



Swarming rate and timing of unmanaged honeybee colonies (*Apis mellifera carnica*) in a forest environment

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Abstract

Investigating the life history of social insect colonies and the demography of their populations are important for their conservation, but data collection is challenging. There is a growing interest in understanding the population status of wild-living honeybee colonies across Europe, for which it is critical to collect data on survival and natality rates. Although survival rates can be investigated through regular inspections of wild nests, the accurate quantification of natality rates (i.e., the number of swarms produced per colony per year) remains a significant challenge. Using digital weight scales, we remotely monitored the natural swarming behavior of ten unmanaged *Apis mellifera carnica* colonies housed in static-volume hives (45 L) in a forest region of southern Germany. During the 2019 season, between mid-May and late June, we recorded 17 swarming events, averaging 1.7 swarms per colony. Our observations offer a reference point for the timing, frequency, and size of honeybee swarms that helps us understand the natural reproductive patterns of wild-living honeybees in a temperate forest environment.

Keywords Honeybee reproduction rates · Population demography · Natality rate · Weight scale analysis · Wild-living honeybees · Free-living honeybees · Swarming behavior

Introduction

Understanding the life-history strategy of species and studying the demography of their populations are core tasks in conservation biology. For an analysis of the viability of a given population, two variables are key: survival and natality. Survival (or its converse, mortality) refers to the proportion of the population that endures from one year to the next, influenced by factors such as predation, diseases, and other environmental conditions (e.g., weather conditions or resource availability). Conversely, natality encompasses the introduction of new individuals into the population through

reproduction, contributing to population growth (Krebs, 1972).

Although European populations of the western honeybee (*Apis mellifera*) have traditionally been viewed through the lens of apiculture, recent studies have highlighted the existence of wild-living¹ honeybees across various regions (e.g., Oleksa et al., 2013; Kohl and Rutschmann, 2018; Requier et al., 2020; Browne et al., 2020; Bila Dubaić et al., 2021; Rutschmann et al., 2022; Visick and Ratnieks, 2023; Rutschmann et al., 2024). Understanding the population demography of these wild-living colonies is crucial for comprehending their ecological and evolutionary roles (Kohl and Rutschmann, 2018; Kohl et al., 2022). Furthermore, if self-sustaining populations were discovered, these would offer unique insights into how honeybees can persist despite challenges like parasitic pressures which, in turn, could contribute to the development of more sustainable beekeeping practices (Panziera et al., 2022; Requier et al., 2019).

Honeybee colonies are intrinsically long-lived (multiple years), since young queens can supersede old queens as the reproductive individual of the society. In temperate regions, *Apis mellifera* colonies exhibit high site fidelity, allowing

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¹ To describe honeybee colonies that are not managed and chose their nest site on their own, we use the term “wild-living”.

survival rates to be assessed through repeated visits to their nest sites (Seeley, 1978; Oldroyd et al., 1997; Rutschmann et al., 2022; Kohl et al., 2022; Lang et al., 2022; Cordillot, 2024; Rutschmann et al., 2024). Acquiring comprehensive and reliable data on the annual natality rates of honeybee colonies within a population, however, poses a significant challenge (Otis and Wearing-Wilde, 1992).

Colony reproduction is achieved through swarming, a process in which a single colony divides into two or more distinct colonies under favorable environmental conditions. In preparation of this process, the colony raises several young queens. Before these new queens emerge, the old queen leaves with more than 70% of the colony's worker bees as the so-called prime swarm (Seeley, 1985; Winston, 1991). Subsequently, several afterswarms—each with at least one of the newly emerged young queens and another portion of the workers—may leave the original nest site to found independent colonies (Otis and Wearing-Wilde, 1992; Seeley, 1985; Winston, 1991). The annual natality rate in honeybees is thus defined as the frequency of swarms (prime swarm plus afterswarms) per colony per year.

Although a queen change during the swarming season (which is typically indicative of a swarming event) can be detected using genetic maternity tests of workers, or, in case of colonies managed in hives by directly monitoring the presence of color-marked queens, these methods fall short in accurately determining the number of afterswarms (Otis and Wearing-Wilde, 1992). Therefore, prior studies on annual swarming rates required continuous observation of nest entrances (Otis and Wearing-Wilde, 1992), a method that is notably time-consuming.

