Dung burial by roller dung beetles (Coleoptera: Scarabaeinae): An individual and specif c-level study

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Abstract. Dung beetles (Coleoptera: Scarabaeinae) mediate many ecological functions that are important to maintain the ecosystem functioning of terrestrial environments. Although a large amount of literature explores the dung beetle-mediated ecological processes, little is known about the individual contribution from distinct species. Here, we aimed to examine the intra and interspecif c variations in dung burial rates performed by two roller dung beetle species (*Canthon smaragdulus* Fabricius, 1781 and *Canthon sulcatus* Castelnau, 1840). Furthermore, we evaluated the relationship between dung beetle biomass and dung burial rates. We set up a laboratorial experiment with three treatments (two males, two females, and a couple) and 10 replicates per treatment for each dung beetle species, and dung burial rates were measured after exposing 100 g of mixed pig and human excrement for 48 hours. Our results demonstrate that dung burial rates of males, females, and couples within each species do not differ. However, *C. smaragdulus* individuals performed a larger dung burial than *C. sulcatus* individuals did. In addition, we found no effect of individual biomass on the amount of dung burial on intra and interspecif c levels. These f ndings highlight the need for further research considering that distinct species, even from the same genus, may perform different rates of ecological processes, as well as about the importance for considering the beetle biomass when measuring their ecological functions. We call for studies to f ll in the knowledge gap about the individual species' contribution to the maintenance of different dung beetle-mediated ecological processes.

Key words: Scarabaeidae, ecological function, dung burial

Introduction

Anthropogenic disturbances have caused wide-spread species extinctions, with further consequences on ecosystem processes and ecological services triggered by organisms (Hooper *et al.*, 2005; Kremen *et al.*, 2007). Much of the debate about the impacts of biodiversity loss on ecosystem functioning has thus far focused on higher organizational levels by considering the role of species groups instead of those from individual species and organisms (Loreau *et al.*, 2001; Bellard

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et al., 2012). However, understanding the functional role of organisms and individual species becomes crucial to further predict their ecological importance and contribution to ecosystem functioning processes (Covich *et al.*, 1999; Thrush *et al.*, 2006).

Despite the almost exponential increase in research on ecological functions and further benefts to human well-being and environmental maintenance, contributions at individual or even specifc levels remain poorly studied. On the other hand, functional groups have been used to better represent the relationship between functional diversity and ecosystem functioning (Slade *et al.*, 2007; Nichols *et al.*, 2013a; Griffths *et al.*, 2016a). Overall, studies have shown that the loss of both taxonomic and functional diversities leads to negative effects on ecosystem functioning (Hooper *et al.*, 2005), but ecological processes in perturbed ecosystems can also be buffered by functionally redundant species (Barragán *et al.*, 2011). Few studies, however, have explored the individual contribution and the infuence of the sex of organisms on ecosystem processes; interspecifc variation and species intraspecifc differences have also been poorly considered (Wohlfahrt *et al.*, 1999; Crutsinger *et al.*, 2006; Carvalho *et al.*, 2018).

Dung beetles (Coleoptera: Scarabaeidae: Scarabaeinae) are a recognized bioindicator group used for ecological investigations (Barlow *et al.*, 2016; França *et al.*, 2016a, b) and for research on ecosystem functioning (Braga *et al.*, 2013; Griffths *et al.*, 2015, 2016a). These detritivore beetles mediate a variety of relevant ecological processes, such as secondary seed dispersal, nutrient cycling, soil bioturbation, and parasite suppression (reviewed by Nichols *et al.*, 2008). The ecological functions performed by these insects mainly result from their nesting and resource allocation strategies. Among dung allocation strategies, roller (or telecoprid) dung beetles are those that make dung balls and roll them for distances ranging from a few centimeters to several meters. Once the nesting place is reached, the balls are buried in tunnels and chambers that have been built (Halffter and Matthews, 1966; Halffter and Edmonds, 1982), which benefts the soil environment and plant productivity (Bang *et al.*, 2005; Yamada *et al.*, 2007; Griffths *et al.*, 2016a).

