria, Croatia, Czech Republic, Denmark, Estonia, Finland, France, Germany, Hungry, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Macedonia, Moldova, Montenegro, Netherlands, Norway, Poland, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Ukraine and the United Kingdom (England) (Bank and Neubert, 2017; Pollard, 1975).

*Helix pomatia* has been introduced to multiple European countries (Pollard 1975), as well as Brazil (Rumi, Sánchez and Ferrando, 2010), Canada (Forsyth and Kamstra, 2019), the Russian Federation (Egorov, 2015), and the United States (Dees, 1970; Imai and Hennessey, 1999). In Japan, one farm is commercially producing *H. pomatia* (Nippon News, 2016).

*Helix pomatia* has been detected at the border by the department, on non-commodity goods. However, numbers are significantly lower compared with other hitchhiking snail species. Due to its popularity as a gourmet food, *H. pomatia* is more likely to be imported illegally for commercial purposes (Twomey, 2017), as has been the case in other countries (Forsyth and Kamstra, 2019).

*Helix pomatia* inhabits mild coastal areas, as well as mountainous regions with continental climates (Błoszyk, Kacprowicz and Książkiewicz-Parulska, 2016; Forsyth and Kamstra 2019). Habitats include forest, shrublands, grasslands abutting woods, gardens and vineyards. *H. pomatia* is also common in gardens, parks, roadside verges/banks, and cemeteries, and is often found under debris; juveniles are reported to climb vegetation (Błoszyk, Kacprowicz and Książkiewicz-Parulska, 2016; Forsyth and Kamstra, 2019; Korábek, Juřičková and Petrusek, 2016; Pollard, 1975).

*Helix pomatia* has been referred to as a polyphagous generalist; information on specific hosts is limited. In England, *H. pomatia* was observed feeding on a range of plants with a preference for the knapweeds *Centaurea* *scabiosa* and *C. nigra* (Pollard, 1975). In Romania, *H. pomatia* has been observed to attack *Dhalia variablilis* (dahlia), *Lilium candidum* (lily), *Phaseolus vulgaris* (beans), *Lactuca sativa* (lettuce), and to a lesser extent *Phlox drummondii*, *Hosta plantaginea*, *Pisum sativum* (pea) (Călin *et al*., 2014). Tluste and Birkhofer (2023) reported a preference for *Urtica diocia* (common nettle) in German populations of *H. pomatia*. The authors also identified that plant litter constituted a significant proportion of the snail’s diet (Tluste and Birkhofer, 2023), in contrast to Pollard (1975) who stated that the bulk of *H. pomatia* diet consisted of fresh plants.

*Helix pomatia* is reported to be invasive in the United States (Purdue University, 2024), and Canada (Harrap, 2021) though there are no specific claims of damage to plants or crops found in the literature. Elmslie (2005) suggests that *H. pomatia* is less of an agricultural pest compared to *C. aspersum* due to smaller egg clutch size, also observing that there are no reports of rapid spread of *H. pomatia*.

*Helix pomatia* is claimed to be the most famous of the world’s edible snails, thanks to its popularity in French cuisine (Elmslie, 2005). *H. pomatia* are more difficult to cultivate than *C. aspersum* and other edible snails. To meet the demand, many snails are collected from the wild (Rygało-Galewska, Zglińska and Niemiec, 2022). This has resulted in a decrease in populations in the native range, leading many European countries to regulate wild harvesting and/or protect the species (Łysak 1999; Pollard, 1975; Welch and Pollard, 1975).

##### Morphological similarities

*Cornu aspersum* can be distinguished from *H. pomatia* by the smaller shell, different pigmentation patterns and shell structure (Forsyth and Kamstra 2019). However, there is significant variation in shell patterns noted within populations of both species (Korábek, Juřičková and Petrusek, 2016; Kougiagka *et al*., 2022; Madec, Bellido and Guiller, 2003; Neubert, 2014; Welter-Schultes, 2012); morphological identification alone may not be sufficient to determine identity.

The morphological identification of snail species that exhibit wide variation in shell complicates identification, especially in juvenile specimens (Blacket *et al*., 2016). The geographic range of *C. aspersum* and *H. pomatia* overlap.

In summary, *H. pomatia* is a quarantine pest for Australia. It has the ability to establish and spread demonstrated by established populations in countries, both native and introduced, with climates similar to areas of Australia. *H. pomatia* has consequences for plant health, feeding on a range of ornamental and agricultural hosts that are present in Australia.

### Conclusion

From a plant health perspective, the subspecies *C. aspersum maximum* does not meet the IPPC definition of a quarantine pest for Australia. The available evidence suggests that *C. aspersum* maximum is very similar to *C. aspersum aspersum*, which is already present and widespread in Australia, and any impacts on plant health and the environment are likely to be commensurate with, or less than, those of *C. aspersum aspersum*.

## Biosecurity import risk assessment

### *Angiostrongylus* spp.

#### Background

*Angiostrongylus* spp. are parasitic nematodes with an intermediate host (snails) and a definitive host (vertebrates). Some Angiostrongylus spp. use rodents as definitive hosts but most of these species display a high degree of host specificity (Anderson, 2000). Nematodes of this genus can cause potentially life-threatening diseases in several accidental host animal species, including humans (Colella *et al*., 2016a). There are 4 species of *Angiostrongylus* (*A. vasorum*, *A. abstrusus*, *A. cantonensis*, *A. chabaudi*) known to use *C. aspersum* as an intermediate host. It is not yet known if *A. mackerrasae* can infect *C. aspersum*, but it is possible given its similarities to other *Angiostrongylus* species (Aghazadeh *et al*., 2015a). *Angiostrongylus* spp. are common in wild snails (Anderson, 2000; Barcante *et al*., 2003; Colella *et al*., 2016b; Elsheikha *et al*., 2018; Bowman *et al*., 2002).

*Angiostrongylus* spp. infection is known as angiostrongylosis in vertebrate animals, including humans who consume raw snails. The disease primarily causes cardiopulmonary signs, but can also lead to eosinophilic meningitis, eosinophilic encephalitis and ocular angiostrongylosis (Wang *et al*., 2012; Sawanyawisuth and Sawanyawisuth, 2008; Sawanyawisuth *et al*., 2013). Clinical signs from A. vasorum can be highly variable with cardiopulmonary disorders (e.g. bronchopneumonia, cardiac dysfunction) being most common, followed by coagulopathies, then neurological presentations. Other non-specific signs have been reported such as ocular and gastrointestinal signs. Eosinophilic meningitis can also be associated with lifelong neurological impairments and well as loss of vision (da Silva and Morassutti, 2021). The variability in clinical presentations can make it challenging to differentiate from other canine diseases on initial presentation. There is a long prepatent period of 1 to 4 months (Di Cesare *et al*., 2014). Cases with severe clinical signs, such as marked pulmonary hypertension, coagulopathies or neurological signs, are associated with a fatal outcome in up to 30% of cases. Conversely, in the absence of severe clinical signs 95% of cases have a positive outcome (Chapmen *et al*., 2004; Willesen *et al*., 2007).

While disease is rare in Australia, eosinophilic meningitis caused by *A. cantonensis* has been found in a patient who ingested two garden slugs as part of a dare (Senanayake *et al*., 2003). In parts of the world where disease is more common, occurrence is also attributed to direct ingestion of the intermediate host as part of the diet (Alicata and Brown, 1962) or from ingestion of products contaminated with infective larvae in secretions from infected intermediate hosts (Hughes and Biggs, 2002). In Taiwan cases of eosinophilic meningitis were attributed to contaminated raw vegetable juice (Tsai *et al*., 2004). Cases of angiostrongylosis in humans have occurred with the accidental consumption of infected snails on produce (Yeung *et al*., 2013).

*Angiostrongylus cantonensis* is present in Australia (Aghazadeh *et al*., 2015a; Chan *et al*., 2015; Prociv and Carlisle, 2001), as is *A. mackerrasae* (Bhaibulaya, 1968; Aghazadeh *et al*., 2015b). *A. abstrusus* has been found in Australia (Anderson, 2000, Barrs *et al*., 2008), though an extensive study of its range does not appear to have occurred. *A. vasorum* was only identified in Australia in a dog imported from the United Kingdom (Tebb *et al*., 2007) and *A. chabaudi* has not been detected in Australia yet.

Little is known about *A. chabaudi*. Gherman *et al*, 2016 state it rarely infects felids. In contrast, Varcasia *et al*, 2014 citing Biocca (1957), stated wildcats may have high burdens of *A. chibaudi*. Whether wildcats (a different subspecies to the domestic cat) are a reservoir of infection for domestic cats in various parts of the world has not been clarified to date. Additional studies to determine its distribution and impact on the health of wild and domestic cats have been recommended (Varcasia *et al*., 2014).

#### Technical information

##### Agent properties

*Angiostrongylus* spp. are part of the phylum Nematoda, Class Chromadorea, Order Strongylida, family Angiostrongylidae (GBIF, 2022a). *A. vasorum*, *A. abstrusus*, *A. cantonensis*, and *A. chabaudi*, are known to infect *C. aspersum* as an intermediate host. *Helicarion* spp. appear to be the most important intermediate hosts of *A. mackerrasae*, at least in South-East Queensland and is theorised to be instrumental in the spread of the nematode south (Prociv *et al*., 2000)

All 5 *Angiostrongylus*spp. have similar lifecycles involving the cardiovascular system and/or lungs in the definitive host. The L1 larvae are shed by the definitive host and are either ingested by or penetrate the cuticle of the intermediate snail host (Anderson, 2000). The location of the larvae within the intermediate host varies depending on the species. *A. abstrusus* larvae are mostly found in the foot and viscera of the snail (Colella *et al*., 2015). *A. cantonensis* larvae are found in the foot, lung, muscle, liver and connective tissue of the snail (HeXiang *et al*., 2009; Modry *et al*., 2021). The period of larval development within mollusc species can vary depending on temperature and location (Alicata, 1965; Anderson, 2000; HeXiang *et al*., 2009). The larvae undergo 2 moults in the snail where they develop into an infective third stage (L3). The definitive (and accidental) hosts become infected when they directly ingest the infected snail (Cowie, 2013). They can also become indirectly infected through ingestion or contact with items contaminated by L3 containing secretions from the infected snail (Hughes and Biggs, 2002), noting L3 larvae persist in water and on moist soils (Alicata, 1965; Modry *et al*., 2021).

##### Epidemiology

*Angiostrongylus* spp. require intermediate and definite hosts to be present in the same environment in order to complete their lifecycle. Each *Angiostrongylus* species has different specific intermediate and definitive hosts (Table 4).

Table 4 *Angiostrongylus species* that use *C. aspersum* as a host, their intermediate and definitive hosts, their presence in Australia and in snail farms

| Species | Intermediate host | Alternative intermediate host | Definitive host | Accidental or Paratenic host | Present in Australia | References |
| --- | --- | --- | --- | --- | --- | --- |
| *Angiostrongylus vasorum* | Snail (range of terrestrial and aquatic)  *C. aspersum*  *Omalonyx matheroni*  *Subulina octona*  *Arionater*  *A. rufus*  *Biomphalaria glabrata*  *Bradybaena similaris*  *Deroceras leave*  *Laevicaulus alte*  *Prosoples javanicum* | Slug  Frog  *Rana temporaria* may also act as intermediate host | Canines  *Canis lupus familiaris*  *Cerdocyon thous*  *Ducicyon azarae*  *D. vetulus*  *Fennecus zerda*  *Vulpes vulpes* | Paratenic: lizards, mice, rats | No – only found in one imported dog | (Rosen *et al*., 1970; Bolt *et al*., 1993; Tebb *et al*., 2007; Ferdushy and Hasan, 2010; Bessa *et al*., 2000; Mozzer *et al*., 2011; Di Cesare *et al*., 2015)  (Ferdushy *et al*., 2009; Morgan *et al*., 2005; Grewal *et al*., 2003) |
| *Angiostrongylus abstrusus* | Snail  *C. aspersum*  *Agriolimax laevis*  Snails of multiple genera | Slug | Felines  *Felis catus*  *Acinonyx jubatus*  *Panthera onca*  *Felis concolor*  *Panthera leo*  *Panthera tigris altaica* | - | Yes | (Hobmaier and Hobmaier, 1935; Mackerras, 1957; Anderson, 2000; Bjork *et al*., 2000; Gonzalez *et al*., 2007; Barrs *et al*., 2008; Thiengo *et al*., 2008; Moskvina, 2018; Di Cesare *et al*., 2013) |
| *Angiostrongylus cantonensis*  Different strains exist | Snail  *C. aspersum*  *Achatina fulica*  *Bellamya ingallsiana*  *Bradybaena similaris*  *Cipangopaludina chinensis*  *Derceras leave*  *Euglandina rosea*  *Girasia peguensis*  *Indoplanorbis exustus*  *Laevicaulus alte*  *Macrochlamys resplendens*  *Microparmarion malayanus*  *Opeas javanicum*  *Pupina complanate*  *Pila ampullaceal*  *P. scutate*  *Quantula striata*  *Subulina octona*  *Vaginalus plebeius*  *Veronicella alte* | Slugs  *Girasia peguensis*  *Microparmarion malayanurn*  *Deroceras leave*  *Vaginalus plebeius*  *Veronicella alte*  Frogs, experimentally  *Xenopus laevis*  *Rana chensinensis* | Rodents  24 rat species with *Rattus norvegicus* and *R. rattus* identified as responsible for greatest distribution | Low host specificity  Accidental hosts: humans, Australian native fauna, domesticated and wild animals | Yes | (Alicata, 1965; Bhaibulaya, 1991; Anderson, 2000; Aghazadeh *et al*., 2015a; Chan *et al*., 2015; Stokes *et al*., 2007; Kim *et al*., 2014; Colella *et al*., 2015) |
| *Angiostrongylus mackerrasae* | Snail  *Helicarion* spp. | Slugs | Rodents  *Rattus fuscipes*  *Melomys cervinipes*  *R. leucopus* | Accidental host: *Pteropus alecto* | Yes, thought to be endemic to Australia | (Prociv *et al*., 2000; Mackie *et al*., 2013) |
| *Angiostrongylus chabaudi* | Snail  *C. aspersum* | - | Felines | - | No | (Colella *et al*., 2015; Colella *et al*., 2017) |

The main definitive hosts are rodents for *A. cantonensis* and *A. mackerrasae*, are felines for *A. abstrusus* and *A. chabaudi*, and canines for *A. vasorum*. A wide variety of species can act as paratenic hosts, which are optional hosts that have eaten infected snails. Paratenic hosts include terrestrial crabs, freshwater shrimp, frogs, fish and seas snakes. Their importance as a source of infection is not well understood (Prociv *et al*., 2000).

The importation of infected snails such as *Achatina fulica* into areas previously considered infection-free is thought to have facilitated the spread of the parasite (particularly *A. cantonensis*) and thus plays an important role in the epidemiology of the disease (Colella *et al*., 2015). *Achatina (Lissachatina) fulica* was introduced in Brazil in the 1980s for commercial purposes (escargot farming) and is now widespread, and has become a pest of public health and veterinary importance (Thiengo *et al*., 2008). *Pomacea canaliculata* was introduced into Taiwan from Argentina for use in commercial snail farms and ended up spreading throughout eastern Asia (Lv *et al*., 2011). Two species, *Achatina fulica* and *P. canaliculate,* are thought to be closely associated with angiostrongylosis in China. These were first imported into mainland China in 1931 and 1981 respectively, and have rapidly extended their geographic ranges. They are now listed as invasive species in China (Lv *et al*., 2009). The snail *P. canaliculata* was introduced into China in 1981 and has become the key intermediate host for *A. cantonensis* in China (Lv *et al*., 2011).

Global warming, changes in phenology of mollusc intermediate hosts and movements of wild reservoirs have been implicated in spreading mollusc-borne parasites, including *A. vasorum* (Di Cesare *et al*., 2015). *C. aspersum* is endemic in most regions of the world and is thought to be increasing the range of any parasites that it is harbouring (Di Cesare *et al*., 2015).

##### Diagnosis

The diagnosis of *Angiostrongylus*spp. in a snail occurs post-mortem. The shell is removed, and body minced before being digested in a solution of hydrochloric acid and pepsin. This digestive fluid is then filtered and spun in a centrifuge. Larvae can be counted and examined for identifying characteristics under a light microscope (Di Cesare *et al*., 2013; Cardillo *et al*., 2018; Giannelli *et al*., 2014). The larvae can also have their DNA sequenced using PCR to provide a definitive identification (Giannelli *et al*., 2014; Colella *et al*., 2017).

##### Treatment

No treatment options for snails were found. Various treatments are available for humans (Evans-Gilbert *et al*., 2014; Senanayake *et al*., 2003). Sporadic cases of *A. cantonensis* in Australia occur and medical treatment may not prevent morbidity or death.

Prevention in dogs and cats includes avoiding ingestion of infected intermediate hosts (e.g. snails, slugs, molluscs), and the use of routine prophylactic anthelmintics. The monthly use of spinosad/milbemycin oxime, imidacloprid 10%/moxidectin 2.5% has been shown to effectively eliminate larvae thereby preventing clinical angiostrongylosis.

In infected definitive felid hosts, several drugs are available to eliminate parasites. Fenbendazole (20–25 mg/kg, oral, once a day for 5 days, or 50 mg/kg, oral, once a day for 15 days), moxidectin (1 mg/kg, 1–3 applications topically in combination with imidacloprid), abamectin (0.3 mg/kg, subcutaneous, repeated in 2 weeks) and eprinomectin have all been identified as effective treatment of *A. abstrusus* in cats (Baydar and Kaya, 2021).

For *A. vasosum*, several drugs are available to eliminate worms in infected canid definitive hosts. Monitoring for anaphylaxis through treatment is advised (Elsheikha *et al*., 2018). A range of anthelmintic protocols have been described for treatment: a spot-on formulation containing imidacloprid 10% and moxidectin 2.5% has an efficacy of 85.2 % after one single application; the use of 25 mg/kg fenbendazole orally once per day for 20 days has an efficacy of 91.3%; and the use of 0.5 mg/kg milbemycin oxime given orally once a week for 4 weeks has a 84.4% efficacy (Di Cesare and Traversa, 2014). In addition to anthelmintics, infected dogs may require supportive treatment according to the presenting clinical signs. This could include oxygen supplementation for respiratory compromise, transfusion of whole blood for coagulopathies, and ACE inhibitors and diuretics for cardiac dysfunction (Di Cesare and Traversa, 2014)

The sudden killing of these worms (as occurs with the use of levamisole at 7.5 mg/kg s/c q24 h for 2 days, then 10 mg/kg s/c q24 h for 2 days) can result in a severe allergic reaction, so concurrent treatment with corticosteroids to suppress potential anaphylaxis due to sudden release of worm antigens in those cases is recommended.

##### Occurrence

*Angiostrongylus abstrusus* and *A. cantonensis* are found globally, including cases within Australia (Barcante *et al*., 2003; Colella *et al*., 2016a; Elsheikha *et al*., 2018).

While *A. abstrusus* has been recorded in Australia, an extensive study of its range does not appear to have occurred. It has the potential to be found throughout the country

*Angiostrongylus cantonensis* is endemic to temperate and tropical regions (York *et al*., 2014; Anderson, 2000). Its current range includes Southeast Asia, Japan, the Pacific Islands, and Australia, as well as parts of Central and South America, the United States and the Caribbean (Wang *et al*., 2008; Dard *et al*., 2017; Qvarnstrom *et al*., 2007; Lv *et al*., 2011;Evans-Gilbert *et al*., 2014; Stockdale Walden *et al*., 2017). In Australia, *A. cantonensis* has been found in Sydney, Brisbane and in forests around Jervis Bay, NSW (Aghazadeh *et al*., 2015a; Chan *et al*., 2015; Stokes *et al*., 2007). The full distribution of *A. cantonensis* within Australia is not known.

There is a patchy distribution of *A. vasorum* in many parts of the world, including tropical, sub-tropical and temperate regions (e.g. Europe, Africa, South America, Asia) (Colella *et al*., 2016a; Elsheikha *et al*., 2018). Clear evidence exists of both an increase in the number of cases within known endemic foci (e.g. Denmark, France, UK) and the appearance of new foci in several regions that were previously free of infection (Morgan and Wall, 2009; Morgan *et al*., 2010; Lurati *et al*., 2015). During the 1990s in the UK, *A. vasorum* was largely confined to dogs living in the south-east and south-west of England and south Wales. From 2008, the disease has become more common and has been identified as far north as Scotland (Elsheikha *et al*., 2018). *A. vasorum* has also been found in *C. aspersum* in Scotland, indicating that this was not just a matter of movement of an infected definitive host (Helm *et al*., 2015). *A. vasorum* is not present in Australia and has only been found in one imported dog (Tebb *et al*., 2007)

*Angiostrongylus chabaudi* has so far only being identified in several European countries in wild and domestic cats (Colella *et al*., 2017). It is known to be present in Italy and Greece (Morgan *et al*., 2021). It has not yet been found in Australia.

#### Current biosecurity measures

For imported dogs and cats, the current generic biosecurity measures for internal parasites for Group 3 countries are that: a government approved veterinarian must treat the dog twice with an internal parasite treatment effective against internal parasites (nematodes and cestodes). The two treatments must be administered at least 14 days apart and within 45 days before export. The second treatment must be given within five (5) days before export.

There are no specific biosecurity measures in place for *Angiostrongylus* spp.

#### Risk assessment

*Cornu aspersum* is present in most regions of the world and is a vector of *Angiostrongylus*spp. and other parasites (Di Cesare *et al*., 2013). Climate change is likely increasing the range of both intermediate and definitive hosts and may further increase expansion of the associated parasitic disease (Morgan and Wall, 2009). The presence of *A. cantonensis* in the Australian environment has been raised as a possible issue of concern for future reintroductions of native birds or mammals into areas known to have infected intermediate or definitive hosts. Fatal cases of infection have been reported in flying foxes (*Pteropus* spp.) and tawny frogmouths (*P. strigoides*) (Prociv and Carlisle, 2001; Barrett *et al*., 2002; Monks *et al*., 2005; Spratt, 2005; Ma *et al*., 2013).

Entry and exposure assessment:

* *Angiostrongylus* spp. are parasitic nematodes that utilise an intermediate host (snails) and a definitive host (vertebrates).
* *Angiostrongylus* spp. have been detected in snails and produce sold for human consumption.
* There are 4 species of *Angiostrongylus* (*A. vasorum*, *A. abstrusus*, *A. cantonensis*, *A. chabaudi*) known to utilise *C. aspersum* as an intermediate host. *A. mackerrasae* may also infect *C. aspersum*.
* *A. vasorum*, *A. abstrusus*, *A. cantonensis* and *A. chabaudi* have similar lifecycles involving the cardiovascular system and/or lungs in the definitive host. The L1 larvae are shed by the definitive host and are either ingested by or penetrate the cuticle of the intermediate snail host.
* Most *Angiostrongylus* spp. display a high degree of host-specificity, except for *Angiostrongylus* species using rodents as definitive hosts. The definitive hosts for *A. vasorum* are canines and for *A. chabaudi* are felines.
* *Angiostrongylus cantonensis, A. abstrusus*, and *A. mackerrasae* have been found in Australia and therefore will not be considered further in this risk assessment.
* *Angiostrongylus vasorum* has a wide geographic distribution and is known to occur in dog populations in Africa, Europe, and North and South America.
* Prevalence estimates for *A. vasorum* in infected countries are not readily available, but are likely to vary considerably between regions and within a population based on exposure to infected intermediate or paratenic hosts.
* *Angiostrongylus vasorum* has spread across Europe since the 1990s and is now broadly endemic. It is likely that spread of the infected intermediate snail hosts have contributed (not simply movement of definitive hosts). Affected countries includes countries proposed for sourcing of snails for breed stock in Australia.
* To determine if a snail is infected with any *Angiostrongylus* spp., the snail must first be euthanised. No treatment options for snails were found.
* A possible pathway for introduction of *A. vasorum* to Australia is via infected snails (as the intermediate host) if there is exposure to the L1 larvae shed by infected canids (as the definitive host). Entry of infected snails onto a snail farm may occur where the security of the farm is not sufficient to prevent entry and exit of snails from the external environment, or fattening or grow-out stages are situated outdoors. It is noted that many, if not all, commercial snail farms have external grow out stages.
* There is a long history of importation of dogs from countries where canine pulmonary angiostrongylosis due to infection with *A. vasorum*, is endemic. Available records indicate only one case of canine pulmonary angiostrongylus in Australia in 2007. The entry likelihood via a potentially infected definitive host is still considered to be very low (Department of Agriculture, 2013).
* The prevalence of *A. chabaudi* and where it occurs in the world is unclear. *A. chabaudi* has so far only being identified in wild and domestic cats in several European countries. It is known to be present in Italy and Greece. At least one paper has called for further research and greater awareness of this parasite.

Based on this information, the likelihood of entry of *A. vasorum* or *A. chabaudi* associated with the importation of infested farmed *C. aspersum* and exposure of potential hosts in Australia was estimated to be moderate.

This takes into account that no snail farm, even well established commercial facilities, could be considered ‘closed’ systems from the biosecurity perspective. *A. vasorum* appears to be endemic in a number of European countries and *A. chabaudi* has also been reported. A closed system does not allow for snail trade or the cryptic nature of snails which may enter and leave undetected within even very well-established facilities with certification of their premises and procedures.

Consequence assessment:

* *Angiostrongylus* spp. cause angiostrongylosis in vertebrate animals, including humans, in the form of respiratory and cardiac disease, as well as eosinophilic meningitis, eosinophilic encephalitis and ocular angiostrongylosis.
* The definitive (and accidental/paratenic) hosts become infected when they directly ingest the infected snail host. They can also become infected indirectly through ingestion or contact with items contaminated by L3 containing secretions shed from the infective intermediate host.
* In the UK in the 1990s, *A. vasorum* was largely confined to dogs living in the south-east and south-west of England and south Wales, but from 2008, the disease has become more common and has been identified as far north as Scotland. This was not likely to be just a matter of movement of an infected definitive host (dogs), as *A. vasorum* was also found in *C. aspersum* in Scotland. A similar scenario could unfold in Australia.
* Adverse health outcomes may occur for Australian dogs, with significant flow on impacts to their owners, if the intermediate host (snail) is imported in large volumes over time, spreads widely, and is carrying exotic *Angiostrongylus* spp.
* *A. chabaudi* has not been detected in Australia. There is general uncertainty about the prevalence and potential impacts on domestic cats. It can cause cardio‑pulmonary disease in felids. Similarly to dogs, adverse health outcomes may eventuate in cats if the intermediate host (snail) is imported in large volumes over time and is carrying *A. chibaudi.*
* Treatment options for definitive hosts, and humans exist but are not always successful in preventing morbidity and mortality in the occasional cases of angiostrongylus reported in Australia due to endemic *Angiostrongylus* spp.