We here demonstrate the use of digital weight scales to remotely and accurately monitor honeybee colony swarming events. We gathered data from ten colonies that were kept in hives with a static volume of approximately 45 L in a forest region in Germany. By avoiding beekeeping manipulations, we ensured that our observations reflected near natural behavior. The method not only enhances previous research techniques but also adds to our understanding of the natural timing and frequency of swarming and, consequently, the population dynamics of wild-living honeybee populations in a central European forest environment.

Methods

Study area

This study was conducted in the Steigerwald, a low mountain forest region in southern Germany. The details of our experimental setup are described in Rutschmann et al. (2023). The managed forested area covers approximately 165 km² and primarily consists of beech trees (*Fagus sylvatica*, 44%),

followed by oak (*Quercus* spp., 21%), Scots pine (*Pinus sylvestris*, 13%), spruce (*Picea abies*), and hornbeam (*Carpinus betulus*) (Mergner and Kraus, 2020). Surrounding grassland areas feature many orchard meadows, and the most prevalent crops include cereals, corn, and oilseed rape.

Honeybee colonies, weight scales, and statistical analysis

In July 2018, we selected 12 equally sized *Apis mellifera carnica* colonies, each featuring a 1-year-old queen obtained from the Bavarian State Research Centre for Viticulture and Horticulture (LWG). These queens represent a typical lineage of *Apis mellifera carnica* commonly found in Germany. The colonies were housed in single 10-frame Zander hive boxes whose volume of approximately 45 L matches the natural cavity volume preferences of temperate *Apis mellifera* (Seeley and Morse, 1976). We placed the colonies at various sites inside the forest or along its edges. The average altitude of the hive locations was 403 m a.s.l. (range: 354–439). Spatial independence was achieved by placing neighboring colonies around 2 km or more apart, as detailed in Rutschmann et al. (2023). Each hive was set up on a weight scale (Capaz BEE HIVE SCALE GSM 200) that recorded hive weight once every hour and forwarded data via mobile communication.

The only beekeeping interventions conducted included treatments against varroosis using formic acid in August 2018 (1 year prior to the swarming season). Additionally, to ensure adequate overwintering resources, colonies that were deemed to have insufficient honey stores had one or two largely empty frames replaced with full honeycombs sourced from other colonies. During preliminary data analysis, we addressed potential weight artifacts arising from hive maintenance manipulations. Due to the malfunctioning of weight scales at two locations, our investigation was limited to recordings from 10 of the 12 initially selected colonies. Swarm departures were identified as abrupt weight losses exceeding 500 g within an hour. This threshold (500 g) was chosen as a compromise to avoid missing small swarms while minimizing false detections caused by environmental or random fluctuations in colony weight, based on previous studies on swarming behavior, which reported mean swarm weights well above 1 kg (Fell et al. 1977; Burgett and Morse 1974). We calculated swarm weight (as a measure of swarm size) by comparing hive weights the hour before and after the swarm's departure. To estimate the average number of bees in swarms, we considered the empirical mean weight of European worker honeybees in swarms of 129.5 mg (Otis, 1982). Although the weight of swarming bees varies depending on the level of engorgement with honey prior to departure and the proportion of drones in swarms (Otis, 1982), using an average value should provide

reasonable approximations for the *average* number of bees in the swarms of our study. All swarms issued throughout the 2019 season were recorded. To evaluate the weather conditions during swarming events, we retrieved average daily temperatures and precipitation data from nearby weather stations, provided by Agrarmeteorologie Bayern (www.wetter-by.de).

