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Comparative analysis of experimental testing procedures for the elicitation of rescue actions in ants
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Abstract

Rescue behaviour is observed when one individual provides help to another individual in danger. Most reports of rescue behaviour concern ants (Formicidae), in which workers rescue each other from various types of entrapment. Many of these entrapment situations can be simulated in the laboratory using an entrapment bioassay, in which ants confront a single endangered nest mate entrapped on a sandy arena by means of an artificial snare. Here, we compared numerous characteristics of rescue actions (contact between individuals, digging around the entrapped individual, pulling at its body parts, transport of the sand covering it and biting the snare entrapping it) in \textit{Formica cinerea} ants. We performed entrapment tests in the field and in the laboratory, with the latter under varying conditions in terms of the number of ants potentially engaged in rescue actions and the arena substrate (marked or unmarked by ants’ pheromones). Rescue actions were more probable and pronounced in the field than in the laboratory, regardless of the type of test. Moreover, different test types in the laboratory yielded inconsistent results and showed noteworthy variability depending on the tested characteristic of rescue. Our results illustrate the specifics of ant rescue actions elicited in the natural setting, which is especially important considering the scarcity of field data. Furthermore, our results underline the challenges related to the comparison of results from different types of entrapment tests reported in the available literature. Additionally, our study shows how animal behaviour differs in differing experimental setups used to answer the same questions.

Key words: ants, entrapment bioassay, \textit{Formica cinerea}, rescue behaviour, animal behaviour

Altruistic behaviour in animals draws high scientific interest (Pennisi 2005). A form of altruistic behaviour, rescue, is a behaviour of one individual helping a related individual in grave danger, not bringing any direct reward to the rescuer, except for the benefits from reciprocal altruism and kin selection (Nowbahari and Hollis 2010). Numerous cases of nonhuman animals rescuing each other can be found in the literature. In capuchin monkeys, males rescue females from harassment by distracting the aggressors (Vogel and Fuentes-Jiménez 2006). In bottlenose dolphins, injured members of
the group receive help in reaching the water surface (Siebenaler and Caldwell 1956). In birds, flock members remove flight preventing bird-catcher tree seeds from each other’s bodies (Hammers and Brouwer 2017). By far, the most commonly studied in this context among all vertebrates are rats due to the vast opportunities to study their rescue behaviour in terms of ensuring appropriate laboratory conditions (Bartal et al. 2011, Vasconcelos et al. 2012, Silberberg et al. 2014, Silva et al. 2020). Notably, however, most reports of rescue behaviour concern animals with a much less complex nervous system than that in vertebrates, namely, ants (e.g., Czechowski et al. 2002, Nowbahari et al. 2009, Taylor et al. 2013, Miler 2016). However, the similarities and differences in ant rescue actions observed under different field and laboratory setups are entirely unstudied. Using several well-known tests to examine the same behaviour may help uncover the role of context in animal behaviour (Watanabe 2012, Scharf and Martin 2013). It has been suggested, for example, that relatively similar tests devoted to the study of boldness in crabs lead to different responses of tested individuals (Watanabe 2012). Moreover, as showed by Scharf and Martin (2013), frequencies of same-sex mating in insects and arachnids are higher in the laboratory conditions than in the field. Feeding strategy in desert gerbils, and their response to predation risk, also differs strikingly between the natural and laboratory contexts (Ovadia et al. 2001). Plausibly, the response of ants to a risky situation, such as another individual in need of rescue, can also vary between contexts.

The simplest forms of ant rescue behaviour, such as helping a nest mate move from under pebbles, were documented for the first time by Belt (1874). Afterwards, similar observations of ants’ digging behaviour around individuals covered by sand were reported several times by different authors (Lafleur 1940, Blum and Warter 1966, Hangartner 1969, Markl 1965, Spangler 1968, Wilson 1958). More advanced forms of rescue behaviour were found by Czechowski et al. (2002) in sand-dwelling ants (Formica sanguinea, Formica fusca and Formica cinerea) trying to set imperilled individuals free from traps of predatory antlions. Other recent examples of risky rescue operations included those described in Matabele ants Megaponera analis saving wounded nest mates after direct confrontation with termites (Frank et al. 2017) and in weaver ants Oecophylla smaragdina and harvester ants Veromessor pergandei trying to save endangered individuals who became stuck on spider webs (Uy et al. 2018, Kwapich and Hölldobler 2019). Rescue may be considerably more widespread in nature than these reports demonstrate (Nowbahari and Hollis 2010).

