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Peroral Echinococcus multilocularis egg inoculation in Myodes glareolus, Mesocricetus auratus and Mus musculus (CD-1 IGS and C57BL/6j)

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Echinococcus multilocularis transmission predominantly occurs in Europe between the red fox (Vulpes vulpes) and various species of rodent intermediate hosts. We infected 3 species of rodent, Myodes glareolus (n = 47), Mesocricetus auratus (n = 11) and outbred Mus musculus (CD-1 IGS (n = 9)) with an E. multilocularis egg suspension that contained 100 eggs with viable oncospheres and performed post mortem examination 6, 8 (M. glareolus) and 10 weeks post inoculation (wpi). C57BL/6j mice (n = 4) were used as positive controls as they have been shown to exhibit macroscopic liver lesions 4 wpi. To the best of our knowledge, this is the first study to experimentally assess susceptibility in the ostensibly competent host M. glareolus. Lesions were only detected in 2 of 47 M. glareolus (4.3%) at 8 and 10 wpi and although both contained protoscolices (1675 at 8 wpi and 88 at 12 wpi) the low percentage of infected animals brings into question their role as transmitters of the parasite. Significant differences were observed between inbred and outbred mice with E. multilocularis infection in the former demonstrating increased establishment (p ≤ 0.0001) and growth (p ≤ 0.0001). No lesions were found in all 11 M. auratus.

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1. Introduction

Studies of the fox tapeworm, Echinococcus multilocularis have focused on naturally infected human and animal populations for disease mapping and risk assessment whilst experimental work is often conducted in mouse models intended for medical benefit (Dematteis et al., 2003; Eckert and Deplazes, 2004; Vuitton and Gottstein, 2010). Although experimental studies have identified profound differences in the susceptibility of the definitive carnivore hosts (Kapel et al., 2006) very little information exists on experimental infections of these tapeworms in their naturally occurring intermediate hosts, although such studies would clarify which host species play a key role in the transmission, why they are physiologically suited for parasite establishment and growth, and which minimum infectious doses would be required in natural settings across various relevant species. In addition to its ecological value, such data would constitute novel information for risk assessment and prevention.

In Europe, Arvicolidae species of rodents serve as intermediate hosts although the parasite is capable of more or less normal development in small mammal species from several families. Thus, the range of intermediate hosts that may be susceptible seems to be wider as compared to that of the definitive hosts. Even rodents not sympatric with Echinococcus multilocularis may establish metacestodes with protoscolices when experimentally inoculated (Thompson and Lymbery, 1995). That said, the growth and persistence of metacestodes varies between species and genus (Ohyayashi et al., 1971) and thus the geographical distribution of intermediate hosts species ought to affect transmission dynamics.

Experimental E. multilocularis infection in rodents can be achieved via various routes. Oral inoculation of E. multilocularis eggs is referred to as primary infection, whereas secondary inoculation involves the injection of metacestode homogenates or oncospheres intraperitoneally (IP), intrahepatically (IH), subcutaneously (SC) or intravenously (IV). Although secondary inoculation bypasses the early gastrointestinal exposure responsible for oncosphere activation and development, and thus provides a more narrow view of
2. Materials and methods

2.1. Experimental inoculation

The *E. multilocularis* eggs used for inoculation were isolated from worms in naturally infected foxes from the city of Zurich and the surrounding area, during the official Swiss hunting season. Eggs were tested for viability by the sodium hypochlorite (s-h) resistant test (Deplazes et al., 2005). In brief, the percentage viability of the eggs was determined to be the number of eggs with intact oncospheres after the s-h solution had been applied. Animals were anesthetized with isoflurane and the egg suspension containing approximately 100 viable *E. multilocularis* eggs was administered via gavage. This was calculated as follows: the total number of eggs per ml divided by the percentage viability (via s-h resistant test) to determine the percentage of viable eggs per ml. The number of viable eggs per ml was then used to calculate the volume of egg suspension that would contain 100 viable eggs. Animals were inoculated on different days with the s-h test conducted prior to each inoculation round and the volume of egg suspension adjusted accordingly. During the period of inoculation eggs were stored at 4 °C to maximise viability (Veit et al., 1995) in 1% penicillin-streptomycin solution.

Animals were housed in a safety facility (Biosafety Level 2++) approved by the Danish Working Environment Authority. Journal no. 20120014119/21 at the Department of Plant and Environmental Sciences (University of Copenhagen, Denmark), under experimental license no. 2012-15-2934-00150. All animals were imported under permission from the Danish Agrifish Agency (CVR: 29979812, No. 1013624417).