Here, we used a laboratorial experiment to investigate the intra and interspecifc contributions of two roller dung beetle species (*Canthon (Goniocanthon) smaragdulus* Fabriciuys, 1781 (Nunes *et al.*, 2018) and *Canthon sulcatus* Castelnau, 1840) to the dung burial function. *Canthon (Peltecanthon) sulcatus* is a diverse genus of dung beetle, distributed throughout the American continent (Vaz-de-Mello, 2000; França *et al.*, 2016c). However, the ecology and behaviour of this genus is poorly known (Vaz-de-Mello *et al.*, 2014). In particular, we evaluated the relationship between the dung beetle individuals'

biomass and the dung burial rates they performed, as this function is considered to be one of the most important dung beetle-mediated ecological processes. Dung consumption and burial can lead to other crucial incidental functioning processes (Nichols and Gardner, 2011; França *et al.*, 2018), such as secondary seed dispersal (Santos-Heredia and Andresen, 2014; Griffths *et al.*, 2015), microbial transport across the soil surface (Slade *et al.*, 2016), and soil bioturbation (Nichols *et al.*, 2008; Braga *et al.*, 2013). We assessed how dung beetle biomass would infuence dung burial rates, as this functional trait has been shown to infuence dung beetle-mediated ecological processes (Gregory *et al.*, 2015). We hypothesized that (1) males, females, and couples of the same species remove dung at different rates; (2) different species remove dung at different rates; and (3) there is a positive relationship between individual biomass and the amount of dung burial on intra and interspecifc levels.

Materials and methods

Study region

We sampled individuals of the dung beetle species *C. smaragdulus* and *C. sulcatus* in January 2015, during the rainy season, at Rio Doce State Park, in the state of Minas Gerais, southeastern Brazil (19°48 18 –19°29 24 S and 42°38 30 – 42°28 18 W). This state park has an area of approximately 36,000 ha and is considered one of the largest preserved Atlantic forest areas within the state of Minas Gerais (Coelho and Ribeiro, 2006). Although it presents a mosaic of vegetation categories, it is considered a tropical rainforest domain, classifed as submontane semidecidual stationary forest (Antunes, 1986). The region presents a transition of mesothermic humid tropical and mesothermic rainy tropical climate (Antunes, 1986), with a mean temperature between 20 °C and 22 °C and a well-defned dry season from April to September and a rainy season between October and March (Antunes, 1986).

Sampling design

We used 25 pitfall traps (25 × 5.2 cm), placed linearly 25 m apart in a transect within a forest fragment. Pitfall traps were made by cutting the neck off from 2-L plastic bottles. The tapering part of the bottle was cut off and placed upside down on the top of the bottle, forming a funnel, which allowed the beetles to be trapped while also preventing them from escaping. Traps were buried with their openings at the ground level. We placed a small amount of soil and leaf litter inside each pitfall trap (below the funnel) to accommodate and avoid stressing the trapped dung beetles. To attract the

dung beetles, we placed a small container with 25 g of pig dung above each trap and we used a plastic lid supported by bamboo sticks as a rain cover. All traps remained in the f eld for 72 hours and dung beetles were collected every 24 hours, when the traps were rebaited. Living dung beetles collected were kept in plastic containers (15 × 9.4 cm) half f lled with a humidif ed mixture of soil (70%) and sand (30%).

After f eldwork, live dung beetles were transferred to the Insect Ecology and Conservation Laboratory at the Federal University of Lavras (Brazil), where containers were kept within a controlled room (humidity 65% ± 10%, temperature 26 °C ± 1 °C, and 12 h/ 12 h light/ dark photoperiod). The beetles were reared following the methodology of Favila (1993).

Selection of individual dung beetles

Prior to the experiment, individual dung beetles were f rst separated by species and then by sex. Due to the absence of sexual dimorphism in *C. smaragdulus* specimens, we opted to separate males from females by observing their reproductive behaviour, as described by Belles and Favila (1983) and Favila (2001). Males made dung balls with resources that were provided to them. We then observed that they stood on the ground, raising their bodies while keeping both hind legs extended, which was interpreted as the moment when pheromones are emitted. When a female approached, sexual recognition was made through contact between heads and antennae. The female then climbed onto the resource ball and the male rolled it to a location where copulation occurred. Here, we observed the male approaching the female and mounting her. The female remained in a favourable position for him to attempt copulation. We used this attempt by the male to copulate with the female to consider them as a couple and to identify each sex. All individuals that failed to display reproductive behaviour were excluded from the trial.