Ant societies are thought to largely dominate the Earth (Hölldobler and Wilson 1990). One of the essential components of their success, beyond advanced nest construction, foraging behaviour and complex division of labour, seems to be rescue behaviour (Andras et al. 2020). The potential meaning of this altruistic behaviour to the overall success of the colony is suggested by two previous studies. As shown by Frank et al. (2017), rescue behaviour of Megaponera analis ants prevents mortality in 32% of cases and allows for up to 28.7% higher colony size. In a study conducted by Kwapich and Hölldobler (2019), Veromessor pergandei ants not only saved endangered nest mates that became stuck in spider webs but also removed the webs to reduce future risks and ensure better foraging performance, which largely determined the success of the colony. Further studies devoted to rescue in ant species with well-known ecology will surely allow to assess the value of rescue actions for ants’ success in nature (Taylor et al. 2013).

To date, although only 20 species have been studied out of over 16000 known species of ants (Bolton 2020), rescue has already been demonstrated in several different subfamilies, i.e., Dolichoderinae, Formicinae, Myrmicinae and Ponerinae. With some exceptions in which rescue was studied in an ecological context in the field (e.g., Czechowski et al. 2002, Frank et al. 2017, Uy et al. 2018), it was most frequently triggered in the laboratory with the use of the so-called “entrapment bioassay”, the most universal type of rescue test in ants so far (Hollis and Nowbahari 2013, Miler et al. 2017b). In the bioassay, ants confront a single endangered nest mate entrapped on a sandy arena by means of an
artificial snare, as was described for the first time by Nowbahari et al. (2009) and then subsequently adapted by multiple authors (e.g., Taylor et al. 2013, Andras et al. 2020).

This test is thought to provide the laboratory setup for drawing conclusions on entrapment situations in which ant workers of various species may find themselves in the field. However, several alternative versions of the entrapment bioassay can be found in the literature. The main variations include the type of substrate used on the test arena (i.e., marked or unmarked by ants’ pheromones), the number of potential rescuer ants (i.e., one or five) and the method used to code behaviour during analysis (i.e., time or interval recording procedures). Thus, despite the recognized universality of the entrapment bioassay, it is unclear how comparable are the results of various independent studies utilizing its different versions. Moreover, considering the overall scarcity of comparable data on ant rescue behaviour in the natural setting (Czechowski et al. 2002, Hollis and Nowbahari 2013, Taylor et al. 2013), it is also unclear how well each of these versions corresponds to field-based rescues.

Here, we decided to compare numerous characteristics of rescue actions in *Formica cinerea* workers, one of the species most often chosen in research on ant rescue behaviour (Czechowski et al. 2002, Miler 2016, Miler and Kuszweska 2017, Miler et al. 2017a, 2017b, Turza et al. 2020), under field- and laboratory-based entrapment bioassays of various types. Furthermore, we conducted comparisons of the data obtained using various coding procedures. We hypothesized that (1) rescue actions would be more pronounced in the field than in laboratory tests, (2) laboratory tests would yield overall similar rescue actions, and (3) coding procedures would not severely affect the conclusions. Regarding the first hypothesis, data from experiments conducted in the field may differ drastically from the results of similar experiments performed under laboratory conditions (e.g., Calisi and Bentley 2009). In the case of second and third hypothesis, many variations of the entrapment bioassay and coding procedures can be found in the literature, all regarded as appropriate for testing all rescue components (e.g., Nowbahari et al. 2009, Tailor et al. 2013, Miler 2016).

Materials and Methods

Tests of rescue behaviour were performed in the field near Klucze (Błędowska Desert, Poland, 50°21′22″N 19°31′03″E) and in the laboratory in Kraków (Institute of Environmental Sciences, Jagiellonian University, Poland) in July 2020. *F. cinerea* ants, which are sand-dwelling ants naturally exposed to different types of entrapment, were selected as the model organism. According to Czechowski et al. (2012), *F. cinerea* ants often create extensive polycalic colonies (i.e., multi-nest colonies between which workers move freely without aggression). Therefore, we decided to test two independent polycalic colonies of *F. cinerea* ants from two areas (southern and northern parts of the Błędowska Desert) 2 km apart and well separated geographically. First, we performed tests in the field, and then we collected ants from the two focal colonies and tested them in different laboratory tests.

