Evolution of sex differences in cooperation: the role of trade-offs with dispersal

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Statement of authorship

PC-L and AJY designed the study. PC-L, AMB, XH, TR, JEY, DLC, LW and AJY contributed to long-term field data collection. PC-L, NB, CR, MN and AJY set up Encounternet system. PC-L performed statistical analysis with input from NB, XH, MN and AJY. PC-L and AJY wrote the first draft of the manuscript, and all authors contributed substantially to revisions.

Data accessibility statement

All R scripts and datasets needed to reproduce the analyses presented in this paper are available at: https://github.com/PabloCapilla/Sex_Diffs_cooperation_dispersal. Should the manuscript be accepted, a DOI to this data repository will be provided.

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Abstract

Explaining the evolution of sex differences in cooperation remains a major challenge. Comparative studies highlight that the more philopatric sex tends to be more cooperative than the more dispersive sex, but we do not understand why. Here we test the two leading evolutionary explanations for this association for the first time in a wild cooperative bird, using a novel automated radio-tracking study. Our findings provide no support for the widely invoked ‘philopatry hypothesis’, which proposes that the more philopatric sex helps more as it gains a greater direct fitness benefit from cooperation. Instead, our findings support the lesser known ‘dispersal trade-off hypothesis’, which proposes that the more dispersive sex helps less because investments in extra-territorial prospecting for dispersal opportunities trade-off against cooperation. This simple explanation for the evolution of sex differences in cooperation could apply widely across taxa as it does not require that cooperation yields a direct fitness benefit.

Introduction

The evolution of sex differences in cooperation remains a major evolutionary puzzle (Clutton-Brock *et al.* 2002; Cockburn 1998; Downing *et al.* 2018; Fenner *et al.* 2023; Williams & Hale 2007; Woolfenden & Fitzpatrick 1978). In many cooperatively breeding species, in which offspring of both sexes delay dispersal from their natal group and help to rear future generations of their parents' young, one sex contributes significantly more than the other to cooperative care within the natal group (Canestrari *et al.* 2005; Clutton-Brock *et al.* 2002; Fenner *et al.* 2023; Williams & Hale 2007; Zöttl *et al.* 2016). These sex differences in natal cooperation are of particular interest as they occur in the absence of sex differences in relatedness to recipients (as helpers of both sexes are still residing within their natal groups), leaving it unlikely that kin selection alone can explain their evolution (Clutton-Brock *et al.* 2002; Downing *et al.* 2018; Fenner *et al.* 2023; Williams & Hale 2007; Woolfenden & Fitzpatrick 1978). Recent comparative analyses have highlighted that, across cooperative breeders of this kind, sex differences in contributions to cooperative care while within the natal group are predicted by the species' sex difference in dispersal; resident offspring of the more philopatric sex tend to contribute more to natal cooperation than resident offspring of the more dispersive sex (Downing *et al.* 2018; Fenner *et al.* 2023). However, it is not clear why the more philopatric sex contributes more to natal cooperation, as the two major evolutionary explanations proposed for this pattern have not been teased apart (Clutton-Brock *et al.* 2002; Downing *et al.* 2018; Fenner *et al.* 2023; Williams & Hale 2007; Young *et al.* 2005).

The leading hypothesis for the evolution of sex differences in natal cooperation, the 'Philopatry hypothesis', proposes that offspring of the more philopatric sex help at higher rates (while both sexes reside within the natal group) because, by staying in their natal group for longer they stand to gain a greater direct fitness benefit from cooperation (Clutton-Brock *et al.* 2002; Downing *et al.* 2018; Woolfenden & Fitzpatrick 1978). This hypothesis therefore requires that cooperation yields a direct fitness benefit and that the accrual of this benefit is to some extent contingent upon remaining within the natal group. For example, the more philopatric sex may be more likely to ultimately breed within the natal territory and may thus stand to gain a greater downstream direct benefit from cooperating to rear additional group members that may ultimately help them back (Clutton-Brock *et al.* 2002; Cockburn 1998; Downing *et al.* 2018; Woolfenden & Fitzpatrick 1978). A recent comparative study of the cooperative birds in which both sexes delay dispersal and help has found support for this idea (Downing *et al.* 2018); species with more female-biased probabilities of natal breeding showed more female-biased helper contributions to cooperative care while within the natal group. These

findings are consistent with the key prediction of the Philopatry hypothesis and could therefore constitute rare evidence that direct fitness benefits have played a widespread role in the evolution of helping (Downing et al. 2018; see also Kingma 2017). However, these findings are also consistent with an alternative less-studied hypothesis which also predicts that the more philopatric sex should help more, regardless of whether helping yields a direct benefit; the 'Dispersal trade-off hypothesis'.

The Dispersal trade-off hypothesis proposes that offspring of the more dispersive sex help at lower rates (while both sexes reside within the natal group) because all individuals face a trade-off between investments in natal cooperation and activities that promote dispersal from the natal group, such as extra-territorial prospecting for dispersal opportunities (Williams & Hale 2007; Young *et al.* 2005). Thus, while the Philopatry hypothesis requires that cooperation yields a downstream direct fitness benefit, the Dispersal trade-off hypothesis does not. In many social species offspring prepare for dispersal by conducting extra-territorial prospecting forays from their natal group, which are thought to yield information on dispersal opportunities in the surrounding population (Doerr & Doerr 2005; Kingma *et al.* 2016b, a; Williams & Hale 2007; Woolfenden & Fitzpatrick 1978; Young *et al.* 2005, 2007). Investments in prospecting could trade-off against natal cooperation because the two activities cannot be carried out at the same time, and because costs and physiological changes associated with prospecting could reduce the expression of helping (Kingma *et al.* 2016b; Maag *et al.* 2022; Ridley *et al.* 2008; Young *et al.* 2005; Young & Monfort 2009). Frequent prospectors could also reduce their costly contributions to helping in order to maintain competitive phenotypes that reduce the risks and/or increase the likely success of prospecting (Heinsohn & Legge 1999). However, little is known about prospecting due to the difficulty of monitoring the cryptic, fast and often long-distance movements that prospectors make (Doerr & Doerr 2005; Kingma *et al.* 2016b; Maag *et al.* 2022; Williams & Hale 2007; Young *et al.* 2005). One study of male meerkats, *Suricata suricatta*, whose frequent extra-territorial forays yield extra-group matings (Young *et al.* 2007), suggests that investments in such forays may indeed trade-off against cooperation (Young *et al.* 2005). But whether prospecting for dispersal opportunities *per se* (which may occur at lower rates than mating forays and could entail different costs) can also trade off against natal cooperation, as envisaged under the Dispersal trade-off hypothesis, is not yet known.