To test whether swarm category (prime or afterswarm) has a statistically significant effect on swarm weight, we performed a linear mixed-effects model analysis using the lme4 package in R (Bates et al., 2015). Weight was the response variable, swarm category was the predictor, and colony ID was included as a random effect to account for variability among colonies. Additional models incorporating Julian date as a fixed effect were tested but did not improve the explanatory power of the model (Figure SII). Residuals of the models were inspected with DHARMA package (Hartig and Hartig, 2017). For inference, a Type II ANOVA was performed using the Anova function from the car package (Fox et al., 2012). All statistical analyses were performed

using R software (version 4.3.1; R Core Team, 2016). For data processing and graphical representation of the results, we utilized ‘tidyverse’, ‘ggplot2’, ‘patchwork’, and ‘ggpol’ (Wickham, 2017, 2016; Pedersen, 2020; Tiedemann, 2020).

Results

We recorded a total of 17 swarming events across ten colonies of *Apis mellifera carnica* during the 2019 swarming season, resulting in an average of 1.7 (SD: ± 0.8) swarms per colony (1 prime swarm and 0.7 afterswarms) (Fig. 1A). All colonies swarmed at least once: five colonies issued a single swarm; three colonies produced two swarms (a prime swarm and one afterswarm); and two colonies gave rise to three swarms (a prime swarm and two afterswarms). Swarms departed between May 17 and June 27, and the median swarming date was May 30 (Fig. 1B). The interval between prime swarms and afterswarms averaged 9 ± 3 days (range: 6–13 days, $N=5$). For the two colonies that produced