Entrapment tests in the field

The testing procedure was similar to those of Hollis and Nowbahari (2013) and Taylor et al. (2013). For each test, using clean forceps, one ant (the victim) was captured close to the entrance to the nest and then inserted into a nylon thread loop so that the thread was located between its thorax and abdomen (i.e., on the petiole) and then tied to a small piece of filter paper (2 cm in diameter). The ant was placed within a plastic ring (7 cm in diameter) that was previously positioned on an ant-free area 5-10 cm away from the nest entrance. The plastic ring was smeared with fluon (Sigma Aldrich, Germany) on the outside to prevent nearby ants from entering the test area before the test began. Then, the
plastic ring was removed, and a 5 minute video recording started. After the test ended, the tested ant worker was not released but placed in a temporary container. The procedure was repeated 50 times for each of the two colonies (a total of 100 tests under field conditions). A new nylon thread and filter paper were used for each test. Forceps were sterilised after each test in 98% ethanol to avoid the transfer of cuticular hydrocarbons between tested ants. The position of the plastic ring was changed for each test but was always within 5-10 cm of the nest entrance. All tests were performed during two days: on 4.07.2020 (23°C, 48% relative humidity) and on 05.07.2020 (25°C, 52% relative humidity) (one day per colony), between 9 AM and 6 PM. After all tests for a given colony were completed, the tested ants were untied and released, with the exception of a few individuals collected for later taxonomic identification. The taxonomic identity of ants was confirmed in the laboratory with reference to a taxonomic key (Czechowski et al. 2012).

**Entrapment tests in the laboratory**

On a day after the entrapment tests in the field were completed, we collected approximately 1000 active *F. cinerea* foragers from each of the two colonies previously tested in the field and transported them to the laboratory where they were housed separately in plastic boxes (28 × 15 × 6 cm) at a constant temperature of 24°C, 40-60% relative humidity and a 12:12 day:night cycle, each with an attached foraging arena of the same size. Both the nest part and the foraging arena were half-filled with sand from the original habitat. Ants were provided with ad libitum water and protein-carbohydrate-vitamin-mineral food recommended by Czechowski and Pisarski (1992) in their foraging arena. The edges of the plastic boxes were smeared with fluon (Sigma-Aldrich, Germany) to prevent escape of ants. After 2-days habituation period, during which time the ants were allowed to move freely inside both boxes and accumulate pheromones on the substrate (Heyman et al. 2017), the experimental part started.

The laboratory simulations of entrapment followed standard protocols (e.g., Nowbahari et al. 2009, Miler 2016). In each test, using clean forceps, one ant (the victim) was captured on the foraging arena and inserted into a nylon thread loop so that the thread was located between its thorax and abdomen (i.e., on the petiole) and then tied to a small piece of filter paper (2 cm in diameter). Depending on the type of test, the victim was placed into a plastic ring (7 cm in diameter) on an ant-free area of the foraging arena or inside a test cup (7 cm in diameter, 8 cm high) half-filled with dry (unmarked) sand. Immediately after, the potential rescuer(s), was(were) introduced into the test area (Table 1). Then, a 5 minute video recording started. No ant was used twice. A new nylon thread and filter paper were used for each test. Forceps were sterilised after each test in 98% ethanol. In the case of tests on the substrate marked by ants’ pheromones (M1 & M5), the position of the plastic ring was changed for each test within 5-10 cm of the entrance to the foraging arena. In the case of tests on the substrate unmarked by ants’ pheromones (NM1 & NM5), unmarked sand in the test cup was renewed for each test. The procedure was repeated 50 times for each test type and colony (a total of 400 tests under laboratory conditions). The order of testing was counterbalanced by test type and colony. Tests were performed over several days, always between 9 AM and 6 PM. See Figure 1 for the scheme of the experiment.

**Analysis of the recordings**

Recorded videos were analysed using BORIS software, which enables coding behaviour of animals in an accurate and quantitative way (Friard and Gamba 2016). The videos were analysed in two ways, namely, using the time and interval recording procedures described by Martin and Bateson (2007). For the time recording procedure, each occurrence of the
given behaviour was recorded, together with information about the time the behaviour started and its total duration of occurrence. For the interval recording procedure, each test (lasting 5 minutes in total) was divided into 10 s intervals (in total 30 intervals), and behaviours were noted for each interval, providing information on the number of intervals in which a given behaviour occurred (e.g., 10 out of 30). Both procedures were previously used in ant rescue behaviour research (e.g., Nowbahari et al. 2012, Miler et al. 2017b).