As the Philopatry and Dispersal trade-off hypotheses for the evolution of sex differences in cooperation *both* predict that helpers of the more philopatric sex should contribute more to cooperative care while within the natal group (consistent with the findings of comparative studies; Downing et al. 2018; Fenner et al. 2023),

discriminating between them now requires innovative field studies to test whether the different underlying processes assumed by each hypothesis are acting as required to explain the observed sex differences in cooperation.

Here, we combine long-term data on life-histories and cooperative behaviour with a high-resolution automated radio-tracking study of prospecting to test both the predictions and core assumptions of the Philopatry and Dispersal trade-off hypotheses, using wild cooperatively-breeding white-browed sparrow weavers, *Plocepasser mahali*, as a model system. White-browed sparrow weavers (hereafter 'sparrow weavers') are cooperative birds that live in social groups and defend year-round territories across the semi-arid regions of sub-Saharan Africa. Within each group, a single dominant male and female monopolise reproduction and their offspring of both sexes delay dispersal, becoming non-breeding subordinates that cooperatively help to feed future generations of nestlings produced by the breeding pair (Harrison *et al.* 2013a). As offspring of both sexes never breed while subordinate (Harrison *et al.* 2013a), the only route to natal breeding is via inheritance of the natal dominant (breeding) position. The primary route to dominance for both sexes, however, is via dispersal to other groups (Harrison *et al.* 2014). Dispersal is typically local in both sexes but genetic analyses and field observations suggest that males disperse further than females from their natal to breeding sites and thus that females may be the more philopatric sex (Harrison *et al.* 2014); a rare reversal of the typical avian sex-bias in dispersal (Greenwood 1980). Here we establish whether there are also clear sex biases in dispersal incidence and natal breeding position inheritance; the dispersal traits most relevant to the rationale of the focal hypotheses. Subordinates of both sexes conduct extra-territorial prospecting forays prior to dispersal from their natal group, which are commonly met with aggression from territory holders (Harrison *et al.* 2014; York *et al.* 2019). The rarity of extra-group parentage by subordinates (<1% of paternity; Harrison et al. 2013a) suggests that the primary function of these forays is to promote dispersal.

Subordinate female sparrow-weavers cooperatively feed offspring at substantially higher rates than subordinate males (Capilla-Lasheras *et al.* 2021); a pattern rarely documented among birds (Williams & Hale 2007). Here we demonstrate that this marked sex difference in cooperation is evident among subordinates within their natal groups, in the absence of a sex difference in relatedness to recipients (Figure 1). Helper contributions are known to increase the overall rates at which nestlings are fed (Capilla-Lasheras *et al.* 2021) and may yield indirect fitness benefits by reducing environmentally-induced variance in the reproductive success of related breeders (Capilla-Lasheras *et al.* 2021), lightening the workloads of related breeders

(Capilla-Lasheras *et al.* 2023; O'Callaghan 2021) and enhancing the quality of their offspring (Wood 2017). Helping could also conceivably yield direct fitness benefits, but as subordinate immigrants rarely help at all (Walker *et al.* 2021) it seems unlikely that these benefits are substantial (see discussion for more detail). Helping behaviour could, for example, yield a downstream direct fitness benefit to helpers if they inherited the dominant (breeding) position in their natal group and were then helped by offspring that they had previously helped to rear; the mechanism commonly envisaged in the Philopatry hypothesis (Downing *et al.* 2018; Woolfenden & Fitzpatrick 1978).

First, we demonstrate that the sex differences in dispersal and cooperation in sparrow-weaver societies satisfy the predictions of both the Philopatry and Dispersal trade-off hypotheses: this species shows a rare reversal of the typical avian sex biases in both dispersal incidence and natal cooperation. We then test for the underlying processes assumed by the Philopatry and Dispersal trade-off hypotheses to establish whether either hypothesis can explain why this rare example of female-biased philopatry in birds is accompanied female-biased natal cooperation. Regarding the Philopatry hypothesis, we test whether female helpers are indeed more likely than males to inherit the breeding position within their natal group; the most commonly invoked mechanism by which the more philopatric sex could gain a greater downstream direct benefit from cooperation (Clutton-Brock *et al.* 2002; Downing *et al.* 2018). Regarding the Dispersal trade-off hypothesis, we use a high-resolution automated radio-tracking study of extra-territorial prospecting to test its two key assumptions: first, that male helpers (the more dispersive sex) prospect at higher rates or over greater distances or for longer durations than females; second, that investments in prospecting from the natal group trade off against concurrent investments in natal cooperative care.

Methods

Study system and life-history monitoring

This work was conducted as part of the long-term field study of white-browed sparrow weavers in Tswalu Kalahari Reserve in the Northern Cape Province of South Africa (27°16'S, 22°25' E). Fieldwork was carried out from September to May between 2007 and 2016 inclusive. All social groups within a study area of approximately 1.5 km^2 were monitored each year; approximately 40 groups, each defending a small year-round territory. Throughout the study period, all birds in the study population were fitted with a single metal ring, three colour rings for identification (under SAFRING license 1444) and genotyped using 10 microsatellite loci (see Supplementary text A in Capilla-Lasheras et al. [2021] and Harrison et al. [2013a] for

genotyping protocols and relatedness calculations). The birds could be reliably caught from their woven roost chambers at night using custom-made sweep nets, and were then returned to their chambers to pass the remained of the night on completion of any processing. Social groups were easily distinguishable as all group members foraged together within their territory, defended it against neighbouring groups, and roosted within woven chambers in a single tree or cluster of trees close to the territory centre (Harrison *et al.* 2013a; Walker *et al.* 2016). Each sparrow weaver group contains a single dominant (reproductive) pair and a variable number of non-breeding subordinates (Harrison *et al.* 2013a). Dominants are easily distinguished from subordinates because they display a distinct set of behaviours (Walker *et al.* 2016), and the sexes are easily distinguished after the first six months of life because the sub-species that we study (*P. mahali mahali*) exhibits a beak colour sexual dimorphism (Leitner *et al.* 2009). Group compositions were assessed every week in the morning throughout each field season (Capilla-Lasheras *et al.* 2021), and were confirmed by periodic captures of all group members from their roost chambers at night. We also sought to confirm the dominance status of birds on all such visits. Once a bird acquired a dominant position they retaining dominance for life. Dominance tenures typically ended with the bird's immediate disappearance from the study population; a scenario that likely reflects their death, given the very local nature of dispersal in this species and the relative rarity with which unmarked adults enter our study population from outside (Harrison *et al.* 2014). On rare occasions dominants did transfer into a dominant position in another group, but on no occasion was a dominant bird known to subsequently become subordinate (either in their own group or any other). Dispersal was deemed to have occurred when an individual that was previously resident in one group (i.e., foraging with the group in the day and roosting on their territory at night) transferred to become resident in another group. All offspring delaying dispersal from their natal group and those that do disperse typically make just one or two dispersal movements in their lifetimes (Harrison *et al.* 2013a, 2014). During our field observations from October 2011 onwards, we also noted any visits to groups by birds that were currently resident in other groups (i.e., prospecting birds) were also noted. Wherever possible we noted the colour-ring combinations of these prospecting birds, but aggression from the territory residents often left this impossible (see results). Throughout each field season, we also checked every territory for nests (conspicuous woven structures with a different shape to roosting chambers) every one or two days, and wherever nests were present we check them for the presence of newly laid eggs. When new eggs were found we monitoring egg laying through to clutch completion and the fate of the brood through to fledging (Harrison *et al.* 2013a).