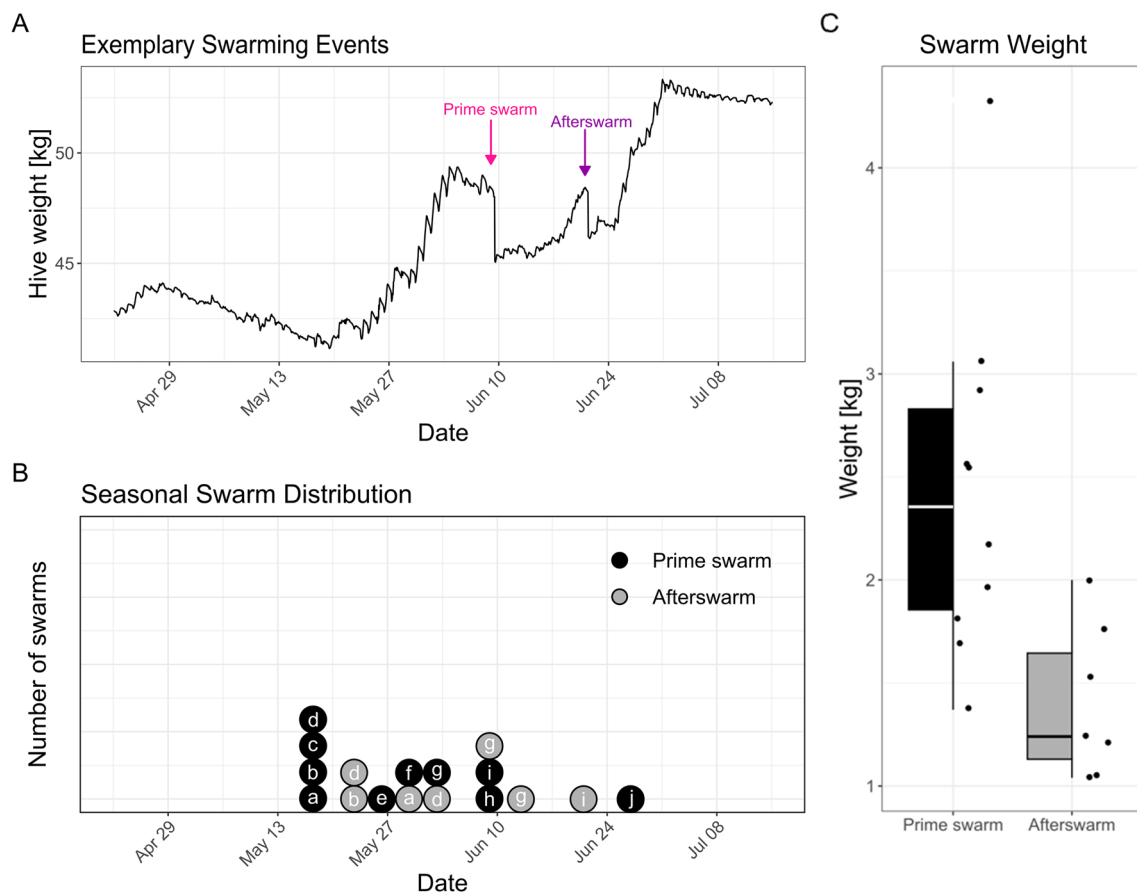


Fig. 1 Overview of swarming activity in a forest environment and its characteristics. **A** Example of hive weight recordings from colony “i” during the swarming period, where arrows indicate the occurrence of a prime swarm (pink) and an afterswarm (violet). **B** Temporal distribution of swarms throughout the season, with letters representing the different parental colonies. **C** A boxjitter plot displays the weight of the recorded prime swarms and afterswarms.

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two afterswarms, the interval between the first and second afterswarm was 4 and 8 days. The median departure time for swarms was 11:00 AM (range: 8:30 AM to 5:30 PM CEST; UTC + 2). Swarming usually occurred when it was getting warmer, following a period of relative cold, and importantly, on days without rain (Fig. 2).

We observed notable variation in swarm weight, with an overall mean and SD weight of 2.01 ± 0.86 kg (range: 1.04–4.33 kg). This variability stemmed mainly from the significant differences between prime swarms, averaging 2.44 ± 0.86 kg, and afterswarms, at 1.40 ± 0.37 kg (Fig. 1C) ($\chi^2 = 21.276$, $df = 1$, $p < 0.001$). Assuming that the average weight of workers in swarms is 129.5 mg, we estimated an average of 18,857 bees in prime swarms and 10,844 bees in afterswarms.

Discussion

Our study provides insights into the natural swarming behavior and the natality rates of unmanaged *Apis mellifera carnica* colonies in a German forest environment. It offers a critical reference point for ongoing and future demographic investigations of wild-living honeybees in Europe.

Comparison with other studies

The observed average of 1.7 swarms falls within the overall range of 0–4 swarms per colony and year recorded for temperate honeybees (reviewed by Winston, 1991). However, a comparison between individual studies reveals notable variations between swarming rates. This highlights the impact of

geographical location and subspecies differences on swarming frequency.

In Kansas, USA, Winston (1980) recorded 3.6 daughter colonies per year from a combination of *Apis mellifera carnica* and *Apis mellifera ligustica* colonies, thus a significantly higher natality rate than reported here. In British Columbia, Canada, Lee and Winston, (1987) observed an annual reproductive rate of 2.2 swarms. Seeley (1978, 2017) reported that the probability for colonies to enter a swarming cycle was around 0.9 in the Arnot forest in New York State. Since this number does not account for afterswarms, it resembles our observations more closely. A low propensity to swarm was reported by Fries et al. (2006) for colonies on Gotland, an island in the Baltic Sea, where the proportion of reproducing colonies (with at least one swarm) varied between 0 and 60% in six observation years. Similarly, Le Conte et al. (2007) documented variable swarming rates in untreated colonies resistant to *Varroa destructor* in France, with an average rate of $41.50 \pm 9.94\%$ across 6 years.

Otis and Wearing-Wilde (1992) pointed to the significant contribution of afterswarms to the overall number of swarms produced in a population, and our finding of an average of 0.7 afterswarms per colony supports that conclusion. Five out of ten colonies in our study produced more than the prime swarm. This shows that it is important to use methods that can detect afterswarms to accurately estimate natality rates.

In regions warmer than temperate zones (e.g., tropical or subtropical regions), natality rates for other subspecies have been found to be substantially higher (Winston et al. 1983). This underscores the significant role local environmental conditions (such as climate) and biological factors play in shaping reproductive strategies of honeybees (Ruttner, 1988;