We obtained data on the dependent variables common to both procedures, which included the number of tests with contact with the victim, the number of tests with any rescue actions among those tests with contact as well as the number of rescuers and the latency to the first episode of rescue in those tests with rescue. Additionally, we obtained data on the duration of (the number of intervals with) contacts in tests with at least one contact as well as the duration of (the number of intervals with) rescue actions and the durations of (the number of intervals with) selected rescue categories in tests with at least one rescue action. Operational definitions of contact and various rescue categories considered in this study were described in previous research (Table 2).

Statistics
Statistical analyses were performed using the statistical programming language R (R Core Team 2020). We used generalized linear mixed models (lme4 package) with binomial distribution, logit link function and two factors, including random ‘colony’ (N vs. S) and fixed ‘type of test’ (F vs. M1 vs. M5 vs. NM1 vs. NM5). Colonies “N” and “S” refer to the parts of the Błędowska desert from where they have been collected, i.e., northern and southern part, respectively. Types of test include “F”, which refers to the field entrapment tests, and other four types, which refer to laboratory entrapment tests. “M” and “NM” refer to marked and unmarked substrate and the numbers “1” and “5” refer to the number of freely moving workers (potential rescuers) in the test type. These models were used to compare the number of tests with contact with the victim (1 – contact, 0 – no contact) and to compare the number of tests with any rescue actions (1 – rescue, 0 – no rescue). We used a similar model to compare whether the number of rescuers differed between tests (1 – more than one rescuer, 0 – one rescuer), but in this analysis, M1 and NM1 tests were not included, as there was no possibility for more than one rescuer to occur in those tests (levels of ‘type of test’ included only F vs. M5 vs. NM5). Furthermore, we used generalized linear mixed models (lme4 package) with Poisson distribution, log link function and two factors, random ‘colony’ (N vs. S) and fixed ‘type of test’ (F vs. M1 vs. M5 vs. NM1 vs. NM5), to compare the duration of (the number of intervals with) contact and rescue. Finally, we used generalized zero inflated linear mixed models (glmmTMB package) with Poisson distribution, log link function and two factors, random ‘colony’ (N vs. S) and fixed ‘type of test’ (F vs. M1 vs. M5 vs. NM1 vs. NM5), to compare the durations of (the number of intervals with) rescue categories. Models were chosen based on deviations from normality and homogeneity of variance (stats package) and zero inflation (performance package). In all models, we performed post hoc Tukey comparisons for ‘type of test’ (emmeans package). All figures were produced in R as visualizations of model outputs (sjPlot package).

Results
Contact probability differed between test types ($\chi^2 = 31.34, P < 0.001$). In the field, the probability of contact was as high as ~90% (Figure 2). Only the laboratory test with the use of one potential rescuer on marked substrate (the M1 test type) showed a clearly lower probability of contact (ca. 70%). A more detailed analysis revealed that among tests with at least one contact with the victim, the duration of contact and the number of intervals with contact depended on the type of test ($\chi^2 = 5074.10, P < 0.001, \chi^2 = 377.95, P < 0.001$, respectively) and were both lowest in the M1 test type.
Indeed, contact duration lasted for about 2 minutes in the field, but only for about 40 seconds in the M1 tests, with in-between durations in the other test types (Figure 3A). The analysis of the number of intervals with contact revealed a very similar pattern (Figure 3B).

Rescue probability differed between test types ($\chi^2 = 42.92, P < 0.001$) in such a way that all four types of laboratory tests had lower probabilities of eliciting rescue (with a mean below 50%) than the field tests (~80%) (Figure 4). In those tests in which rescue was observed, the number of rescuers only exceeded one occasionally, although there were significant differences between test types ($\chi^2 = 18.92, P < 0.001$). In the field, the mean probability of observing more than one rescuer was lower than 50%, but in the two laboratory tests with the use of five potential rescuers it was below 20% (Figure 5). Furthermore, latency to the first episode of rescue differed between test types ($\chi^2 = 2555.00, P < 0.001$) and was highest for those conducted on unmarked substrates (the NM1 and NM5 test types). Indeed, in their case the mean latency was over 2 minutes, whereas in the field it was below 1 minute, with in-between results in tests conducted on marked substrates (the M1 and M5 test types) (Figure 6). The results for the duration of rescue and the number of intervals with rescue were consistent in the detected differences between test types ($\chi^2 = 3078.20, P < 0.001$, $\chi^2 = 354.02, P < 0.001$, respectively, Figure 7), namely, being highest in the field tests (with a mean of almost 2 minutes and over a half of all intervals). Among laboratory tests, the duration of rescue and the number of intervals with rescue were closest to that in the M5 test with the use of five potential rescuers on marked substrate (with a mean of almost 1 minute and about 9 intervals). The other test types yielded poorer results, i.e., even lower rescue durations and numbers of intervals with rescue (Figure 7).