Quantifying the contributions of natal subordinates to cooperative provisioning

To determine the rates at which helpers within their natal groups (i.e., all subordinate, and therefore non-breeding, group members still residing within the group in which they born) cooperatively provision the nestlings of the dominant pair, provisioning behaviour was recorded for all breeding attempts throughout the study period in which the nestlings survived to the standard recording days. A video camera was placed on a low tripod below the nest to record the birds entering the nest on the mornings of the 6th to 12th days (inclusive) after the first egg in the focal clutch hatched (days that cover the period of highest nestling demand). At least 5 days before video recording started, we caught every bird in the group apart from the dominant female to confirm group composition and mark each bird's vent feathers with a unique black dye mark to facilitate their identification on provisioning videos. On recording days, video cameras were set up at a standardized time following sunrise to record provisioning for approximately 3 hours per nest per day (yielding a total of 289 days of recording over the 177 monitored broods in groups with one or more helpers within their natal group; median of 2, range 1 - 6, days per brood). Within-nest cameras have confirmed that all nest visits in which the birds are not conspicuously carrying grass (events that were readily excluded from our provisioning rate calculations below), entail the visiting bird carrying a single food item to the nest and delivering it to the chicks (Walker 2015). The videos were subsequently transcribed to determine the timing of each provisioning visit, the duration of the visit (from the bird entering the nest to exiting), and the identity of the provisioning bird (using its unique vent mark and information from its leg rings and bill colouration; which differs between the sexes). Whenever the bill of the provisioning bird was visible on their approach to the nest, we also categorised the size of the food item being delivered; simply 'large' if it was evidently longer than the bird's bill or 'short' if it was not.

We used the resulting provisioning data set to calculate the number of provisioning visits that each group member made during each observation period (termed 'provisioning number' within our methods). In some cases, we were unable to reliably identify all group members, adding uncertainty to our provisioning number estimates. We therefore excluded any provisioning number estimates derived from observation periods in which there were five or more provisioning visits in which the feeder was of uncertain identity (72 rate estimates excluded). To address the residual uncertain in the provisioning number estimates derived from the observation periods containing one to four feeds by birds of uncertain identity, we set each bird's provisioning number estimate for these observation periods to be the integer midway between the lowest and highest possible provisioning number for that bird in that period given any partial identity information that was available for the feeders of uncertain identity. This yielded a final dataset containing 1,338 daily

provisioning number measures for subordinates (helpers) of known sex within their natal groups for whom age was known with less than 180 days of uncertainty.

Quantifying the prospecting behaviour of natal subordinates

We used Encounternet (Mennill *et al.* 2012; Rutz *et al.* 2012) to identify extra-territorial prospecting movements by natal subordinate sparrow weavers. This system is a wireless sensor network formed by an array of stationary radio receivers (termed 'base stations') that log the radio-signals of any animal-mounted transmitters (termed 'tags'; used here in transmit-only mode) within range. The logs collated by the base stations provide information on the tag (and therefore animal) ID detected, the date and time of detection and the signal strength (expressed as Received Signal Strength Indicator [RSSI] values). Between February and April 2017, we fitted Encounternet tags to 32 adult subordinate birds, using a figure-of-eight leg harness made of a stretchable, porous material that was stuck to the tag. The total mass of the tag and harness was ~1.2g (2.6-3.1% of the body mass of the tagged birds at fitting; Figure 3a). The birds were tracked for an average of 37 days (range = 11–59 days) generating a total of 22,518,022 detection logs (individual base-stations log all tags within detection range every 5 seconds). Field observations after tag fitting confirmed that none of the tagged birds showed any signs of affected behaviour or impaired movement, and our results suggest that they still regularly engaged in extra-territorial movements. All tags were retrieved from the birds before the end of April 2017. The tagged birds were all (non-breeding) subordinates that were resident on their natal territory (hereafter, their 'home' territory) at the time of tagging and remained so for the duration of the Encounternet deployment. All tags were set to transmit ID-coded pulses every five seconds. To track the locations of the tags across our study site, we set up an array of 35 Encounternet base station receivers, each placed in the central tree (typically the main roost tree) of 35 contiguous sparrow weaver territories (Figure S1). See Supplementary text 1 for further detail on Encounternet deployment and base station calibration.

We processed Encounternet logs for each tag separately, working through the continuous time series of logs for each tag to estimate the locations of the tagged bird throughout the tag's deployment. A given tag's logs were processed in moving windows of 15 seconds and in time steps of 5 seconds. For each of these 15-second windows, we attempted to assign the tagged bird a 'best estimate' location following a conservative set of rules depicted as a flow diagram in Figure S3 (see also Supplementary text 1). We then processed the resulting time series of 'best estimate' locations for the tagged bird to identify extra-territorial prospecting

forays using a conservative approach that should minimise the chance of false positives (Figure S3 and Supplementary text 1).

Using this dataset, we identified prospecting forays as events that satisfied both of the following criteria: (i) a single 'time period' or set of contiguous time periods for which the tagged bird had been assigned known 'best estimate' locations that were not the tagged bird's home territory for a total of >15 seconds, and (ii) the furthest assigned 'best estimate' location during this contiguous non-home period was >250 metres away from the centre of their home territory (we set this distance threshold to >250m a *priori*, see Supplementary text 1 for further details). This approach was designed to minimise the chance that the bird's territorial interactions with its neighbours while 'at home' are misclassified as extra-territorial forays. While setting the distance threshold at 250m makes sense *a priori* based on the biology of the bird (Supplementary text 1), we verified via a sensitivity analysis that our key finding of a sex difference in prospecting rate was not specific to this threshold setting (Figure S4).