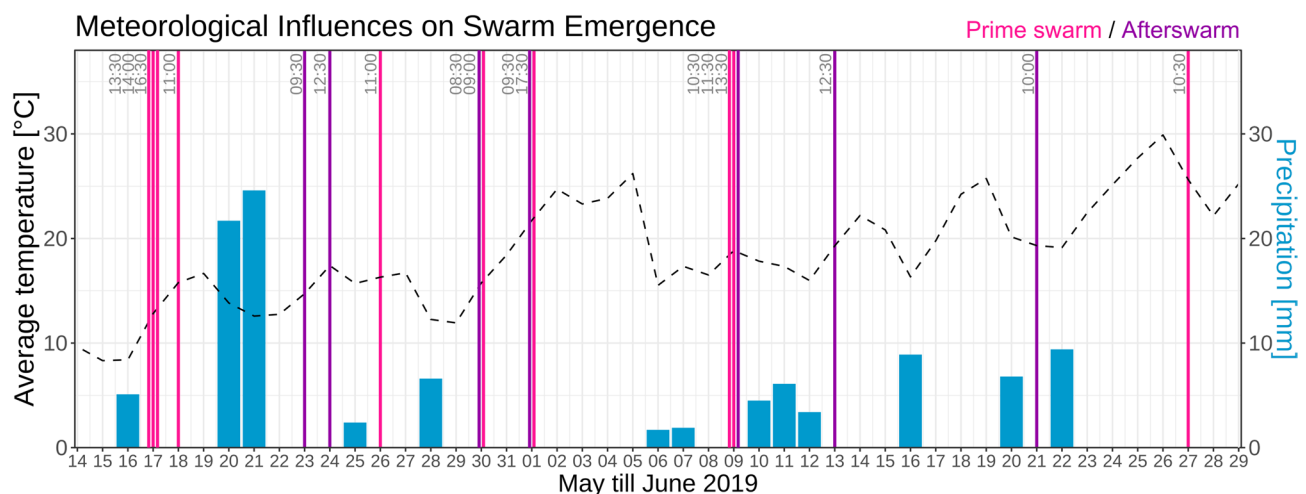


Fig. 2 Average temperature and precipitation in the study area from mid-May to late June 2019. The blue bars represent daily precipitation in millimeters, while the black dashed line represents the daily

average temperature in degrees Celsius. Vertical lines indicate the occurrence of prime swarms (pink) and afterswarms (violet), with the corresponding time of day labeled (CEST; UTC + 2).

Nuru et al., 2002; Strange et al., 2007; Uzunov et al., 2014; Norrström et al., 2021). Additionally, factors such as the queen's age (younger queens exhibit increased pheromone production, thereby more effectively suppressing the development of new queens) and the size of the hive have been demonstrated to influence swarming behavior (Uzunov et al., 2014; Smith et al., 2017). For example, Loftus et al. (2016) showed a clear relationship between hive size and swarming propensity; small hives (42 L) had a higher swarming rate, with 10 out of 12 colonies swarming, compared to larger hives (168 L), where only 2 out of 12 colonies swarmed.

Besides providing data on swarming rate, this study points to the large variation in swarm sizes that exist within a single season and population, noting marked differences between primary swarms and afterswarms. Prior research revealed a broad spectrum of swarm sizes, ranging from as few as 2,400 bees (0.31 kg) to as many as 41,500 bees (5.33 kg), with mean populations reported at 11,800 bees (1.53 kg) by Fell et al. (1977) and 14,000 bees (1.8 kg) by Burgett and Morse (1974). However, it should be noted that estimating the number of workers from swarm weight may be imprecise due to variation in the body mass of individual bees, which can be influenced by the amount of honey carried in their crops (Combs, 1972; Otis, 1982). Additionally, the smaller worker populations and unmated queens associated with afterswarms lower their survival prospects (Lee and Winston, 1987).

Swarming and net reproductive rate

Assessing the self-sustainability of a honeybee population necessitates a thorough understanding of its annual survival and natality rates. The net reproductive rate (R_0), defined for honeybees by the formula $R_0 = s + s \times n$ (Kohl et al., 2022), integrates survival rate (s) and natality rate (n) (i.e., the number of swarms). This index helps in identifying whether populations are diminishing ($R_0 < 1$), maintaining stability ($R_0 = 1$), or experiencing growth ($R_0 > 1$), intrinsically—i.e., without considering immigration from external sources such as managed apiaries (Kohl and Rutschmann, 2024).