Based on this information, the likely consequences of establishment and/or spread of *Angiostrongylus* spp. associated with the importation of *C. aspersum* was estimated to be low.

##### Conclusions

Based on the preceding information, the likelihood of entry of *Angiostrongylus* spp. associated with imports of *C. aspersum* sourced from commercial, government certified snail farms is considered to be moderate and the likely consequences of establishment and/or spread of *Angiostrongylus*spp. is considered low. Using Table 3, the likelihood of entry and exposure (moderate) was combined with the likely consequences of establishment and/or spread (low), which resulted in a risk estimation for *Angiostrongylus* spp. *(A. vasorum, A. chibaudi)* of low risk.

As the overall risk of *Angiostrongylus*spp. associated with the importation of *C. aspersum* from commercial government certified snail farms that produce snails for human consumption is low and therefore does not achieve Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *Angiostrongylus*spp. are required.

#### Risk management measures

**Importation of adult snails**: for the risk of *Angiostrongylus* spp. associated with the importation of live adult garden snails (*C. aspersum)*, to achieve Australia’s ALOP, the snails must be imported into an Approved Arrangement site, where they will be reared in isolation from other snails and the definitive host, thus breaking the lifecycle of the nematode. The next generation of *C. aspersum* would be free of *Angiostrongylus* spp. and would be eligible for release from the Approved Arrangement site.

**Importation of hatchling snails**: these are hatchlings less than 10 days old that have not been released from the hatching environment and have not been exposed to plant substrate. As these hatchling snails have also not been exposed to the definitive hosts for *Angiostrongylus* spp., the risk for hatchling snails will achieve Australia’s ALOP.

### *Brachylaima* spp.

#### Background

*Brachylaima* spp. are parasitic trematodes that have 2 intermediate hosts (snails) and a definitive host (vertebrate). There are four species of *Brachylaima* (*B. aspersae, B. cribbi, B. mascomai* and *B. llobregatensis*) that have been recorded using *C. aspersum* as an intermediate host*.* Brachylaima spp. are common in field populations of *C. aspersum* (Gerard *et al*., 2020; Kose *et al*., 2015) and snail farms (Segade *et al*., 2013; Segade *et al*., 2011). Brachylaima spp. can cause brachylaimiasis in humans. The clinical signs of brachylaimiasis in humans include abdominal pain and recurrent diarrhoea and it has a mortality rate of 5–10% without treatment (Gracenea and Gallego, 2017; Butcher, 2016). Brachylaimaisis is rare but has been found in several patients in Australia through infection with B. cribbi (Butcher, 2016; Butcher *et al*.; 1996, Butcher *et al*., 1998). The trematode would have been ingested with the snails (likely raw or undercooked) or from vegetables that had been in contact with snails (Butcher, 2016).

B. cribbi is present in Australia but the other Brachylaima spp. found in *C. aspersum* have not yet been recorded in Australia. (Gallego *et al*., 2014).

#### Technical information

##### Agent properties

*Brachylaima*spp. are part of the phylum Platyhelminthes, class Trematoda, order Diplostomida and family Brachylaimidae (GBIF, 2022b). Four species are known to occur: *B. aspersae*, *B. cribbi*, *B. mascomai* and *B. llobregatensis* (Butcher and Grove, 2001; Gonzalez-Moreno and Gracenea, 2006; Segade *et al*., 2011; Gracenea and Gonzalez-Moreno, 2002). Recent studies have shown that little is known about *Brachylaima* spp., with new species being discovered in France (Gerard *et al*., 2020). Further snail research elsewhere may uncover more species.

##### Epidemiology

All four *Brachylaima* spp. have a similar lifecycle. *Brachylaima* eggs are ingested by the first snail host, in which cercariae will develop in sporocysts. When they reach maturity, the cercariae exit the host and crawl until they find a second host (another snail). Once inside the second host, they will form metacercariae usually located in the snail kidney. If the snail is ingested by a vertebrate (the definitive host), the trematode may become adults and colonise the intestine where they will mate. Eggs are excreted with the host faeces. If these eggs are ingested by a suitable first intermediate host, the lifecycle continues (Segade *et al*., 2011; Butcher, 2016; Gracenea and Gonzalez-Moreno, 2002).

The intermediate and definitive hosts relevant to the lifecycle depend on the *Brachylaima* spp. (Table 5). The most thorough study on hosts was done in Australia with *B. cribbi*. In Australia, all snail species tested by Butcher and Grove (2005) were found to be second intermediate hosts of *B. cribbi*. The snail species tested included six introduced and two native species (Butcher and Grove, 2005b). This indicates that other snail species not yet tested could also be intermediate hosts. Mammals, birds and lizards were found to be definitive hosts but the authors suggest that more research is needed to fully ascertain host range (Butcher and Grove, 2005b). Birds likely spread the parasite broadly while mice and lizards contribute to local dispersion (Butcher and Grove, 2005b). There is seasonal variation in *B. cribbi* infection in snails, with infection rates being higher in winter and spring, and lower in summer (Butcher and Grove, 2005a).

Table 5 *Brachylaima* species that use *C. aspersum* as a host, their intermediate and definitive hosts, their presence in Australia and in snail farms

| Species | Intermediate host 1 | Intermediate host 2 | Definitive host | Present in Australia | Detected in snail farms | References |
| --- | --- | --- | --- | --- | --- | --- |
| *Brachylaima aspersae* | Snail  *C. aspersum* | Snail  *C. aspersum* | Rodent  mouse (*Mus musculus*) | No | Yes | (Segade *et al*., 2013; Segade *et al*., 2011) |
| *Brachylaima cribbi* | Snail  *C. aspersum*  *Theba pisana*  *Cochlicella acuta*  *Cochlicella barbara*  *Microxeromagna armillata* | Snail  Introduced:  *C. aspersum*  *Theba pisana*  *Cochlicella acuta*  *Cochlicella Barbara*  *Cernuella virgata*  *Microxeromagna armillata*  Native:  *Succinea australis*  *Strangesta gawleri* | Bird:  Emu  (*Dromaius novaehollandiae*)  Chickens  (*Gallus gallus*)  Pigeon  (*Columba livia*)  Little raven  (*Corvus mellori*)  Black bird  (*Turdus merula*)  Starling  (*Sturnus vulgaris*)  Mammal:  Mouse  (*Mus domesticus*)  Sheep  (*Ovis* spp.)  Cat (*Felis catus*)  Reptile:  Shingleback lizards (*Tiliqua rugosa*) | Yes | Unknown | (Butcher *et al*., 1996; Butcher and Grove, 2001; Butcher, 2016; Butcher and Grove, 2005b; Butcher and Grove, 2003) |
| *Brachylaima Ilobregatensis* | Snail  *C. aspersum* | Snail  *C. aspersum*  *Theba pisana* | Rodent  *Crocidura russula*  *Mus spretus*  *Mus musculus* | No | No | (Gonzalez-Moreno and Gracenea, 2006; Gallego *et al*., 2014) |
| *Brachylaima mascomai* | Snail  *Pseudotachea splendida* | Snail  *C. aspersum*  *Theba pisana*  *Otala punctata*  *Pseudotachea splendida* | Rodent  Rattus norvegicus  Rattus rattus  Mus musculus  Crocidura russula  Meriones unguiculatus  Apodemus sylvaticus | No | No | (Gracenea and Gonzalez-Moreno, 2002; Gracenea and Gallego, 2017; Gallego *et al*., 2014) |

The definitive host for *B. aspersae*, is the domestic mouse, *Mus musculus* (Segade *et al*., 2011). All other *Brachylaima* spp. should be considered as non-specific as they have several definitive hosts, usually rodents (Table 5).

Gerard *et al*., 2023 suggest that *C. aspersum* is capable of trapping cercariae (trematode larvae) in its shell, thus possibly reducing the intensity of infestation by these parasites.

##### Diagnosis

To determine if a snail is infected with *Brachylaima* spp., it needs to be dissected to look for the presence of sporocysts and metacercariae (Segade *et al*., 2013; Segade *et al*., 2011). In definitive hosts, dissections can also be used to look for the presence of adults in the gastrointestinal tract and eggs can be found in faeces (Butcher and Grove, 2005b; Butcher and Grove, 2001).

In humans, symptoms are linked to intestinal issues (abdominal pain, diarrhoea, weight loss) (Butcher, 2016). Symptoms can last from one month to one year. Diagnosis involves looking for eggs in faeces (Butcher, 2016; Butcher *et al*., 1996).

##### Treatment

Snails can be treated with praziquantel by mixing it with food (Gallego and Gracenea, 2015).

In humans, brachylaimiasis was successfully treated with a daily dose of 20 mg/kg of praziquantel for three days. The treatment had no side effect (Butcher, 2016).

##### Occurrence

*B. cribbi* has been found in Australia (Victoria, South Australia and Western Australia). The origin of this parasite has not been confirmed. It is unclear whether *B. cribbi* is native or introduced (Butcher and Grove, 2003). *B. aspersae* is currently only found in Spain, while *B. ilobregatensis* is found in Spain and Algeria (Gonzalez-Moreno and Gracenea, 2006; Gallego *et al*., 2014) and *B. mascomai* is found in Spain and South Africa (Gracenea and Gonzalez-Moreno, 2002). In the Republic of Türkiye, a *Brachylaima* species was detected in *C. aspersum* on pasture, with 2% of individuals infected. The *Brachylaima* were not identified at the species level (Kose *et al*., 2015). Recently, Gerard *et al*., (2020) identified a novel *Brachylaima* species on *C. aspersum* in France. This species has yet to be named. Prevalence in 2 wild populations was high with 10% and 73% of snails being infected (Gerard *et al*., 2020).

*Brachylaima* spp. are commonly found in *C. aspersum* sold in markets (Gracenea and Gallego, 2017; Gallego *et al*., 2014). In Spain, the overall percentage of snails infected in markets was over 40%, and closer to 60% in autumn (Gracenea and Gallego, 2017). Studies have shown that imported *C. aspersum* sold in markets in Spain were infected with different *Brachylaima* species (Gallego *et al*., 2014). Imported snails could therefore contribute to the geographic spread of *Brachylaima* if they come in contact with local intermediate and definitive hosts (Gallego *et al*., 2014).

*B aspersae* is prevalent in snail farms and was found in every farm sampled in Spain (Segade *et al*., 2011; Segade *et al*., 2013), with metacercariae found in 10 to 97% of snails (Segade *et al*., 2013). Infections was more common in *C. a. aspersum* than *C. aspersum* *maximum*) (Segade *et al*., 2013). Segade *et al*., (2011) suggested that controlling mice, the definitive host of *B. aspersae*, would help reduce the infection rates of *C. aspersum* on farms (Segade *et al*., 2013).

#### Current biosecurity measures

There are no specific biosecurity measures for *Brachylaima* spp. in Australia.

#### Risk assessment

Entry and exposure assessment:

* *Brachylaima* spp., including species not reported in Australia, may be common in snail farms overseas. *Brachylaima* spp. have been found in snails sold in markets and natural environments.
* *Brachylaima* spp. have been poorly studied; however, at least four species are known to use *C. aspersum* as a host (three of which have not been reported in Australia). More species are likely to be described in the future.
* *C. aspersum* is capable of trapping [cercariae](https://en.wikipedia.org/wiki/Cercariae) ([trematode](https://en.wikipedia.org/wiki/Trematode) larvae) in its shell, thus possibly reducing the intensity of infestation by these parasites.
* *Brachylaima* sporocysts and metacercariae can only be detected by microscopic examination following dissection of the snails.
* Infested wild or field reared snails may introduce the infection to a snail farm. It is unknown whether testing is, or would be, regularly undertaken for this parasite on commercial premises.
* A probable pathway for the introduction of exotic *Brachylaima* spp. is via the importation of parasitised imported snails (infested intermediate host). These snails may then shed metacercaria that infect other intermediate hosts or that may be consumed by definitive vertebrate hosts. The possible impact on Australian wildlife is unknown.

Based on this information, the likelihood of entry of *Brachylaima* spp. associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be high.

Consequence assessment:

* Importation of *C. aspersum* for human consumption has been identified as a risk for the expansion of *Brachylaima* spp. distribution.
* Because detection requires dissection of the snail, an incursion of exotic *Brachylaima*spp. would likely remain undetected until a suitable survey was undertaken, or associated disease presented.
* Some species of *Brachylaima* can infect humans.
* There is potential for serious health impacts of *Brachylaima* infested *C. aspersum* on other definitive hosts (likely rodents). This has not yet been scientifically investigated.
* It is possible that the introduction, spread and establishment of exotic *Brachylaima* spp. could adversely impact native and introduced animals in Australia. There is little information in the available scientific literature to suggest consequences of animal biosecurity significance.

Based on this information, the likely consequences of establishment and/or spread of *Brachylaima*spp. associated with the importation of *C. aspersum* was estimated to be very low from an animal biosecurity perspective.

The consequence assessment rating should be reviewed by relevant experts within the Australian Government Department of Health and Aged Care as this agency has responsibility for estimating the level of adverse impacts on human health that might eventuate associated with importation of this parasite.

##### Conclusions

Based on the preceding information, the likelihood of entry of *Brachylaima*spp. associated with the importation of infested *C. aspersum* sourced from commercial, government certified snail farms is considered to be high and the likely consequences of establishment and/or spread of *Brachylaima*spp. are considered to be very low. Using Table 3, the likelihood of entry and exposure (high) was combined with the likely consequences of establishment and/or spread (very low), which resulted in a risk estimation for *Brachylaima* spp. of very low.

As the overall risk of *Brachylaima*spp. associated with the importation of *C. aspersum* is very low and therefore achieve Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *Brachylaima*spp. are not required.

### *Crenosoma vulpis*

#### Background

*Crenosoma vulpis* identified by Dujardin (1845) is a lungworm nematode that causes chronic respiratory disease in domestic dogs in parts of Europe and in the north-eastern region of North America (Bihr and Conboy, 1999; Maksimov *et al*., 2017; Fuehrer *et al*., 2020; Elsheikha *et al*., 2014). It is considered endemic in red fox populations across these regions. *C. vulpis* needs an intermediate host snail, including *C. aspersum*, and a vertebrate definitive host canid to complete its development (Colella *et al*., 2016b; Fuehrer *et al*., 2020). It is thought that *C. aspersum* plays an important role in infecting animals in the Mediterranean basin and in Austria (Colella *et al*., 2016b). Other snail species, particularly the giant African snail *Achatina fulica*, have been identified as important intermediate hosts of canine lungworm species, including *C. vulpis* (Penagos-Tabares *et al*., 2019).

*C.* *vulpis* has not been detected in Australia.

#### Technical information

##### Agent properties

*C. vulpis* is part of the phylum Nematoda, class Secernentea, order Strongylida, and family Crenosomatidae (Anderson, 2000). Four haplotypes of *C. vulpis* in wild and domestic carnivores were identified in Italy, with only one haplotype being infective to dogs and all four being infective to foxes (Latrofa *et al*., 2015).

The lifecycle of *C. vulpis* starts in the definitive host, a canid. The first-stage larvae (L1) are coughed out of the lungs and swallowed, and then pass through with the faeces. Once in the environment the L1 infects the intermediate host, a snail, via the foot. It will then develop into the infective third-stage larvae (L3) within 17 days (Stockdale and Hulland, 1970). In infected snails, larvae are primarily located in the viscera (69%) and foot (31%) (Colella *et al*., 2016b). L3 *C. vulpis* can also be shed by the snail host and may remain infective in the environment for up to 8 weeks (Robbins *et al*., 2021).

The definitive host becomes infected after ingesting L3 larvae, either from environmental contamination or from ingesting an infected intermediate host (Stockdale and Hulland, 1970; Robbins *et al*., 2021). Canids can be infected through eating or licking contaminated material or surfaces.

Once ingested, the L3 larva penetrates the definitive host’s intestines and will reach its lungs within 20 hours using the hepatic portal system (Anderson, 2000; Elsheikha *et al*., 2018). The larvae colonise the small bronchi and bronchioles of the lung. They moult twice before reaching adulthood (Nevarez *et al*., 2005; Elsheikha *et al*., 2018). Females are ovoviviparous and will start laying larvae 19 days post-infection. Adults are large, measuring 5 to 10 mm in length and can live up to nearly 10 months (Anderson, 2000).

The parasite causes microscopic lesions around the bronchi and bronchioles and to a lesser extent around the alveoli (Nevarez *et al*., 2005). Crenosomosis is a frequent cause of chronic respiratory disease in domestic dogs (Elsheikha *et al*., 2018), although it is rarely fatal (Maksimov *et al*., 2017).

##### Epidemiology

Definitive hosts of *C. vulpis* are foxes (*Vulpes vulpes*), coyotes (*Canis latrans*), dogs (*Canis familiaris*) and badgers (*Meles meles*) (Bihr and Conboy, 1999; Barutzki and Schaper, 2009; Nelson *et al*., 2007; Popiolek *et al*., 2009).

A large range of terrestrial gastropods, including both snails and slugs can serve as the intermediate hosts for *C. vulpis* (Stockdale and Hulland, 1970; Anderson, 2000; Conboy *et al*., 2017). *C. aspersum* is a suitable intermediate host (Colella *et al*., 2016b).

There is seasonal variation in *C. vulpis* infection and distribution is thought to be influenced by precipitation and temperature (Tolnai *et al*., 2015).

##### Diagnosis

The diagnosis in the intermediate host involves killing the snail and digesting it in a solution of hydrochloric acid and pepsin before being spun in a microcentrifuge (Colella *et al*., 2016b). Any nematode larvae can be examined under a light microscope. It is also possible to identify *C. vulpis* larvae using a duplex RT-PCR (Lange *et al*., 2018). However, availability and other details of this test are not clear.

In definitive hosts, diagnoses are made through detection of first-stage larvae in faeces or via transtracheal wash samples.

##### Treatment

There is no treatment for the immediate snail host.

In domestic dogs, treatment includes the administration of Febantel (14 mg/kg, oral, once a day for 7 days) and fenbendazole (25–50 mg/kg, oral, once a day for 3–14 days) or with oral milbemycin oxime (0.5 mg/kg) and moxidectin spot-on treatments with clinical cure reportedly occurring 7–10 days post-treatment with an efficacy of 98–99% (Conboy, 2004; Elsheikha *et al*., 2018).

##### Occurrence

*Crenosoma vulpis* is endemic in red fox populations across Europe and north-east North America and the range of this nematode is thought to be expanding (Bihr and Conboy, 1999) (Maksimov *et al*., 2017;Rinaldi *et al*., 2007). In North America, occurrence primarily occurs in Canada’s Atlantic provinces (New Brunswick, Newfoundland-Labrador, Nova Scotia, PEI), with some cases also reported in Ontario, Quebec, New York and Illinois (Nelson *et al*., 2007; Maksimov *et al*., 2017; Barutzki and Schaper, 2011; Conboy *et al*., 2009).

The expansion of *C. vulpis* is a likely consequence of the red fox carrying the parasite across geographical boundaries and bringing the parasite into contact with domestic dogs and other susceptible wildlife (Latrofa *et al*., 2015).

There is no record of *C. vulpis* in snail farms. However, absence of evidence does not infer evidence of absence. There may have been no incentive to specifically look for this parasite on snail farms to date. In the absence of targeted surveillance, the prevalence of this parasite is undefined.

#### Current biosecurity measures

For imported dogs and cats, the current generic biosecurity measures for internal parasites for Group 3 countries are that: a government approved veterinarian must treat the dog twice with an internal parasite treatment effective against internal parasites (nematodes and cestodes). The two treatments must be administered at least 14 days apart and within 45 days before export. The second treatment must be given within five (5) days before export.

There are no specific biosecurity measures for *C. vulpis* in Australia.

#### Risk assessment

Entry and exposure assessment:

* *C. vulpis* is endemic in red fox populations across Europe and north-east North America and the range of this nematode is thought to be expanding. Coyotes, dogs and badgers are other definitive hosts.
* *C. vulpis* can use *C. aspersum* as an intermediate host.
* The first-stage larvae (L1) are coughed out of the lungs of the canid definitive host, swallowed and are then passed in faeces. Once in the environment the L1 infects the intermediate host, a snail, via the foot. It will then develop into the infective third-stage larvae (L3) in the snail.
* There are no records of *C. vulpis* on snail farms to date. However, this information should receive appropriate weighting considering a general paucity of information and detailed scientific reports assessing the prevalence of pathogens and parasites in farmed snails that may represent a biosecurity (rather than food safety) risk.
* Diagnosis in the snail is a laboratory procedure. It involves killing and digesting the snail in a solution of hydrochloric acid and pepsin before being spun in a microcentrifuge. A species-specific PCR has also been used for diagnosis. However, test specific details are unclear.
* There is no treatment for the immediate snail host.
* A possible pathway for introduction of *C. vulpis* to Australia from farmed *C. aspersum* could be through importation of parasitised snails that have entered a snail farming unit (e.g. at the fattening stage outside) or been introduced into a facility (snail farm). In addition, if parasitised dogs or foxes are allowed access to the snail breeding facilities, and the snails are able to come into contact with the faeces of these canids, the lifecycle could be maintained within the facility. (Note: Depending on the mode of operation of the sourcing facility, this latter scenario with dogs accessing commercial government certified facilities may be more improbable than entry of parasitised snails).

It is also noted that there is a history of importation of dogs from countries where *C. vulpis* is endemic. No cases have been reported in Australian dogs. The risk management applied currently for internal parasites for imported dogs into Australia is considered appropriate, and infections in pet dogs in Australia would be highly likely to have been detected through veterinary investigation over the years.

Based on this information, the likelihood of entry of *C. vulpis* associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be moderate.

Consequence assessment:

* *C. vulpis* has not yet been detected in Australia.
* *C. vulpis* is a lungworm nematode that causes chronic respiratory disease in domestic dogs in parts of Europe and in the north-eastern region of North America. Crenosomosis is reported as a common cause of chronic respiratory disease in domestic dogs, although it is rarely fatal.
* Terrestrial gastropods, including both snails and slugs, serve as the intermediate hosts for *C. vulpis*. *C. aspersum* has been identified as a suitable intermediate host.
* Canids can be infected by eating intermediate hosts with L3 forms of this parasite. L3 *C. vulpis* can be shed by the gastropod intermediate host and remain infective in the environment for up to eight weeks. The canine hosts could become infected through eating or licking contaminated material or surfaces or eating parasitised snails.
* Since foxes and dogs are definitive hosts of this parasite, the presence of feral foxes and dogs in Australia could result in a rapid spread of this parasite (introduced by parasitised *C. aspersum* and potentially extending to *C. aspersum* populations already present in Australia) from agricultural to urban areas and natural areas. There is also a significant population of working and assistance dogs in rural and regional areas overlapping with populations of foxes and feral dogs. This parasite appears to prefer cold temperatures so this could limit its spread in parts of northern Australia.
* The introduction of this parasite into Australia would impact domestic dog and dingo populations resulting in chronic respiratory disease of a new origin. There is a large companion dog owning population in Australia, a number of whom could be impacted by introduction of a parasite that could cause significant disability in their animals as well as expense related to veterinary investigations and treatment. The impact on the working ability of working dogs on animal production could also be significant for primary producers.

Based on this information, the likely consequences of establishment and/or spread of *C. vulpis* associated with the importation of *C. aspersum* was estimated to be lowfrom an animal biosecurity perspective.

##### Conclusions

Based on the preceding information, the likelihood of entry of *C. vulpis* associated with imports of *C. aspersum* from commercial, government certified snail farms is considered to be moderate and the likely consequences of establishment and/or spread of *C. vulpis* is considered low. Using Table 3, the likelihood of entry and exposure moderate was combined with the likely consequences of establishment and/or spread (low), which resulted in a risk estimation for C. vulpis of low risk.

As the overall risk of *C. vulpis* associated with the importation of *C. aspersum* is low and therefore does not achieve Australia’s ALOP with respect to animal biosecurity risks, specific biosecurity measures for *C. vulpis* are required.

#### Risk management measures

**Importation of adult snails**: for the risk of *C. vulpis* associated with the importation of live garden snails (*C. aspersum)*, to achieve Australia’s ALOP, the snails must be imported into an Approved Arrangement site, where they will be reared in isolation from other snails and the definitive host, thus breaking the lifecycle of the nematode. The next generation of *C. aspersum* would be free of *C. vulpis* and eligible for release from the Approved Arrangement site.

**Importation of hatchling snails**: these are hatchlings less than 10 days old that have not been released from the hatching environment and have not been exposed to plant substrate. As these hatchling snails have also not been exposed to the definitive hosts for *C. vulpis*, the risk for hatchling snails will achieve Australia’s ALOP.

### *Dicrocoelium dendriticum*

#### Background

*Dicrocoelium dendriticum* (Rudolphi, 1819)*,* formerly known as *Distoma dendriticum,* is commonly known as the lancet liver fluke. It is a trematode that infests herbivorous and omnivorous mammals including humans. It requires two intermediate hosts, a snail and an ant (Ismail and Gurelli, 2018). The definitive hosts are often ruminants such as sheep and cattle, but can include rabbits, pigs and dogs (Kahl *et al*., 2021). *D. dendriticum* infection is known as dicrocoeliasis and can infect many vertebrate species, including humans. Infection with *D. dendriticum* in humans has been associated with consuming raw liver of infected animals. However, infections are rare and often spurious (Schweiger, 2008). Most infections are subclinical and any clinical signs are typical of liver disease, as caused by other liver flukes (Jeandron *et al*., 2011).

This parasite is reported as problematic in northern and southern Europe, northern Africa, western and eastern America and Asia (Arbabi *et al*., 2021). It is unclear whether this parasite is in Australia. According to the online Global Biodiversity Information Facility database (GBIF), *D. dendriticum* is not present in Australia (GBIF, 2022c). The Atlas of Living Australia database only includes one species of *Dicrocoelium* (*D. antechini*) (Cribb, 1992). This parasite has been reported to infect people in the Republic of Türkiye, Iran, Canada and Kyrgyzstan (Jeandron *et al*., 2011). Several references state that it is present in Australia, but the source of this information is unclear. For example, according to Otranto and Traversa, 2003, it is found in some focal points in Australia, but no references are provided for this statement. Noting that cattle and sheep may show very large numbers of these parasites in the bile ducts it is probable that surveillance activities would have detected its presence if it were in Australia.