For each foray, we estimated 'foray duration' as the time elapsed between the focal bird's first 'best estimate' location outside their home territory and their subsequent first 'best estimate' location back on their home territory. We estimated 'foray distance' as the linear distance between the bird's home territory base station (approximately the centre of their home territory) and the furthest 'best estimate' location that they were assigned during the foray. These 'foray distances' could therefore markedly underestimate the true distance travelled during the foray, because (i) the bird could have travelled on a complex path to and from the outer point on the foray, taking in many groups, and (ii) the furthest 'best estimate' location could be much closer to home than the true furthest location visited by the bird during the foray, as the former is constrained to be the location of a base station within our array. If prospecting birds did travel beyond the boundary of our receiver array, such forays would still be detected (despite having potentially long periods of 'unknown' locations while far from the array) if the bird passed within receiving range of a base station >250m from their 'home' base station, either on their outbound or return journey. The final dataset included extra-territorial forays for 27 tagged birds (Supplemenaty text 1).

Statistical methods

We employed a full model approach, evaluating the statistical importance of each predictor using likelihood-ratio tests (LRT). If not statistically significant, interactive terms were removed from initially-specified full models to ease the interpretation of the effects of non-interactive terms. All non-interactive fixed effect

predictors were retained within the full model regardless of their significance. All statistical analyses were carried out using R 4.3.1. (R Core Team 2023). Unless otherwise stated, statistical models were fitted using the R package 'lme4' (v1.1.29; Bates et al. 2015). Over-dispersion and zero-inflation of GdLMMs were checked by simulating scaled model residuals in the R package 'DHARMa' (v0.4.6; Hartig 2018).

1. Modelling the age-specific dispersal probabilities of natal subordinates

We investigated whether sparrow-weaver subordinates within their natal groups exhibit a sex difference in the age-specific probability of dispersal from their natal group, by fitting a binomial generalised linear mixed model (GdLMM) to data reflecting whether or not they dispersed within a given year-long age window of their lives as a natal subordinate. All individuals hatched into our population between September 2007 and April 2014 that reached six months of age contributed to our analysis (individuals under six months of age very rarely disperse) and we utilised group census data up until April 2016 to determine if and when they dispersed (giving all individuals a minimum of 2 years of monitoring in which to disperse). The lives of each of the focal individuals were divided into year-long 'age windows' from birth onwards. Each year-long age window that the individual started as a subordinate within its natal group (and before April 2016) contributed a line of data to the data set, and was flagged to reflect whether they were (1) or were not (0) observed to have dispersed away from their natal group to another study group within our population during that year of life. Our analysis therefore focussed on known dispersal events (see Supplementary text 2 for a discussion of unknown dispersal events). This approach resulted in a data set of 812 year-long windows for 338 individuals (178 males and 160 females) monitored as natal subordinates within 36 social groups. The full model contained the following fixed effect predictors: the subordinate's sex, age class during the focal year-long age window (as a 4-level categorical variable: <1 year old; 1-2 years old, 2-3 years old, and > 3 years old; this last age class therefore captured any year-long age windows in which the focal individual was 3 years or older at the start of the age window), and the interaction between its sex and age class. Breeding season of hatching, individual ID and natal group ID were included as random effect intercepts.

2. Modelling the cooperative provisioning traits of natal subordinates

To investigate whether there is a sex difference in the nestling provisioning (helping) rates of natal subordinates, we fitted a zero-inflated negative binomial GdLMM using the R package 'glmmTMB' (v1.1.7; Brooks et al. 2017). We modelled provisioning rate (feeds / hour) by fitting the birds' 'provisioning number' measure as the response term (the number of provisioning visits it conducted during a given ~3 hour observation period; see above), with the duration of the observation period fitted as an offset. We included

the following fixed effect predictors in the full model: the sex of the bird, their age (a 4-level factor: < 1 year, 1 year – 2 years, 2 years – 3 years, and > 3 years), the interaction between sex and age (given the possibility that the magnitude of sex differences changes with age), brood size (as a continuous variable, 1 to 3 nestlings) and brood age (a 7-level factor, 6 to 12-day old, given the potential for non-linear effects). Breeding year, social group ID, clutch ID and individual ID were included as random intercepts. The model was fitted to 1,338 observations of provisioning number by 314 different natal subordinates feeding 177 broods in 38 social groups. We found evidence for zero-inflation (zero-inflation parameter ± SE = -1.605 ± 0.181) and over-dispersion (dispersion parameter = 2.87), which were accounted for by our zero-inflated negative binomial model. To investigate whether there is a sex difference in natal subordinate provisioning visit durations (time spent within the nest chamber, from entry to exit, during which the provisioning bird interacts with the brood), we fitted a linear mixed model to the provisioning visit durations of the 5,034 provisioning events that natal subordinates conducted during the observation periods monitored above. The full model included the same fixed and random effect predictors as the analysis of provisioning rate described above. Provisioning visit duration was 'log+1' transformed to fulfil the assumption of normality in model residuals. To investigate whether there is a sex difference in the size of the food items that natal subordinates provisioning to broods, we fitted a binomial GdLMM to the binomial prey item size information (large or small) collected from the 1,325 provisioning events in which the focal bird's bill was visible prior to entering the nest (see above). We analysed the probability that the prey item was large, with a full model that included the same fixed and random effect predictors as the analysis of provisioning rates described above.

3. Modelling the probability of acquiring dominance within the natal group

We investigated sex differences in the probability that natal subordinates inherit a dominant (breeding) position within their natal group using binomial GdLMMs, As the probability of inheriting natal dominance (and any sex difference therein) may change with a natal subordinate's age, we conducted different models each focussing on the dominance inheritance prospects of subordinates of a given age: one model at 0.5 years of age, one at 1 year of age, one at 2 years of age and one at 3 years of age. The data set for each model comprised all birds that were known to be subordinate within their natal group at the focal age, and modelled whether or not they ultimately became dominant within their natal group. Subordinates were only included in the analysis if dominance turnovers were monitored within their natal group for a minimum of 2 years after the date on which the subordinate reached the focal age. Sex was included as a fixed effect predictor in these models. Breeding season of hatching and natal group ID were included as random intercepts. We also verified that similar patterns were reflected in a simple contingency table test for a sex

difference in the proportion of all dominance acquisition events for a given sex in which the winner took dominance within their natal group. As subordinates could also conceivably gain a downstream direct fitness benefit from natal helping if they take a dominance position via 'territorial budding' from the natal territory (i.e., founding a new group on territory previously held by their natal group), we then repeated our modelling analysis above including within the 'natal dominance inheritance events' events in which the focal bird did not inherit dominance within the natal *group* but instead became dominant by founding a new group via probable 'territorial budding'. Using the 88 dispersal events in the dataset, we used binomial GdLMMs to explain the probability of dispersing into subordinate or dominant positions, including sex as a fixed effect predictor, and breeding season of hatching and natal group ID as random intercepts. We analysed dominance tenure length using a linear mixed model, including sex as a fixed effect and, breeding season of hatching and natal group ID as random intercepts. This model used data from 66 dominance tenures (32 males and 32 females) acquired by individuals hatched in the study population.

