



Unexpected sensitivity of the highly invasive spider *Mermessus trilobatus* to soil disturbance in grasslands

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Abstract The dwarf spider *Mermessus trilobatus* (Araneae: Linyphiidae), native to North America, has expanded its range over large parts of Europe within less than fifty years. It is notable for occurring in a wide range of mostly agricultural habitats, while most other invasive spiders in Europe are associated with human buildings. As in other invasive invertebrates and plants, the tremendous colonisation success of *Mermessus trilobatus* might be related to anthropogenic habitat disturbance. Here we aim to test if the

invasion success of *Mermessus trilobatus* in Europe is associated with high tolerance towards soil disturbance. We sampled spiders from eight grasslands experimentally disturbed with superficial soil tillage and eight undisturbed grasslands without tillage. Opposite to our expectation, *Mermessus trilobatus* densities decrease sharply with soil disturbance. This is in contrast to several native species such as *Oedothorax apicatus*, which becomes more abundant in the fields after superficial soil tillage. Our study suggests that invasion success of *Mermessus trilobatus* is not connected to a ruderal strategy. The ecological and evolutionary processes behind

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colonisation success of *Mermessus trilobatus* need to be further investigated.

Keywords Araneae · Disturbance · Habitat preference · Invasibility · Linyphiidae · *Oedothorax apicatus*

Introduction

Despite their essential role in ecosystems (Michalko et al. 2019; Nyffeler and Birkhofer 2017), invasions by spiders have only recently started to receive scientific attention (Nentwig 2015). One of the most widespread alien spider species in Europe is the North American dwarf spider *Mermessus trilobatus* (Araneae: Linyphiidae), formerly known as *Eperigone trilobata* (Milledge 1987; Nentwig 2015; Nentwig and Kobelt 2010; Schmidt et al. 2008). It was first detected in Europe in the late 1970s in the Upper Rhine valley near Karlsruhe in South-West Germany (Dumpert and Platen 1985). The species has undergone a largely concentric range expansion and has been recorded in numerous other countries since 1990, such as Austria, Belgium, Croatia, Czech Republic, France, Great Britain, Hungary, Italy, the Netherlands, Poland, Slovakia, Slovenia, Switzerland, and Ukraine (Hirna 2017). To our knowledge, this rapid spread makes *M. trilobatus* currently the most invasive (sensu Richardson et al. 2000) spider in Europe.

Mermessus trilobatus has mostly been collected in open habitats within agricultural landscapes and can be among the most abundant spider species there (Schmidt et al. 2008). Its occurrence in agricultural lands suggests that the invasion success of *M. trilobatus* could be based on a ruderal strategy, whereby it would benefit from reduced competition with native species in disturbed habitats (Elton 1958). Lab experiments confirm that *M. trilobatus* is a poor competitor due to its slightly smaller body size compared to native spiders living in the same habitats (Eichenberger et al. 2009). Furthermore, *M. trilobatus* might benefit from post-disturbance resource influxes to the habitat (e.g. from decomposing plant material), or from altered structure and habitat opening (Lear et al. 2020).

Here we aim to test if *Mermessus trilobatus* benefits from soil disturbance in one of its preferred habitats, perennial hay meadows. We compare its abundance to

native linyphiid spiders in replicated experimentally disturbed and control grassland sites, expecting that *M. trilobatus* abundances increase after disturbance.