Four species/strains of rodent were experimentally inoculated with 100 viable *E. multilocularis* eggs:

- **Myodes glareolus**

  Female (*n* = 23) and male (*n* = 24) *M. glareolus* were obtained from Institute of Environmental Sciences, Jagiellonian University Kraków, Poland. All animals were 56 days old at inoculation (DAI). Animals were euthanized at 6 wpi (Female *n* = 4, Male *n* = 4), 8 wpi (Female *n* = 15, Male *n* = 16) and 10 wpi (Female *n* = 4, Male *n* = 4). The animals euthanized at 8 wpi were a control group of a separate study investigating *E. multilocularis* infection in relation to basal metabolic rate (BMI). The conditions that these rodents were exposed to (housing, nutrition, and *E. multilocularis* infection) were precisely the same as the 6 and 10 wpi animals and were thus included. All animals were inoculated between 20/04/2015 and 29/04/2015.

- **Mus musculus** (CD-1® IGS)

  Male (*n* = 3) and female (*n* = 6) CD-1 animals were obtained from Charles River Germany. Animals were 56 DAI. All animals were euthanized 6 wpi. The original study design was for 12 animals to be inoculated but a shortage of eggs meant that it was not possible for 3 male animals to be inoculated. Animals were inoculated 1 month after the *M. glareolus* and C57BL/6j mice on 26/05/2015. As such it was intended to also inoculate an additional two C57BL/6j mice. The shortage of eggs precluded this but in the interest of the 3 R’s (Russell et al., 1959) it was decided to proceed with the inoculations.

  - **Mesocricetus auratus**

    Female (*n* = 5) and male (*n* = 6) *Mesocricetus auratus* were obtained from Charles River France.

    Animals were 56 DAI. These animals were euthanized 6 wpi (Female *n* = 3, Male *n* = 3) and 10 wpi (Female *n* = 2, Male *n* = 3). Animals were inoculated 07/04/2014. These animals were inoculated in the same period as the Woolsey et al., 2015b study that demonstrated heavy *E. multilocularis* infection in *Microtus arvalis* (which were inoculated with the same egg suspension spanning dates before and after 07/04/2014).

    - **Mus musculus** (C57BL/6j)

      Female (*n* = 4) *M. musculus* were obtained from Charles River Germany in March 2015. Mice were 42 DAI. All animals were inoculated 20/04/2015.
2.2. Animal euthanasia

All animals were anesthetised with isoflurane and then euthanized with CO2 (gradual fill). Rodents were removed from the CO2 once they were observed to cease breathing. Cervical dislocation was performed subsequently as a precaution. Rodents were weighed and their livers were removed and lesions counted by eye and measured along their length (longest dimension). Protoscolecom quantification was conducted as described by (Burlet et al., 2011). Internal liver lesions were investigated and counted and measured by palpating the liver between two clear plastic sheets. Although more accurate determination of internal lesions would have been obtained by slicing the organ, this would have reduced the accuracy of protoscolecom enumeration, which was deemed more important to understanding the transmission potential than lesion number (as this parameter is a more robust determinant of the animal’s potential to transmit the parasite). All other organs (with the exception of the brain) were inspected for any metacestode growth.

2.3. Statistical analysis

The number of metacestodes, their size and species susceptibility (defined here as the number of all inoculated animals per species displaying E. multilocularis metacestodes in the liver), was run in a multiple linear regression analyses against species and sex at 6 wpi. Single rodents were found infected at 8 and 10 wpi respectively, negating statistical analysis at these time points. All analyses was conducted in R (R Core Team, 2014) and differences considered significant when \( p < 0.05 \).

3. Results

At 6 wpi 4/4 C57BL/6j and 5/9 CD-1 (3 female, 2 male) mice had developed visible E. multilocularis infections. No metacestodes were observed in M. auratus or M. glareolus at 6 wpi. Initial C57BL/6j susceptibility was significantly greater than M. auratus \( (p < 0.0001) \), M. glareolus \( (p < 0.0001) \) and CD-1 \( (p < 0.0001) \). No rodents exhibited protoscolecis at 6 wpi. Sex was not found to be significant (Fig 1).

No CD-1 outbred mice harboured metacestodes larger than 1 mm. Three C57BL/6j harboured metacestodes larger than this \( (>1 \text{ mm} < 2 \text{ mm}) \) and this was significant \( (p < 0.0001) \).