If we assume that our finding of 1.7 swarms per colony and year, based on observations from a single year, represented the average annual reproductive rate in our study system, a wild honeybee population would require a minimum annual survival rate of 37% to achieve population sustainability ($R_0 = 1$). (It has to be emphasized that winter survival is only one component of annual survival and that the annual survival rate refers to the average of all colonies, new swarms/founders, and established colonies combined.) Notably, such an annual survival rate is considerably higher than the observed rates of 10.6% reported by Kohl et al. (2022) and 12% by Rutschmann et al. (2024), underscoring the urgent need for further investigations into the factors

affecting survival and reproduction to inform conservation and management strategies.

The role of landscape in the timing of swarming

Resource availability on the landscape scale plays a pivotal role in the survival of wild-living honeybee colonies (Rutschmann et al., 2022; Kohl et al., 2023) and the establishment of new swarms (Seeley, 1978; Seeley and Visscher, 1985; Otis and Wearing-Wilde, 1992; Seeley, 2017). A comparison with two other phenological data sets also suggests that the timing of swarming is affected by landscape context. Henneken et al. (2012), utilizing a crowdsourced dataset of 1,335 swarming events in Germany during 2011, found that swarming began in April with a first peak in early May. In addition, a predominantly urban dataset from 2019, the same year as our study, which included 77 swarm observations (55 from Munich and 22 from across Germany), reported swarming as early as 17 April (Rutschmann et al., 2024). While the median swarming date in the urban setting was around 19 May 2019 (Remter, Roth, Rutschmann, personal communication), the median swarming date in this study was 30 May 2019. The delayed timing of swarms in our study region might be explained by inferior foraging conditions in forest-dominated compared to agro-urban landscapes. We know from a study on foraging behavior in the same study region that, especially in April and May during an important phase of colony weight gain, honeybees find superior foraging opportunities in open areas such as grassland and cropland (Rutschmann et al. 2023). Colonies nesting in forests might also face cooler temperatures and fewer hours of sunlight compared to colonies in non-forest landscapes and thus might show delayed development in spring due to fewer hours available for foraging (Czekońska et al., 2023). An additional possible influence on the delayed onset of swarming observed in forest settings could be the lack of stimulative feeding practices that are commonly used in urban/agricultural settings by many beekeepers. Additionally, the temporary placement of pollen traps for 1–3 days each month during the foraging season as part of a different study may also have contributed to a slight delay.

Future directions and methodological advancements

The adoption of weight scales for the surveillance of swarming events introduces a non-invasive, efficient approach to studying the reproductive patterns of honeybees. We encourage the broader application of this technology to detect swarming in weight monitoring initiatives, such as McMinn-Sauder et al. (2024), Johannesen et al. (2022), Lecocq et al. (2015), Kuchling et al. (2018), Komasilova et al. (2023), or Prešern et al. (2019). Combining weight monitoring with

weather data, as demonstrated in our study, opens up exciting possibilities for predicting swarming events. Future research could explore integrating weight monitoring with other remote colony monitoring methods, such as temperature, vibration, and acoustic or video surveillance, to further enhance the precision and scope of swarming monitoring systems (Zacepins et al., 2016; Ramsey et al., 2020; Minaud et al., 2024).

While our study focused on a limited number of colonies ($N=10$) observed during a single season, future research with a larger sample size, spanning multiple years, could explore differences in swarming across various landscapes, climatic regions, altitudes, and subspecies. Additionally, an important area for future investigation is the survival chances of primary and afterswarms. Such studies would deepen our understanding of the factors influencing honeybee reproduction and survival.

Supplementary Information The online version contains supplementary material available at <https://doi.org/10.1007/s00040-025-01028-y>.

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Author contributions BR conceived the study, analyzed and visualized the data, and wrote the first draft of the manuscript. PLK provided assistance in the field. All authors reviewed and edited the manuscript.

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Data availability The data that support the findings of this study are openly available in Zenodo at <https://doi.org/10.5281/zenodo.14814245>.

Declarations

Conflict of interest The authors have declared no conflict of interest.

Ethical approval No honeybees were harmed during this study, which used non-invasive monitoring methods. Ethical approval was therefore not required.

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