Rescue categories (digging, pulling, sand transport and snare biting) in terms of both the duration and number of intervals all depended on the type of test (Table 3, Figure 8). Digging and pulling behaviours were most pronounced in the field and their levels seemed to be closest in the M5 test type conducted in the laboratory with the use of five potential rescuers on marked substrate (Figure 8). Sand transport was generally very rare in all types of tests (Figure 8). Interestingly, snare biting seemed even more pronounced in the M5 test type in the laboratory than in the field tests (Figure 8). In the analysis of rescue categories, differences between test types were more visible when analysed in terms of the duration than number of intervals. In other words, the duration was a better measure than the number of intervals for the fine-scale analysis of behavioural categories. This was especially clear for snare biting, in which there were differences between types of tests in the duration (highest in the M5 test type, lower in the F and NM1 test types, lowest in the M1 and NM5 test types, Figure 8) but less so in the number of intervals (similar in the F, M1, M5, and NM1 test types and lower in the NM5 test type, Figure 8).

Discussion

Our results reveal for the first time the similarities and differences in rescue actions elicited in the field and various laboratory tests conducted on ants. We compared different versions of the entrapment bioassay, considered to be a highly universal test of rescue behaviour in ants (Hollis and Nowbahari 2013, Miler et al. 2017b). Our first hypothesis, that (1) rescue actions would be more pronounced in the field than in laboratory tests, was confirmed. This illustrates the difficulty of reproducing appropriate rescue contexts in the laboratory. Our second hypothesis, that (2) laboratory tests would yield overall similar rescue actions, was not confirmed. Although we expected considerable similarities between the results of the different laboratory test types, and these results were indeed always closer to each other than to those conducted in the field, we detected some marked differences. Our third hypothesis, that (3) coding procedures would not severely affect the conclusions, was only partly confirmed. On the one hand, major issues of interest, such as...
the assessment of the contact and rescue proneness in different types of tests, were generally unaffected by the coding procedure. On the other hand, however, conclusions on a fine-scale analysis of different rescue categories differed depending on the procedure, with generally higher resolution of the analysis focused on duration rather than on intervals (and, thus, more detailed conclusions in the former case).

In this study, we measured contact with the victim and examined its importance in the context of later elicitation of rescue behaviour. Laboratory tests with the use of one potential rescuer on marked substrate (the M1 test type) were least effective in eliciting contact with the victim (Figure 2). Moreover, contacts within this M1 test, expressed as either the duration or number of intervals, differed the most from contacts in the field (Figure 3). It is reasonable to expect that contact generally impacts the occurrence of rescue because it is a prerequisite of this more advanced behaviour (Taylor et al. 2013, Silberberg et al. 2014). However, the probability of rescue was much lower in all laboratory tests compared to the field (Figure 4). Indeed, this probability in the laboratory was approximately 50%, which was in line with earlier results obtained on Formica cinerea ants (Miler and Kuszewska 2017) and much lower than that in the field (~80%).

This general difference between the field and the laboratory might stem from the fact that the field study, in contrast to artificial conditions, provides many more natural environmental cues and close proximity of the whole nest, leading to a more natural behaviour and stimulation of risky behaviour (Czechowski et al. 2002, 2012, Hollis and Nowbahari 2013). Based on these results alone, it is obvious that whenever possible, field tests should be preferred when studying ant rescue behaviour. Of note, however, the present results do not mean that laboratory tests are useless. All types of tests triggered rescue actions and may be suitable for many studies on rescue behaviour. In general, however, tests might be enhanced under laboratory conditions by, for example, collecting full ant colonies (including queen and brood), which would provide more natural conditions and probably intensify the occurrence of rescue behaviours (Nowbahari et al. 2009).