Canthon sulcatus specimens present sexual dimorphism, which allowed us to separate individuals by observing their anterior tarsal claws: males present bidentate tarsal claws, whereas the female's anterior tarsal claw is simple dentate. We further dissected 10 individuals of each sex from both species to conf rm our sex classif cation. The high number of matches (100%) conf rmed that our techniques were precise for sexing this dung beetle species.

After sexing the beetles, we separated 10 couples, 20 males, and 20 females of each species, and placed them in separate plastic containers. There were 10 containers with one couple in each container, 10 containers with two males each, and 10 containers with two females each.

Experimental design for dung burial evaluation

To evaluate dung burial capacity (f rst hypothesis), three treatments were performed with two individuals of the same species: males (MM), females (FF), and distinct-sex couples (MF). A total of 10 replicates were performed for each treatment, totalling 30 replicates for each species (for the second hypothesis). Each repetition comprised a bucket with 5 kg of a humidif ed substrate, prepared with 70% soil and 30% sand. Two individuals of each respective treatment (MM, FF, and MF) of one of the species and 100 g of pig dung mixed with human excrement (80:20 pig-to-human ratio, following Marsh *et al.*, 2013) were placed in each bucket, which was then covered by canvas to prevent any beetle from escaping. Before being placed in a bucket, the beetle biomass was weighed with a Shimatzu AY220 balance scale (Shimadzu Corporation, Kyoto, Japan) accurate to within ± 0.0001 g to further analyse it (third hypothesis). Beetles were not fed for 7 days prior to the experiment, and beetle-mediated dung burial was quantif ed 48 hours after the beginning of the experiment.

We prepared 10 additional buckets following the same methodology above, but without placing any beetle, to account for any humidity loss/ gain that could inf uence the measurement of dung removal rates. In each of these 'humidity control' buckets, the changes in humidity from the dung were calculated and the mean value was subtracted from the amount of dung buried by the beetles in each replicate. The positions of the buckets, both those corresponding to the treatments and those of the humidity control, were randomized to avoid any inf uence of bucket placement.

Statistical analysis

All data were analysed using the software R (R Core Team, 2017). A general linear model assuming quasi-Poisson distribution was used to examine whether the biomass varied among treatments within each species (males, females, and couples) or between species (*C. smaragdulus* and *C. sulcatus*). The average biomass of the two individuals of each bucket was considered the dependent variable, and the treatment levels and the species of beetle were the explanatory variable. We used general linear models (GLMs) assuming quasi-Poisson distribution to examine whether the dependent variable (dung burial rates) varied among treatments (three levels: MM, FF, and MF) and the two *Canthon* species (*C. smaragdulus* and *C. sulcatus*) (f rst and second hypothesis). Biomass was used as a covariable in the model, because it could inf uence the response variable (third hypothesis). On both models when signif cant differences were detected in formed

interactions, we performed a contrast analysis to determine which categories were different in relation to the response variable. All models were submitted to residual analysis to verify the adequacy of distribution errors (Crawley, 2002, 2013). The quasi-Poisson distribution was performed as recommended to deal with overdispersed data (Ver Hoef and Boveng, 2007). We performed Shapiro–Wilk tests of normality with the response variable and model residuals in the *shapiro.test* function in *stats* package (R Core Team, 2017). The signif cance level of the analysis was alpha = 0.05.

Results

Biomass in the intra and interspecif c levels

There were no signif cant differences in mean biomass when we considered the interaction of treatment with *C. smaragdulus* (MM pairs 0.241 ± 0.012 g; FF pairs 0.246 ± 0.007 g; 0.230 ± 0.011 g) (Fig. 1A) and with *C. sulcatus* (MM pairs 0.149 ± 0.004 g; FF pairs 0.163 ± 0.006 g; MF pairs 0.162 ± 0.006 g) ($F_{2,54} = 1.033$; $P = 0.363$) (Fig. 1B). However, the mean biomass of individuals differed between species ($F_{1,56} = 152.235$, $P < 0.01$). *Canthon smaragdulus* individuals (0.239 ± 0.006 g) had greater biomass than *C. sulcatus* individuals (0.156 ± 0.04 g) (Fig. 1C).