Only 2/47 M. glareolus and no M. auratus developed metacestodes. At 8 wpi, one male M. glareolus developed an infection with a single mass of metacestode material \( (1.1 \times 1.2 \times 1 \text{ cm}) \) located in the liver with 1675 protoscolecis. At 10 wpi, one female M. glareolus was found infected with a single metacestode mass in the liver \( (2 \times 0.9 \times 0.6 \text{ cm}) \) with 88 protoscolecis.

4. Discussion

The infection rate of 4.3% in M. glareolus is of special interest considering that under very similar experimental conditions, M. arvalis and Microtus agrestis both exhibited much greater susceptibility to the parasite (95.2% and 88.9% of animals developing infection respectively) and many more metacestodes per rodent \( (27.5 \pm 6.63 \text{ S.D.} \) and \( 23.9 \pm 15.3 \text{ S.D.} \) respectively) at 6 wpi (Woolsey et al., 2015a, 2015b). Comparisons of these data with the current study can be seen in Figs. 2 and 3. Two M. glareolus were infected and both harboured protoscolecis, so the species is clearly capable of transmitting the parasite and in the literature there are numerous references to this species being infected with protoscolecis in Europe e.g. (Eckert, 1998; Osterman Lind et al., 2011; Liccioli et al., 2013). Prevalence of the parasite in M. glareolus has been observed as high, 10.3% (6/58) (Reperant et al., 2009) but is generally reported as low with prevalence values of 4.3% (1/23) (Hanosset et al., 2008) and 2.4% (2/83, but with 108 000 protoscolecis in one animal) (Stieger et al., 2002) in central Europe. It would be prudent however to consider this limited susceptibility in relation to Microtus spp. in assessing the competence of this species in parasite transmission. Limited susceptibility in this species is verified by the 100% infection rate in the C57BL/6j mice used as positive controls. The lack of susceptibility in this species is of even greater interest considering that in Japan, the transmission of E. multilocularis is largely based upon the grey-sided vole (Myodes rufocanus) (Saitoh and Takahashi, 1998) suggesting that susceptibility to this parasite is perhaps not determined by genus. It would have been desirable to see data on lesion size in M. glareolus at 6 wpi in order to determine some idea regarding the metacestode rate of growth but unfortunately no animals had lesions at this endpoint. To ensure infections at all time points when working with this species much larger cohort sizes will be needed.

Lending support to the limited role played by this species in parasite transmission is the finding that variations in E. multilocularis prevalence in foxes was not associated with M. glareolus but was with Microtus spp. in the Swiss canton of Grisons, even though M. glareolus constituted the second highest prey item in the study (Tanner et al., 2006). Considering the role of intermediate host populations being highly important to variations in definitive host prevalence (Raoul et al., 2015), this is of great interest.

The C57BL/6j mice were significantly more susceptible to the parasite than the CD-1 outbred mice. C57BL/6j mice were not found to harbour protoscolecis at 16 wpi after oral infection with 200 eggs (Matsumoto et al., 2010) and thus, considering the reduced susceptibility of CD-1 to the inbred strain it is unlikely that the outbred mice would have produced protoscolecis either. Although there are a limited number of examples of this species harbouring the infection in the wild e.g. (Leiby et al., 1970; Leiby and Kritsky, 1972; Pětavy et al., 1990), these data are highly indicative of M. musculus.
not playing any significant role in *E. multilocularis* transmission. Although outbred, these CD-1 mice obtained from Charles River originate from a very small number of animals (2 males and 7 females), which represents a significant bottleneck, and thus caution should be exercised when interpreting these results as representative of wild type *M. musculus*. Furthermore, only 4 C57BL/6j mice were used in this study but their inbred nature should mitigate the small sample size. In previous studies using the same methodology similar infection dynamics were observed in this strain (Woolsey et al., 2015a, 2015b).

The CD-1 mice were included in this study as a representative of the family Muridae. Based on trapping studies aiming to determine *E. multilocularis* prevalence, it is clear that *Apodemus* spp. are caught a great deal more frequently than *M. musculus* (Takeuchi-Storm et al., 2015). This is due to habitat overlap between this species and species widely regarded to be key drivers of *E. multilocularis* transmission in Europe whereas *M. musculus* are found in villages, away from these species (Giraudoux et al., 2003). Due to this, *Apodemus* spp. would constitute a much better representative of the Muridae family. That said, if protoscolices had been found in mice in this study it would have made this species potentially important considering the urbanisation of foxes (Deplazes et al., 2004).