Methods

Field characteristics and sampling

The experiment was conducted in 16 permanent hay meadows in the Canton of Bern, Switzerland, in 2008 (Table S2 in supplementary material). All grassland sites belonged to the same community type and were situated 0.5–50 km from each other. The treatments were randomly assigned to the 16 grassland sites. In each grassland, one plot of 240 m² was used. Eight plots were superficially tilled with a rotary tiller (Figure S1 and Figure S2 in supplementary material) in the first half of April, creating soil and ground surface disturbance (disturbed fields). The vegetation was left to decay. The other eight grasslands served as a control and were mown instead of tilled also in the first half of April, and the mown grass was left to decay (undisturbed fields in the following). Disturbance with the rotary tiller had profound effects, killing part of the vegetation and loosening the soil surface, but still leaving sufficient perennial plants alive for continuous vegetation cover. By contrast, mowing only shortened the vegetation at an early growing stage, which is common practice in this grassland type and was required for a plant introduction experiment reported elsewhere (Kempel et al. 2013), but did not affect the ground surface. The sites received the same set of plant species with variable propagule pressure at the beginning of May for the plant introduction experiment. Most adults of *M. trilobatus* are found in summer (Arachnologische Gesellschaft 2020). Thus, the spiders were sampled in late June to early July, 1–2 months after the disturbance event, which meant that the immediate impact was over, but that the vegetation was still different between disturbed and undisturbed sites. The sown plants were hardly visible at the time of sampling and were therefore unlikely to have affected the spiders in the field. We sampled spiders with a vacuum sampler with an 11 cm diameter nozzle (modified STIHL SH85 blower; Stihl, Waiblingen, Germany). It was lowered 150 times per meadow, each time over a different location, resulting in a sampled area of 1.4 m² per meadow, except for

two undisturbed plots with 200 times each, or 1.9 m² (Table S2 in supplementary material). Densities per square metre were analysed to account for this difference in sampling effort. By lowering the nozzle until just above the ground, both the vegetation and ground surface was sampled (Sanders and Entling 2011). All samples were transferred in ethanol (70%) for further identification in the lab.

Study species

All spiders were identified to species level with the aid of a stereomicroscope (Table S1 in supplementary material). Linyphiid species were identified using “The Spiders of Great Britain and Ireland” by Roberts (1987) and “Spiders of Europe” online key (Nentwig et al. 2020). The non-linyphiid spiders were identified with “Collins Field Guide: Spiders of Great Britain and Northern Europe” by Roberts (1995), names following the World Spider Catalog (Nentwig et al. 2020). To reduce the effects of rare species, we used only species present in at least half of the plots in each treatment group (at least 4). We ended up with eight linyphiids: the invasive species *Mermessus trilobatus* and seven native species, namely, *Agyneia rurestris*, *Erigone atra*, *Erigone dentipalpis*, *Oedothorax apicatus*, *Oedothorax fuscus*, *Pelecopsis parallela* and *Tenuiphantes tenuis*. These are all small (< 3 mm) spider species that live among vegetation close to the ground surface. They represent a gradient in hunting strategies, with *A. rurestris*, *M. trilobatus* and *T. tenuis* being obligatory builders of horizontal sheet webs; *E. atra*, *E. dentipalpis* and *P. parallela* capturing prey both within and outside webs; and *O. apicatus* and *O. fuscus* being free hunters (ME, personal observation; Cordoso et al. 2011).

Statistical analysis

We calculated the number of individuals per square meter in each field. We modelled the number of individuals per spider species fitting a multivariate generalized linear model (MvGLM) from *mvabund* package in R 3.6.1 (R Core Team 2019; Wang et al. 2012). We used a negative binomial distribution as the most flexible and appropriate for count data (O’Hara and Kotze 2010). We analysed soil disturbance (disturbed, undisturbed) as a fixed predictor with the “*anova.manyglm*” function with correction for

multiple tests using the “*p.uni*” function (test = “LR”) with 100,000 permutations.

Results

Mermessus trilobatus individuals were found in half of the disturbed and in 7 out of 8 undisturbed sites. Community composition of spiders was significantly affected by soil disturbance (Dev = 22.71; *P* = 0.02). Opposite to our expectations, *M. trilobatus* densities were reduced almost 90% after disturbance (Dev = 9.451; *P* = 0.003), and none of the native species showed a comparable decline (Fig. 1). In undisturbed grasslands, *M. trilobatus* was the most abundant spider together with *Erigone dentipalpis*. Densities of *O. apicatus* were approx. 13-fold higher in disturbed than in undisturbed meadows (Dev = 5.099; *P* = 0.03). The other six native linyphiids showed no significant response to the disturbance treatment (Fig. 1).