Dung burial rates in the intra and interspecif c levels

We did not f nd signif cant differences in mean dung buried when we considered the interaction of treatment with *C. smaragdulus* (MM pairs 19.91 ± 7.22 g; FF pairs 17.63 ± 4.84 g; MF couples 18.82 ± 3.52 g; mean ± SE) and with *C. sulcatus* (MM pairs 11.74 ± 2.28 g; FF pairs 8.84 ± 2.94 g; MF couples 7.95 ± 1.23 g; mean ± SE) ($F_{2,53} = 0.49$; $P = 0.61$) (f rst hypothesis). However, supporting our second hypothesis, we found a difference in the amount of dung buried by species, where *C. smaragdulus* (18.78 ± 3.03) buried more dung than *C. sulcatus* individuals (8.07 ± 1.38; mean ± SE) ($F_{1,58} = 7.12$, $P < 0.01$) (Fig. 2).

Relationship between individual biomass and the amount of dung burial on intra and interspecif c levels

No signif cant effect on the amount of dung buried was observed when considering the interaction between the biomass of the beetles, treatments within each species, and species ($F_{2,48} = 0.05$, $P = 0.95$). We found no effect on the amount of dung buried when we considered the interaction between biomass and *C. smaragdulus* and *C. sulcatus* ($F_{1,52} = 0.21$, $P = 0.65$) (third hypothesis).
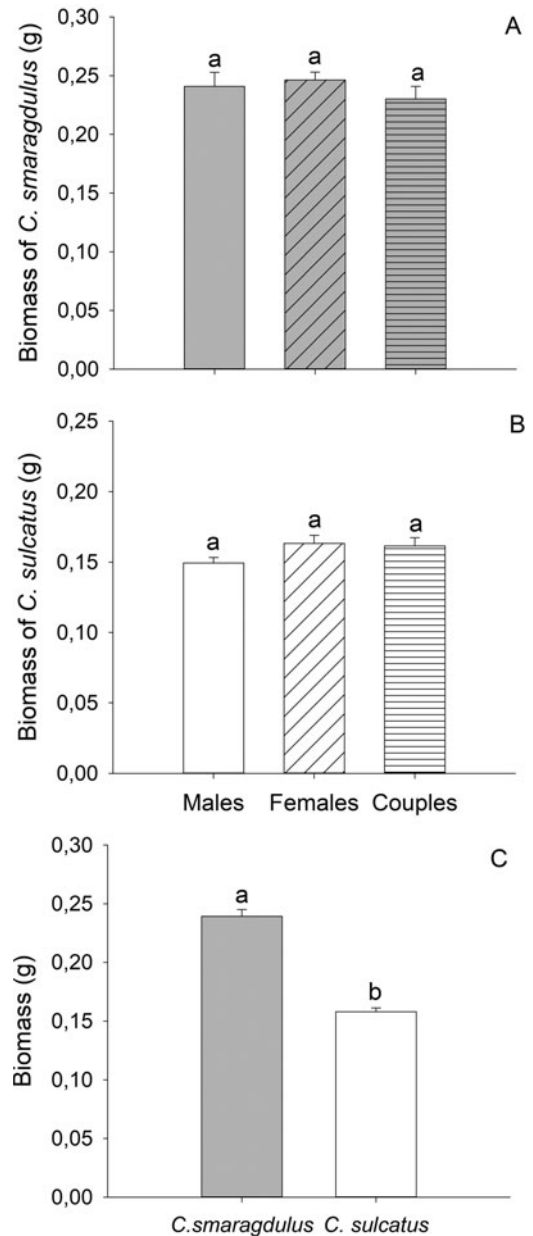

Fig. 1. Mean and standard error of the biomass of individuals (A) males, females, and couples of *Canthon smaragdulus*, (B) males, females, and couples *C. sulcatus*, and (C) of *C. smaragdulus* and *C. sulcatus*. Different letters indicate statistical difference between treatments as indicated by GLM at 5% signif cance, *n* = 60.