It was previously suggested that at least five nest mates (potential rescuers) must be present to trigger any rescue action in the laboratory (Nowbahari et al. 2009, 2012). Our data showed that, even in tests with multiple potential rescuers, the number of ants actually performing rescue was rarely more than 1, even in the field (Figure 5). Notably, the maximum number of active rescuers did not exceed three in the laboratory and five in the field, although more than one rescuer in any test was rare. One possible explanation for this is related to energy limitations as ants might have evolved ways to prevent the elicitation of potentially costly rescue behaviours simultaneously in many individuals (Nowbahari and Hollis 2010). In many cases, we observed active rescuers to cease their rescue activity during a test to interact with other freely moving workers, possibly exchanging signals and changing behaviour of both the rescuer and the other surroundings ants (Mallon and Franks 2000, Pratt et al. 2001, McLeman et al. 2002).

Of note, when considering latency to the first episode of rescue in our study, it clearly depended on the familiarity of the substrate (Figure 6). Marked substrate enabled ants to identify the rescue context faster and react appropriately to the trapped individual. Also, when analysing more detailed data, i.e., the duration of rescue and the number of intervals with rescue, the laboratory test with the use of five potential rescuers on marked substrate (the M5 test type) differed the least from that performed in the field (Figure 7). Thus, the use of laboratory tests performed on familiar substrates and/or with the use of several potential rescuers should be considered in further studies especially if, for instance, the total number of tests to be performed is to be low and cannot be increased. Doing so might facilitate rescue observations, even in its more advanced forms, such as pulling or snare biting (Figure 8).

Snare biting is considered to be precision rescue, i.e., the ability to target the object holding the victim directly (Nowbahari et al. 2009, 2012). Indeed, a study by Hollis and Nowbahari (2013) showed that some species can perform rescue actions with digging, sand transport and pulling, but not with precision rescue behaviour. These differences
between species suggest that rescue behaviour categories (or at least precision rescue behaviour) are not necessarily common among ants. Indeed, ant species may differ strikingly in their rescue behaviour. For instance, ants show division of labour, influenced by age and/or body size (Hölldobler and Wilson 1990), and different species might differ in the type of workers responding to rescue-associated stimuli. As pointed out by Nowbahari et al. (2012), temporal polyethism can regulate the expression of rescue behaviour. In Cataglyphis cursor ants, which are characterized by this type of polyethism, older foragers both rescue and are being rescued more frequently than younger nurses. However, polyethism is not present in all species of ants (Traniello 1978), and for those in which it occurs, it is diverse in its forms (Hölldobler and Wilson 1990). Similarly, rescue in ants can be under genetic control (Andras et al. 2020). Apparently, in Cataglyphis cursor ants, 34% of the variation in propensity for rescue behaviour is explained by paternity. This means that the relatedness of workers within a colony, which often differs drastically between ant species, might affect the ease of rescue elicitation. Indeed, Nowbahari et al. (2009) showed that the behaviour of Cataglyphis cursor rescuers is preferentially aimed at providing help to more related individuals. This means that multifaceted analyses of rescue actions in ants are indispensable.

Here, we also indicate the need for careful interpretation of results depending on the examined behavioural component of rescue (Figure 8). In this context, we found pronounced differences between both the test types and coding procedures. In the case of coding procedures, differences between test types in some behaviours were more difficult to detect when measured by interval coding procedure than by time coding procedure (compare, e.g., differences between tests in digging when analysed via time duration, Figure 8a, and interval duration, Figure 8e, or differences between tests in pulling when analysed via time duration, Figure 8b and interval duration, Figure 8f). When analysing the results, one has to keep in mind to fit the resolution of the coding method to the amount of gathered data. In the case of test types, on the other hand, in line with what was mentioned above, the M5 test type with the use of five potential rescuers on marked substrate was closest to field tests in behavioural categories of digging, pulling and snare biting (Figure 8). It needs to be stressed that choosing a category to study is highly important and may significantly affect the conclusions. In different test types, the expression of behavioural categories may differ. Moreover, the detected behavioural categories may differ between species. Here, for example, the most common behaviour was pulling the victim (Figure 8), which was in line with previous results obtained on Formica cinerea ants (Miler 2016). Interestingly, in the case of Cataglyphis cursor, digging around the victim is considered to be most common rescue category (Nowbahari et al. 2009). In a further study, it would be interesting to examine the differences in the intensity of behavioural categories in different ant species more closely. Such differences may suggest that rescue actions evolved in various contexts, depending on the selective pressure of the environment and ecology of given species (Hollis and Nowbahari 2013).