4. Modelling the prospecting behaviour of natal subordinates

To investigate whether there is a sex difference in the daily rate of extra-territorial prospecting (number of forays per day) we employed a Poisson GdLMM. To investigate whether there is a sex difference in mean 'foray duration' and mean 'foray distance' (see above) we employed LMMs. In each of these three models, we included the same set of fixed effect predictors: age of the individual (as a continuous variable), sex, provisioning status of their home group (2-level factor [yes/no] to capture whether their home group was provisioning nestlings at the time), and the interaction between sex and provisioning status. In all three models we included bird ID and home territory ID as random intercepts. In the model of the daily rate of prospecting, we also included observation ID as a random intercept, to account for overdispersion (Harrison 2014). All three models used data from the 27 individuals with working tags (see above; 13 males and 14 females).

To investigate the existence of a trade-off between cooperative nestling provisioning and prospecting, we tested whether these traits were correlated at the individual level. For this analysis, we used the Encounternet data from the 13 tagged subordinates (6 males and 7 females) whose prospecting rates were quantified during a nestling feeding period within their group. For these birds we therefore had daily estimates of their prospecting rate, as well as estimates of their nestling provisioning rate from those mornings in which provisioning observations were also conducted (see above). To estimate a proxy for each bird's general level of prospecting effort, we calculated the bird's prospecting rate over a given number of

days preceding each day of provisioning observations (see below). The full model included this proxy for the focal bird's prospecting effort as the main fixed effect predictor of interest in a GdLMM of daily provisioning rate (implemented using a Poisson process to model the number of nestling feeds conducted by a given bird during each observation watch and including the duration of the observation watch as an offset). We also included subordinate sex, brood age (as a continuous predictor in this case due to lower sample size), brood size and the interaction between prospecting rate and subordinate sex as fixed effects. Group ID and bird ID were included as random effect intercepts. The sample size for the analysis was 43 daily measures of provisioning rate for the 13 tagged subordinates. The analyses presented within the main paper use the focal bird's prospecting rate calculated over a window of three days prior to the focal daily measure of their provisioning rate, as we thought it plausible *a priori* that any energetic or stress-related costs of prospecting might accumulate and be evident over this timescale. Recognising that the choice of three days is somewhat arbitrary though, we confirmed through sensitivity analyses that the observed negative covariance between prospecting rate and provisioning rate was still evident when prospecting rate was calculated over a range of time window lengths (Figure S2).

Results

1. Testing the Predictions of the Philopatry and Dispersal trade-off Hypotheses

Consistent with the predictions of both hypotheses, sparrow weavers show a significant male-bias in dispersal incidence coupled with a significant female-bias in contributions to natal cooperation. Modelling the age-specific probability of dispersal from the natal group revealed that subordinate natal males are significantly more likely to disperse than subordinate natal females across all age classes (Figure 1a, Table S1). There was also evidence that dispersal probability increased with age (Figure 1a, Table S1), but no evidence that the sex difference in dispersal probability changed with age (Table S1). The Philopatry and Dispersal trade-off hypotheses would therefore both predict that subordinate females (the more philopatric sex) should help at higher rates than subordinate males (the more dispersive sex) while within the natal group.

Subordinate females did indeed cooperatively feed broods at significantly higher rates than subordinate males while within the natal group (Figure 1b, Table S2). Cooperative provisioning rates also increased initially with advancing age before decreasing slightly (Figure 1b, Table S2). There was no indication that the magnitude of the sex difference in cooperation changed with age (Table S2). Brood size and brood age both positively predicted cooperative provisioning rates (Table S2). During their provisioning visits, subordinate

females also spent slightly but significantly longer than subordinate males within the nest chamber (Figure 1c, Table S3), and were just as likely as subordinate males to provision nestlings with food items that were large (Figure 1d, Table S4). These sex differences in the cooperative provisioning rates of subordinates within the natal group occurred in the absence of sex differences in their relatedness to the nestlings being fed (Figure 1e, Table S5).