This group of trematodes usually have wide host range. *Dicrocoelium dendriticum* has only been recorded in one study on *C. aspersum* (Ismail and Gurelli, 2018).

#### Technical information

##### Agent properties

The genus *Dicrocoelium* (Rudolphi, 1819) is in the phylum Platyhelminthes, class Trematoda, order Plagiorchiida and in family Dicrocoeliidae.

The adults are semitransparent, 6–12 mm length and 1.5–2.5 mm width, with a small oral sucker and a larger ventral sucker. The eggs are dark brown, 35–45 µm in length and 22–30 µm wide (Kahl *et al*., 2021)

Adults reside in the gall bladder and bile ducts of the definitive host. Embryonated eggs are shed via bile to the intestine and then are excreted in faeces into the environment. The first intermediate host, snails, feed on faeces consuming the eggs. The egg will hatch into a miracidium, then migrates through the intestinal wall into the hepatopancreas. The parasite remains inside the snail for 3 to 4 months while developing from miracidium to cercariae. The cercariae migrate to the respiratory tract and are expelled through the respiratory movements once coated in slime. The second intermediate host, an ant, will consume the ball of slime with the cercariae. Some of the cercariae will migrate into the suboesophageal ganglion causing a shift in ant behaviour where the ants cling to the vegetation. This increases the chance of being consumed by the definitive host. The remaining cercariae encyst in the abdomen of the ant. Once in the definitive host, the metacercariae excyst as juvenile fluke which follow the bile duct to the liver to mature into adult fluke thus completing the lifecycle (Kahl *et al*., 2021).

##### Epidemiology

*D. dendriticum* requires two suitable intermediate hosts co‑occurring in the environment to complete its lifecycle: a snail and an ant. Over 90 mollusc species, mostly from the Helicidae and Geomitridae families, can be intermediate hosts (Otranto and Traversa, 2003; Sanchez *et al*., 2021). *Cernuella virgata* and *Cochlicella acuta* are commonly first intermediate hosts (Mitchell *et al*., 2017; Fasanella *et al*., 1995). These two introduced species are now widespread in Australia and are major pests of grain crops (Baker, 2008). *D. dendriticum* has been detected in *C. aspersum* in only one study to date. This study was conducted in Türkiye (Kose *et al*., 2015).

*D. dendriticum* is less pathogenic than the liver fluke *Fasciola hepatica* (Cengiz *et al*., 2010). It’s pathology is often masked by the pathological effects of other parasitic infections of ruminants (Arbabi *et al*., 2011). Unlike other flukes, *D. dendriticum* doesn’t migrate within the host so doesn’t cause tissue damage or bleeding. This helps explain the lack of clinical symptoms in infected animals (Kahl *et al*., 2021). However, Arbabi *et al*. (2018) observed that infection can cause slow development and impaired fertility in animals, which can reduce product quality and rate of production. It could also increase costs of anthelmintic treatments and costs of stock replacement (Arbabi *et al*., 2018).

In humans, *D. dendriticum* infection can cause abdominal pain, weight loss and chronic diarrhoea. Patients are often asymptomatic. Heavier infections can cause symptoms including anaemia, constipation, and eosinophilia (Cengiz *et al*., 2010).

Egg excretion in sheep in the Mediterranean regions is seasonal, peaking during winter (Otranto and Traversa 2002). The eggs are very resistant, and they can survive for up to 20 months on pastures with the right conditions (dry and calcareous soils) (Otranto and Traversa, 2003).

##### Diagnosis

The diagnosis of *D. dendriticum* in humans is through the FLOTAC technique which identifies the presence of eggs in the stool.

It can be diagnosed in ruminants at slaughterhouses through the presence of worms in the bile duct and gall bladder, swollen livers, whitish spots and scarring on liver and cholangitis (Arbabi *et al*., 2011).

Diagnosis in the intermediate hosts involves dissection followed by either microscopic morphological examination or molecular analysis using PCR. The PCR molecular analysis has a higher sensitivity (Mitchell *et al*., 2017).

##### Transmission

Definitive hosts are infected through ingestion of the second intermediate host (the ant). The trematode inside the ant intermittently changes its behaviour so it clings to vegetation, increasing its chance of being eaten by the definitive host (Kahl *et al*., 2021). The ant will attach to vegetation in cooler temperatures, but will detach if temperature exceeds 20°C. The ant then returns to normal worker activities but will reattach to the same place once the temperature cools down (Martin-Vega *et al*., 2018).

Terrestrial snails enter hibernation during colder months. Transmission of *D. dendriticum* is higher in warmer months when snails are more active (Ismail and Gurelli, 2018).

##### Treatment

There is no treatment for intermediate hosts.

There are several treatments available for the definitive hosts. A field study showed that the use of praziquantel at 50 mg/kg in sheep was 95.9% effective against *D. dendriticum* (Akkaya, Deniz and Sezen 2006). Another study on merino sheep using 5% and 15% suspension of netobimin at a dose rate of 20 mg/kg showed good efficacy with 90.80% and 91.50% respectively (Senlik *et al*., 2008). In another study, a double administration of micronized albendazole had higher efficacy than a single treatment against *D. dendriticum* (Bosco *et al*., 2015). Khanjari *et al*., (2010) recommends preventative measures through treating sheep with albendazole and triclabendazole before autumn (Khanjari *et al*., 2010).

##### Occurrence

*D.* *dendriticum* occurs in northern and southern Europe, northern Africa, western and eastern America and Asia (Manga-Gonzalez, 2001). Distribution reporting may be confounded by linking to fasciolosis distribution.

Prevalence in livestock varies. Infection rates of cattle and sheep can reach 37.5% and 46% respectively in Northern France, or 80% overall in Poland (Tarry, 1969). Infection rates of up to 100% are reported in sheep located in the Mediterranean and Middle East (Jeandron *et al*., 2011). In a more recent study, 21.1% (sheep) and 7.0% (goats) were infected with *D. dendriticum* in southern Germany (Alstedt *et al*., 2022).

*D. dendriticum* was not observed in 4,680 sheep imported from Australia to Jordan (Maraqa *et al*., 2005).

#### Current biosecurity measures

There are no specific biosecurity measures for *D. dendriticum* to manage risk of entry of this parasite into Australia.

#### Risk assessment

Entry and exposure assessment:

* *D.* *dendriticum* occurs in northern and southern Europe, northern Africa, western and eastern America and Asia. Its prevalence in livestock varies.
* *D. dendriticum* was detected in *C. aspersum* in one study overseas (l., 2015). *Cernuella virgata* and *Cochlicella acuta* are the common first intermediate hosts, and these species are now widespread in Australia.
* The status of *D.* *dendriticum* on snail farms is unknown.
* The life cycle requires two suitable intermediate hosts co-occurring in the environment: a snail and an ant, and definitive hosts which could be ruminants or humans.
* Diagnosis of infestation in the snail intermediate host requires dissection of the snail.
* A possible pathway for introduction of *D. dendriticum* to Australia from farmed *C. aspersum* could be through importation of parasitised snails that have entered a breeding and rearing facility (snail farm) undetected, or been deliberately introduced into an insecure breeding and rearing source facility; for example, from sourcing from other breeders or from a fattening stage in the external environment.

Based on this information, the likelihood of entry of *D.* *dendriticum* associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be low.

This takes into account that no snail farm, even well-established commercial facilities, could be considered ‘closed’ systems from a biosecurity perspective. A closed system does not allow for snail trade or the cryptic nature of snails which may enter and leave undetected within even well-established facilities that are regulated through certification of their premises and procedures.

Consequence assessment:

* It is unclear whether *D. dendriticum* is present in Australia. The available evidence suggests it is exotic as surveillance activities, including routine abattoir inspections, would likely detect it.
* The introduction of *D. dendriticum* to Australia could have adverse consequences for the agriculture industry. As seen in Iran, it is possible that infection could slow development, and impair fertility in livestock, which can reduce product quality and rate of production. It could also increase costs associated with anthelmintic treatments and costs of stock replacement. There is a lack of clinical symptoms in grazing animals.
* *D. dendriticum* can infect humans so may be pose a public health risk.
* Two intermediate snail hosts are already widespread in Australia (*C. virgata* and *C. acuta*), therefore *D. dendriticum* already has some suitable intermediate hosts present. Australia has very diverse ants. Once introduced, it is likely *D. dendriticum* could spread geographically.
* The level of direct economic loss due to partial or total condemnation of livers infected with infected with *D. dendriticum* was found to be equivalent to the loss due to liver fluke (*Fasciola hepatica*) infection of sheep, goats and cattle in a 2018 Iranian study (Arabi *et al*., 2018)

Based on this information, the likely consequences of establishment and/or spread of *D. dendriticum* associated with the importation of *C. aspersum* were estimated to be low from an animal biosecurity perspective. However, the consequence assessment rating should be reviewed by relevant experts within the Australian Government Department of Health and Aged Care for consideration of potential adverse impacts on human health that might eventuate associated with importation of this parasite.

##### Conclusions

Based on the preceding information, the likelihood of entry of *D. dendriticum* associated with imports of *C. aspersum* from commercial government certified snail farms is considered to be low, and the likely consequences of establishment and/or spread of *D. dendriticum* is considered low. Using Table 3, the likelihood of entry and exposure (low) was combined with the likely consequences of establishment and/or spread (low), which resulted in a risk estimation for *D. dendriticum* of very low risk.

As the overall risk of *D. dendriticum* associated with the importation of *C. aspersum* from commercial, government certified snail farms is very low and therefore achieves Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *D. dendriticum* are not required.

### *Phasmarhabditis* spp.

#### Background

*Phasmarhabditis* *hermaphrodita* (synonym= *Pellioditis hermaphrodita* and *P. californica* are soil nematodes that are facultative parasites of gastropods. They are also able to live and reproduce in slug faeces and other organic material in the environment (Rae *et al*., 2009). *Phasmarhabditis* *hermaphrodita* has been well studied but comparatively little is known about *P. californica* (Mc Donnell *et al*., 2020).

Several slug and snail species are susceptible to *P. hermaphrodita* and *P. californica*, including *C. aspersum* (Williams and Rae, 2015; Andrus *et al*., 2020; Grannell *et al*., 2021). *P. hermaphrodita* is lethal to snails and slugs when associated with the bacterium *Moraxella osloensis* and is sold as a biocontrol agent across Europe as ‘Nemaslug’ (Holley, 2020).

There is no record of *Phasmarhabditis*spp. in Australia (Mc Donnell *et al*., 2020).

#### Technical information

##### Agent properties

*Phasmarhabditis* spp. are in the Phylum Nematoda, class Chromadorea, order Rhabditida and family Rhabditidae. *Phasmarhabditis hermaphrodita* can parasitise several mollusc families including Limacidae, Agriolimacidaem, Arionidae, Milacidae, Vaginulidae and Helicidae (Holley, 2020).

*Phasmarhabditis hermaphrodita* has a mutualistic relationship with bacteria carried in its intestines (Hapca *et al*., 2007). The slug or snail becomes infected when encountering the infective juvenile stage (dauer larvae) in the environment. The nematode larvae enter the dorsal integumental pouch of the snail and then penetrate the shell cavity (Rae *et al*., 2007). Once inside the shell cavity, the bacteria are released, producing an endotoxin (Grewal and Grewal, 2003). This leads to an accumulation of fluid, causing swelling in the shell cavity and death within 4–21 days after the initial infection (Rae *et al*., 2007). After death of the host, the nematodes distribute themselves amongst the entirety of the corpse to feed on the decaying body (Rae *et al*., 2007; Tan and Grewal, 2001a). They can also reproduce on slug faeces or other rich bacterial substrates (Rae *et al*., 2007). New generations of dauer larvae will be produced and will migrate to the soil in search of their new host (Cutler and Rae, 2020). Dauer larvae are nonfeeding and rely on energy reserves until they infect a new host (Grewal and Grewal, 2003).

##### Epidemiology

Mortality of the host is caused by the bacteria, and not the nematode. Nematodes free from the bacteria were found to be non-pathogenic when injected directly into the host, while the bacteria killed the host without the nematode (Tan and Grewal, 2001b). Some snail species have developed resistance to *P. hermaphrodita* infection including *Cepaea nemoralis*, *Oxychilus helveticus*, *Disus rotundatus* and *Clausilia bidentata* (Williams and Rae, 2015). Resistance is also size and age dependent. Juvenile *C. aspersum* are susceptible but adults seem more resistant (Williams and Rae, 2015). The same resistance pattern was found with *P. californica* (Grannell *et al*., 2021).

A single host carcase can support the production of thousands of nematodes. Fifty nematodes feeding on the carcase can result in the production of between 15,000–40,000 infectious dauer larvae (Morris *et al*., 2018). These nematodes are also able to complete their life cycle in nonparasitic conditions, thus maximising their ability to survive in the environment (Tan and Grewal, 2001b).

Dauer larvae are sensitive to ultraviolet light, high temperatures and desiccation (El-Danasoury and Iglesias-Pineiro, 2017). However, they can survive relatively tough environmental conditions due to their thick cuticle and closed openings (Grewal and Grewal, 2003). The nematode is killed at temperatures over 35°C (Rae *et al*., 2007). It is thought that the nematodes manipulate the spatial behaviour of the host to ensure the host dies in the soil, hence protecting against high temperature and escaping predation. The nematodes requires a few days to complete their life cycle inside the cadaver, where under normal circumstances the cadaver would be consumed by predators within a matter of hours (Pechova and Foltan, 2008).

##### Diagnosis

The diagnosis of *Phasmarhabditis* spp. is performed on dead snails, historically through dissection, which is time consuming (Wilson *et al*., 2012). Wilson *et al* 2012 developed a new technique that was equally effective. This involved decapitating the host and incubating it for a week, allowing the development of the dauer larvae into adults.

##### Transmission

Transmission occurs in the soil when an infective larvae encounter a host. The larvae are attracted to mucus, faeces and volatile cues of snails and slugs (Andrus *et al*., 2020). Optimum temperature for growth and transmission is 17°C, but transmission can occur at temperatures as low as 5°C (Rae *et al*., 2007). Infection causes a great reduction in host movement and feeding, meaning environmental dispersal is minimal (Rae *et al*., 2007).

##### Occurrence

*P. hermaphrodita* was first discovered in Germany and believed to be European in origin (Howe *et al*., 2020). It is commercialised as Nemaslug and sold widely over Europe (14 countries) (MacMillan *et al*., 2006). It has also been found in New Zealand, Chile, Iran, Egypt and the western United States. It is unknown whether these populations are naturally occurring or illegally introduced with the biological control agent, Nemaslug (Nemaslug is not available for sale in NZ or USA). *P hermaphrodita*’s potential impacts on gastopods native to these areas is unknown (Howe *et al*., 2020). *P. hermaphrodita* has not been recorded in Australia. Australian average temperatures are higher than the optimum temperature of 15°C for transmission, and therefore it may not spread as much as in other countries (Charwat and Davies, 2001).

*P. californica* has been isolated from the U.K., U.S.A., New Zealand and Canada (Grannell *et al*., 2021), but its native range remains unknown (Mc Donnell *et al*., 2020).

#### Current biosecurity measures

There are no specific biosecurity measures for *Phasmarhabditis*spp. in Australia.

#### Risk assessment

Entry and exposure assessment:

* *Phasmarhabditis* spp. can infect *C. aspersum*.
* Infected gastropods can take up to three weeks to die from infection. It is possible that an infected snail would be moved while not showing symptoms. Adults seem more resistant and therefore they could carry the parasite without showing signs.
* The prevalence of *Phasmarhabditis* spp. in snail farms is unknown. It has been detected in various countries but its distribution is not well-described.
* Transmission occurs when infective larvae in the soil encounter a host (snail or slug).
* In a commercial government certified snail farm, it would be expected that loss of breeding stock (discovery of snail carcases) would be investigated, the infection detected, and suitable action taken to protect stock.
* Only dead snails can be diagnosed with *Phasmarhabditis* spp.

Based on this information, the likelihood of entry of associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be very low.

Consequence assessment:

* There is no record of *Phasmarhabditis* spp. in Australia.
* *Phasmarhabditis* spp. can infect many slugs and snails.
* Environmental spread is limited due to decreased movement of the infected host. It is unlikely to move rapidly from any source of introduction.
* Average temperatures in Australia may be too high to support populations of this nematode in some areas, as optimum temperatures of infection are reported as being 15–17°C; however, transmission can occur at temperatures as low as 5°C.
* The introduction of this nematode to Australia could adversely impact native snails and slugs.

Based on this information, the likely consequences of establishment and/or spread of *Phasmarhabditis*spp. associated with the importation of *C. aspersum* was estimated to be negligible from an animal biosecurity perspective.

##### Conclusions

Based on the preceding information, the likelihood of entry of *Phasmarhabditis*spp. associatedwith imports of *C. aspersum* sourced from commercial, government certified snail farms is considered to be very low and the likely consequences of establishment and/or spread of *Phasmarhabditis*spp. is considered negligible. Using Table 3, the likelihood of entry and exposure (very low) was combined with the likely consequences of establishment and/or spread (negligible), which resulted in a risk estimation for *Phasmarhabditis* spp. of negligible.

As the overall risk of *Phasmarhabditis*spp.associated with the importation of *C. aspersum* is negligible and therefore achieves Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *Phasmarhabditis*spp. are not required.

The consequence assessment rating should be reviewed by relevant experts within the Australian Government Department of Climate Change, Energy the Environment and Water as this agency has responsibility for estimating the level of adverse impacts on native slugs and snails that might eventuate associated with import.

### *Riccardoella limacum*

#### Background

*Riccardoella limacum* is a parasitic mite of snails. It is common on *C. aspersum* (Turk and Phillips, 1946) and can be found in snail farms (Segade *et al*., 2013). It has not been recorded in Australia.

#### Technical information

##### Agent properties

*Riccardoella limacum* (Acari: Prostigmata: Ereynetidae) is a parasitic mite of snails. Little information exists on *Riccardoella*spp. in general as they are small, live inside hosts and are therefore inconspicuous. Two species have been confused in past studies. However, a study showed that one species, *R. limacum* infects snails, while the other species *R. oudemans* infect slugs (Graham *et al*., 1993). *R. limacum* can infect several snail species, with *C. aspersum* being one of the most common hosts (Turk and Phillips, 1946).

*R. limacum* is a hematophagous mite, feeding on its host blood while living in the mantle cavity. Eggs are laid in the host’s lung cavity and hatch within eight to 12 days (Baker, 1970). There are three nymphal stages (Baker, 1970). Newly hatched nymphs have six legs and move fast allowing dispersal (Turk and Phillips, 1946). Adults are 0.4 mm long (Baker, 1970). Mating occurs inside the host and males are present for a short time only (Turk and Phillips, 1946). The total life cycle of *R. limacum* takes 19 to 23 days to complete at temperatures between 20.1–25.1°C (Baker, 1970). Therefore, several generations can occur in a year. There is a seasonal pattern in mite numbers, which peak during autumn (reviewed in Baur and Baur, 2005). Adult mites rarely survive winter in the host (Baker, 1970) and infections decrease over the host hibernating period (Haeussler *et al*., 2012). Eggs appear to be the overwintering stage with nymphs hatching in spring (Baker, 1970; Haeussler *et al*., 2012). The number of mites increases with host size with larger individuals having higher infestation load (Schupbach and Baur, 2010b).

##### Epidemiology

While *R. limacum* does not directly kill its hosts, several studies have found that infestation impact snail health. These impacts can be subtle. In a study where healthy *Arianta arbustorum* snails were infested with *R limacum*, shell growth and reproduction (growth of albumen glands and reaching sexual maturity) were not affected by the mite, but they ingested less food (Wacker, 2008). In the same snail species, less eggs were produced when more mites were present in the host (Schupbach and Baur, 2008b). Several studies showed that snail mortality in infested individuals was higher in winter (Wacker, 2008; Shupbach and Baur, 2008b; Hausller *et al*., 2012) and mortality increased with parasite load (Schupbach and Baur, 2010b).

Susceptibility to *R. limacum* is dependent on the snail family with some families being more resistant (Schupbach and Baur, 2010b). Intensity of infection was found to vary between field populations with some populations harbouring higher numbers of *R. limacum* per individual (Baur and Baur, 2005).

##### Diagnosis

The definitive diagnosis of *R. limacum* infestation is through dissection when the mite is found inside the host. While it is possible to observe mites in snails by looking through the respiratory pore, this is not a reliable technique and mite eggs cannot be detected (Baur and Baur, 2005). Dissection is therefore a necessary part of diagnosis. In a recent study on *Riccardoella tokyoensis*, it was shown that snails did not show any sign of inflammation after being artificially infected. (Waki *et al*., 2021). It is impossible to assess infestation based on snail health only.

##### Transmission

Transmission occurs when snails come in contact with each other (Schupbach and Baur, 2010a) during courtship and mating, but they can also use mucus to infect new hosts (Schupbach and Baur, 2008a). Fresh mucus is most effective for transmission (Schupbach and Baur, 2008a). Transmission is influenced by the infested snail’s parasite load. The heavier the infestation, the more transmission will occur (Schupbach and Baur, 2010a).

##### Occurrence

Parasitic *Riccardoella*spp. mites may be easily missed due to their small size and because they live within their hosts. Therefore, they could be more widely distributed than currently thought. *R. limacum* is currently widespread in Europe. The greatest height it was found in Europe was at 1290 metres in Switzerland (Baur and Baur, 2005).

*R. limacum* is commonly found in snail farms. Segade *et al*., (2013) found this mite species to be infecting predominantly *C. aspersum aspersum* (Segade *et al*., 2013). In one of the farms sampled, close to 60% of individuals were infected by mites. Mites were less prevalent in *C. aspersum maximum* (Segade *et al*., 2013).

*R.* *oudemans* is widespread in Australia however *R. limacum* has not been recorded in Australia to date (B. Halliday, personal communication).

#### Current biosecurity measures

There are no specific biosecurity measures for *R. limacum* in Australia.

#### Risk assessment

Entry and exposure assessment:

* *R. limacum* is widespread in Europe and it is commonly found in snail farms.
* The hosts for these parasites seem to be restricted to snails and slugs.
* As *R. limacum* is found within the mantle cavity of the snail, it is hard to detect, and dissection is necessary for diagnosis. Infested hosts may seem healthy based on visual inspection.
* Transmission occurs when snails come in contact with each other. The mites may also use snail mucus to infect new hosts. Any opportunity for entry of wild snails or field reared snails onto snail farms represents a potential entry point for this pathogen onto the farm.
* A probable pathway for the introduction of *R. limacum* is from infested imported snails transmitting *R. limacum* to local *C. aspersum* populations, which is one of the most common hosts. However, *R. limacum* can infest several snail species.

Based on this information, the likelihood of entry of *R. limacum* associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be high.

Consequence assessment:

* *R. limacum* has not been reported in Australia. A related species, *R. oudemans* is present and infests slugs.
* *R. limacum* has become widespread in Europe which suggests it is an efficient parasite in terms of ability to spread.
* Several studies have found that infestation with *R. limacum* impacts snail health, although such impacts may be subtle. Reported adverse effects for affected snails include lower food consumption, lower egg production, and increased mortality.
* Detrimental impacts on the existing population of *C. aspersum* might widely be viewed favourably by the Australian public, given its pest status. However, *R. limacum* may also infest Australia’s native snails, of which there are some 2,500 species (Stanisic *et al*., 2022). This could result in potentially significant adverse outcomes for the ongoing viability of these species.
* Infestation of native snails (and possibly native slugs) could have wider implications for Australia’s biodiversity and conservation efforts.

Based on this information, the likely consequences of establishment and/or spread of *R. limacum* associated with the importation of *C. aspersum* was estimated to be very low.

The consequence assessment rating should be reviewed by relevant experts within the Australian Government Department of Climate Change, Energy the Environment and Water as this agency has responsibility for estimating the level of adverse impacts to native fauna that might eventuate associated with importation of this parasite.

##### Conclusions

Based on the preceding information, the likelihood of entry of *R. limacum* associated with imports of *C. aspersum* sourcedfrom commercial, government certified snail farms is considered to be high and the likely consequences of establishment and/or spread of *Riccardoella*spp. are considered to be very low. Using Table 3, the likelihood of entry and exposure (high) was combined with the likely consequences of establishment and/or spread (very low), which resulted in a risk estimation for *R. limacum* of very low.

As the overall risk of *R. limacum* associated with the importation of *C. aspersum* is very low and therefore achieve Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *R. limacum* are not required.

### *Tetrahymena* spp.

#### Background

*Tetrahymena rostrata* and *Tetrahymena limacis* are free-living, ciliated protozoans that can also be facultative parasites of terrestrial snails and slugs (Haites *et al*., 2021). Some studies suggest that the gastropod host is not harmed or injured when infected (Michelson, 1971), while others found that high parasitic loads in gastropods are often fatal once the ciliates enter internal organs, particularly the kidneys (Zhang and Vdacny, 2021).

*T. rostrata* is present in Australia and has been isolated from the egg of the grey field slug *Deroceras reticulatum* (Watt *et al*., 2021).

*T. rostrata* is being considered for its potential as a biocontrol agent for *D. reticulatum* (Watt *et al*., 2021).

#### Technical information

##### Agent properties

*Tetrahymena* spp. are part of the phylum Ciliophora, class: Oligohymenophorea, order Hymenostomatida and family Tetrahymenidae. These protozoans have four developmental stages: trophonts, tomonts, tomites and theronts (Haites *et al*., 2021).

The natural habitat for *T. rostrata* is diverse and can include soil, leaf litter and moss (Zhang and Vdacny, 2021). This species can survive harsh conditions and can persist in the environment for long periods (Wilson *et al*., 1998). As *T. rostrata* can go locally extinct due to the death of the host or exhaustion of its food source it must migrate frequently to find suitable environments to live (Kaczanowski *et al*., 2016).

Infection of *T. rostrata* is largely concentrated in the nephridium or kidneys of the host (Segade *et al*., 2009). Other areas of infection include the pulmonary chamber, muscle, albumen gland, hepatopancreas and gonads (Haites *et al*., 2021). Maximum densities in hosts can be very high. The maximum recorded by Segade *et al* (2009) was 6300 ciliates for petit gris *C. a. aspersum* and 15,700 ciliates for *C. aspersum maximum* (Segade *et al*., 2009).