Rescue behaviour plausibly plays an essential role in ecological success of ants by increasing individual survival and, ultimately, benefiting the ant colony. However, studies assessing the benefits of rescue actions in ants are generally scarce (Frank et al. 2017, Kwapić and Hölldobler 2019). As mentioned in the Introduction, rescue actions at least in some cases decrease mortality of individuals (Frank et al. 2017). How important each and every individual is for the colony is demonstrated by Kwapić and Hölldobler (2019), which calculated that the loss of five individuals in a colony of Veromessor pergandei seed harvester ants causes the loss of 65700 seeds per year. This suggests that rescue actions translate into real benefits. Here, during field entrapment tests, F. cinerea ants carrying resources (i.e., nest material or prey item) were observed several times to abandon it and start a rescue action of the entrapped nest mate (F. Turza, personal observation). Attempts to measure the benefits of rescuing a nest mate in F. cinerea would contribute to a better understanding of its generally high propensity for rescue behaviour.
In further research, it would be interesting to examine rescue propensity in the field, depending on how far away from the nest the entrapped individual needs help. This would supplement our current suggestions that field studies yield highest probabilities of rescue actions due to familiarity of the surroundings. Such research could also compare rescue propensity on and off the foraging trails (see Kwapich and Hölldobler 2019) or in areas with a few or many nest mates. In addition, connection between rescue and activity cycle seems worthwhile of investigation too (Fujioka et al. 2017, Kay et al. 2018). It could provide information on the specificity of rescue behaviour, which may contribute to a better understanding of the unknown variability in the expression of such behaviour among Formicidae (Hollis and Nowbahari 2013, Miler et al. 2017b).

Data from experiments utilizing laboratory entrapment tests (Nowbahari et al. 2009, 2012, 2016, Miler and Kuszewska 2017, Miler et al. 2017a, 2017b, Duahoo et al. 2017, Andras et al. 2020) may strongly underestimate rescue proneness in comparison to field conditions. Therefore, regarding ant species tested only in the laboratory, that showed no rescue behaviour, such as Camponotus korthalsiae, Anoplolepis gracilipes and Myrmica ruginodis (Miler et al. 2017b), or that showed low-level rescue behaviour, such as Camponotus aethiops (Nowbahari et al. 2016), Formica polyctena and Iridomyrmex anceps (Miler et al. 2017b), their inability to perform rescue actions should be considered carefully. Even species showing high levels of rescue behaviour in laboratory tests, such as Cataglyphis cursor (Nowbahari et al. 2009, 2012, 2016, Duahoo et al. 2017, Andras et al. 2020) and Formica cinerea (Miler and Kuszewska 2017, Miler et al. 2017a, 2017b), can potentially be even more prone to express rescue behaviour than so far thought. Overall, the choice of a test variant in a study should be carefully considered depending on the specifics of the research questions. Here, we agree with Duahoo et al. (2017) that ants, despite common belief, are not ”the hardwired reflex automatons” and might be highly sensitive to the behavioural context. Our data supports a growing number of studies showing that methods of assessing animal behaviour, even if differing only slightly, may lead to significantly different results (Ovadia et al. 2001, Watanabe 2012, Scharf and Martin 2013).

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**Authors’ contributions**
K.M. conceived the study; F.T. and K.M. designed the study; F.T. collected the data; F.T. and K.M. analyzed the data and wrote the manuscript.

**Data availability**
The datasets generated and analyzed during the current study are available in the supplementary Excel file.