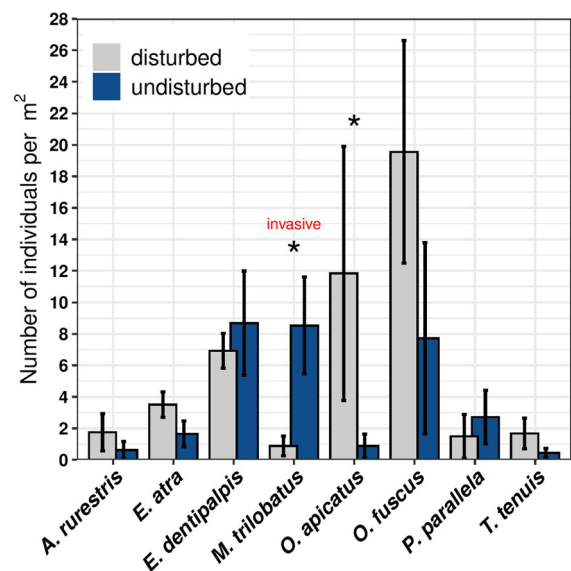


Fig. 1 Response of spiders to soil disturbance. The number of individuals per 1 m² for all 8 spider species are illustrated. Spiders were sampled from 8 meadows after soil tillage (disturbed) and 8 meadows without tillage (undisturbed). Mean ± SE are presented, with significant differences marked with asterisk. Invasive species: *Mermessus trilobatus* (Dev = 9.451; *P* = 0.003); Native species: *Agyneia rurestris* (Dev = 0.968; *P* = 0.39), *Erigone atra* (Dev = 2.909; *P* = 0.12), *Erigone dentipalpis* (Dev = 0.283; *P* = 0.61), *Oedothorax apicatus* (Dev = 5.099; *P* = 0.03), *Oedothorax fuscus* (Dev = 1.127; *P* = 0.21), *Pelecopsis parallela* (Dev = 0.194; *P* = 0.64), and *Tenuiphantes tenuis* (Dev = 2.681; *P* = 0.22)

Discussion

Opposite to our expectations, our results suggest that the highly invasive spider *M. trilobatus* is more sensitive to soil disturbance than sympatric native European species. One of the native species, *O. apicatus*, even increases in abundance in the disturbed grassland sites. The increase of *O. apicatus* in disturbed grassland does not come as a surprise since they are adapted to live and even overwinter in annual crop fields with little vegetation cover (Mestre et al. 2018; Schmidt and Tscharrntke 2005). Furthermore, since mainly cursorial spiders show avoidance behaviour towards intraguild predators like ants (Mestre et al. 2020), *O. apicatus* may benefit from soil disturbance which destroys ant nests. By contrast, the webs of *M. trilobatus* can protect them against predators (Blackledge et al. 2003). *Mermessus trilobatus* uses webs for prey capture (ME, personal observation). The destruction of these webs during disturbance represents a disadvantage. However, native obligatory web builders like *A. rurestris* and *T. tenuis* (ME, personal observation; Cardoso et al. 2011) are not sensitive to disturbance, so the hunting mode cannot fully explain the decline of *M. trilobatus*. Thus, other factors such as microclimate, prey availability, or competition with the better disturbance-adapted native species (Eichenberger et al. 2009) are potential mechanisms behind the sensitivity of *M. trilobatus* to disturbance but require further study. From an evolutionary perspective, the reduced adaptation of *M. trilobatus* to soil disturbance compared to European species may be related to the much more recent spread of annual cropping systems in its native North American range, and thus reduced time to co-evolve with intensive land-use.

Irrespective of the mechanisms, the decline of *M. trilobatus* after disturbance raises the question of how it can nevertheless be so successful in European agricultural landscapes. Importantly, the short-term decline of *M. trilobatus* observed here should not be mistaken for a general avoidance of disturbed habitats. Most (86%) of the specimens in Germany have been recorded from grasslands, which depend on regular disturbance of the vegetation layer, i.e. mowing or grazing, in this climatic region. *Mermessus trilobatus* is rarely found both in completely undisturbed habitats such as forests (2.4% of individuals), but also in highly disturbed annual crops (1.3% of individuals)

(Arachnologische Gesellschaft 2020). This avoidance of habitats with cultivated soil is in line with the results found in the current experiment.