*T limacis* protozoans were also concentrated in the kidneys but in heavy infestations were also found in the body wall and mantle shield. In heavy infections the ciliates can also replace the normal tissue structure (Michelson, 1971).

##### Epidemiology

*Tetrahymena* spp. have several host species. *Tetrahymena limacis* can use hosts from the Arionidae, Milacidae and Limacidae slug families and from the Bradybaenidae, Daudebardiidae, Helicidae (which includes *C. aspersum*), Hygromiidae, Succineidae, Vitrinidae and Zonitidae snail families. *Tetrahymena rostrata* is known to infect species such as *C. aspersum, Chochilcopa lubrica,* and more commonly, *Deroceras reticulatum* (Kaczanowski *et al*., 2016). It has been found in seven slug species in Agriolimacidae and Arionidae families and seven snail species in the Oleacinidae and Zonitidae families (Van As and Basson, 2004).

In *C. aspersum*, high *T. rostrata* density causes mantle swelling and severe renal pathology (Segade *et al*., 2009). It is reported that snails infected with *T. rostrata* have slower growth rates, lower feeding rates and decreased fecundity. Heavily parasitised *C. aspersum* suffer significant damage to the renal epithelium alongside deleterious effects such as the inability to retract their body back into their shell and mantle collar swelling (Zhang and Vdacny, 2021). Infected slugs will also have reduced tentacle mobility (Haites *et al*., 2021).

Prevalence of *Tetrahymena*spp. is significantly reduced after hibernation (Segade *et al*., 2013).

##### Diagnosis

*T. rostrata* is identifiable in the kidneys of infected snails such as *C. aspersum.* Diagnosis therefore requires dissection of the snail (Haites *et al*., 2021). It is assumed this also applies to *T. limacis.*

##### Transmission

The host becomes infected by trophonts. It is likely that snails or slugs are also infected by the free-swimming, excysted cells (theronts) that penetrate the host's urethras and renal organs, as mucus of snails enhances T. rostrata excystment (Kaczanowski *et al*., 2016). Snails can also undergo secondary self-infections by excysted cells which outcompete the remaining older cells (Kaczanowski *et al*., 2016).

Transmission of *T. limacis* is faecal-oral, with host becoming infected when ingesting *T. limacis.* Directinfection into the respiratory chamber also seems possible in *D. reticulatum* (Michelson, 1971).

##### Occurrence

The first discovery of *T. rostrata* occurred in New Zealand and today it can be found in the USA, North America, Europe and Australia (Kaczanowski *et al*., 2016). The distribution in Australia has not been defined.

*T. rostrata* is common in snail farms (Segade *et al*., 2009; Segade *et al*., 2013). A study into parasite prevalence in mixed system-based heliciculture farms in North-West Spain found *T. rostrata* in approximately 3% of *C. aspersum* populations involved in the study. However, infection rate can be higher with 12.6% of juveniles being infected (Segade *et al*., 2013).

A Spanish study found that *T. limacis* had an overall prevalence of 0.5% in heliciculture systems., with higher prevalence in juveniles with 29.4% of individuals infected (Segade *et al*., 2013).

#### Current biosecurity measures

There are no specific biosecurity measures for this species in Australia.

#### Risk assessment

Entry and exposure assessment:

* *T. rostrata* is known to be present in Australia; however, available information for *T. rostrata* is likely to be largely applicable to the related species, *T. limacis.*
* *T. limacis* are free-living, ciliated protozoans that are also facultative parasites of terrestrial snails and slugs.
* *T. limacis* has an overall prevalence of 0.5% in heliciculture systems., with a higher prevalence in juveniles with 29.4% of individuals infected.
* Transmission of *T. limacis* is faecal-oral, with the host becoming infected when ingesting T. limacis.
* Diagnosis requires dissection of the snail.

There is a paucity of relevant scientific information on *T. limacis*; however, based on this information, the likelihood of entry of associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be very low.

Consequence assessment:

* The status in Australia of *T. limacis* is unknown. The related species, *T. rostrata* is here.
* The impact of *Tetrahymena*spp. on their mollusc hosts is unknown; but there is a possibility that in heavy infections the ciliates could replace the normal tissue structure in parts of the snail or slug.
* The fact that *T. rostrata* is being considered for its potential as a biocontrol agent for *D. reticulatum* would suggest significant harm to the host; in this case the grey field slug*.*
* The full host range is unknown.
* *Tetrahymena* spp. could also have an impact on snail farms if initial stock is infected.
* The introduction of new species of these protozoans to Australia could adversely impact native slugs and snails.

Based on this information, the likely consequences of establishment and/or spread of *T. limacis* associated with the importation of *C. aspersum* was estimated to be negligible from an animal biosecurity perspective.

##### Conclusions

Based on the preceding information, the likelihood of entry of *T. limacis* associated with imports of *C. aspersum* sourced from commercial, government certified snail farms is considered to be very low and the likely consequences of establishment and/or spread of *T. limacis* is considered negligible Using Table 3, the likelihood of entry and exposure (very low) was combined with the likely consequences of establishment and/or spread (negligible), which resulted in a risk estimation for *T. limacis* of negligible**.**

As the overall risk of *T. limacis* associated with the importation of *C. aspersum* is negligible and therefore achieves Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *T. limacis* are not required.

The consequence assessment rating should be reviewed by relevant experts within the Australian Government Department of Climate Change, Energy the Environment and Water as this agency has responsibility for estimating the level of adverse impacts on native slugs and snails that might occur.

### *Troglostrongylus brevior*

#### Background

*Troglostrongylus brevior* (Gerichter, 1948) is a nematode lungworm that uses gastropods as intermediate hosts and cats as definitive hosts. It can cause serious broncho‑pulmonary disease in cats which are the definitive hosts and is often fatal in kittens (Traversa *et al*., 2014). It was initially considered as a parasite of wild cats, but its prevalence in domestic cats appears to be increasing. This is difficult to be certain of as it is believed that *T. brevior* infection has frequently been historically misidentified as *A. abstrusus* (Traversa *et al*., 2014). Several gastropods serve as intermediate hosts including *C. aspersum* (Brianti *et al*., 2013). *T. brevior* has not been detected in Australia.

#### Technical information

##### Agent properties

*T. brevior* is a nematode in the class Chromadorea, order Strongylida and family Crenosomatidae (GBIF, 2022). First stage larvae (L1) are released into the environment through faeces of the definitive host, cats. The larvae penetrate the integument of molluscs, the intermediate hosts. Inside the snail they develop until they become infective L3 larvae. The definitive host will become infected by eating the intermediate host or indirectly by eating a paratenic host that has ingested an infected snail (rodents, birds, reptiles, and amphibians) (Traversa *et al*., 2014). Kittens may also acquire infection through vertical transmission and ingestion of L3 stage larvae in their mother’s milk (Brianti *et al*., 2013).

Once inside the digestive system of the definitive host, the L3 larvae penetrate the intestinal tract and enter the blood steam or lymphatic system to eventually reach the lungs where they will mature to adults (Brianti *et al*., 2013). Adults live in the bronchi and bronchioles where they will reproduce (Traversa *et al*., 2014). Females release eggs in the lungs. These will hatch into L1 larvae, exit the respiratory system to be swallowed back by the host and travel down the digestive system until they are released in the environment through faeces for the lifecycle to recommence (Traversa *et al*., 2014). The lifecycle takes approximately 28 days to complete.

##### Epidemiology

Several gastropods serve as intermediate hosts of *T. brevior* including *Agriolimax*spp., *Helicella*spp., *Helix*spp., *Monacha*spp.and *Theba*spp. (Brianti *et al*., 2013)*.* These authors also note *C. aspersum* as an intermediate host.

Feline symptoms can range from subclinical to life threatening and include dyspnoea, mucoid-purulent nasal discharge, sneezing, depression, anorexia (Giannelli *et al*., 2014), as well as irreversible pulmonary hypertension and death (Morelli *et al*., 2020). Severity of infection is a result of the large body size of the nematodes and their location in the trachea, bronchi, and bronchioles (Brianti *et al*., 2013). Prevalence is higher in kittens (85.7%) than in adults (64.3%) (Falsone *et al*., 2014). There is limited data on prevalence and potential for co‑infections of cat metastrongylid species in Europe.

Seasonal patterns of transmission in southern Europe have been observed. A decrease in temperature favours the development of *T. brevior* in the snail, therefore infection rates in snails are highest in autumn and winter. The highest infection rates in cats are seen in spring when gastropods exit hibernation (Morelli *et al*., 2020).

##### Diagnosis

The diagnosis of *T. brevior* infection in the intermediate gastropod host involves dissection as for other lungworms.

Diagnosis in the definitive host require microscopic analysis of faecal samples following the ZnSO4 flotation and Baermann method. Diagnosis can be complicated by the non-specific nature of the clinical signs and an absence of clinical signs in some infected animals (Jefferies *et al* 2010 citing Travers and Guglielmini 2008).

##### Transmission

*T. brevior* infects the snail host as L1 larvae when they come in contact in the environment with faeces carrying an egg burden from an infected feline. Transmission between snails is also possible with L3 larvae being shed in the snail mucus (Brianti *et al*., 2021). Cats can acquire the infection directly through ingestion of infected intermediate hosts or indirectly through ingestion of paratenic hosts (birds, amphibians, reptiles or rodents), which have ingested the infected snails (Morelli *et al*., 2020). Research has also shown kittens can acquire infection through lactation (Brianti *et al*., 2013).

##### Treatment

In a pilot study, *T. brevior* infection was successfully treated in cats with Advocate® spot-on solution for cats (Bayer Animal Health GmbH) which contains 1% w/v moxidectin and 10% w/v imidacloprid (Diakou *et al*., 2019).

There are no known treatments for *T. brevior* in intermediate hosts.

##### Occurrence

*Troglostrongylus brevior* has been reported in the Mediterranean Basin, Eastern Europe and South America (Morelli *et al*., 2020). In Europe it has been recorded in Spain, Italy, Greece, Bulgaria, Romania and Cyprus (Deak *et al*., 2017). All of Mediterranean Europe provides suitable conditions for this parasite (Diakou *et al*., 2015). Infection rates can be very high in wild felids. In Italy, 71.4% of wildcats were shown to be infected (Falsone *et al*., 2014).

#### Current biosecurity measures

For imported dogs and cats, the current generic biosecurity measures for internal parasites for Group 3 countries are that: a government approved veterinarian must treat the dog twice with an internal parasite treatment effective against internal parasites (nematodes and cestodes). The two treatments must be administered at least 14 days apart and within 45 days before export. The second treatment must be given within five (5) days before export.

There are no specific biosecurity measures for *T. brevior* in Australia.

#### Risk assessment

Entry and exposure assessment:

* *T. brevior* may use *C. aspersum* as an intermediate host and infections may be increasing in domestic cats in Europe based on detection of a number of cases in domestic cats over the last decade. However, there is little information available for clear statements about prevalence.
* Domestic cats are known to eat snails but it can be assumed this is not their preferred diet. Therefore, detection of cases may argue for an increasing prevalence in the snail host and/or infested potential paratenic hosts. It may also reflect increased research effort.
* *T. brevior* has not been reported on snail farms. As with *C. vulpis*, this information should receive appropriate weighting considering a general paucity of information and detailed scientific reports assessing the prevalence of pathogens and parasites in farmed snails that may represent a biosecurity (rather than food safety) risk.
* Diagnosis in the snail is a laboratory procedure and requires dissection.
* There is no treatment available for snails.

Based on this information, the likelihood of entry of *T. brevior* associated with the importation of infested farmed *C. aspersum*, and exposure of potential hosts in Australia was estimated to be low.

Consequence assessment:

* *T. brevior* has not been detected in Australia in intermediate hosts or definitive hosts.
* Several gastropods serve as intermediate hosts of *T. brevior,* including *C. aspersum*. *T. brevior* would likely have ample intermediate, paratenic and definitive hosts (domestic and feral cats) in Australia if it were introduced. Once introduced, there is potential for this parasite to establish and spread.
* *T. brevior* can infect domestic cats and may be fatal in kittens.
* Feline symptoms can range from subclinical to life threatening and include dyspnoea, mucoid-purulent nasal discharge, sneezing, depression and anorexia, as well as irreversible pulmonary hypertension and death. Severity of infection is variable and some infected animals may show no clinical signs.
* Cats can be diagnosed through clinical signs and a faecal examination. Diagnosis can be complicated by the non-specific nature of the clinical signs in affected animals as well as the possibility of a complete absence of clinical signs in infected animals.
* Cats can be successfully treated based on limited studies.
* Impact on the domestic (companion) cat population in Australia, if *T. brevior* was introduced in the snail, is anticipated to be minimal. There is potential for kittens to succumb and incur treatment costs.

Based on this information, the likely consequences of establishment and/or spread of *T. brevior* associated with the importation of *C. aspersum* was estimated to be low.

##### Conclusions

Based on the preceding information, the likelihood of entry of *T. brevior* associated with imports of *C. aspersum* sourced from commercial, government certified snail farms is considered to be low and the likely consequences of establishment and/or spread of *T. brevior* is considered low. Using Table 3, the likelihood of entry and exposure (low) was combined with the likely consequences of establishment and/or spread low), which resulted in a risk estimation for *T. brevior* of very low.

As the overall risk of *T. brevior* associated with the importation of *C. aspersum* is very low and therefore achieves Australia’s ALOP with respect to animal biosecurity risks, additional biosecurity measures for *T. brevior* are not required.

### *Cornu aspersum* as a vector for plant pathogens

This section examines the potential for garden snails, including *C. aspersum*, to vector microorganisms that are not pathogenic to snails but may have an impact on plant life and health.

#### Relevant vectored plant pathogens

The potential transmission of plant disease agents by snails, including *C. aspersum*, has received little attention. A number of plant pathogenic organisms have been recovered from snails; however, there is no evidence that snails play a key role in the dispersal of these disease agents or are the primary vectors of these organisms:

* El-Hamalawi and Menge (1996) demonstrated the transmission of *Phytophthora citricola* by *C. aspersum* (syn. *Helix aspersa*) to *Persea americana* (avocado) plants and *P. indica* stem cuttings.
* Alvarez et al. (2009) demonstrated *C. aspersum* (syn. *H. aspersa*) successfully transmitted *Phytophthora citrophthora* (Phytophthora branch canker of citrus) to three varieties of citrus plants.
* Hyder, Coffey and Stanghellini (2009) demonstrated successful transmission of the plant pathogen *Phytophthora ramorum* by *C. aspersum* (syn. *H. aspersa*) to *Rhododendron* leaves.

The transmission of Phytophthora by snails has not been directly observed in the field, and transmission studies are often conducted under controlled conditions. Phytophthora are spread naturally via soil, rain splash, water, wind and water and by humans moving infected plant material, soil and equipment (DAWR, 2015; Ristaino and Gumpertz, 2000). These modes of transmission are highly effective in spreading the disease to new hosts.

In contrast to microbial disease, snails are well documented vectoring and transmitting a range of nematodes, and similar organisms, associated with animal and human diseases – similarly, plant related nematodes have also been found inhabiting *C. aspersum*.

* Sanchez (2010) recovered an extensive range of nematodes and bacteria from *C. aspersum* (as *H. aspersa*) collected in California, USA. Nematodes were recovered from 92% of snails, including plant pathogenic species *Aphelenchoides fragariea*, *Heterodera schachtii* and *Xiphinema index*. Thirty-one bacterial species were isolated from snail organ tissues and mucus slime; of these, only one was phytopathogenic – *Xanthomonas campestris* pv. *campestris*. Many of the bacterial species recovered are frequently found in or on plants, soil and water. Some bacteria isolated during the study have been reported to be entomopathogenic such as *Pseudomonas entomophila*, *Serratia marcescens* and *S. proteamaculans*, and several are associated with infections in humans.
* Sanchez (2015) further investigated the role of *C. aspersum* (as *H. aspersa*) in transmitting plant pathogens. Additional plant pathogenic nematodes *Ditylencus dipsaci*, *Mesocriconema discus* and *Pratylenchus vulnus*, as well as fungi *Fusarium solani*, *F. oxysporum* f. sp. *callistephi,* *F. oxysporum* f. sp. *chrysanthemi*, *Mucor hiemalis* and *Rhizoctonia solani*, were recovered from snails collected in California, USA.
* In controlled studies, Sanchez (2015) demonstrated that plant parasitic nematodes survived digestion by *C. aspersum*,and were passed in decreasing numbers in snail faeces for up to eight days (minimal nematodes were recovered after 9 days). Likewise, viable fungi were also recovered from snail faecal pellets.
* Sanchez (2015) went on to demonstrate that root knot nematodes consumed by *C. aspersa*, successfully infected tomato plants, when collected faecal pellets were placed on the base of seedlings and incubated for two months.
* Similarly, Michaud (2019) also demonstrated the consumption, and passing of viable plant nematodes *Meloidogyne javanica* and *Heterodera schachtii* by *C. aspersum*. Egg masses and cysts consumed by the snails, survived digestion and were viable after excretion. The hatched nematodes were infective, causing galls on tomato plants.

Each of these authors point to the potential role that *C. aspersum* may play in distributing plant pathogens within crops/hosts. Thus far, all studies have been conducted under controlled conditions. There are no publications that identify plant pathogenic nematodes being transmitted by snails in either agricultural or environmental settings, and no known plant pathogenic nematodes that rely on snails as the sole method of dispersal, or in contrast to animal pathogenic nematodes, rely on snails as an obligate intermediate host.

#### Risk assessment and management for vectored plant pathogens

There is substantial uncertainty in respect of both: (a) the likelihood that imported *C. aspersum* will vector an individual pathogen; and (b) the likely consequences of introduction within Australia in respect of the impact of each of these pests and diseases may have on plant life and health. In view of this uncertainty, and the potential for significant impact, the biosecurity risk attached to each vectored pest or disease was conservatively estimated as low and, thus requiring risk management if the importation of *C. aspersum* is to achieve Australia’s ALOP.

**Importation of adult snails**: the management measure that has been specified for *Angiostrongylus* spp. and for *Crenosoma vulpis* (Sections 5.1 and 5.3, respectively) is to require that adult snails are imported into an Approved Arrangement site, where they can be reared in isolation from other snails and any other host species. This measure can be extended to manage vectored diseases and pests. Isolation would break the life cycle of plant pathogens. The next generation of *C. aspersum* would then be considered free of any key vectored diseases and pests and will be eligible for release from the Approved Arrangement site.

**Importation of hatchling snails**: these are hatchlings less than 10 days old that have not been released from the hatching environment and have not been exposed to plant substrate. As these hatchling snails have not been exposed to plant substrate, the likelihood that they will vector plant pathogens will be sufficiently low as to achieve Australia’s ALOP.

## Species verification and biosecurity

In this biosecurity import risk review, risk management describes the process of implementing measures to address the underlying requirement for species verification, as well as the biosecurity risks identified in the review as exceeding Australia’s ALOP. In establishing these measures, the negative effects on trade have been minimised.

Before issuing an import permit for this commodity, the department will consider (*inter alia*) certain criteria, including:

* The animal health status of the country
* The effectiveness of the country’s animal and plant health authorities
* Legislative controls over animal and plant health, including biosecurity policies and practices
* The standard of reporting to WOAH and IPPC, of significant disease and pest outbreaks
* The effectiveness of official laboratory services, including compliance with relevant international standards
* The effectiveness of systems for the certification and documentation of products intended for export to Australia.

### Measures for species verification and biosecurity

Biosecurity measures will be required for the importation of live *C. aspersum* for heliciculture within Australia, noting that the scope of this review was extended from *adult C. aspersum* for heliciculture to include the option for the importation of hatchling snails.

**Exporting country**: *C. aspersum* breed-stock must only be sourced from approved countries. Approval will depend on the department undertaking a Competent Authority assessment for this commodity.

**Snail farming establishments**: snail farms seeking to export *C. aspersum* to Australia must be commercial operations and government certified for basic quality standards for food safety, and general hygienic operations. Government certification for relevant biosecurity controls will also be required at the time of export. The current Quality Standard applicable to the commercial operation which is exporting *C. aspersum* to Australia must be linked to a specific import application and will be reviewed as part of the application assessment process. Some modification or amendment of this Quality Standard may be required.

Species verification and biosecurity measures for adult snails

* Adult snails exported to Australia must be certified by the exporter and exporting country as *C. aspersum* only. Certification of species may be based on morphological identification or on an approved molecular method. Approval will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity.
* Adult snails will be directed to an Approved Arrangement on arrival at the Australian border. This will be a biosecure facility and may be a government-supervised quarantine facility or may be a snail farming facility that is managed privately. The privately managed Approved Arrangement and site will include (*inter alia*) site design and security, as well as biosecurity protocols for site maintenance, people and the disposal of waste. The privately managed Approved Arrangement and site will also be subject to the department’s requirements for audit and compliance.
* A subset of the snails within each imported consignment will be subject to species verification by morphological or molecular identification. Morphological Identification involves sacrifice of the selected snails as dissection is required. Dissection and identification must be undertaken by a suitably qualified inspector who has been approved by the department to undertake this task. Identification must be based on a diagnostic key that has been approved by the department for use in this context. Molecular identification will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity. The identification procedure will be described in the Approved Arrangement and the protocol for its application will be based on the anticipated size of consignment.
* Imported adult snails will remain under the Approved Arrangement and within the biosecure facility where they will be bred to the next generation. Throughout this period, the imported snails will be isolated from Australian snails and from the definitive host of both *Angiostrongylus spp*. and *Crenosoma vulpis*. This will ensure that the following generation of *C. aspersum* will be free of *Angiostrongylus spp*. and *Crenosoma vulpis*. Isolation within the biosecure facility will also break the life cycle of vectored plant pathogens. Successive generation(s) of snails will then be eligible for release from the Approved Arrangement and site.
* The imported adult snails will not be eligible for release from the Approved Arrangement site. They will either remain within the site for their natural life or will be euthanised. The remains of imported adult snails will be managed as biosecurity waste.

Species verification and biosecurity measures for hatchling snails

* Hatchling snails are hatchlings less than 10 days old that have not been released from the hatching environment and have not been exposed to plant substrate.
* Hatchling snails must be sourced from establishments that employ either an internal or mixed (internal and external) farming system and must only have received commercially processed livestock feed.
* Hatchling snails must be certified by the exporter and exporting country as *C. aspersum* only. Certification of species may be based on morphological identification of breed stock or on an approved molecular method. Approval will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity.
* Hatchling snails will be directed to an Approved Arrangement on arrival at the Australian border. This will be a government-supervised biosecure facility where the snails can be held while species is verified.
* A subset of the hatchling snails within each imported consignment will be subject to species verification by molecular identification. This will entail the development, validation and standardisation of a molecular procedure that meets the department’s requirements for sensitivity and specificity. The protocol for the application of this procedure will be described in the Approved Arrangement and will be based on the anticipated size of a consignment.
* Having verified the species of hatchling snails within a consignment (as above), the consignment may be released from biosecurity control. It will be a condition of the import permit that imported hatchling snails are for breeding purposes only and must not be used for human consumption.

### Certification for species verification and biosecurity

A veterinary health certification will be required for live garden snails (*C. aspersum*) exported to Australia. This certificate must be signed by an Official Veterinarian. The department may review these conditions if there is reason to believe that the animal health or phytosanitary status of an exporting country has changed.

The certificate must provide details of:

* The packaging of the live garden snails for export to Australia, including details of the labelling
* The addresses and, where applicable, the identification numbers of establishments at which the live garden snails for export to Australia were raised
* The names and addresses of the exporter and the consignee
* Species of the live garden snails for export to Australia within the consignment.

The Official Veterinarian of the source country must certify in English and also in a language understood by the Official Veterinarian of the approved country, if required, that:

* The live garden snails for export to Australia are *C. aspersum*
* The live garden snails for export to Australia have been continuously resident in the exporting country since hatching
* The live garden snails for export to Australia have been sourced from an approved snail farming establishment. To be approved, this must be a commercial establishment that is not incorporating field (wild) snails into their stock. These establishments must be government certified for basic quality standards for food safety and general hygienic operation.

The current Quality Standard document(s) applicable to the commercial operation must be attached to the import application.

* The live garden snails for export to Australia must not be field collected
* The live garden snails for export to Australia were prepared and/or stored at the following establishments <insert list>
* The live garden snails for export to Australia have been subject to effective separation controls to ensure the prevention of inadvertent or deliberate substitution, and the prevention of comingling with, or contamination by, animal material not eligible for export to Australia
* The live garden snails for export to Australia have been prepared for export and packed on <insert date> and the bags, wrappers or packing containers were clean and new
* The live garden snails for export to Australia were not exposed to contamination prior to export
* The live garden snails for export to Australia will be transported in a clean packing container sealed with a seal bearing the number or mark <insert mark>. The container contains only garden snails eligible for entry into Australia.

## Appendices

### Appendix A: hazard identification

Under the World Organisation of Animal Health (WOAH) Code, hazard identification is a categorisation step, identifying biological agents dichotomously as hazards or not. The risk assessment may be concluded if hazard identification fails to identify hazards associated with the importation.

A set of questions distils the key considerations for hazard identification for this biosecurity import risk review, including:

* Are there any pathogens that can infect *C. aspersum* that could be imported along with the snail?
* Are these pathogens present in Australia?
* Can the introduction, establishment and spread of pathogens introduced in imported infected *C. aspersum* which have escaped/been released into the environment, cause adverse outcomes in Australia if a significant outbreak eventuated?

Hazard identification identified 14 pathogens (to genus group) which could infect *C. aspersum*, were exotic to Australia and which could have significant adverse consequences in Australia. Of these, eight pathogens (to genus group) were then retained for further review.