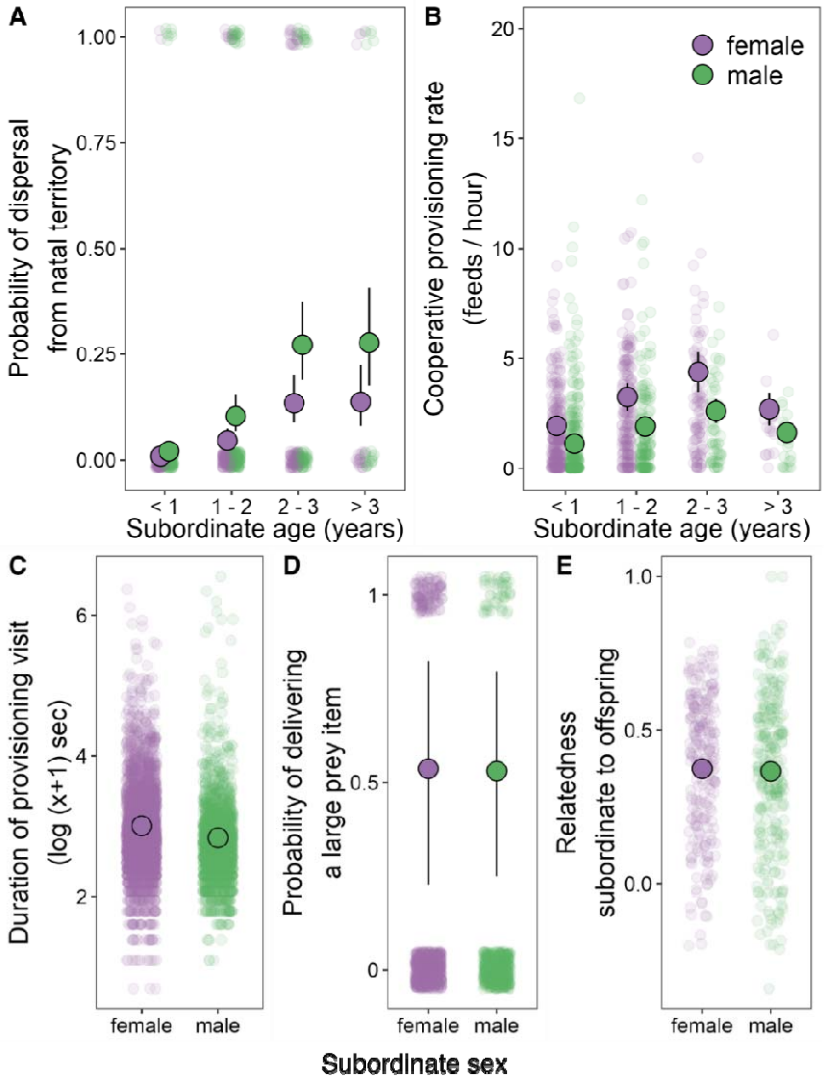

Figure 1. **Sparrow-weavers show a rare reversal of the typical avian sex differences in dispersal and natal cooperation.** Among subordinate birds residing within their natal groups, (**A**) males (green points) showed significantly higher age-specific probabilities of dispersal than females (purple points; Table S1), while (**B**) females showed significantly higher cooperative nestling provisioning rates than males (a sex difference that did not vary with age; Table S2). (**C**) Natal subordinate females also spent slightly but significantly longer in the nest per provisioning visit than males (Table S3), and (**D**) were just as likely as natal subordinate males to feed the nestlings with food items that were large (Table S4). (**E**) These sex differences in natal cooperation occurred in the absence of a sex difference in relatedness to the nestlings being fed (Table S5). Translucent points represent raw data whereas larger points and error bars give mean model predictions ± SE (where error bars cannot be seen, they are smaller than the radius of the point presenting the mean).

2. Testing the Mechanisms Invoked by the Philopatry Hypothesis

Contrary to expectations under the Philopatry hypothesis, females were no more likely than males to obtain a dominant breeding position within their natal group (Figure 2; for all age classes of natal subordinates, Table S6-S9). For natal subordinates under 3 years of age, the probability of inheriting natal dominance was low (<10% on average; Figure 2) and showed no sex difference, yet sex differences in natal cooperation were clear at these ages (Figure 1b). Once subordinates reached 3 years of age within their natal group their probability of inheriting natal dominance markedly increased (to ~ 25%), but their contributions to natal cooperation markedly decreased (Figure 1b). If we also included in these 'natal dominance acquisition' events any dominant breeding positions acquired *outside* the natal group by founding a new group within territory previously held by the natal group (i.e., territorial budding; Woolfenden & Fitzpatrick 1978), we still found no evidence of a sex difference in the probability of natal dominant acquisition in any age class (Table S10-13). Overall, dominance turnover events rarely involved subordinates winning dominance within their natal group, and the sexes did not differ in this regard (just 7 natal dominance wins of 32 total turnovers for males; 9 natal of 32 total for females; $\chi^2_1 = 0.08$, $p = 0.773$); in *both* sexes dominance was typically acquired via dispersal to other groups. Dominance turnover events were also rare in both sexes, as average dominance tenure lengths are long and do not differ between the sexes (mean ± S.E.: males = 2.62 ± 0.32 years, range 0.18 – 5.84 years; females = 2.51 ± 0.33 years, range 0.16 – 7.51 years; Table S14).

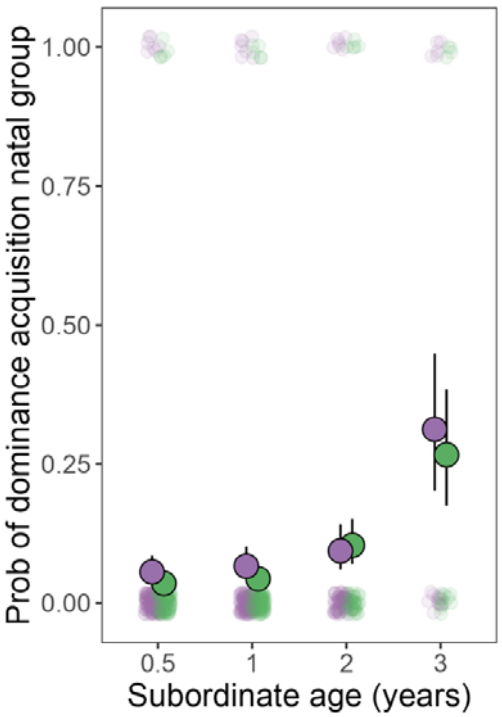

Figure 2. Age-specific probabilities of inheriting a dominant breeding position within the natal group, for subordinates still residing within their natal group at various ages. Subordinate females (purple points; the more cooperative sex; Figure 1b) were no more likely to inherit a natal breeding position than subordinate males (green points; Table S6-S9). Our analyses focussed on subordinates within their natal group at a given focal age and tested for an effect of sex on their probability of inheriting natal dominance at some later time. The solid points show the sex-specific means (± SE) for the focal age class, and the translucent points represent raw data. Subordinates were only included in the analysis if dominance was monitored within their focal group for at least 2 years after the date on which the subordinate reached the focal age.

3. Testing the Mechanisms Invoked by the Dispersal Trade-off Hypothesis

Direct observations of prospecting

On 678 occasions, resident birds were observed visiting other territories. While they were often chased by residents, hampering identification, the visitor was identified for 293 visits, and in 184 (62.8%) of these they were known to be subordinates resident in another group, who must therefore have been prospecting. Of these 184 prospecting forays by subordinates, 115 (62.5%) were conducted by 68 males and 69 (37.5%) were conducted by 51 females, which suggests a male-bias in prospecting incidence given the approximately equal adult sex ratio in our population (115 of 184 forays is a significant deviation from 50%; Binomial test $p < 0.001$).

Automated tracking of prospecting behaviour

To verify this sex difference in prospecting incidence and study the consequences of variation in prospecting rates, we deployed an automated radio-tracking technology to provide continuous high-resolution information on the prospecting movements of 27 radio-tagged subordinate adults residing within their natal groups (13 males and 14 females; Figure 3a). After processing the 22,518,022 detection logs from the base station receiver array (see Methods), we detected a total of 971 extra-territorial prospecting forays. We used a conservative approach to identify forays (see Methods): events in which the signal-strength distribution across the receiver array suggested that the bird's closest base-station was >250m away from the centre of their own territory (where there was also always a base-station). As base-stations were situated in the centres of all territories within the tracking study area and the mean ± S.E. distance between neighbouring territory centres was just 93.7 m ± 4.56 m, these events should typically constitute forays beyond the territory centres of neighbouring groups (see Methods). This approach will minimise the chance that a resident bird's territorial interactions with its neighbours are incorrectly interpreted as extra-territorial prospecting, but may underestimate foray frequency by excluding more local prospecting. These 971 forays principally occurred between 06:00h and 19:00h, with frequencies peaking in the cooler periods towards the start and end of the day (Figure 3b). The tagged bird's estimated daily rates of prospecting varied widely from 0 to 26 forays per day (n = 895 daily rate measures from 27 tagged birds; mean = 1.09 forays / day; median = 1 foray / day). Forays had a mean duration of 18.59 minutes (median = 5.67 minutes; range = 0.42 – 493 minutes). Our best estimate of the round-trip distance travelled per foray had a mean of 720.74 m (median = 647.60 m; range = 502.29 – 2026.15 m), though this is likely an underestimate as our ability to detect longer-distance forays was constrained by the spatial coverage of our receiver array (Figure S1), and as the estimate was

calculated as twice the straight line distance to the bird's furthest location (while the birds could have taken a more complex path).