Possible ecological mechanisms for the success of this species in Europe include the enemy release hypothesis (Roy et al. 2011). Reduced pressure by native predators, parasitoids and pathogens enhances the survival of alien relative to native species. Such potential advantages could be straightforwardly tested experimentally using important enemies of linyphiid spiders such as ants (Hymenoptera: Formicidae) or wolf spiders (Araneae: Lycosidae; Nyffeler 1999). Lastly, it is possible that *M. trilobatus* can spread in its invasive range without being limited by ecological interactions with native species, just as high numbers of native linyphiid spiders are able to coexist in the same habitat.

In summary, our study shows that in contrast to the theory of disturbance-mediated invasion success, *M. trilobatus* does not benefit from soil disturbance. Thus, other potential mechanisms behind its colonisation success remain to be studied, notably its potentially higher reproduction or reduced sensitivity to predators, parasitoids, or pathogens. Given the increasing dominance of invasive spiders in many agricultural (e.g. Hogg et al. 2010) and natural habitats (e.g. Pétiillon et al. 2020) across the globe, further studies on their ecology are strongly encouraged.

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Authors' contributions AK, ME and MvK conceived the idea; AK and MvK designed and performed the disturbance experiment; ME collected the data; NN analysed the data and led the writing of the manuscript. All authors contributed critically to the drafts and gave final approval for publication.

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Availability of data and material The dataset generated during and/or analysed during the current study is available in the Figshare repository, [<https://doi.org/10.6084/m9.figshare.12726998.v1>]

Compliance with ethical standard

Conflicts of interest The authors declare no competing interests.