The results of the hazard refinement process, including the reason for removal or retention of each identified hazard are summarised in Table 6. The diseases retained for review include:

* *Angiostrongylus*spp., including:
  + *Angiostrongylus (Aelurostrongylus) abstrusus*
  + *Angiostrongylus cantonensis*
  + *Angiostrongylus chabaudi*
  + *Angiostrongylus vasorum*
* *Brachylaima*spp., including:
  + *Brachylaima aspersae*
  + *Brachylaima cribbi*
  + *Brachylaima Ilobregatensis*
  + *Brachylaima mascomai*
* *Crenosoma vulpis*
* *Dicrocoelium dendriticum*
* *Phasmarhabditis*spp.
* *Riccardoella limacum*
* *Tetrahumena* spp., including:
  + *Tetrahymena rostrata*
  + *Tetrahymena limacis*
* *Troglostrongylus brevior*

Table 6 Hazard identification

| Agent | Agent species | Known susceptible species | Adverse consequences in Australia | Exotic to Australia | Retained for further review |
| --- | --- | --- | --- | --- | --- |
| *Alloionema appendiculatum*  Nematode | n/a | Snails in the families Helicidae (including *C. aspersum*), Agriolimacidae, Hygromiidae, Succineidae  Slugs in the family Arionidae | Could potentially infect native snails and slugs. | Unknown | No. Refer to Department of Climate Change, Energy, the Environment and Water |
| *Angiostoma aspersae*  Nematode | n/a | *C. aspersum*  Salamanders:  *Lyciasalamandra antalyana*  *Lyciasalamandra luschani* | Little information on this species. Could potentially infect native snails and slugs. | Yes | No  Sparse information available.  Hosts are salamanders.  Very restricted distribution. |
| *Angiostrongylus*spp.  Nematode | *Angiostrongylus (Aelurostrongylus) abstrusus* | Felines  *Felis catus*  *Acinonyx jubatus*  *Panthera onca*  *Felis concolor*  *Panthera leo*  *Panthera tigris altaica* | Yes  Domestic cats are affected by this species. The introduction of new strains/genotypes could have adverse consequences to domestic cats. | No | No |
| *Angiostrongylus cantonensis* | Rodents24 rat species with Rattus norvegicus and *R. rattus* identified as responsible for greatest distribution  Accidental hosts:  Humans, Australian native fauna, domesticated and wild animals | Yes | No | No. However, some details included in chapter on *Angiostrongylus* spp. |
| *Angiostrongylus chabaudi* | Felines | Yes  Domestic cats are affected by this species. | Yes | Yes |
| *Angiostrongylus mackerrasae* | Rodents  *Rattus fuscipes*  *Melomys cervinipes*  *Rattus leucopus* | No | No | No |
| *Angiostrongylus vasorum* | Canines  *Canis lupus familiaris*  *Cerdocyon thous*  *Ducicyon azarae*  *Ducicyon. vetulus*  *Fennecus zerda*  *Vulpes vulpes*  Paratenic hosts:  lizards, mice, rats | Yes  This species infects canids. Domestic dogs, dingoes and foxes would be at risk of infection. | Yes | Yes |
| *Brachylaima*spp.  Trematode | *Brachylaima aspersae* | Rodents  Known definite host:  mouse (*Mus musculus*) | Yes  Little information available on this species. Only one known host. It could potentially infect native rodents and other animals. | Yes | Yes |
| *Brachylaima cribbi* | Birds, mammals, reptiles  Known definitive hosts:  Bird:  emu (*Dromaius novaehollandiae*)  chicken (*Gallus gallus*)  pigeon (*Columba livia*)  little raven (*Corvus mellori*)  black bird (*Turdus merula*)  starling (*Sturnus vulgaris*)  Mammal:  mouse (*Mus domesticus*)  sheep (*Ovis* spp.)  cat (*Felis catus*)  Reptile:  Shingleback lizard (*Tiliqua rugosa*) | Yes, but already present.  The origin of this species has not been established. It is therefore possible that new strains / genotypes are introduced. | No | Yes |
| *Brachylaima Ilobregatensis* | Rodents  Known definite hosts:  *Crocidura russula*  *Mus spretus*  *Mus musculus* | Yes  Little information available on this species. Known hosts are all rodents. It could potentially infect native rodents and other animals. | Yes | Yes |
| *Brachylaima mascomai* | Rodents  Known definitive host:  *Rattus norvegicus*  *Rattus rattus*  *Mus musculus*  *Crocidura russula*  *Meriones unguiculatus*  *Apodemus sylvaticus* | Yes  Little information available on this species. Known hosts are all rodents. It could potentially infect native rodents and other animals. | Yes | Yes |
| *Crenosoma vulpis*  Nematode | n/a | Fox (*Vulpes vulpes*)  Coyote (*Canis latrans*),  Dogs (*Canis lupus familiaris*)  Badgers (*Meles meles*) | Yes  Domestic dogs, dingoes and foxes would be at risk of infection. | Yes | Yes |
| *Cryptobia helicogenae*  Protozoan | n/a | Snails  *C. aspersum*  *Helix pomatia* | Yes  Could potentially infect native snails and slugs. | Yes | No  Sparse information available. |
| *Dicrocoelium dendriticum*  Trematode | n/a | Livestock including sheep, cattle, llama  Rabbits, pigs, dogs  Humans | Yes  The role of *C. aspersum* as an intermediate host is unclear. This parasite has a broad host range, can impact livestock and can also infect humans. It could potentially infect native animals. | Yes | Yes |
| *Phasmarhabditis*spp.  Nematode | *Phasmarhabditis hermaphrodita* | Snails in the family Helicidae (including *C. aspersum*)  Slugs in the families Limacidae, Agriolimacidae, Arionidae, Milacidae, Vaginulidae | Yes  Could potentially infect native snails and slugs. | Yes | Yes |
| *Phasmarhabditis californica* | *C. aspersum* | Yes  Little information on this species. Could potentially infect native snails and slugs. | Yes | Yes |
| *Rhabditis maupasi*  Nematode  10 | n/a | *C. aspersum* | No, but could potentially infect native snails and slugs. | Unknown | No  Sparse information, no evidence of animal biosecurity impact, referred to Department of Health and Aged Care as *C. aspersum* for human consumption can carry parasite. |
| *Riccardoella limacum*  Mite | n/a | *C. aspersum*  *Arianta arbustorum* | Yes  Could potentially infect native snails and slugs | Yes | Yes |
| *Tetrahymena* spp.  Protozoan | *Tetrahymena limacis* | Slugs in the Arionidae, Milacidae and Limacidae families  Snails in the Bradybaenidae, Daudebardiidae, Helicidae, Hygromiidae, Succineidae, Vitrinidae and Zonitidae families. | Yes  Could potentially infect native snails and slugs. | Unknown | Yes. |
| *Tetrahymena rostrata* | Snails and slugs  Snails in the Helicidae (including *C. aspersum*) Oleacinidae and Zonitidae  Slugs in the Agriolimacidae and Arionidae families | Yes  Could potentially infect native snails and slugs. | No | As part of *Tetrahymena* spp. chapter. |
| *Tetratrichomonas limacis*  Protozoan | n/a | Snails and slugs  *C. aspersum*  *Limax flavus*  *Deroceras agreste*  *Limax maximus*  *Helix pomatia*  *Otala lacteal* | No information but  could potentially infect native snails and slugs. | Unknown | No |
| *Troglostrongylus brevior*  Nematode | n/a | Felids | Yes  Domestic cats would be at risk of infection. | Yes | Yes |

n/a: not applicable.

### Appendix B: species of garden snail

Table 7 Comparison of snail species in Australia and worldwide

| Snail species | Morphology | Origin | Present in Australia | Is this species farmed? | Is this species field collected? | Comments | References |
| --- | --- | --- | --- | --- | --- | --- | --- |
| *Cornu aspersum*  *= Helix aspersa aspersa*  Common garden snail  Petit gris | Shell size: 25–35 mm diameter x 25–35 mm height  Shell colour: light brown to dark brown with darker coloured bands and yellow fleck  Shell shape: round and globose  Whorls: 4–5  Large and rounded aperture  Closed umbilicus | Mediterranean region  Now also found in:  Africa  Asia  Europe  North America  Oceania  South America | Yes | Yes  Sizes of farms vary from cottage industry or hobbyists to large scale commercial production.  No commercial farms in Australia, only cottage industry. | Yes  Farm raised in countries where it has been intentionally introduced such as eastern Europe, South America and Asia.  Field collected, due to overharvesting and agricultural chemicals has become rare to find in the wild in places like the Republic of Türkiye. | *Helix* snails constitute 70% of world market (*Helix aspersa* 40%).  Most suitable and easily grown species to farm in Italy.  Invasive pest of agricultural crops in the Americas, south-eastern parts of Australia, New Zealand, South Africa and elevated areas (>1,000 m).  High change of wild caught snails being contaminated with heavy metals presenting risk to humans. | (Conte, 2015; Begg, 2003; Blacket *et al*., 2016; Begg, 2006; Nordsieck, 2022a; Murphy, 2001; CABI, 2015) |
| *Helix aspersa maxima*  *= Cornu aspersum maximum*  Gros gris | Shell size: 45 mm diameter  Shell colour: similar to *C. aspersum aspersum*  Weight: 20–30 g  Bigger than *C. aspersum aspersum* | North Africa | No | Yes  Commercial | No | Growing conditions have been adapted to support breeding of these snails in Europe. | (Aude *et al*., 2019; CABI, 2015) |
| *Helix pomatia*  Roman snail, Burgundy snail, escargot de Bourgogne | Shell size: 38–50 mm or 1/3 size of snail  Shell colour: brown with 4–5 light brown bands  Shell shape: round spiral  Whorls: 4–5 | Central and Southeast Europe  Now also found in:  North America,  South America  Asia | No | Yes  160 farms in Sardinia (Italy) for *Helix* genus under the support of the International Institute of Snail Farming (2,000 members).  9,800 kg produced in Italy 200.  Sizes of farms vary from cottage industry or hobbyists to large scale commercial production. | Yes  Field collected, but due to overharvesting and agricultural chemicals has become rare to find in the wild.  Difficult to breed on a large scale due to artificial conditions of a farm.  Commercial.  Exploitation prohibited in Western Europe (still occurring in Eastern Europe). | Most popular in Europe and North America.  Meat is considered most juicy and tasty.  In places like Central Romania, 50–150 families at a time make collection an annual even.  *Helix* snails constitute 70% of world market (*Helix pomatia* 28%). | (Animalia, 2022; Ligaszewski *et al*., 2007; Ligaszewski and Pol, 2021; Ligaszewski *et al*., 2009; Conte, 2015; Thompson and Cheney, 1996) |
| *Helix lucorum*  Turkish snail | Shell size: 35–60 mm width x 25–45 mm height  Shell colour: brown tip with white spherical spiral bands  Shell shape: heavily depressed  Weight: 20–25 g | Eastern Black Sea region through Asia Minor  Southern Romania and Bulgaria  Italy  Introduced in Austria | No | No | Yes  Field collected in the Republic of Türkiye – regional standards suggested such as size, diameter, weight and time of year (after copulation). | Most popular in Europe and North America.  *Helix* snails constitute 70% of world market (*Helix lucorum* 22%). | (Nordsieck, 2022b; Conte, 2015; Neubert, 2011) |
| *Helix cincta* | Shell size: 37 ± 4 mm diameter  Shell colour: light brown with thin bands  Shell shape: compressed last turn with rough surface  Short brown peristome | The Republic of Türkiye  North East Mediterranean from Italy to the Middle East | No | No | Yes | Not consumed in the Republic of Türkiye but a main export item. | (Yildirim *et al*., 2004; Giusti *et al*., 2015) |
| *Cantareus apertus*  *= Helix adanensis*  *= Cornu apertus*  Green snail | Shell size: 15–30 mm diameter  Shell colour: olive green (juveniles), light brown shell (adults), no banding  Whorls: 4–5  Aperture extremely large relative to body (larger aperture than *C. aspersum*) | Mediterranean region (Europe and North Africa)  Introduced to Australia (WA) and North America | Yes  Declared pest | No | Yes | Not consumed in the Republic of Türkiye but a main export item. | (Blacket *et al*., 2016; Yildirim *et al*., 2004) |
| *Cepaea nemoralis*  English garden snail | Shell size: 20–26mm diameter  Shell colour: yellow, pink or brown with 1–5 bands  Distinguished from *C. hortensis* by its brown apertural lip (opening of the shell) | Central and Western Europe  Introduced to North America | No | No | Yes | Ease of culture, near worldwide availability and hardiness has led to them being cultured for human consumption. | (Dees, 1970; Pearce *et al*., 2010; Rosin *et al*., 2013; Ozgo and Schilthuizen, 2012; Whitson, 2005) |
| *Cepaea hortensis*  Smaller banded snail | Shell size: 20 mm diameter  Shell colour: yellow to brown with 1–5 thin chestnut brown stripes  Distinguished from *C. nemoralis* by its white apertural lip (opening of the shell) | Central and Northern Europe, extending more north than *C. nemoralis*  North America | No | No | Yes | Least popular of the large European snails. | (Cowie and Jones, 1987; Pearce *et al*., 2010; Dees, 1970; Thompson and Cheney, 1996) |
| *Eobania vermiculata =Massylaea vermiculata*  Chocolate-banded snail | Shell size: 22–35mm diameter  Shell colour: white, green or yellow with darker chocolate colour bands  Whorls: 5–6 | Mediterranean region  The Republic of Türkiye | Unsure  Introduced to South-eastern Australia; appears locally extinct.  On National Priority Pests list 2019. | Yes  4,420 kg produced in Italy 2002.  Sizes of farms vary from cottage industry or hobbyists to large scale commercial production. | Yes  collected | Quite difficult to raise in captivity.  Important export item for the Republic of Türkiye.  8.5% world market. | (Conte, 2015; Yildirim *et al*., 2004) |
| *Otala lactea*  Milk snail | Shell size: 27.5–36 mm diameter x 16–25 mm high  Shell colour: light brown (almost white) with speckled dark brown lines along the shell whorls  Shell shape: non globular, slightly depressed | Northern Africa  Spain  South America | No |  | Yes | None | (White-McLean, 2022b; Dees, 1970) |
| *Otala punctata* | Shell size: 33–39 mm wide x 20–24 mm high  Shell colour: dark brown with lots of white flecks along whorls | Mediterranean  South Africa  North America | No |  | Yes | None | (White-McLean, 2022b; Barbara and Schembri, 2008) |
| *Theba pisana*  White garden snail | Shell size: 12–25 mm width x 9–20 mm height  Shell colour: creamy white shell with diverse range of pale-dark brown markings. Uninterrupted spiral bands, dotted lines or small radial smudges | Mediterranean  Atlantic coast of Europe  Southwest England and Wales  Introduced to Australia and USA | Yes |  | Yes | Serious agricultural pest worldwide, and in Australia, particularly in South Australia.  Mainly consumed in the Iberian Peninsula (Spain and Portugal).  4,000 tonnes per year consumed in Portugal.  Growing and large economic market.  Important export item for the Republic of Türkiye. | (Cowie *et al*., 2009; Dees, 1970; Leonard, 2003; Caetano *et al*., 2021; CABI, 2020; ALA, 2022) |
| *Achatina achatina* Common African snail | Shell size: 180 mm length x 90 mm diameter  Body size: 300 mm length and 250 mm height (largest snail species worldwide)  Shell colour: brown with striped pattern ‘tiger snail’  Shell shape: conical | West Africa | No | Yes  Commercial  Cottage industry | Yes  18.2% of farmers in Nigeria obtain snails from wild.  Very few farmers (<4%) start with eggs or hatchlings.  Market makes up 30% of snails that are farmed. | Snails purchased from the market are hand-picked from their natural habitat.  Preferred source of snails for consumers as it is widely believed they taste better.  Snails are taken from wild before reaching sexual maturity – threatens survival.  High change of wild caught snails being contaminated presenting risk to humans.  Hardest species to farm due to stable conditions it is accustomed to in the wild but by far most popular in West Africa (2nd and 3rd most popular *Archachatina marginata* and *Achatina fulica* respectively*).* | (Chah and Inegbedion, 2013; Nyoagbe *et al*., 2016; Ngenwi *et al*., 2010; Cobbinah, 1993) |
| *Achatina (Lissachatina) fulica*  Giant African land snail | Shell size: 200 mm length x 120 mm diameter  Shell colour: brown with darker bands across spiral  Weight: 250 g | East Africa | No  An outbreak occurred in QLD in 1977 but was quickly eradicated.  A single animal was also found in QLD in 2004.  On National Priority Pests list 2019. | Yes  Commercial  Cottage industry | Yes  18.2% of farmers in Nigeria obtain snails from wild.  Very few farmers (<4%) start with eggs or hatchlings.  Market makes up 30% of snails that are farmed. | Snails purchased from the market are hand-picked from their natural habitat.  Preferred source of snails for consumers as it is widely believed they taste better.  Snails are taken from wild before reaching sexual maturity – threatens survival.  High change of wild caught snails being contaminated presenting risk to humans.  Causes considerable economic damage to crops. | (Chah and Inegbedion, 2013; Nyoagbe *et al*., 2016; Government, 2019; Cobbinah, 1993) |
| *Archachatina marginata* | Shell size: up to 210 mm length x 130 mm in diameter  Shell colour: brown to yellow, striated ‘woven’ look  Shell shape: more rounded than other African species  Weight: 500 g | African rainforest belt / West Africa | No | Yes  Commercial  Cottage industry | Yes  18.2% of farmers in Nigeria obtain snails from wild.  Very few farmers (<4%) start with eggs or hatchlings.  Market makes up 30% of snails that are farmed.  Largely rudimentary and its demand is met by hunting from their habitat in the wild. | Snails purchased from the market are hand-picked from their natural habitat.  Preferred source of snails for consumers as it is widely believed they taste better.  Snails are taken from wild before reaching sexual maturity – threatens survival.  High change of wild caught snails being contaminated presenting risk to humans.  Highest yield out of the three main species farmed in Nigeria. | (Chah and Inegbedion, 2013; Nyoagbe *et al*., 2016; Jimoh and Akinola, 2020; White-McLean, 2022a) |
| *Elona quimperiana*  Quimper snail | Shell size: 30 mm diameter  Shell colour: translucent  Shell shape: flat and smooth  Slightly reflected apertural margin | France  Spain | No | No | Yes | None | (Raven, 2022) |
| *Sphincterochila candidissima* | Shell size: 13.3–20.5 mm  Shell colour: pure white | Mediterranean  Spain | No | No | Yes | None | (Moreno-Rueda, 2008; Yanes and Fernandez-Lopez-de-Pablo, 2017) |
| *Iberus alonensis* | Shell size: 30 mm diameter  Shell colour: creamy light brown |  | No | No | Yes | Only in Spain  Threatened  Unlikely important | (Thompson and Cheney, 1996) |

## Glossary

| Term | Definition |
| --- | --- |
| ALOP | Appropriate Level of Protection |
| Approved Arrangement | Approved arrangements, previously Quarantine Approved Premises and Compliance Agreements, are voluntary arrangements entered into with the Department of Agriculture, Fisheries and Forestry.  These arrangements allow operators to manage biosecurity risks and/or perform the documentary assessment of goods in accordance with departmental requirements, using their own sites, facilities, equipment and people, and without constant supervision by the department and with occasional compliance monitoring or auditing.  Further detail at [Approved Arrangements](https://www.agriculture.gov.au/biosecurity-trade/import/arrival/arrangements). |
| Approved Arrangement site | A site where a biosecurity industry participant undertakes biosecurity activities covered by an Approved Arrangement. For biosecurity industry participants that undertake mobile biosecurity activities, the biosecurity industry participants Approved Arrangement site is the location from which the mobile activities are based.  Further detail at [Approved Arrangement site](https://www.agriculture.gov.au/biosecurity-trade/import/arrival/arrangements/glossary). |
| Approved Country | An approved country is one assessed by the Department of Agriculture, Fisheries and Forestry based on relevant factors such as history of trade and knowledge of the competent authority, history of reporting to WOAH for notifiable animal diseases, history of compliant trade to Australia, and verification activities which may have been undertaken historically. |
| BIRA | Biosecurity import risk analysis |
| DAFF | Australian Government Department of Agriculture, Fisheries and Forestry |
| DCCEEW | Australian Government Department of Climate Change, Energy, the Environment and Water |
| EPBC Act | Environment Protection and Biodiversity Conservation Act 1999 |
| Heliciculture | Synonymous with heliculture and means the science of snail rearing (or farming) |
| IPPC | International Plant protection Convention |
| ISPM | International Standards for Phytosanitary Measures |
| ISPM 11 | Pest Risk Analysis for Quarantine Pests |
| The department | Australian Government Department of Agriculture, Fisheries and Forestry |
| WOAH | The World Organisation for Animal Health |

## References

Abo Bakr, Y, Eshra, EH and Hussein, HI (2007). Calotropis procera glycosides are more effective on Eobania vermiculata (Müller) than methomyl and other plant glycosides, Journal of Agricultural Science 32(12): 10519-27

Aghazadeh, M, Reid, SA, Aland, KV, Restrepo, AC, Traub, RJ, McCarthy, JS and Jones, MK (2015a). A survey of Angiostrongylus species in definitive hosts in Queensland, International Journal for Parasitology: Parasites and Wildlife 4(3): 323-8

Aghazadeh, M, Traub, RJ, Mohandas, N, Aland, KV, Reid, SA, McCarthy, JS and Jones, MK (2015b). The mitochondrial genome of Angiostrongylus mackerrasae as a basis for molecular, epidemiological and population genetic studies, Parasites and Vectors 8: 473

AJOT (2017). Philadelphia CBP slows invasive snail mail, American Journal of Transportation, available at http://www.ajot.com/news/philadelphia-cdp-slows-invasive-snail-mail, accessed 13 March 2024.

Akkaya, H, Deniz, A and Sezen, A (2006). Effect of praziquantel on Dicrocoelium dendriticum in naturally infected sheep, Medycyna Weterynaryjna 62(12): 1381-1382

ALA (2022a). Cornu aspersum (Müller, 1774), Atlas of Living Australia, accessed 05-06-2022.

ALA (2022b). Dicrocoelium (Dujardin, 1845), Atlas of Living Australia, accessed 27-05-2022.

ALA (2022c). Theba pisana (Müller, 1774), Atlas of Living Australia, accessed 07-06-2022.

Alicata, JE and Brown, RW (1962). Observations on the method of human infection with Angiostrongylus cantonensis in Tahiti, Canadian Journal of Zoology 40: 755-760

Alicata, JE (1965). Biology and Distribution of the Rat Lungworm, Angiostrongylus cantonensis, and its Relationship to Eosinophilic Meningoencephalitis and other Neurological Disorders of Man and Animals, Advances in Parasitology 3: 223-248

Al-Khayat, JA (2010). First record of five terrestrial snails in the State of Qatar, Turkish Journal of Zoology 34(4): 539-545

Allendorf, FW, Leary, RF, Spruell, P and Wenburg, JK (2001). The problems with hybrids: setting conservation guidelines. Trends in Ecology and Evolution 16: 613-622

Alstedt, U, Voigt, K, Jager, MC, Knubben-Schweizer, G, Zablotski, Y, Strube, C and Wenzel, C (2022). Rumen and Liver Fluke Infections in Sheep and Goats in Northern and Southern Germany, Animals 12(7), available at ARTN 876 10.3390/ani12070876

Alvarez, LA, Gramaje, D, Abad-Campos, P and García-Jiménez, J (2009). Role of the Helix aspersa snail as a vector of Phytophthora citrophthora causing branch cankers on clementine trees in Spain, Plant Pathology 58: 956-963

Amr, ZS, Baker, MA and Katbeh-Bader, A (2019). Massylaea vermiculata (O.F. Müller, 1774): a serious land snail pest introduced to Jordan, Jordan Journal of Natural History 6(6): 60-61

Anderson, RC (2000). The superfamily Megastrongyloidea, in Nematode parasites of vertebrates: their development and transmission, 2nd Ed, CABI Publishing, Wallingford, 129-172

Andrus, P, Ingle, O, Coleman, T and Rae, R (2020). Gastropod parasitic nematodes (Phasmarhabditis sp.) are attracted to hyaluronic acid in snail mucus by cGMP signalling. Journal of Helminthology 94, available at ARTN e9 10.1017/S0022149X18000986

Animalia (2022). Helix pomatia [Online]. Available at https://animalia.bio/helix-pomatia [Accessed 30-05-2022]

Apostolou, K, Staikou, A, Sotiraki, S and Hatziioannou, M (2021). An Assessment of Snail-Farm Systems Based on Land Use and Farm Components. Animals 11, available at: https://www.mdpi.com/2076-2615/11/2/272

Arbabi, M, Dalimi, A, Ghafarifar, F and Moghadam, MF (2011). Prevalence and intensity of Dicrocoelium dendriticum in sheep and goats of Iran, Research Journal of Parasitology 6(5,): 160-167

Arbabi, M, Hadad, A, Hooshyar, H, Akbari, H and Mashkani, SMH (2021). Maintenance of liver fluke, Dicrocoelium dendriticum, outside the body of its native host, International Archives of Health Sciences 8(3): 201-105

Arbabi, M, Nezami, E, Hooshyar, H and Delavari, M (2018). Epidemiology and economic loss of fasciolosis and dicrocoeliosis in Arak, Iran, Veterinary World 11(12): 1648-1655

Arbabi, M, Nezami, E, Hooshyar, H, Delavari, M (2018). Epidemiology and economic loss of fasciolosis and dicrocoeliosis in Arak, Iranian Veterinary World 11(12): 1648-1655

Arnaud, JF, Madec, L, Bellidio, A and Guiller, A (1999). Microspatial genetic structure in the land snail Helix aspersa (Gastropoda : Helicidae), Heredity 83: 110-119

Aubry, S, Labaune, C, Magnin, F, Roche, P and Kiss, L (2006). Active and passive dispersal of an invading land snail in Mediterranean France. Journal of Animal Ecology 75: 802-813

Aude, D, Marion, B, Raphael, C and Colet, JM (2019). Proton Nuclear Magnetic Resonance (H-1 NMR) profiling of isolated organs in the snail Helix aspersa maxima, Ecological Indicators 105: 177-187

Aude, D, Marion, B, Raphael, C and Colet, JM (2019). Proton Nuclear Magnetic Resonance (H-1 NMR) profiling of isolated organs in the snail Helix aspersa maxima, Ecological Indicators 105: 177-187

AUSVEG (2021). Slugs and Snails [Online]. Available: https://ausveg.com.au/biosecurity-agrichemical/crop-protection/overview-pests-diseases-disorders/slugs-and-snails/

Baker, RA (1970). Studies on Life History of Riccardoella-Limacum (Schrank) (Acari-Trombidiformes), Journal of Natural History 4(4): 511

Bank, RA and Neubert, E (2017). Checklist of the land and freshwater Gastropoda of Europe, MolluscaBase, available at https://docplayer.net/131138986-Molluscabase-checklist-of-the-land-and-freshwater-gastropoda-of-europe-ruud-a-bank-eike-neubert-last-update-july-16-th-2017.html

Barbara, N and Schembri, PJ (2008). The status of Otala punctata (Muller, 1774): a recently established terrestrial gastropod in Malta, Bollettino Malacologico 44(5-8): 101-107

Barbara, N and Schembri, PJ (2008). The status of Otala punctata (Muller, 1774): a recently established terrestrial gastropod in Malta. Bollettino Malacologico 44: 101-107

Barcante, TA, Barcante, JM, Dias, SR and Lima Wdos, S (2003). Angiostrongylus vasorum (Baillet, 1866) Kamensky, 1905: emergence of third-stage larvae from infected Biomphalaria glabrata snails, Parasitology Research 91(6): 471-475

Barker, GM and Watts, C (2002). Management of the invasive alien snail Cantareus aspersus on conservation land. Available at: https://citeseerx.ist.psu.edu/document?repid=rep1&type=pdf&doi=d2c718a99a3354ba3bf004d38cb5f6bc404960fa

Barrett, JL, Carlisle, MS and Provic, P (2002). Neuro-angiostrongylosis in wild black and grey-headed flying foxes (Pteropus spp.), Australian Veterinary Journal 80: 554-558

Barrs, VR, Swinney, GR, Martin, P and Nicoll, RG (2008). Concurrent Aelurostrongylus abstrusus infection and salmonellosis in a kitten, Australian Veterinary Journal 77(4): 229-32

Barutzki, D and Schaper, R (2009). Natural Infections of Angiostrongylus vasorum and Crenosoma vulpis in dogs in Germany (2007-2009), Parasitology Research 105: S39-S48

Barutzki, D and Schaper, R (2011). Results of parasitological examinations of faecal samples from cats and dogs in Germany between 2003 and 2010, Parasitology Research 109: S45-S60

Bashê, SK and Al-Qassab, SE (2024). Morphological and molecular identification of some terrestrial snails: first report, Zanco Journal of Pure and Applied Sciences 36(1): 94-104

Baur, A and Baur, B (2005). Interpopulation variation in the prevalence and intensity of parasitic mite infection in the land snail Arianta arbustorum, Invertebrate Biology 124(3): 194-201

Baydar, E and Kaya, F (2021). Case report: Aelurostrongylus abstrusus infection and radiographic findings in a kitten, Kocatepe Veterinary Journal, available at: https://dergipark.org.tr/tr/download/article-file/1726058

Begg, S (2006). Free-range Snail Farming in Australia, Australian Government Rural Industries Research and Development Corporation publication 06/104, RIRDC

Begg, S (2003). Farming Edible Snails – Lessons from Italy, Australian Government Rural Industries Research and Development Corporation publication 03137, RIRDC

Benbellil-Tafoughalt, S, Sahnoune, M, De Vaufleury, A and Moali, A (2009). Effects of Temperature and Photoperiod on Growth and Reproduction of the Land Snail Helix Aperta Born (Gastropoda, Pulmonata). Revue D Ecologie-La Terre Et La Vie, 64, 207-219.