Consistent with our direct observations above, subordinate males showed significantly higher daily rates of prospecting from their natal groups than subordinate females (Figure 3e, Table S17). Prospecting rates were modestly but not significantly lower during nestling provisioning periods (i.e., when cooperative care occurs) than at other times (Figure 3e, Table S17). The magnitude of the sex difference in prospecting rate was not affected by whether the bird's social group was provisioning nestlings at the time (Table S17). There was no evidence of an effect of subordinate sex (or nestling provisioning periods) on either the duration (Figure 3c, Table S18) or estimated distance of forays (Figure 3d, Table S19). Older subordinates conducted forays of shorter duration (Table S18) but age did not predict prospecting rate (Table S17) or distance (Table S19).

Focussing on 13 radio-tagged subordinates (6 males and 7 females) whose forays were tracked through a nestling provisioning period, we found evidence suggestive of a trade-off between prospecting and cooperative provisioning. An individual's prospecting rate over the three-day window prior to us quantifying cooperative contributions to nestling provisioning negatively predicted its cooperative provisioning rate (Figure 3f, Table S20). We found no evidence that either the intercept or gradient of this apparent trade-off regression line differed between the sexes (i.e., there was no sex effect on provisioning rate while accounting for prospecting rate and no significant sex x prospecting rate interaction; Table S20), though the sample size for this comparison was small. The existence of a negative correlation between prospecting and cooperative provisioning rate was also robust to the choice of time window over which we calculated prospecting effort (Figure S2).

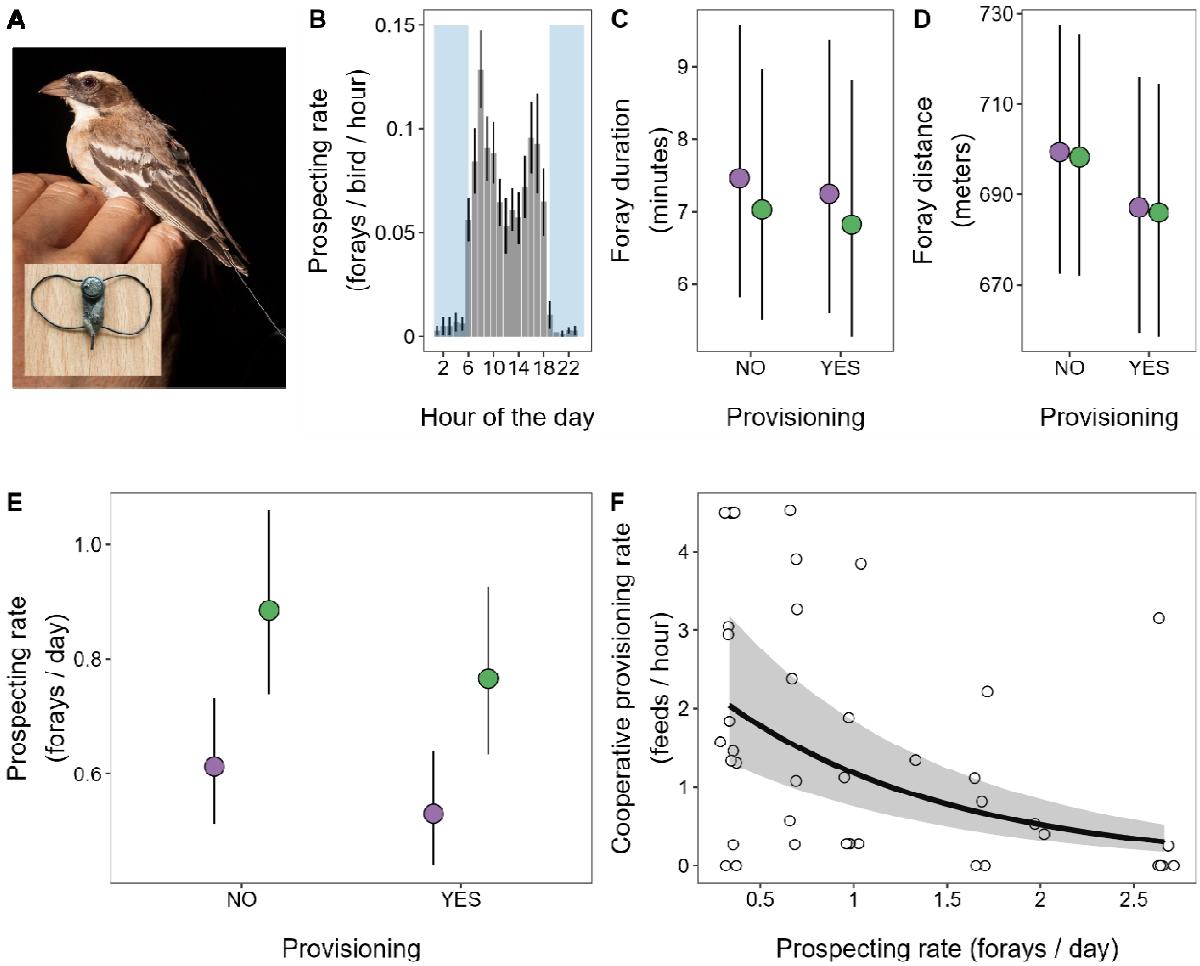

Figure 3. Extra-territorial prospecting behaviour and its trade-off with cooperation. (**A**) A subordinate female (females have pale bills) fitted with an Encounternet radio-tag (pictured inset). The tag is fitted with leg loops and sits on the bird's lower back. The aerial can be seen sitting along the tail feathers. (**B**) Circadian variation in prospecting behaviour; mean hourly prospecting rates per bird per hour of the day (mean ± SE; from the 27 radio-tracked birds). The shaded area illustrates night hours (using the local civil twilight time on March 11th 2017, the middle date of our tracking study). The rare forays at night could reflect birds being flushed from their roost chambers by predators. We found no evidence of sex differences in either the (**C**) mean duration of forays (Table S18) or (**D**) mean estimated round-trip distance travelled on forays (Table S19). (**E**) Subordinate males have a significantly higher daily prospecting rate than subordinate females, both inside and outside peak periods of cooperative nestling provisioning (Table S17). (**F**) Individual variation in prospecting rate negatively predicted cooperative provisioning rate for the 13 radio-tagged birds with prospecting data during a nestling provisioning period (Table S20; the plot shows >13 data points as we have repeated measures per individual). There was no evidence of a sex difference in the intercept or gradient of this regression line (Table S20). Male and female model estimates in **C**, **D** and **E** illustrated in green and purple colour respectively. **C**, **D**, **E** and **F** present mean model predictions ± SE.