It should be noted that the CD-1 mice were inoculated approximately 1 month after the *M. glareolus* and C57BL/6j mice and although the s-h resistance test was conducted prior to inoculation of the outbred mice, recent studies have demonstrated that this primarily tests the maturity of the eggs, not necessarily their infectivity (Federer et al., 2015) and this may have declined in the time between inoculations. As such it would have been ideal to have an additional group of C57BL/6j mice inoculated at the same time for a more robust comparison of susceptibility between these two strains and results need to be considered in this context. It does seem however that even in highly susceptible species, inoculated with 100–1000 eggs, maximum establishment in the liver is < 35 (Woolsey et al., 2015a, 2015b) and thus the egg dose used in this study was much higher than any possible establishment. It therefore seems likely that the CD-1 mice received enough infective eggs for a robust comparison as eggs have been demonstrated to survive for 78 days in summer conditions (Veit et al., 1995) far less favourable than the ideal storage conditions of the eggs in this study.

The infections observed in *M. glareolus* and both strains of mouse are indicative of two distinct mechanisms pertaining to *E. multilocularis* infection dynamics in the host; i) establishment of the oncosphere and ii) its persistence and subsequent growth. Clearly, the establishment of the oncosphere in the mouse is much more successful than in *M. glareolus* however, growth of the metacestode if it does establish in this latter species is far more prolific, with protoscolices appearing 8 wpi. This may permit the distinction of two types of host in relation to *E. multilocularis* infection, those that have the physiological/morphological/immunological profile to permit establishment of the oncosphere but not its proliferation in the liver and those that are refractory to oncosphere invasion but permit rapid growth if it does occur. *M. arvalis* and *M. agrestis*, which along with high metacestode establishment and growth (Woolsey et al., 2015a, 2015b) would clearly represent a third type which posses both of these ‘attributes’. Of course, there may be a range of issues that result in the lack of establishment of the oncosphere in the liver such as the physio/chemical conditions in the stomach or the morphology of the small intestine, however the two positive *M. glareolus* in this study would indicate that such aspects are not relevant in this species.

The mechanisms behind the increased resistance of the outbred...
strain to *E. multilocularis* infection are impossible to establish with the data obtained in this study. In order to elucidate such mechanisms, assessment of various immune parameters, such as varying cytokine profiles, would have been useful. Investigating such aspects in ecologically relevant Cricetidae species in the manner that has been achieved with various mouse strains (Vuitton and Gottstein, 2010) is a clear avenue for prospective study. If the cause does lie in differing immune mechanisms it would not be surprising that outbred mice demonstrate greater resistance to infection considering that inbreeding is thought to reduce host immunity as a result of decreased heterozygosity and inbreeding increases homozygosity (Carrington et al., 1999). Indeed, it was found in song sparrows (*Melospiza melodia*) that a greater inbreeding coefficient decreased the effectiveness of cell mediated immune response (Reid et al., 2003), precisely the type associated with resistance to *E. multilocularis* (Emery et al., 1996, 1997; Vuitton and Gottstein, 2010). It should be noted however that in the context of oncosphere establishment and metacestode growth, it is the latter that has been demonstrated to be inhibited by cell mediated immune responses and the aspects governing establishment in the liver are far less investigated (not least due to the lack of studies utilising primary infection).

*Mesocricetus auratus* appears not to be of relevance in transmission as they were not susceptible to *E. multilocularis*, but it is not possible to determine whether this is a consequence of the oncosphere failing to establish in the liver or another mechanism by which the parasite was prevented from reaching the organ. Furthermore, it should be considered that with a greater number of animals an infection might have been observed. Considering *M. glareolus* in this study this is particularly relevant with only 2 infections resulting from 47 inoculations.

In conclusion, clear differences between rodent susceptibility to *E. multilocularis* transmission have been demonstrated. This methodology has the potential to shed light on parasite transmission dynamics that cannot be obtained through trapping studies or secondary infections in non-ecologically relevant species. In Europe, there ought to be a clear emphasis on conducting such studies on *Arvicola* and *Apodemus* spp. and assessment of cytokine and endocrine profiles in all species aiming to determine why these species manifest such differences in response to the parasite. Furthermore, early stages of parasite-host interaction are still unclear and this present model could be used to compare e.g. intestinal dynamics in susceptible/refractory species.

**Conflict of interest**

The authors declare no conflict of interest.

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