Benfradj, N, Vettraino, AM, Tomassini, A, Bruni, N, Vannini, A and Boughalleb-M'Hamdi, N (2018). Citrus gummosis incidence and role of ants (Lasius grandis) and snails (Helix aspersa) as vectors of the disease in Tunisia, Forest Pathology, 48(3): e12423

Bessa, EC, Lima, WDS, Daemon, E, Cury, MC and Araujo, JLDB (2000). Desenvolvimento biológico de Angiostrongylus vasorum (Baillet) Kamensnky (Nematoda, Angiostrongylidae) em Subulina octona Bruguiere (Molusca, Subulinidae) em condições de laboratório, Revista Brasileira de Zoologia 17(1): 29-41

Bezemer TM and Knight, KJ (2001). Unpredictable responses of garden snail (Helix aspersa) populations to climate change. Acta Oecologica-International Journal of Ecology 22: 201-208

Bhaibulaya, M (1968). A new species of Angiostrongylus in an Australian rat, Rattus fuscipes, Parasitology 58(4): 789-799

Bhaibulaya, M (1991). Snail borne parasitic zoonoses: Angiostrongyliasis, Southeast Asian Journal of Tropical Medicine and Public Health 22: 189-93

Bihr, T and Conboy, GA (1999). Lungworm (Crenosoma vulpis) infection in dogs on Prince Edward Island, Canadian Veterinary Journal-Revue Veterinaire Canadienne 40(8): 555-559

Biocca, E (1957). Angiostrongylus chabaudi n. sp. parassita del cuore e dei vasi polmonari del gatto selvatico (Felis silvestris), Reale Accademia dei Lincei 22: 526-532

Bjork, KE, Averbeck, GA and Stromberg, BE (2000). Parasites and parasite stages of free-ranging wild lions (Panthera leo) of northern Tanzania, Journal of Zoo and Wildlife Medicine 31(1): 56-61

Blacket, MJ, Shea, M, Semararo, L and Malipatil,MB (2016). Introduced Helicidae Garden Snails in Australia: Morphological and Molecular Diagnostics, Species Distributions and Systematics. Records of the Australian Museum 68: 99-116

Błoszyk, J, Kacprowicz, J and Książkiewicz-Parulska, Z (2016). Notes on the activity of the Roman snail (Helix pomatia L.), Folia Malacologica 24(2): 103-106

Bolt, G, Monrad, J, Frandsen, F, Henriksen, P and Dietz, HH (1993). The common frog (Rana temporaria) as a potential paratenic and intermediate host for Angiostrongylus vasorum, Parasitology Research 79: 428-30

Bosco, A, Rinaldi, L, Salamina, V, Santaniello, M, Morgoglione, ME, Guariglia, I, Cappelli, G, Scala, A and Cringoli, G (2015). Field trial on the efficacy of albendazole micronised (single and double treatment) against Dicrocoelium dendriticum in naturally infected sheep: A new strategy for the control of dicrocoeliosis, Small Ruminant Research 126: 2-5

Bößneck, U (2011). New records of freshwater and land molluscs from Lebanon: (Mollusca: Gastropoda and Bivalvia), Zoology in the Middle East 54(1): 35-52

Bowman, DD, Hendrix, CM, Lindsay, DS and Barr, SC (2002). ‘Metastrongyloidea’, Feline Clinical Parasitology 1: 272-173

Brianti, E, Gaglio, G, Napoli, E, Falsone, L, Giannetto, S, Latrofa, MS, Giannelli, A, Dantas-Torres, F and Otranto, D (2013). Evidence for direct transmission of the cat lungworm Troglostrongylus brevior (Strongylida: Crenosomatidae), Parasitology 140(7): 821-824

Brianti, E, Varcasia, A and Otranto, D (2021). Troglostrongylus brevior, Trends in Parasitology 37(6): 569-570

Butcher, AR and Grove, DI (2001). Description of the life-cycle stages of Brachylaima cribbi n. sp (Digenea : Brachylaimidae) derived from eggs recovered from human faeces in Australia, Systematic Parasitology 49(3): 211-221

Butcher, AR and Grove, DI (2003). Field prevalence and laboratory susceptibility of southern Australian land snails to Brachylaima cribbi sporocyst infection, Parasite 10(2): 119-125

Butcher, AR and Grove, DI (2005a). Seasonal variation in rates of sporocyst and metacercarial infection by Brachylaima cribbi in helicid and hygromiid land snails on the Yorke Peninsula, South Australia, Australian Journal of Zoology 53(6): 375-382

Butcher, AR and Grove, DI (2005b). Second intermediate host land snails and definitive host animals of Brachylaima cribbi in southern Australia, Parasite 12(1): 31-37

Butcher, AR (2016). Children, snails and worms: the Brachylaima cribbi story, Microbiology Australia 37(1): 30-33

Butcher, AR, Parasuramar, P, Thompson, CS and Grove, DI (1998). First report of the isolation of an adult worm of the genus Brachylaima (Digenea : Brachylaimidae), from the gastrointestinal tract of a human, International Journal for Parasitology 28(4): 607-610

Butcher, AR, Talbot, GA, Norton, RE, Kirk, MD, Cribb, TH, Forsyth, JRL, Knight, B and Cameron, AS (1996). Locally acquired Brachylaima sp (Digenea: Brachylaimidae) intestinal fluke infection in two South Australian infants, Medical Journal of Australia 164(8): 475-478

Cabaret, J, Morand, S, Aubert, C and Yvore, P (1988). Snail Farming - a Survey of Breeding Management, Hygiene and Parasitism of the Garden Snail, Helix-Aspersa Muller, Journal of Molluscan Studies 54: 209-214

CABI. 2015. Cornu aspersum (common garden snail) [Online]. Available: https://www.cabi.org/isc/datasheet/26821 [Accessed 30-05-2022 2022].

CABI. 2015. Cornu aspersum (common garden snail) [Online]. Available: https://www.cabi.org/isc/datasheet/26821 [Accessed 30-05-2022 2022].

CABI. 2020. Theba pisana (white garden snail) [Online]. Available: https://www.cabi.org/isc/datasheet/62094 [Accessed].

Caetano, D, Miranda, A, Lopes, S, Paiva, J, Rodrigues, A, Videira, A and Almeida, CMM (2021). Nutritional and toxicity profiles of two species of land snail, Theba pisana and Otala lactea, from Morocco, Journal of Food Composition and Analysis, doi: 10.1016/j.jfca.2021.103893

Călin, M, Oana Cristea, T, Ambarus, S, Brezeanu, C, Brezeanu, PM, Prisecaru, M and Şova, GF (2014). The behavior of some plant species to snails attack, Studii şi Cercetări: Biologie 23(2): 31-33

Cardillo, NM, Ercole, M, Farina, F, Pasqualetti, M, Loiza, Y, Perez, M, Bonboni, A and Ribicich, M (2018). Larval development of Aelurostrongylus abstrusus in experimentally infected Rumina decollata snails, Veterinary Parasitology 251: 50-55

Catalogue of Life (2020). Pellioditis hermaphrodita (A. Schneider, 1859)

Catalogue of Life (2022a). Cornu aspersum (O. F. Müller, 1774), Catalogue of Life, accessed 04-06-2022

Catalogue of Life (2022b) Dicrocoelium Dujardin, 1845, accessed 27-05-2022

Catalogue of Life (2022c) Tetrahymena, accessed 25-05-2022

Çelik, MY, Duman, MB, Sariipek, M, Goran, GU and Karayucel, S (2018). Growth and mortality rates of Cornu aspersum: organic snail culture system, Black Sea region, Journal of Agricultural Sciences 25: 189-196

Çelik, MY, Duman, MB, Sariipek, M, Goran, GU and Karayucel, S (2020). Comparison of Proximate and Amino Acid Composition between Farmed and Wild Land Snails (Cornu aspersum Muller, 1774), Journal of Aquatic Food Product Technology 29: 383-390

Çelık, MY, Duman, MB, Sariıpek, M, Uzun Gören, G, Kaya Öztürk, D and Karayücel, S (2018). Effect of shell height on the reproductive success and survival of Cornu aspersum (O.F. Müller, 1774), AACL Bioflux 11(2): 525-532

Cengiz, ZT, Yilmaz, H, Dulger, AC and Cicek, M (2010). Human infection with Dicrocoelium dendriticum in Turkey, Annals of Saudi Medicine 30(2): 159-161

Chan, JM and Inegbedion, G (2013). Characteristics of snail farming in Edo South Agricultural Zone of Edo State, Nigeria, Tropical Animal Health and Production 45(2): 625-631

Chan, D, Barratt, J, Roberts, T, Lee, R, Shea, M, Marriott, D, Harkness, J, Malik, R, Jones, M, Aghazadeh, M, Ellis, J and Stark, D (2015). The Prevalence of Angiostrongylus cantonensis/mackerrasae Complex in Molluscs from the Sydney Region, PLoSOne 10(5), available at ARTN e0128128 10.1371/journal.pone.0128128

Charwat, SM and Davies, KA (2001). Nematodes: Biocontrol Agents of Helicid Snails, Rural Industries Research and Development Corporation publication 01/03, RIRDC

Chevallier, H (1980). Cornu aspersum (common garden snail), CABI Digital Library, doi: 10.1079/cabicompendium.26821

Chevallier, H (1977). La variabilité de l'Escargot Petit-gris Helix aspersa Müller, (Variability of the Petit-gris snail Helix aspersa Müller), Bulletin Du Muséum National D'histoire Naturelle 3(448): 425-442

Civeyrel, L and Simberloff, D (1996). A tale of two snails: Is the cure worse than the disease? Biodiversity and Conservation 5: 1231-1252

Clark, SA (2004). Native snails in an urban environment - conservation from the ground up. Urban Wildlife: More Than Meets the Eye: 78-81

Cobbinah, JR (1993). Snail farming in West Africa, Wageningen, The Netherlands, available at: https://hdl.handle.net/10568/63667

COL (2022). Cornu aspersum (O. F. Müller, 1774). Catalogue of Life

Colella, V, Cavalera, MA, Deak, G, Tarallo, VD, Gherman, CM, Mihalca, AD and Otranto, D (2017). Larval development of Angiostrongylus chabaudi, the causative agent of feline angiostrongylosis, in the snail Cornu aspersum, Parasitology 144 (14): 1922-30

Colella, V, Giannelli, A, Brianti, E, Ramos, RAN, Cantacessi, C, Dantas-Torres, F and Otranto, D (2015). Feline lungworms unlock a novel mode of parasite transmission, Scientific Reports 5, available at: https://www.nature.com/articles/srep13105

Colella, V, Lia, RP, Premont, J, Gilmore, P, Cervone, M, Latrofa, MS, D'Anna, N, Williams, D and Otranto, D (2016a) Angiostrongylus vasorum in the eye: new case reports and a review of the literature, Parasites and Vectors 9: 161

Colella, V, Mutafchiev, Y, Cavalera, MA, Giannelli, A, Lia, RP, Dantas-Torres, F and Otranto, D (2016b). Development of Crenosoma vulpis in the common garden snail Cornu aspersum: implications for epidemiological studies, Parasites and Vectors 9, available at: https://parasitesandvectors.biomedcentral.com/articles/10.1186/s13071-016-1483-8

Colomba, MS, Gregorini, A, Liberto, F, Reitano, A, Giglio, S and Sparacio, I (2015). The genus Erctella Monterosato, 1894: new molecular evidence (Pulmonata Stylommatophora Helicidae), Biodiversity Journal 6(1): 401-411

Conboy, G (2004). Natural infections of Crenosoma vulpis and Angiostrongylus vasorum in dogs in Atlantic Canada and their treatment with milbemycin oxime. Veterinary Record 155(1): 16-18

Conboy, G, Guselle, N and Schaper, R (2017). Spontaneous Shedding of Metastrongyloid Third-Stage Larvae by Experimentally Infected Limax maximus. Parasitology Research 116: S41-S54

Conboy, G, Hare, J, Charles, S, Settje, T and Heine, J (2009). Efficacy of a Single Topical Application of Advantage Multi (R) (=Advocate (R)) Topical Solution (10% Imidocloprid+2.5% Moxidectin) in the Treatment of Dogs Experimentally Infected with Crenosoma vulpis. Parasitology Research 105: S49-S54

Conte, R (2015). Heliciculture: purpose and economic perspectives in the European community, The Official Journal of the Institute of Science and Technology, available at: https://www.researchgate.net/publication/275030005\_Heliciculture\_purpose\_and\_economic\_perspectives\_in\_the\_European\_community

Cowie, RH and Jones, JS (1987). Ecological interactions between Cepaea nemoralis and Cepaea hortensis: competition, invasion but no niche displacement, Functional Ecology 1: 91-97

Cowie, RH (2001). Decline and homogenization of Pacific faunas: the land snails of American Samoa, Biological Conservation 99: 207-222

Cowie, RH, Dillon, RT, Robinson, DG and Smith, JW (2009). Alien non-marine snails and slugs of priority quarantine importance in the United States: A preliminary risk assessment, American Malacological Bulletin 27: 113-132

Cowie, RH, Hayes, KA, Tran, CT and Meyer, WM (2008). The horticultural industry as a vector of alien snails and slugs: widespread invasions in Hawaii. International Journal of Pest Management 54: 267-276

Cowie, RH and Jones, JS (1987). Ecological interactions between Cepaea nemoralis and Cepaea hortensis: competition, invasion but no niche displacement, Functional Ecology 1(2): 91-97

Cowie, RH (2013). Pathways for transmission of Angiostrongyliasis and the risk of disease associated with them, Hawai’i Journal of Medicine and Public Health 72(6): 70-74

Cowie, RH, Dillon, RT, Robinson, DG and Smith, JW (2009). Alien non-marine snails and slugs of priority quarantine importance in the United States: A preliminary risk assessment, American Malacological Bulletin 27(1-2): 113-132

Cowlishaw, RM, Andrus, P and Rae, R (2019). An Investigation into Nematodes Encapsulated in Shells of Wild, Farmed and Museum Specimens of Cornu Aspersum and Helix Pomatia, Journal of Conchology 43: 385-392

Cribb, TH (1992). The Brachylaimidae (Trematoda, Digenea) of Australian Native Mammals and Birds, Including Descriptions of Dasyurotrema N-G and 4 New Species of Brachylaima, Systematic Parasitology 22(1): 45-72

Cutler, J and Rae, R (2020). Pathogenicity of wild and commercial Phasmarhabditis hermaphrodita exposed to the pestiferous slug Deroceras invadens, Journal of Invertebrate Pathology 174: 107435

Czarnoleski, M, Labecka, AM and Kozłoeski, J (2016). Thermal plasticity of body size and cell size in snails from two subspecies of Cornu aspersum, Journal of Molluscan Studies 82(2): 235-243

da Silva, A and Morassutti, A (2021) Angiostrongylus spp. (Nematoda; Metastrongyloidea) of global public health importance, Research in Veterinary Science 135: 397-403

Dahirel, M, Ansart, A and Madec, L (2014). Stage- and weather-dependent dispersal in the brown garden snail Cornu aspersum, Population Ecology 56: 227-237

Dahirel, M, Vardakis, M Ansart, A and Madec, L (2016). Density-dependence across dispersal stages in a hermaphrodite land snail: insights from discrete choice models, Oecologia 181: 1117-1128

Daniell, A (1994). The impact of terrestrial molluscs on native vegetation in South-eastern Australia, The Victorian Naturalist 111(6): 218-222

Danilova, IS (2022). Heliceculture as a new promising direction of agriculture in Ukraine, (in Ukrainian), Scientific Messenger of LNU of Veterinary Medicine and Biotechnologies, Series: Agricultural Sciences 24(97): 44-47

Dard, C, Piloquet, JE, Qvarnstrom, Y, Fox, LM, M'Kada, H, Hebert, JC, Mattera, D and Harrois, D (2017). First Evidence of Angiostrongyliasis Caused by Angiostrongylus cantonensis in Guadeloupe, Lesser Antilles, American Journal of Tropical Medicine and Hygiene 96(3): 692-697

DAWE (2020). Exotic invasive snails, Department of Agriculture, Water and Energy (ed), available at: https://www.agriculture.gov.au/biosecurity-trade/pests-diseases-weeds/plant/giant-african-snail

DAWR (2015). Final review of policy: importation of Phytophthora ramorum host propagative material into Australia, Department of Agriculture and Water Resources, Canberra, available at https://www.agriculture.gov.au/biosecurity-trade/policy/risk-analysis/memos/ba2015-19

Deak, G, Ionica, AM, Mihalca, AD and Gherman, CM (2017). Troglostrongylus brevior: a new parasite for Romania, Parasites and Vectors 10, doi: 10.1186/s13071-017-2551-4

Dedov, IK, Mitov, PG, Zapryanov, L, Georgiev, D and Gashtarov, V (2022). Distribution of the invasive land snail Eobania vermiculata (O.F. Müller, 1774) (Gastropoda: Helicidae) in Bulgaria, Acta Zoologica Bulgarica 74(4): 611-622

Dees, LT (1970). Edible land snails in the United States, In: United States Department of the Interior, FAWS (ed), Washington, DC, USA, available at: https://pubs.usgs.gov/publication/rp91

Dekle, GW and Fasulo, TR (2021). Brown garden snail, Cornu aspersum (Müller, 1774) (Gastropoda: Helicidae), EENY-240, University of Florida, Institute of Food and Agricultural Sciences, available at: https://entnemdept.ufl.edu/creatures/misc/gastro/brown\_garden\_snail.htm

Department of Agriculture (2013). Importation of dogs and cats and their semen from approved countries: Final Policy Review, Australian Government Department of Agriculture, Canberra, Australia, available at: https://www.agriculture.gov.au/biosecurity-trade/policy/risk-analysis/animal

Department of Agriculture (2019). National priority plant pests 2019, Australian Government Department of Agriculture, Canberra, Australia, available at: https://www.agriculture.gov.au/biosecurity-trade/pests-diseases-weeds/plant/national-priority-plant-pests-2019.

Desouky, MMA and Busais, S (2012). Phylogenetic relationships of the land snail; Eobania vermiculata (Müller, 1774) from Egypt and Saudi Arabia. A combined morphological and molecular analysis, The Journal of Basic and Applied Zoology 65(2): 144-151

Di Cesare, A, Crisi, PE, Bartolini, R, Iorio, R, Talone, T, Filippi, L and Traversa, D (2015). Larval development of Angiostrongylus vasorum in the land snail Helix aspersa, Parasitology Research 114(10): 3649-3655

Di Cesare, A, Crisi, PE, Di Giulio, E, Veronesi, F, di Regalbono, AF, Talone, T and Traversa, D (2013). Larval development of the feline lungworm Aelurostrongylus abstrusus in Helix aspersa, in English), Parasitology Research 112(9): 3101-3108

Di Cesare, A, di Regalbono, AF, Tessarin, C, Seghetti, M, Iorio, R, Simonato, G and Traversa, D (2014). Mixed infection by Aelurostrongylus abstrusus and Troglostrongylus brevior in kittens from the same litter in Italy, Parasitology Research 113(2): 613-618

Diakou, A, Di Cesare, A, Barros, LA, Morelli, S, Halos, L, Beugnet, F and Traversa, D (2015). Occurrence of Aelurostrongylus abstrusus and Troglostrongylus brevior in domestic cats in Greece, Parasites and Vectors 8, doi: 10.1186/s13071-015-1200-z

Diakou, A, Morelli, S, Dimzas, D, Di Cesare, A, Capelli, G, Parrinello, C, Pollmeier, M, Schaper, R and Traversa, D (2019). Efficacy of a moxidectin/imidacloprid spot-on formulation (Advocate (R)) for the treatment of Troglostrongylus brevior in naturally infected cats in a field study in Greece, Parasites and Vectors 12(1), doi: 10.1186/s13071-019-3760-9

Drozd, L, Ziomek, M, Szkucik, K, Paszkiewicz, W, Mackowiak-Dryka, M, Belkot, Z and Gondek, M (2017). Selenium, copper, and zinc concentrations in the raw and processed meat of edible land snails harvested in Poland, Journal of Veterinary Research 61(3): 293-298

Dupont-Nivet, M, Guiller, A and Bonnet, JC (1997). Genetic and environmental variability of adult size in some stocks of the edible snails, Helix aspersa, Journal of Zoology 241(4): 757-765

Dupont-Nivet, M, Mallard, J, Bonnet, JC and Blanc, JM (2001). Evolution of genetic variability in a population of the edible snail, Helix aspersa Müller, undergoing domestication and short-term selection, Heredity 87: 129-135

Egorov, R (2008). Treasure of Russian Shells: illustrated catalogue of the recent terrestrial molluscs of Russia and adjacent regions, Supplement 5, available at: https://www.nhbs.com/treasure-of-russian-shells-supplement-5-illustrated-catalogue-of-the-recent-terrestrial-molluscs-of-russia-and-adjacent-countries-book

El-Danasoury, H and Iglesias-Pineiro, J (2017). Performance of the slug parasitic nematode Phasmarhabditis hermaphrodita under predicted conditions of winter warming, Journal of Pesticide Science 42(1-2): 62-66

Elhamalawi, ZA and Menge, JA (1996). The role of snails and ants in transmitting the avocado stem canker pathogen, Phytophthora citricola, Journal of the American Society for Horticultural Science, 121: 973-977

Ellijimi, C, Ben Hammouda, M, Othman, H, Moslah, W, Jebali, J et al. (2018). Helix aspersa maxima mucus exhibits antimelanogenic and antitumoral effects against melanoma cells, Biomedicine and Pharmacotherapy 101: 871-880

Ellstrand, NC and Schierenbeck, KA (2000). Hybridization as a stimulus for the evolution of invasiveness in plants? Euphytica 148: 35-46

Elmslie, LJ (2005). Snail collection and small-scale production in Africa and Europe, in Ecological implications of minilivestock: potential of insects, rodents, frogs and snails, Paoletti, MG (ed), CRC Press

Elsheikha, HM, Holmes, SA, Wright, I, Morgan, ER and Lacher, DW (2014). Recent advances in the epidemiology, clinical and diagnostic features, and control of canine cardio-pulmonary angiostrongylosis, Veterinary Research 45(1): 92

Elsheikha, HM, Wright, I and McGarry, J (2018). Parasites of the Respiratory System, in HM Elsheikha (ed) Parasites and Pets: A Veterinary Nursing Guide, CABI, London, pp. 33-51

El-Wakil, HB, Banaja, AEA and Amer, SAM (2011). Morphometric and genetic insignts for three terrestrial snails in Taif Province of Saudi Arabia, World Applied Sciences Journal 14(4): 546-551

Eshra, ESH (2013). Survey and distribution of terrestrial snails in fruit orchards and ornamental plants at Alexandria and EL-Beheira governorates, Egypt, Alexandria Science Exchange Journal 34(2): 242-248

Evans-Gilbert, T, Lindo, JF, Henry, S, Brown, P and Christie, CD (2014). Severe eosinophilic meningitis owing to Angiostrongylus cantonensis in young Jamaican children: case report and literature review, Paediatrics and International Child Health 34(2): 148-152

Facon, B, Jarne, P, Pointer, JP and David, P (2005). Hybridisation and invasiveness in the freshwater snail Melanoides tuberculate: hybrid vigour is more important than increase in genetic variance. Journal of Evolutionary Biology 18: 524-535

Falsone, L, Brianti, E, Gaglio, G, Napoli, E, Anile, S, Mallia, E, Giannelli, A, Poglayen, G, Giannetto, S and Otranto, D (2014). The European wildcats (Felis silvestris silvestris) as reservoir hosts of Troglostrongylus brevior (Strongylida: Crenosomatidae) lungworms, Veterinary Parasitology 205(1-2): 193-198

FAO (2019). International Standards for Phytosanitary Measures (ISPM) no. 11: Pest risk analysis for quarantine pests, Secretariat of the International Plant Protection Convention, Food and Agriculture Organization (FAO) of the United Nations, Rome, Italy, available at https://www.ippc.int/en/core-activities/standards-setting/ispms/