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| Type of test                                                                 | Detailed procedure                                                                                                                                 |
|------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------|
| One victim and potential rescuer(s) on marked substrate under field conditions (F) | The plastic ring is placed on an ant-free area 5-10 cm away from the nest entrance (i.e., sand marked by ant pheromones). Then, the victim is placed in the centre of the ring. Then, the plastic ring is removed. |
| One victim and one potential rescuer on marked substrate under laboratory conditions (M1) | The plastic ring is placed on an ant-free area of the foraging arena attached to the nest (i.e., sand marked by ant pheromones). Then, the victim is placed in the centre of the ring. Then, a randomly chosen nest mate from the foraging arena is placed into the ring. |
| One victim and five potential rescuers on marked substrate under laboratory conditions (M5) | The plastic ring is placed on an ant-free area of the foraging arena attached to the nest (i.e., sand marked by ant pheromones). Then, the victim is placed in the centre of the ring. Then, 5 randomly chosen nest mates from the foraging arena are placed into the ring. |
| One victim and one potential rescuer on unmarked substrate under laboratory conditions (NM1) | The victim is placed into the plastic cup filled with dry (unmarked) sand. Then, a randomly chosen nest mate from the foraging arena is placed inside the cup. |
| One victim and five potential rescuers on unmarked substrate under laboratory conditions (NM5) | The victim is placed into the plastic cup filled with dry (unmarked) sand. Then, 5 randomly chosen nest mates from the foraging arena are placed inside the cup. |

Table 1. Detailed procedures for the different types of tests conducted under field and laboratory conditions.
| Behaviour      | Operational definition                                                                 |
|---------------|----------------------------------------------------------------------------------------|
| **Contact**   | The ant touches any part of the body of the victim, using antennae                      |
| **Digging**   | The ant stands in front of the victim and repels sand backward, using legs             |
| **Pulling**   | The ant grabs any part of the body of the victim and drags it backwards, using mandibles|
| **Sand transport** | The ant picks up a pebble near the victim and moves it away, using mandibles          |
| **Snare biting** | The ant tugs on the nylon thread holding the victim, using mandibles                   |

Table 2. Definitions of rescue categories used in the study (based on Hollis and Nowbahari 2013).
| Behaviour      | Procedure      | $\chi^2$  | p value |
|---------------|----------------|-----------|---------|
| Digging       | Time recording | 497.01    | < 0.001 |
|               | Interval recording | 98.73   | < 0.001 |
| Pulling       | Time recording | 1836.90   | < 0.001 |
|               | Interval recording | 242.60   | < 0.001 |
| Sand transport| Time recording | 9.84      | 0.043   |
|               | Interval recording | 26.45   | < 0.001 |
| Snare biting  | Time recording | 179.26    | < 0.001 |
|               | Interval recording | 24.53   | < 0.001 |

Table 3. Results of the models for the duration of (the number of intervals with) rescue categories depending on the test type (F vs. M1 vs. M5 vs. NM1 vs. NM5).
Figure 1. Scheme of the experiment.
Figure 2. Probability of contact in different types of tests (field entrapment tests, “F”, and four types of laboratory entrapment bioassays). See Table 1 for differences between the “M1”, “M5”, “NM1” and “NM5” tests. Dots represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0125.
Figure 3. Duration of contacts (panel A) and number of intervals with contact (panel B) in different types of tests (field entrapment tests, “F”, and four types of laboratory entrapment bioassays). See Table 1 for differences between the “M1”, “M5”, “NM1” and “NM5” tests. Dots represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0125.
Figure 4. Probability of rescue in different types of tests (field entrapment tests, “F”, and four types of laboratory entrapment bioassays). See Table 1 for differences between the “M1”, “M5”, “NM1” and “NM5” tests. Dots represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0125.

Figure 5. Probability of observing more than one rescuer in different types of tests (field entrapment tests, “F”, and two types of laboratory entrapment bioassays). See Table 1 for differences between the “M5” and “NM5” tests. Dots
represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0250.

Figure 6. Latency to first episode of rescue in different types of tests (field entrapment tests, “F”, and four types of laboratory entrapment bioassays). See Table 1 for differences between the “M1”, “M5”, “NM1” and “NM5” tests. Dots represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0125.
Figure 7. Duration of rescue (panel A) and number of intervals with rescue (panel B) in different types of tests (field entrapment tests, “F”, and four types of laboratory entrapment bioassays). See Table 1 for differences between the “M1”, “M5”, “NM1” and “NM5” tests. Dots represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0125.
Figure 8. Duration of selected rescue categories (panels A-D) and number of intervals within rescue categories (panels E-H) in different types of tests (field entrapment tests, “F”, and four types of laboratory entrapment bioassays). See Table 1 for differences between the “M1”, “M5”, “NM1” and “NM5” tests. Dots represent model predictions, and whiskers indicate estimated 95% confidence intervals. Small letters above upper whiskers indicate significance with Tukey post hoc contrasts after Bonferroni correction < 0.0125.