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References

- Arachnologische Gesellschaft (2020) Atlas of the European Arachnids, accessed at <https://atlas.arages.de>, on 09.04.2020
- Blackledge TA, Coddington JA, Gillespie RG (2003) Are three-dimensional spider webs defensive adaptations? *Ecol Lett* 6:13–18. <https://doi.org/10.1046/j.1461-0248.2003.00384.x>
- Cardoso P, Pekár S, Jocqué R, Coddington JA (2011) Global patterns of guild composition and functional diversity of spiders. *PLoS ONE* 6:e21710. <https://doi.org/10.1371/journal.pone.0021710>
- Core Team R (2019) R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna
- Dumpert K, Platen R (1985) Zur Biologie eines Buchenwaldbodens. 4. Die Spinnenfauna. *Carolinea* 42:75–106
- Eichenberger B, Siegenthaler E, Schmidt-Entling MH (2009) Body size determines the outcome of competition for webs among alien and native sheetweb spiders (Araneae: Linyphiidae). *Ecol Entomol* 34:363–368. <https://doi.org/10.1111/j.1365-2311.2008.01085.x>
- Elton CS (1958) The ecology of invasions by animals and plants. Springer, Boston. <https://doi.org/10.1007/978-1-4899-7214-9>
- Hirna A (2017) First record of the alien spider species *Mermessus trilobatus* (Araneae: Linyphiidae) in Ukraine. *Arachnol Mitt* 54:41–43. <https://doi.org/10.5431/aramit5409>
- Hogg BN, Gillespie RG, Daane KM (2010) Regional patterns in the invasion success of Cheiracanthium spiders (Miturgidae) in vineyard ecosystems. *Biol Invasions* 12:2499–2508. <https://doi.org/10.1007/s10530-009-9659-1>
- Kempel A, Chrobock T, Fischer M, Rohr RP, van Kleunen M (2013) Determinants of plant establishment success in a multispecies introduction experiment with native and alien species. *PNAS* 110:12727–12732. <https://doi.org/10.1073/pnas.1300481110>
- Lear L, Hesse E, Shea K, Buckling A (2020) Disentangling the mechanisms underpinning disturbance-mediated invasion. *Proc R Soc B* 287:20192415. <https://doi.org/10.1098/rspb.2019.2415>
- Mestre L, Schirmel J, Hetz J, Kolb S, Pfister SC, Amato M, Sutter L, Jeanneret P, Albrecht M, Entling MH (2018) Both woody and herbaceous semi-natural habitats are essential for spider overwintering in European farmland. *Agric Ecosyst Environ* 267:141–146. <https://doi.org/10.1016/j.agee.2018.08.018>
- Mestre L, Narimanov N, Menzel F, Entling MH (2020) Non-consumptive effects between predators depend on the foraging mode of intraguild prey. *J Anim Ecol* 89:1690–1700. <https://doi.org/10.1111/1365-2656.13224>
- Michalko R, Pekár S, Dul'a M, Entling MH (2019) Global patterns in the biocontrol efficacy of spiders: a meta-analysis. *Glob Ecol Biogeogr* 28:1366–1378. <https://doi.org/10.1111/geb.12927>
- Millidge AF (1987) The erigonine spiders of North America. Part 8. The Genus *Eperigone* Crosby and Bishop (Araneae, Linyphiidae). *Am Mus Novit* 2885:1–75
- Nentwig W (2015) Introduction, establishment rate, pathways and impact of spiders alien to Europe. *Biol Invasions* 17:2757–2778. <https://doi.org/10.1007/s10530-015-0912-5>
- Nentwig W, Kobelt M (2010) Spiders (Araneae). Chapter 7.3. *BioRisk* 4:131–147. <https://doi.org/10.3897/biorisk.4.48>
- Nentwig W, Blick T, Bosmans R, Gloor D, Hänggi A, Kropf C (2020) Spiders of Europe. Online at <https://www.araneae.nmbe.ch>, accessed July 2008
- Nyffeler M (1999) Prey selection of spiders in the field. *J Arachnol* 27:317–324
- Nyffeler M, Birkhofer K (2017) An estimated 400–800 million tons of prey are annually killed by the global spider community. *Sci Nat* 104:30. <https://doi.org/10.1007/s00114-017-1440-1>
- O'Hara RB, Kotze DJ (2010) Do not log-transform count data. *Methods Ecol Evol* 1:118–122. [https://doi.org/10.1111/j.2041-210X.2010.00021.x@10.1111/\(ISSN\)2041-210X](https://doi.org/10.1111/j.2041-210X.2010.00021.x@10.1111/(ISSN)2041-210X). TOPMETHODS
- Pétillon J, Privet K, Roderick GK, Gillespie RG, Price DK (2020) Non-native spiders change assemblages of Hawaiian forest fragment kipuka over space and time. *NeoBiota* 55:1–9. <https://doi.org/10.3897/neobiota.55.48498>
- Richardson DM, Pyšek P, Rejmánek M, Barbour MG, Panetta FD, West CJ (2000) Naturalization and invasion of alien plants: concepts and definitions. *Divers Distrib* 6:93–107. <https://doi.org/10.1046/j.1472-4642.2000.00083.x>
- Roberts MJ (1987) The Spiders of Great Britain and Ireland: Linyphiidae and check list. Harley Books, United Kingdom
- Roberts MJ (1995) Collins Field Guide: Spiders of Great Britain and Northern Europe. Later prt. edition. ed. HarperCollins, New York
- Roy HE, Lawson Handley L-J, Schönrogge K, Poland RL, Purse BV (2011) Can the enemy release hypothesis explain the success of invasive alien predators and parasitoids? *Bio-control* 56:451–468. <https://doi.org/10.1007/s10526-011-9349-7>

- Sanders D, Entling MH (2011) Large variation of suction sampling efficiency depending on arthropod groups, species traits, and habitat properties: variation in arthropod suction sampling. *Entomol Exp Appl* 138:234–243. <https://doi.org/10.1111/j.1570-7458.2010.01094.x>
- Schmidt MH, Tscharrntke T (2005) The role of perennial habitats for Central European farmland spiders. *Agric Ecosyst & Environ* 105:235–242. <https://doi.org/10.1016/j.agee.2004.03.009>
- Schmidt MH, Rocker S, Hanafi J, Gigon A (2008) Rotational fallows as overwintering habitat for grassland arthropods: the case of spiders in fen meadows. *Biodivers Conserv* 17:3003–3012. <https://doi.org/10.1007/s10531-008-9412-6>
- Wang Y, Naumann U, Wright ST, Warton DI (2012) mvabund - an R package for model-based analysis of multivariate abundance data. *Methods Ecol Evol* 3:471–474. <https://doi.org/10.1111/j.2041-210X.2012.00190.x>

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