Discussion

Our study provides empirical evidence that different dung beetle species remove different amounts of dung, therefore supporting previous research demonstrating that bigger beetles may perform larger amounts of dung beetle-mediated ecological functions (Gregory *et al.*, 2015). Dung consumption
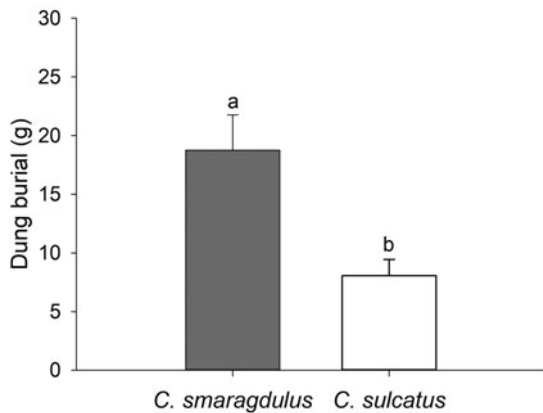
Fig. 2. Mean and standard error of the amount of dung buried by *Canthon smaragdulus* and *C. sulcatus*. Different letters indicate statistical difference between treatments as indicated by GLM at 5% signif cance, $n = 60$.

and burial by dung beetles are key processes for the occurrence of other ecological functions, such as soil bioturbation from the construction of tunnels, seed dispersal, and the increase in plant productivity (Nichols *et al.*, 2008; Braga *et al.*, 2013; Griff ths *et al.*, 2015; Santos-Heredia *et al.*, 2016; Braga *et al.*, 2017).

We found no differences in dung burial rates by males, females, or couples within each species. Previous research has shown that dung beetle sex is an important driver for tunnel depth (Gregory *et al.*, 2015) and that dung burial behaviour may change between sexes (Emlen, 1997). However, the resource offered was incompletely buried in either of our replicates. This demonstrates that the amount of dung offered in this experiment may be too large, and that there was no competition between the individuals placed within each bucket. While we concede that beetles in this experiment did not reallocate the resource as quickly as they usually do (Hanski and Cambefort, 1991), future studies are needed to evaluate the effect of different densities of beetle individuals on dung burial rates.

Our results also demonstrate that the biomass of males, females, and couples of each species had no relation to dung burial within the examined species. Larger intraspecif c variation in biomass within a species increases the probability of relationships between the individual biomass and the functions they perform (Braga, unpublished observations). If this variation in intraspecif c biomass is low (or does not exist), the relationship of biomass to ecological function will most likely not exist. This is a likely explanation for the patterns found here, as there was no intraspecif c variation in the biomass from males, females, and distinct-sex couples in this experiment.

Individuals of *C. smaragdulus* performed more dung burial than *C. sulcatus*. Although *C. smarag-*

dulus species has a higher biomass than *C. sulcatus*, our results demonstrate that the biomass of the individuals of each species had no direct relation to dung burial rates. In studies with dung beetles at the community level, the biomass of individuals in the community has had a relation to the amount of dung buried by the beetles (Andresen, 2002; Slade *et al.*, 2007; Nichols *et al.*, 2013b). Communities that have dung beetles with greater biomass have a higher capacity for dung burial (Horgan, 2001; Anduaga, 2004). Although this relationship is valid in studies with dung beetle communities, it did not exist in the species we worked with. The biomass of individuals will only be an ecologically considerable trait for dung burial in those species with large biomass and presenting a high capacity for dung burial, like those of the genus *Dichotomius* (Andresen, 2003; Anduaga, 2004; Braga, unpublished observations). This demonstrates the need for research using other factors, such as different functional traits and environmental contexts in which the beetles are found (Nichols *et al.*, 2013b; Griff ths *et al.*, 2015, 2016b).

Our work addressed the individual contribution of two roller dung beetle species to dung burial rates and demonstrated that each species may contribute differently. In addition, the dung burial rates were indirectly related to the biomass of each species, but the larger species performed higher dung burial rates. Thus, it is important to understand the behaviour and ecology of the species and their inf uence on the capacity for dung burial, one of the main ecological functions performed by dung beetles (Nichols *et al.*, 2008). Further studies are needed to evaluate the contribution of other individual species of dung beetle for dung burial. We believe that f ne-scale data based on individuals can be useful for better understanding and valuating the ecological function provided by dung beetles. Further studies are needed to evaluate the contribution of other individual species of dung beetle for dung burial and the other ecological processes they mediate.

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