Fasanella, A, Lia, R and Giangaspero, A (1995). Cernuella-Virgata (Mollusca, Gastropoda, Pulmonata) Intermediate Snail Host for Dicrocoelium-Dendriticum (Rudolphi, 1819) in Apulia Region, Italy, (in French), Parasite 2(3): 331-333

Ferdushy, T and Hasan, MT (2010). Angiostrongylus vasorum: the 'French Heartworm', Parasitology Research 107(4): 765-771

Ferdushy, T, Kapel, CM, Webster, P, Al-Sabi, MN and Gronvold, J (2009). The occurrence of Angiostrongylus vasorum in terrestrial slugs from forests and parks in the Copenhagen area, Denmark, J Helminthol 83(4): 379-383

Février, J, Russo, J and Madec, L (2009). Intraspecific variation in life history traits of a land snail after a bacterial challenge, Journal of Zoology 277(2): 149-56

Fevrier, Y, Russo, J and Madec, L (2009). Intraspecific variation in life history traits of a land snail after a bacterial challenge, Journal of Zoology 277: 149-156

Forsyth, RG and Kamstra, J (2019). Roman snail, Helix pomatia (Mollusca: Helicidae), in Canada, The Canadian Field-Naturalist 133(2): 156-159

Frankham, R (2005). Invasion biology - Resolving the genetic paradox in invasive species. Heredity 94: 385-385

Freitas, FAO, Carrara, ER, Ladeira, G, Lourenҫo, M, Leôncio, T, Miranda, C, César, FL and de Genova Gaya, L (2023). Heritability and genetics correlations for body weight in escargots, Acta Scientiarum: Animal Sciences 45, doi: 10.4025/actascianimsci.v45i1.58130

Fuehrer, HP, Morelli, S, Bleicher, J, Brauchart, T, Edler, M, Eisschiel, N, Hering, T et al. (2020). Detection of Crenosoma spp., Angiostrongylus vasorum and Aelurostrongylus abstrusus in Gastropods in Eastern Austria, Pathogens 9(12), doi: 10.3390/pathogens9121046

GACC (2019). Huangpu Customs seizes alien invasive snails at Xingang Port, General Administration of Customs of the People's Republic of China (GACC), 6 May 2019, available at http://english.customs.gov.cn/Statics/cc31370c-91a9-4eec-a52c-e855da134172.html

Gallego, L and Gracenea, M (2015). Praziquantel efficacy against Brachylaima sp metacercariae (Trematoda: Brachylaimidae) parasitizing the edible landsnail Cornu aspersum and its HPLC-MS/MS residue determination, Experimental Parasitology 157: 92-102

Gallego, L, Gonzalez-Moreno, O and Gracenea, M (2014). Terrestrial Edible Land Snails as Vectors for Geographic Dissemination of Brachylaima Species, Journal of Parasitology 100(5): 674-678

Garefalaki, ME, Kalyva, S, Janicke, T and Staikou, A (2017). Intraspecific variation in reproductive characters is associated with the strength of sexual selection in the hermaphroditic land snail Cornu aspersum, Behavioral Ecology and Sociobiology 71, doi: 10.1007/s00265-017-2383-4

Gerard, C, Ansart, A, Decanter, N, Martin, MC and Dahirel, M (2020). Brachylaima spp. (Trematoda) parasitizing Cornu aspersum (Gastropoda) in France with potential risk of human consumption, Parasite 27, doi: 10.1051/parasite/2020012

Gerard, C, De Tombeur, Y, Dahirel, M and Ansart, A (2023). Land snails can trap trematode cercariae in their shell: Encapsulation as a general response against parasites? Parasite 30, doi: 10.1051/parasite/2023001

Gherman, CM, Ionica, AM, D'Amico, G, Otranto, D and Mihalca, AD (2016). Angiostrongylus chabaudi (Biocca, 1957) in wildcat (Felis silvestris silvestris, S) from Romania, Parasitology Research, 115(6): 2511-2517

Giannelli, A, Ramos, RAN, Annoscia, G, Di Cesare, A, Colella, V, Brianti, E, Dantas-Torres, F, Mutafchiev, Y and Otranto, D (2014). Development of the feline lungworms Aelurostrongylus abstrusus and Troglostrongylus brevior in Helix aspersa snails, Parasitology 141(4): 563-569

Giusti, F, Fiorentino, V and Manganelli, G (2015). A Neotype for Helix Cincta Muller, 1774 (Gastropoda, Pulmonata, Helicidae), Journal of Conchology 42: 209-212

Glasheen, PM, Burks, RL, Campos, SR and Hayes, KA (2020). First evidence of introgressive hybridization of apple snails (Pomacea spp.) in their native range, Journal of Molluscan Studies 86: 96-103

Gomot, A and Pihan, F (1997). Comparison of the bioaccumulation capacities of copper and zinc in two snail subspecies (Helix). Ecotoxicology and Environmental Safety 38: 85-94

Gomot, A (1997). Dose-dependent effects of cadmium on the growth of snails in toxicity bioassays, Archives of Environmental Contamination and Toxicology 33: 209-216

Gomot-De Vaufleury, A and Borgo, R (2001). Experimental hybridization between two sub-species of snails (Helix aspersa aspersa and Helix aspersa maxima): consequences for fertility, survival and growth, Invertebrate Reproduction and Development 40(2-3): 217-226

Gomot-De Vaufleury, A and Pihan, F (2002). Methods for toxicity assessment of contaminated soil by oral or dermal uptake in land snails: Metal bioavailability and bioaccumulation, Environmental Toxicology and Chemistry 21: 820-827

Gonzalez, P, Carbonell, E, Urios, V and Rozhnov, VV (2007). Coprology of Panthera tigris altaica and Felis bengalensis euptilurus from the Russian far east, Journal of Parasitology 93(4): 948-950

Gonzalez-Moreno, O and Gracenea, M (2006). Life cycle and description of a new species of brachylaimid (Trematoda : Digenea) in Spain, Journal of Parasitology 92(6): 1305-1312

Gracenea, M and Gallego, L (2017). Brachylaimiasis: Brachylaima Spp. (Digenea: Brachylaimidae) Metacercariae Parasitizing the Edible Snail Cornu Aspersum (Helicidae) in Spanish Public Marketplaces and Health-Associated Risk Factors, Journal of Parasitology 103(5): 440-450

Gracenea, M and Gonzalez-Moreno, O (2002). Life cycle of Brachylaima mascomai n. sp (Trematoda : Brachylaimidae), a parasite of rats in the Llobregat Delta (Spain), Journal of Parasitology 88(1): 124-133

Gracenea, M and Gallego, L (2017). Brachylaimiasis: Brachylaima Spp. (Digenea: Brachylaimidae) Metacercariae Parasitizing the Edible Snail Cornu Aspersum (Helicidae) in Spanish Public Marketplaces and Health-Associated Risk Factors, Journal of Parasitology 103: 440-450

Graham, FJ, Ford, JB and Runham, NW (1993). Comparison of 2 Species of Mites of the Same Genus, Riccardoella Associated with Mollusks, Acarologia 34(2): 143-148

Grannell, A, Cutler, J and Rae, R (2021). Size-susceptibility of Cornu aspersum exposed to the malacopathogenic nematodes Phasmarhabditis hermaphrodita and P. californica, Biocontrol Science and Technology 31(11), doi: 10.1080/09583157.2021.1929072

Grewal, PS, Grewal, SK, Tan, L and Adams, BJ (2003). Parasitism of molluscs by nematodes: Types of associations and evolutionary trends, Journal of Nematology 35(2): 146-156

Grewal, SK and Grewal, PS (2003). Effect of osmotic desiccation on longevity and temperature tolerance of Phasmarhabditis hermaphrodita (Nematoda : Rhabditidae), Journal of Parasitology 89(3): 434-438

Grigore, S (2021). First record of Eobania vermiculata (O.F. Müller, 1774) (Gastropoda: Eupulmonata: Helicidae) in Romania, Folia Malacologica 29(1): 51-53

Guiller, A and Madec, L (2010). Historical biogeography of the land snail Cornu aspersum: a new scenario inferred from haplotype distribution in the Western Mediterranean basin, BMC Evolutionary Biology 10, doi: 10.1186/1471-2148-10-18

Guiller, A, Coutellec-Vreto, MA, Madec, L and Deunff, J (2001). Evolutionary history of the land snail Helix aspersa in the Western Mediterranean: preliminary results inferred from mitochondrial DNA sequences, Molecular Ecology 10(1): 81-87

Guiller, A, Coutellec-Vreto, MA, Madec, L and Deunff, J (2001). Evolutionary history of the land snail Helix aspersa in the Western Mediterranean: preliminary results inferred from mitochondiral DNA sequences, Molecular Ecology 10: 81-87

Guiller, A, Martin, MC, Hiraux, C and Madec, L (2012). Tracing the Invasion of the Mediterranean Land Snail Cornu aspersum aspersum Becoming an Agricultural and Garden Pest in Areas Recently Introduced, PLoSOne 7(12), doi: 10.1371/journal.pone.0049674

Gural-Sverlova, N and Gural, R (2021). Cornu aspersum (Gastropoda: Helicidae) in Western Ukraine with an overview of introduced species of land molluscs from this area, Malacologica Bohemoslovaca 20: 123-135

Gürelli, G (2017). Importance of Land Snails in Dicrocoeliosis Epidemiology, Turkiye Parazitolojii Dergisi 41(3): 169-172

Haeussler, EM, Piza, J, Schmera, D and Baur, B (2012). Intensity of parasitic mite infection decreases with hibernation duration of the host snail, Parasitology 139(8): 1038-1044

Haites, RE, Watt, AE, Russell, DA and Billman-Jacobe, H (2021). Infection of Slugs with Theronts of the Ciliate Protozoan, Tetrahymena rostrata, Microorganisms 9(9), doi: 10.3390/microorganisms9091970.

Hapca, S, Crawford, J, Rae, R, Wilson, M and Young, I (2007). Movement of the parasitic nematode Phasmarhabditis hermaphrodita in the presence of mucus from the host slug Deroceras reticulatum, Biological Control 41(2): 223-229

Harrap, L (2021). 1 way to deal with invasive snails in Revelstoke is...to eat them?, Revelstoke Review, 26 June 2021, available at: https://www.revelstokereview.com/community/1-way-to-deal-with-invasive-snails-in-revelstoke-is-to-eat-them-4296298

Hayes, KA, Cowrie, RH, Thiengo, SC and Strong, EE (2012). Comparing apples with apples: clarifying the identities of two highly invasive Neotropical Ampullariidae (Caenogastropoda), Zoological Journal of the Linnean Society 166: 723-753

Hazir, A and Ulusoy, MR (2012). Adana ve Mersin illeri şeftali ve nektarin alanlarında saptanan zararlıar ile predatör ve parazitoit türler, (Pest, parasitoid and predator species determined in peach and nectarine orchards in Adana and Mersin provinces), Türkiye Biyolojik Mücadele Derneǧi 3(2): 157-168

Helm, J, Roberts, L, Jefferies, R, Shaw, SE and Morgan, ER (2015). Epidemiological survey of Angiostrongylus vasorum in dogs and slugs around a new endemic focus in Scotland, Veterinary Record 177(2), doi: 10.1136/vr.103006

Herbert, DG (2010). The introduced terrestrial mollusca of South Africa, SANBI Biodiversity Series 15, South Africa National Biodiversity Institute, Pretoria, ISBN: 978-1-919976-56-3

Hetherington, S (2009). Integrated pest management for Australian apples and pears. Apple and Pear Australia Limited, available at: https://www.dpi.nsw.gov.au/\_\_data/assets/pdf\_file/0009/321201/ipm-for-australian-apples-and-pears-complete.pdf

HeXiang, L, Shan, L, Ling, H and XiaoNong, Z (2009). Establishment and observation of the life cycle of Angiostrongylus cantonensis in a laboratory setting, Journal of Pathogen Biology 4(11): 836-839

Hobmaier, M and Hobmaier, A (1935). Intermediate hosts of Aelurostrongylus abstrusus of the cat, Proceedings of the Society for Experimental Biology and Medicine 32(9): 1641-1647

Holland, KD, McDonnell, MJ and Williams, NSG (2007). Abundance, species richness and feeding preferences of introduced molluscs in native grasslands of Victoria, Australia, Austral Ecology 32(6): 626-634

Holley, M (2020). Phasmarhabditis hermaphrodita is not the only slug killing nematode, IOBC-WPRS Bulletin 150: 152-156

Hughes, AJ and Biggs, BA (2002). Parasitic worms of the central nervous system: an Australian perspective, Internal Medicine Journal 32: 541-553

Hyder, N, Coffey, MD and Stanghellini, ME (2009). Viability of oomycete propagules following ingestion and excretion by fungus gnats, shore flies, and snails, Plant Disease 93(7): 720-726

Iglesias, J and Castillejo, J (1999). Field observations on feeding of the land snail Helix aspersa Müller, Journal of Molluscan Studies 65: 411-423

Imai, E and Hennessey, RD (1999). Importation and interstate movement of live, edible land snails: Cantareus apertus (Born), Cryptomphalys aspersus (Müller), Eobania vermiculata (Müller), Helix pomatia Linné, and Otala lactea (Müller) (Pulmonata: Helicidae): Qualitative Pest Risk Analysis, United States Department of Agriculture, Riverdale (MD) USA

Ismail, FMAK and Gurelli, G (2018). First report of natural infection of Xerolenta obvia (Pulmonata, Mollusca) by Dicrocoeliidae (Digenea) larval stages in Turkey, Kastamonu University Journal of Forestry Faculty 18(3): 272-278

Jeandron, A, Rinaldi, L, Abdyldaieva, G, Usubalieva, J, Steinmann, P, Cringoli, G and Utzinger, J (2011). Human Infections with Dicrocoelium dendriticum in Kyrgyzstan: The Tip of the Iceberg? Journal of Parasitology 97(6): 1170-1172

Jess, S and Marks, RJ (1998). Effect of temperature and photoperiod on growth and reproduction of Helix aspersa var. maxima, Journal of Agricultural Science 130(3): 367-372

Jiang, X, Zheng, P, Soto, I, Haubrock, PJ, Chen, J and Ji, L (2022). Global economic costs and knowledge gaps of invasive gastropods, Ecological Indicators 145, doi: 10.1016/j.ecolind.2022.109614

Jimoh, OA and Akinola, MO (2020). Reproductive performance of laying snails (Archachatina marginata) fed on roughages and different concentrate mixes. Bulletin of the National Research Centre 44: 1-6

Johnstone, R (2021). Managing snails in citrus orchards [Online]. Western Australia Department of Primary Industries and Regional Development. Available: https://www.agric.wa.gov.au/citrus/managing-snails-citrus-orchards?nopaging=1 [Accessed].

Kaczanowski, A, Brunk, CF and Kazubski, SL (2016). Cohesion of Clonal Life History, Senescence and Rejuvenation Induced by Autogamy of the Histophagous Ciliate Tetrahymena rostrata, Protist 167(5): 490-510

Kahl, A, von Samson-Himmelstjerna, G, Krucken, J and Ganter, M (2021). Chronic Wasting Due to Liver and Rumen Flukes in Sheep, Animals 11(2), doi: 10.3390/ani11020549

Kahraman, T, Kirişik, M and Kahraman, MU (2020). Determination of pests and beneficial species in avocado orchards in Antalya Province, Horticultural Studies 37(2): 150-154

Karaman, BJ (2006). Former investigations of the fauna of snails (Mollusca, Gastropoda) in Bosnia and Herzegovina, Natura Montenegrina, Podgorica 5: 55-66

Khanjari, A, Partovi, R, Abbaszadeh, S, Nemati, G, Bahonar, A, Misaghi, A, Akhondzadeh-Basti, A, Alizadeh-Ilanjegh, A, Motaghifar, A (2010). A retrospective survey of fasciolosis and dicrocoeliosis in slaughtered animals in Meisam abattoir, Tehran, Iran (2005-2008), paper presented at Veterinary Research Forum, available at: https://www.researchgate.net/publication/229020888\_A\_Retrospective\_Survey\_of\_Fasciolosis\_and\_Dicrocoeliosis\_in\_Slaughtered\_Animals\_in\_Meisam\_Abattoir\_Tehran\_Iran\_2005-2008

Kim, JR, Hayes, KA, Yeung, NW and Cowie, RH (2014). Diverse gastropod hosts of Angiostrongylus cantonensis, the rat lungworm, globally and with a focus on the Hawaiian Islands, PLoSOne 9(5), doi: 10.1371/journal.pone.0094969

Koene, JM (2006). Tales of two snails: sexual selection and sexual conflict in Lymnaea stagnalis and Helix aspersa, Integrative and Comparative Biology 46: 419-429

Korábek, O, Juřičková, L and Petrusek, A (2016). Splitting the Roman snail Helix pomatia Linnaeus, 1758 (Stylommatophora: Helicidae) into two: redescription of the forgotten Helix thessalica Boettger, 1886, Journal of Molluscan Studies 82: 11-22

Kose, M, Eser, M, Kartal, K and Bozkurt, MF (2015). Infections of Larval Stages of Dicrocoelium dendriticum and Brachylaima sp in Brown Garden Snail, Helix aspersa, in Turkey, Korean Journal of Parasitology 53(5): 647-651

Kougiagka, E, Gkafas, GA, Exadactylos, A and Hatziioannou, M (2022). Morphology and genetic structure profile of farmed snails Cornu aspersum aspersum and Cornu aspersum maximum in Greece, Sustainability 14(23), doi: 10.3390/su142315965

Lange, MK, Penagos-Tabares, F, Hirzmann, J, Failing, K, Schaper, R, Van Bourgonie, YR, Backeljau, T, Hermosilla, C and Taubert, A (2018). Prevalence of Angiostrongylus vasorum, Aelurostrongylus abstrusus and Crenosoma vulpis larvae in native slug populations in Germany, Veterinary Parasitology 254: 120-130

Latrofa, MS, Lia, RP, Giannelli, A, Colella, V, Santoro, M, D'Alessio, N, Campbell, BE, Parisi, A, Dantas-Torres, F, Mutafchiev, Y, Veneziano, V and Otranto, D (2015). Crenosoma vulpis in wild and domestic carnivores from Italy: a morphological and molecular study, Parasitology Research 114(10): 3611-3617

Leonard, E (2003). Bash’Em Burn’Em Bait’Em - Integrated snail management in crops and pastures, South Australian Research and Development Institute, GRDC Project Code: UOA2205-005RTX

Ligaszewski, M and Pol, P (2017). Calcium, crude ash and crude protein content in the visceral sac and shell of the edible garden snail (Cornu aspersum synonym Helix aspersa) farmed in confined conditions, Wiadomości Zootechniczne 4: 3-9

Ligaszewski, M and Pol, P (2021). Reproduction of the Roman Snail (Helix Pomatia L) from a Local Natural Population in Farm Conditions and in a Natural Habitat, Annals of Animal Science, 21(2): 693-708

Ligaszewski, M, Lysak, A and Mach-Paluszkiewicz, Z (2007). Reproductive performance of Helix pomatia (Gastropoda : Pulmonata : Helicidae) and survival of its hatchlings under farm conditions, American Malacological Bulletin 22(1-2): 1-6

Ligaszewski, M, Surowka, K and Stekla, J (2009). The shell features of Cornu aspersum (synonym Helix aspersa) and Helix pomatia: Characteristics and comparison, American Malacological Bulletin 27(1-2): 173-181

Ligaszewski, M and Pol, P (2017). Calcium, crude ash and crude protein content in the visceral sac and shell of the edible garden snail (Cornu aspersum synonym Helix aspersa) farmed in confined conditions, Wiadomości Zootechniczne 4: 3-9

Lurati, L, Deplazes, P, Hegglin, D and Schnyder, M (2015). Seroepidemiological survey and spatial analysis of the occurrence of Angiostrongylus vasorum in Swiss dogs in relation to biogeographic aspects, Veterinary Parasitology 212(3-4): 219-226

Lv, S, Zhang, Y, Liu, HX, Hu, L, Yang, K, Steinmann, P, Chen, Z, Wang, LY, Utzinger, J and Zhou, XN (2009). Invasive snails and an emerging infectious disease: results from the first national survey on Angiostrongylus cantonensis in China, PLoS Neglected Tropical Diseases 3(2): e368

Lv, S, Zhang, YI, Steinmann, P, Yang, G-J, Yang, KUN, Zhou, X-N and Utzinger, J (2011). The emergence of angiostrongyliasis in the People’s Republic of China: the interplay between invasive snails, climate change and transmission dynamics, Freshwater Biology 56(4): 717-734

Lydeard, C, Cowie, RH, Ponder, WF, Bogan, AE, Bouchet, P, Clark, SA, et al. (2004). The global decline of nonmarine mollusks, Bioscience 54: 321-330

Łysak, A (1999). Significance of Helix farming for protection of Helix pomatia, Folia Malacologica 7(4): 267-270

Ma, G, Dennis, M, Rose, K, Spratt, D and Spielman, D (2013). Tawny frogmouths and brushtail possums as sentinels for Angiostrongylus cantonensis, the rat lungworm, Veterinary Parasitology 192(1): 158-165

Mackerras, MJ (1957). Observations on the life history of the cat lungworm Aelurostrongylus abstrusus (Railliet, 1898)(Nematoda: Metastrongylidae), Australian Journal of Zoology 5: 188-195

Mackie, JT, Lacasse, C and Spratt, DM (2013). Patent Angiostrongylus mackerrasae infection in a black flying fox (Pteropus alecto), Australian Veterinary Journal 91(9): 366-367

Mackowiak-Dryka, M, Szkucik, K and Pyz-Lukasik, R (2020a). Snail eggs as a raw material for the production of a caviar substitute, Journal of Veterinary Research 64:543-547

Mackowiak-Dryka, M, Szkucik, K and Pyz-Lukasik, R (2020b). Fatty acid profile in edible eggs of snails from the Cornu genus, Journal of Veterinary Research 64: 137-140

MacMillan, K, Blok, V, Young, I, Crawford, J and Wilson, MJ (2006). Quantification of the slug parasitic nematode Phasmarhabditis hermaphrodita from soil samples using real time qPCR, International Journal for Parasitology 36(14): 1453-1461

MacPherson, CN (2005). Human behaviour and the epidemiology of parasitic zoonoses. International Journal for Parasitology 35: 1319-1331

Madec, L, Bellido, A and Guiller, A (2003). Shell shape of the land snail Cornu aspersum in North Africa: unexpected evidence of a phylogeographical splitting, Heredity 91(3): 224-231

Madec, L, Guiller, A, Coutellec-Vreto, MA and Desbuquois, C (1998). Size-fecundity relationships in the land snail Helix aspersa: preliminary results on a form outside the norm, Invertebrate Reproduction and Development 34(1): 83-90

Maksimov, P, Hermosilla, C, Taubert, A, Staubach, C, Sauter-Louis, C, Conraths, FJ, Vrhovec, MG and Pantchev, N (2017). GIS-supported epidemiological analysis on canine Angiostrongylus vasorum and Crenosoma vulpis infections in Germany, Parasites and Vectors 10, DOI 10.1186/s13071-017-2054-3

Manea, D, Ienciu, AA, Stef, Pet, I, Smuleac, L et al. (2021). The "Sandwich" System: A Potential Solution for Protecting Overwintering Cornu aspersum Snails Reared in Semi-Intensive Heliculture Farms in Colder Climates. Animals, 11, doi: 10.3390/ani11051420

Maraqa, A, Amr, Z, Rifai, L and Al-Melhim, W (2005). An abattoir survey of liver and lung helminthic infections in local and imported sheep in Jordan, Turkish Journal of Veterinary and Animal Sciences 29(1): 1-2

Martin-Vega, D, Garbout, A, Ahmed, F, Wicklein, M, Goater, CP, Colwell, DD and Hall, MJR (2018). 3D virtual histology at the host/ parasite interface: visualisation of the master manipulator, Dicrocoelium dendriticum, in the brain of its ant host, Scientific Reports, vol. 8, article number: 8587

Mc Donnell, RJ, Colton, AJ, Howe, DK and Denver, DR (2020). Lethality of four species of Phasmarhabditis (Nematoda: Rhabditidae) to the invasive slug, Deroceras reticulatum (Gastropoda: Agriolimacidae) in laboratory infectivity trials, Biological Control (150), doi: 10.1016/j.biocontrol.2020.104349

Michaud, A (2019). The ingestion and defecation of viable root-knot nematode and cyst nematode eggs by Cornu aspersum, Master of Science Thesis, University of California, Davis

Michelson, EH (1971). Distribution and Pathogenicity of Tetrahymena-Limacis in Slug Deroceras-Reticulatum, Parasitology 62, doi: 10.1017/S003118200007133x

Mienis, HK, Rittner, O and Vaisman, S (2016). Information concerning Eobania vermiculata, I. On the presence of this exotic species in Israel (Mollusca, Gastropoda, Helicidae), Triton 34: 29-36

Milinsk, MC, Padre, RD, Kayashi, C de Souza, NE and Matsushita, M (2003). Influence of diets enriched with different vegetable oils on the fatty acid profiles of snail Helix aspersa maxima, Food Chemistry 82: 553-558

Mitchell, G, Cuthill, G, Haine, A, Zadoks, R, Chaudhry, U, Skuce, P and Sargison, N (2017). Evaluation of molecular methods for the field study of the natural history of Dicrocoelium dendriticum, Veterinary Parasitology 235: 100-105

Modry, D, Feckova, B, Putnova, B, Manalo, SM and Otranto, D (2021). Alternative pathways in Angiostrongylus cantonensis (Metastrongyloidea: Angiostrongylidae) transmission, Parasitology 148(2): 167-173

Mohammed, GR (2015). Incidence of land snails inhabiting different vegetation at some governorates in north-east of delta Egypt, Journal of Plant Protection and Pathology 6(6): 899-907

Mona, MH, Desouky, MM, Mohamadeen, AM, Salama, WM, Fahmy, AM (2019). Heliculture of garden snail (Helix aspersa): Effect of different food items on growth performance and biochemical composition, Egyptian Journal of Experimental Biology 15: 161-169

Monks, DJ, Carlisle, MS, Carrigan, M, Rose, K, Spratt, D, Gallagher, A and Prociv, P (2005). Angiostrongrylus cantonensis as a cause of cerebrospinal disease in a yellow-tailed black cockatoo (Calyptorhynchus funereus) and two tawny frogmouths (Podargus strigoides), Journal of Avian Medicine and Surgery 19(4): 289-293

Morelli, S, Traversa, D, Colombo, M, Raue, K, Strube, C, Pollmeier, M and Di Cesare, A (2020). The effect of the hibernation on the larval development of Troglostrongylus brevior in the land snail Cornu aspersum, Veterinary Parasitology 282, doi: 10.1016/j.vetpar.2020.109123

Moreno-Rueda, G (2008). The colour white diminishes weight loss during aestivation in the arid-dwelling land snail Sphincterochila (Albea) candidissima, Iberus 26: 47-51

Morgan, ER and Wall, R (2009). Climate change and parasitic disease: farmer mitigation? Trends in Parasitology 25(7): 308-313

Morgan, ER, Jefferies, R, van Otterdijk, L, McEniry, RB, Allen, F, Bakewell, M and Shaw, SE (2010). Angiostrongylus vasorum infection in dogs: Presentation and risk factors, Veterinary Parasitology 173(3-4): 255-261

Morgan, ER, Shaw, SE, Brennan, SF, De Waal, TD, Jones, BR and Mulcahy, G (2005). Angiostrongylus vasorum: a real heartbreaker, Trends in Parasitology 21(2): 49-51

Morris, A, Green, M, Martin, H, Crossland, K, Swaney, WT, Williamson, SM and Rae, R (2018). A nematode that can manipulate the behaviour of slugs, Behavioural Processes 151: 73-80

Moskvina, TV (2018). Current knowledge about Aelurostrongylus abstrusus biology and diagnostic, Annals of Parasitology 64(1): 3-11

Mozzer, LR, Montresor, LC, Vidigal, TH and Lima, WS (2011). Angiostrongylus vasorum: Experimental Infection and Larval Development in Omalonyx matheroni, Journal of Parasitology Research 2011: 178748

Mumladze, L and Paposhvili, N (2016). A new addition to the malacofauna of Georgia – Eobania vermiculata is replenishing its range, Proceedings of the Institute of Zoology 25: 153-155

Murphy, B (2001). Breeding and Growing Snails Commercially in Australia, RIRDC Publication No. 00/188, Australian Government Rural Industries Research and Development Corporation

Nelson, TA, Gregory, DG, Burroughs, C and Laursen, JR (2007). Prevalence of lungworms in Illinois coyotes, Transactions of the Illinois State Academy of Science 100(1): 89-95

Nemaslug (2016). Nemaslug [Online]. Available: http://nemaslug.co.uk/using-nemaslug

Neubert, E (2014). Revision of Helix Linnaeus, 1758 in its eastern Mediterranean distribution area, and reassignment of Helix godetiana Kobelt, 1878 to Maltzanella Hesse, 1917 (Gastropoda, Pulmonata, Helicidae), Contributions to Natural History 26: 1-200

Neubert, E (2011). Helix lucorum. The IUCN Red List of Threatened Species 2011 ed

Nevarez, A, Lopez, A, Conboy, G, Ireland, W and Sims, D (2005). Distribution of Crenosoma vulpis and Eucoleus aerophilus in the lung of free-ranging red foxes (Vulpes vulpes), Journal of Veterinary Diagnostic Investigation 17(5): 486-489

Ngenwi, AA, Mafeni, JM, Etchu, KA and Oben, FT (2010). Characteristics of snail farmers and constraints to increased production in West and Central Africa, African Journal of Environmental Science and Technology 4(5): 274-278

Nippon News (2016). Japanese farmer breeding rare escargot de Burgogne snails, 25 May 2016, available at https://www.nipponnews.net/features/food-features/japanese-farmer-breeding-rare-escargot-de-burgogne-snails/, accessed 19 March 2024

Nordsieck, R (2022a). The Brown Garden Snail, The Living World of Molluscs

Nordsieck, R (2022b). The Turkish Snail, The Living World of Molluscs

Notton, D (2006). Eobania vermiculata in the UK, Mollusc World 11: 6

Nyoagbe, LA, Appiah, V, Nketsia-Tabiri, J, Larbi, D and Adjei, I (2016). Evaluation of African giant snails (Achatina and Archachatina) obtained from markets (wild) and breeding farms, African Journal of Food Science 10(7): 94-104

Otranto, D and Traversa, D (2003). Dicrocoeliosis of ruminants: a little known fluke disease, Trends in Parasitology 19(1): 12-15

Ozgo, M and Schilthuizen, M (2012). Evolutionary change in Cepaea nemoralis shell colour over 43 years, Global Change Biology 18(1): 74-81

Páll-Gergely, B, Fehér, Z and Čejka, T (2020). New records of the Mediterranean land snail Massylaea vermiculata (OF Müller, 1774) in Hungary and Slovakia, Folia Malacologica 28(4): 337-41

Pearce, TA, Olori, JC and Kemezis, KW (2010). Land Snails from St. Elzear Cave, Gaspe Peninsula, Quebec: Antiquity of Cepaea Hortensis in North America, Annals of Carnegie Museum 79(1): 65-78

Pechova, H and Foltan, P (2008). The parasitic nematode Phasmarhabditis hermaphrodita defends its slug host from being predated or scavenged by manipulating host spatial behaviour, Behavioural Processes 78(3): 416-420

Penagos-Tabares, F, Lange, MK, Velez, J, Hirzmann, J, Gutierrez-Arboleda, J, Taubert, A, Hermosilla, C and Gutierrez, JJC (2019). The invasive giant African snail Lissachatina fulica as natural intermediate host of Aelurostrongylus abstrusus, Angiostrongylus vasorum, Troglostrongylus brevior, and Crenosoma vulpis in Colombia, PLoS Neglected Tropical Diseases 13(4), doi: 10.1371/journal.pntd.0007277

Pollard, E (1975). Aspects of the ecology of Helix pomatia L., Journal of Animal Ecology 44(1): 305-329

Popiolek, M, Jarnecki, H and Luczynski, T (2009). A record of Crenosoma vulpis (Rudolphi, 1819) (Nematoda, Crenosomatidae) from Eurasian badger (Meles meles L.) from Poland, Wiadomooeci Parazytologiczne 55(4): 437-439

Prociv, P and Carlisle, MS (2001). The spread of Angiostrongylus cantonensis in Australia, Southeast Asian Journal of Tropical Medicine and Public Health 32: 126-128

Prociv, P, Spratt, DM and Carlisle, MS (2000). Neuro-angiostrongyliasis: unresolved issues, International Journal for Parasitology 30(12-13): 1295-303

Puizina, J, Fredotović, Ž, Šamanić, I, Šušnjara, T, Kekez, L, Cukrov, D and Pleslić, G (2013). Phylogeography of the land snail Eobania vermiculata (OF Müller, 1774) (Gastropoda: Pulmonata) along the Croatian coast and islands, Journal of Entomology and Zoology Studies 1(4): 23-31

Purdue University (2024). Pest Tracker, Purdue University, West Lafayette (IN) USA, available at https://www.pesttracker.org/, accessed 2024

Qvarnstrom, Y, Sullivan, JJ, Bishop, HS, Hollingsworth, R and da Silva, AJ (2007). PCR-based detection of Angiostrongylus cantonensis in tissue and mucus secretions from molluscan hosts, Applied and Environmental Microbiology 73(5): 1415-1419

Rae, R, Verdun, C, Grewal, P, Robertson, JF and Wilson, MJ (2007). Biological control of terrestrial molluscs using Phasmarhabditis hermaphrodita - progress and prospects, Pest Management Science 63(12): 1153-1164

Rae, RG, Robertson, JF and Wilson, MJ (2009). Chemoattraction and Host Preference of the Gastropod Parasitic Nematode Phasmarhabditis Hermaphrodita, Journal of Parasitology 95(3): 517-526

Raven, JH (2022). Juvenile shells of Elona quimperiana (Férussac, 1821) (Gastropoda: Elonidae), Revista de Malacologia Iberica 3: 48-49

Regnier, C, Fontaine, B and Bouchet, P (2009). Not Knowing, Not Recording, Not Listing: Numerous Unnoticed Mollusk Extinctions, Conservation Biology 23: 1214-1221

Rinaldi, L, Calabria, G, Carbone, S, Carrella, A and Cringoli, G (2007). Crenosoma vulpis in dog: first case report in Italy and use of the FLOTAC technique for copromicroscopic diagnosis, Parasitology Research 101(6): 1681-1684

Ristaino, JB and Gumpertz, ML (2000). New frontiers in the study of dispersal and spatial analysis of epidemics caused by species in the genus Phytophthora, Annual Review of Phytopathology 38: 541-76

Robbins, W, Conboy, G, Greenwood, S and Schaper, R (2021). Infectivity of gastropod-shed third-stage larvae of Angiostrongylus vasorum and Crenosoma vulpis to dogs, Parasites and Vectors 14(1), doi: 10.1186/s13071-021-04802-6

Robinson, DG, Redmond, L and Hennessey, R (1999). Importation and Interstate Movement of Live, Edible Land Snails: Cantareus apertus (Born), Cryptomphalus aspersus (Müller), Eobania vermiculata (Müller), Helix pomatia Linné, and Otala lactea Muller (Pulmonata: Helicidae). USDA APHIS PPQ SS

Robinson, DG (1999). Alien invasions: the effects of the global economy on non-marine gastropod introductions into the United States, Malacologia 41(2): 413-38

Ronsmans, J and Van den Neucker, T (2016). A persistant population of the chocolate-band snail Eobania vermiculata (Gastropoda: Helicidae) in Belgium, Belgian Journal of Zoology 146(1): 66-68

Rosen, L, Ash, LR and Wallace, GD (1970). Life History of Canine Lungworm Angiostrongylus-Vasorum (Baillet), American Journal of Veterinary Research 31(1): 131

Rosin, ZM, Kobak, J, Lesicki, A and Tryjanowski, P (2013). Differential shell strength of Cepaea nemoralis colour morphs-implications for their anti-predator defence, Naturwissenschaften 100(9): 843-851

Rumi, A, Sánchez, J and Ferrando, NS (2010). Theba pisana (Müller, 1774) (Gastropoda, Helicidae) and other alien land molluscs species in Argentina, Biological Invasions 12: 2985-2990

Russo, J and Madec, L (2011). Dual Strategy for Immune Defense in the Land Snail Cornu aspersum (Gastropoda, Pulmonata), Physiological and Biochemical Zoology 84(2): 212-221

Russo, J and Madec, L (2011). Dual strategy for immune defense in the land snail Cornu aspersum (Gastropoda, Pulmonata), Physiological and Biochemical Zoology 84(2): 212-221

Rygalo-Galewska, A, Zglinska, K and Niemiec, T (2022). Edible Snail Production in Europe, Animals 12(20), doi: 10.3390/ani12202732

Sakai, AK, Allendorf, FW, Holt, JS, Lodge, DM, Molofsky, J et al. (2001). The population biology of invasive species, Annual Review of Ecology and Systematics 32: 305-332

Sakovich, NJ (1996). An integrated pest management (IPM) approach to the control of the brown garden snail, (Helix aspersa) in California citrus orchards, available at: https://calisphere.org/item/ark:/86071/d2kn84/

Sakovich, NJ (2002). Integrated Management of Cantareus aspersus (Muller)(Helicidae) as a Pest of Citrus in California. In: BARKER, G. M. (ed.) Molluscs as crop pests. Wallingford, United Kingdom: CABI Pubishing

Sanchez, KR (2010). Nematode and bacterial associates of the invasive brown garden snail, Helix aspersa, Masters of Science Thesis, University of California, Davis

Sanchez, O, Robla, J and Arias, A (2021). Annotated and Updated Checklist of Land and Freshwater Molluscs from Asturias (Northern Spain) with Emphasis on Parasite Transmitters and Exotic Species, Diversity-Basel 13(9), doi: 10.3390/d13090415

Sanderson, G and Sirgel, W (2002). Helicidae as pests in Australian and South African grapevines, in GM Barker (ed) Molluscs as Crop Pests, CABI Publishing, Wallingford, United Kingdom, pp. 255-70.

Sawanyawisuth, K and Sawanyawisuth, K (2008). Treatment of angiostrongyliasis, Transactions of the Royal Society of Tropical Medicine and Hygiene 102(10): 990-996

Sawanyawisuth, K, Chindaprasirt, J, Senthong, V, Limpawattana, P, Auvichayapat, N, Tassniyom, S, Chotmongkol, V, Maleewong, W and Intapan, PM (2013). Clinical manifestations of Eosinophilic meningitis due to infection with Angiostrongylus cantonensis in children, Korean Journal of Parasitology 51(6): 735-738

Schultes, F (2014). Species summary for Eobania vermiculata, Animal Base, Gottingen University, available at http://www.animalbase.uni-goettingen.de/zooweb/servlet/AnimalBase/home/species?id=1367.

Schupbach, HU and Baur, B (2008a). Experimental evidence for a new transmission route in a parasitic mite and its mucus-dependent orientation towards the host snail, Parasitology 135(14): 1679-1684

Schupbach, HU and Baur, B (2008b). Parasitic mites influence fitness components of their host, the land snail Arianta arbustorum, Invertebrate Biology 127(3): 350-356

Schupbach, HU and Baur, B (2010a). Contact-based transmission models in terrestrial gastropod populations infected with parasitic mites, International Journal for Parasitology 40(9): 1045-1050

Schupbach, HU and Baur, B (2010b). Within- and Among-Family Variation in Parasite Load and Parasite-Induced Mortality in the Land Snail Arianta arbustorum, a Host of Parasitic Mites, Journal of Parasitology 96(4): 830-832

Schweiger F (2008). *Dicrocoelium dendriticum* infection in a patient with Crohn’s disease, Canadian Journal of Gastroenterology 22: 571-573

Segade, P, Crespo, C, Garcia, N, Garcia-Estevez, JM, Arias, C and Iglesias, R (2011). Brachylaima aspersae n. sp. (Digenea: Brachylaimidae) infecting farmed snails in NW Spain: Morphology, life cycle, pathology, and implications for heliculture, Veterinary Parasitology 175(3-4): 273-286

Segade, P, Garcia-Estevez, J, Arias, C and Iglesias, R (2013). Parasitic infections in mixed system-based heliculture farms: dynamics and key epidemiological factors, Parasitology 140(4): 482-497

Segade, P, Kher, CP, Lynn, DH and Iglesias, R (2009). Morphological and molecular characterization of renal ciliates infecting farmed snails in Spain, Parasitology 136(7): 771-782

Segade, P, Crespo, C, Garcia, N, Garcia-Estevez, JM et al. (2011). Brachylaima aspersae n. sp. (Digenea: Brachylaimidae) infecting farmed snails in NW Spain: Morphology, life cycle, pathology, and implications for heliculture, Veterinary Parasitology 175: 273-286

Senanayake, SN, Pryor, DS, Walker, J and Konecny, P (2003). First report of human angiostrongyliasis acquired in Sydney, Medical Journal of Australia 179(8): 430-431

Senlik, B, Cirak, VY and Tinar, R (2008). Field efficacy of two netobimin oral suspensions (5% and 15%) in sheep naturally infected with Dicrocoelium dendriticum, Small Ruminant Research 80(1-3): 104-106

Sherpa, S, Ansart, A, Madec, L, Martin, MC, Dréano, S and Guiller, A (2018). Redefining the biogeographical scenario of the land snail Cornu aspersum aspersum: natural spatial expansion and human-mediated dispersal in the Mediterranean basin, Molecular Phylogenetics and Evolution 120: 218-232

Smith, BJ (1992). Non-marine Mollusca, vol. 8, Houston, WWK (ed), Australian Government Public Service, Canberra, ISBN-10: 0 644 14598 6

Soares, CM, Hayashi, C and Cocito, IC (2002). Exigência de proteína para o escargot francês, Helix aspersa maxima em fase de crescimento, (Protein requirement for Helix aspersa maxima during the growth phase), Revista Brasileira de Zootecnia 31(2): 835-841

Soes, DM (2014). Een vondst van Eobania vermiculata in Wageningen (A find of Eobania vermiculata in Wageningen), Spirula 398: 84-85

Solem, A (1990). How many Hawaiian land snail species are left? And what can we do for them. Bishop Museum Occasional Papers, 30: 27-40

Spratt, DM (2005). Neuroangiostrongyliasis: disease in wildlife and humans, Microbiology Australia 26(2), CSIRO Publishing, doi: 10.1071/MA05063

Stanisic, J, Potter, D and Stanisic, I (2023). A Guide to Land Snails of Australia, CSIRO Publishing, ISBN: 9781486313525

Stockdale Walden, HD, Slapcinsky, JD, Roff, S, Mendieta Calle, J, Diaz Goodwin, Z, Stern, J, Corlett, R, Conway, J and McIntosh, A (2017). Geographic distribution of Angiostrongylus cantonensis in wild rats (Rattus rattus) and terrestrial snails in Florida, USA, PLoSOne 12(5): e0177910

Stockdale, PH and Hulland, TJ (1970). Pathogenesis, Route of Migration, and Development of Crenosoma-Vulpis in Dog, Pathologia Veterinaria 7(1): 28

Stokes, VL, Spratt, DM, Banks, PB, Pechc, RP and Williams, RL (2007). Occurrence of Angiostrongylus species (Nematoda) in populations of Rattus rattus and Rattus fuscipes in coastal forests of south-eastern Australia, Australian Journal of Zoology 55(3): 177-184

Tan, L and Grewal, PS (2001a). Infection behavior of the rhabditid nematode Phasmarhabditis hermaphrodita to the grey garden slug Deroceras reticulatum, Journal of Parasitology 87(6): 1349-1354

Tan, L and Grewal, PS (2001b). Pathogenicity of Moraxella osloensis, a bacterium associated with the nematode Phasmarhabditis hermaphrodita, to the slug Deroceras reticulatum, Applied and Environmental Microbiology 67(11): 5010-5016

Tarry, DW (1969). Dicrocoelium dendriticum: the life cycle in Britain, Journal of Helminthology 43(3-4): 403-416

Taylor, J (1914). Monograph of the land and freshwater Mollusca of the British Isles, doi: 10.5962/bhl.title.54424

Taylor, JW (1883). Life histories of British helices. Helix (Pomatia) aspersa Müll., Journal of Conchology 4: 88-105

Tebb, AI, Johnson, VS and Irwin, PJ (2007). Angiostrongylus vasorum (French heartworm) in a dog imported into Australia, Australian Veterinary Journal. 85(1-2): 23-28

Texas Invasive Species Institute (2014). Chocolate-band snail, Texas Invasive Species Institute, Huntsville (TX) USA, available at https://tsusinvasives.org/home/database/eobania-vermiculata

Thiengo, SC, Fernandez, MA, Torres, EJ, Coelho, PM and Lanfredi, RM (2008). First record of a nematode Metastrongyloidea (Aelurostrongylus abstrusus larvae) in Achatina (Lissachatina) fulica (Mollusca, Achatinidae) in Brazil, J Invertebr Pathol 98(1): 34-39

Thompson, R and Cheney, S (1996). Raising snails, NAL The Alternative Farming Systems Information Center (ed), US Department of Agriculture, available at: https://www.nal.usda.gov/programs/afsic

Tluste, C and Birkhofer, K (2023). The Roman snail (Gastropoda: Helicidae) is not a generalist herbivore, but shows food preferences for Urtica dioica and plant litter, Journal of Natural History 57: 758-770

Tolnai, Z, Szell, Z and Sreter, T (2015). Environmental determinants of the spatial distribution of Angiostrongylus vasorum, Crenosoma vulpis and Eucoleus aerophilus in Hungary, Veterinary Parasitology 207(3-4): 355-358

Traversa, D, Romanucci, M, Di Cesare, A, Malatesta, D, Cassini, R, Iorio, R, Seghetti, M and Della Salda, L (2014). Gross and histopathological changes associated with Aelurostrongylus abstrusus and Troglostrongylus brevior in a kitten, Veterinary Parasitology 201(1-2): 158-162

Trub, H and Ribi, G (1997). High fecundity of hybrids between the sympatric snail species Viviparus ater and V-contectus (Gastropoda: Prosobranchia), Heredity 79: 418-423

Tsai, HC, Lee, SS-J, Huang, C-K, Yen, C-M, Chen, E-R and Liu, Y-C (2004). Outbreak of eosinophilic meningitis associated with drinking raw vegetable juice in southern Taiwan, American Journal of Tropical Medicine and Hygiene 71(2): 222-226

Turk, FA and Phillips, SM (1946). A Monograph of the Slug Mite - Riccardoella-Limacum (Schrank), Proceedings of the Zoological Society of London 115(3-4): 448-472

Twomey, S (2017). Snail mail: Biosecurity officers intercept escargot from the Ukraine, The Weekly Times, 20 July 2017, available at https://www.weeklytimesnow.com.au/news/national/snail-mail-biosecurity-officers-intercept-escargots-from-the-ukraine/news-story/3760ac2f69694370e21ea531dad47f0d, accessed 20 March 2024.

Twyford, AD and Ennos, RA (2012). Next-generation hybridization and introgression, Heredity 108: 179-189

Ueshima, R, Okamoto, M and Saito, Y (2004). Eobania vermiculata, a land snail newly introduced into Japan, (in Japanese) Chiribotan 35(3): 71-74

USDA (2022). Snails and slugs [Online]. United States Department of Agriculture. Available: https://www.aphis.usda.gov/aphis/ourfocus/planthealth/sa\_import/sa\_permits/plant-pests/sa\_snails\_slugs/ct\_snails\_slugs [Accessed 05-06-2022].

Van As, JG and Basson, L (2004). Ciliophoran (Ciliophora) parasites of terrestrial gastropods, in GM Barker (ed) Natural Enemies of Terrestrial Molluscs CABI Publishing, Wallingford, UK, pp. 559–78

Varcasia, A, Tamponi, C, Brianti, E, Cabras, PA, Boi, R, Pipia, AP, Giannelli, A, Otranto, D and Scala, A (2014). Angiostrongylus chabaudi Biocca, 1957: a new parasite for domestic cats? Parasites and Vectors 7, Article number: 588

Wacker, A (2008). Impact of parasitic mite infection on a terrestrial snail, Invertebrate Reproduction and Development 51(2): 69-75

Waki, T, Ikezawa, H, Umeda, K and Shimano, S (2021). Natural history study on Riccardoella tokyoensis: life history with ontogeny and host distribution records, Experimental and Applied Acarology 83(1): 13-30

Wang, QP, Lai, DH, Zhu, XQ, Chen, XG and Lun, ZR (2008). Human angiostrongyliasis, The Lancet Infectious Diseases 8(10): 621-630

Wang, QP, Wu, ZD, Wei, J, Owen, RL and Lun, ZR (2012). Human Angiostrongylus cantonensis: an update, European Journal of Clinical Microbiology and Infectious Diseases 31(4): 389-395

Watt, A, Young, N, Haites, R, Dunse, K, Russell, D and Billman-Jacobe, H (2021). Intraspecies Variation in Tetrahymena rostrata, Microorganisms 9(10), doi: 10.3390/microorganisms9102100

Welch, JM and Pollard, E (1975). The exploitation of Helix pomatia L., Biological Conservation 8(2): 155-160

Welter-Schultes, F (2012). European non-marine molluscs, a guide for species identification, Planet Poster Editions, ISBN: 9783933922755

White-McLean, J (2022a). Archachatina marginata, Terrestrial Mollusc Tool, available at: https://idtools.org/mollusk/index.cfm?packageID=1178&entityID=8183

White-McLean, J (2022b). Otala spp., Teerestrial Mollusc Tool, available at: https://idtools.org/mollusk/index.cfm?packageID=1178&entityID=8183

Whitson, M (2005). Cepaea nemoralis (Gastropoda, Helicidae): the invited invader, Journal of the Kentucky Academy of Science 66(2): 82-88

Williams, AJ and Rae, R (2015). Susceptibility of the Giant African snail (Achatina fulica) exposed to the gastropod parasitic nematode Phasmarhabditis hermaphrodita, Journal of Invertebrate Pathology 127: 122-126

Williams, EH, Jr and Bunkley-Williams, L (2023). New country geographic record of chocolate-band snail, Eobania vermiculata Müller, for Malta, [epub ahead of print], accessed 20 March 2024

Wilson, MJ, Burch, G, Tourna, M, Aalders, LT and Barker, GM (2012). The potential of a New Zealand strain of Phasmarhabditis hermaphrodita for biological control of slugs, New Zealand Plant Protection 65: 161-165

Wilson, MJ, Coyne, C and Glen, DM (1998). Low temperatures suppress growth of the ciliate parasite, Tetrahymena rostrata, and pathogenicity to field slugs, Deroceras reticulatum, Biocontrol Science and Technology 8(1): 181-184

WOAH (2023). Terrestrial Animal Health Code, World Organisation of Animal Health, Paris, France.

Yanes, Y and Fernandez-Lopez-de-Pablo, J (2017). Calibration of the stable isotope composition and body size of the arid-dwelling land snail Sphincterochila candidissima, a climatic archive abundant in Mediterranean archaeological deposits, Holocene 27(6): 890-899

Yeung, N, Hayes, K and Cowie, R (2013). Effects of Washing Produce Contaminated with the Snail and Slug Hosts of Angiostrongylus cantonensis with Three Common Household Solutions, Hawai‘i Journal of Medicine & Public Health, 72(6, supp 2): 83-86

Yildirim, MZ, Kebapci, U and Gumus, BA( 2004). Edible snails (terrestrial) of Turkey, Turkish Journal of Zoology 28(4): 329-335

York, EM, Butler, CJ and Lord, WD (2014). Global decline in suitable habitat for Angiostrongylus ( = Parastrongylus) cantonensis: the role of climate change, PLoSOne 9(8), doi: 10.1371/journal.pone.0103831

Zak, LF, Yanez-Morales, MD, Alanis-Martinez, I and Gonzalez-Perez, E (2011). New hosts of 16Srl phytoplasma group associated with edible Opuntia ficus-indica crop and its pests in Mexico, African Journal of Microbiology Research 5: 910-918

Zhang, TY and Vdacny, P (2021). A discovery of two new Tetrahymena species parasitizing slugs and mussels: morphology and multi-gene phylogeny of T. foissneri sp. n. and T. unionis sp. n., Parasitology Research 120(7): 2595-2616

Ziȩtek, J, Ziomek, M and Wilczyńska, A (2019). Method of dissecting edible snails of the genus Cornu, Medycyna Weterynaryjna 75(10): 609-612

Zucaro, A, Forte, A, De Vico, G and Fierro, A (2016). Environmental loading of Italian semi-intensive snail farming system evaluated by means of life cycle assessment, Journal of Cleaner Production 125: 56-67