Discussion

Our findings reveal that white-browed sparrow weavers show a rare reversal of the typical avian sex difference in dispersal: males show higher age-specific probabilities of dispersal from their natal group than females, and both population genetics and observed dispersals suggest that males disperse further than females from their natal to breeding sites (Harrison *et al.* 2014). Undetected longer-distance dispersals (beyond the bounds of our study site) may also be more common among males, as the low rate of immigration into our study population also shows a male bias (Harrison *et al.* 2014). Both the Philopatry and Dispersal trade-off hypotheses for the evolution of sex differences in cooperation would therefore predict that sparrow-weavers show female-biased natal cooperation, and our findings confirm this. Natal subordinate females helped to feed nestlings at higher rates than natal subordinate males across the age classes, and spent longer within the nest on their visits. This coupling of rare reversals of the typical avian sex biases in both dispersal and cooperation provides novel support for the predictions of the focal hypotheses, which are otherwise difficult to test given the limited variation in dispersal sex biases within taxonomic classes (Downing *et al.* 2018; Fenner *et al.* 2023; Williams & Hale 2007). Below, we consider the potential for the Philopatry and Dispersal trade-off hypotheses to account for the evolution of female-biased natal cooperation in this species, drawing on our empirical tests of the underling mechanisms assumed by each. Our findings suggest that the mechanisms envisaged in the Philopatry hypothesis are unlikely to be acting in this species, and that the evolution of female-biased cooperation can be more readily explained by the Dispersal trade-off hypothesis. We consider the wider implications of this finding for our understanding of the evolution of sex differences in cooperation and, more broadly, the role that direct fitness benefits have played in the evolution of helping behaviour across cooperative societies.

Among the diverse hypotheses proposed for the evolution of sex differences in cooperation (Cockburn 1998; Fenner *et al.* 2023), the Philopatry and Dispersal trade-off hypotheses are unusual in predicting the evolution of female-biased natal cooperation in sparrow-weaver societies; alternative hypotheses cannot readily explain this pattern. That the observed sex difference in natal cooperation occurs in the absence of a sex difference in average relatedness to recipients suggests that it cannot be attributed to a sex difference in the indirect benefits of cooperation (Clutton-Brock *et al.* 2002; Downing *et al.* 2018; Fenner *et al.* 2023). The heterogamety hypothesis (Cockburn 1998; Fenner *et al.* 2023; Whitney 1976) proposes that effects of the sex chromosomes on patterns of genetic relatedness could leave the heterogametic sex (females in birds) less cooperative, yet we see the opposite pattern here. Similarly, the paternity uncertainty hypothesis

(Charnov 1978; Cockburn 1998; Fenner *et al.* 2023) predicts male-biased cooperation when extra-pair paternity occurs, yet we see the opposite pattern here despite dominant males losing 12-18% of paternity to other dominant males (Harrison *et al.* 2013b). The parental skills hypothesis (Cockburn 1998; Fenner *et al.* 2023) proposes that the sex that invests most in parental care could contribute most to helping if (i) helping yields a direct benefit by improving parenting skills, and (ii) this benefit is larger for the sex that contributes most to parental care. This hypothesis could be relevant to sparrow-weavers as dominant females feed offspring at higher rates than dominant males (Walker *et al.* 2021). However, compelling evidence that helping improves parenting skills remains elusive (Chesterton *et al.* 2023; Cockburn 1998), and as immigrant subordinate sparrow-weavers rarely help at all (Walker *et al.* 2021), it seems unlikely that direct benefits arising via skills acquisition are a major driver of helping in this system. It has also been suggested that the sex that shows higher variance in lifetime reproductive success (LRS) may invest more in helping given its lower chance of securing direct fitness by breeding (Cockburn 1998; Koenig *et al.* 1983). However, male and female sparrow-weavers likely have very similar variances in LRS, as both sexes only breed as dominants and do so in socially monogamous pairs (Harrison *et al.* 2013a) with no evident sex difference in dominance tenure length (see Results). If anything, competition for extra-group paternity among dominant males (Harrison *et al.* 2013b) may leave males with slightly higher variance in LRS than females, but this would lead to the prediction of male-biased natal cooperation, and we observe the reverse.

The Philopatry hypothesis proposes that offspring of the more philopatric sex contribute more to natal cooperation because, by staying in their natal group for longer, they stand to gain a greater downstream direct benefit from natal cooperation (Clutton-Brock *et al.* 2002; Downing *et al.* 2018; Woolfenden & Fitzpatrick 1978). The most commonly invoked mechanism by which the more philopatric sex could gain a greater direct benefit from cooperation is if they were more likely to ultimately breed in their natal group and thus benefit from having helped to increase the size of the natal workforce (Clutton-Brock *et al.* 2002; Downing *et al.* 2018; Woolfenden & Fitzpatrick 1978). However, this mechanism cannot readily explain female-biased natal cooperation in sparrow weavers, as neither sex breeds within their natal group while subordinate (Harrison *et al.* 2013a), and subordinate females were no more likely than males to inherit a dominant breeding position within their natal group or territory. Our findings also give cause to question more broadly whether helping in this species yields any form of direct benefit that is contingent upon inheriting natal dominance. If it did, one might expect the markedly higher chance of inheriting natal dominance beyond three years old (Figure 2) to be accompanied by a concomitant increase in natal cooperation, yet we observe the opposite (Figure 1b). This pattern may instead reflect subordinates showing greater restraint

from costly helping once they become credible contenders for dominance (Cant & Field 2001). As subordinate males and females are equally likely to inherit natal dominance (Figure 2), the observed male-bias in age-specific dispersal probability (Figure 1a) appears to reflect a sex difference in the *strategy* used to acquire dominance via dispersal, rather than in the *incidence* of acquiring dominance via dispersal. Indeed, our data suggest that males take a more complex route to dominance that often involves early dispersal into subordinate positions in other groups prior to dominance acquisition from there, while females more commonly disperse directly into dominant positions elsewhere (see Supplementary text 3).

A second mechanism by which the more philopatric sex might gain a greater downstream direct benefit from natal helping (regardless of whether they inherit natal dominance), is if helping to rear offspring augments natal group size and living in a larger group improves survival while subordinate. Where this is the case, the more philopatric sex may enjoy this downstream direct benefit for longer because it will tend to remain in the natal group for longer (Clutton-Brock *et al.* 2002). However, this mechanism cannot readily account for the female-biased natal cooperation observed here for two reasons. First, recent analyses strongly suggest that subordinate sparrow-weavers suffer markedly *lower* survival rates in larger social groups (O'Callaghan 2021), which would leave the more philopatric sex suffering a greater downstream *cost* (rather than benefit) from cooperative group augmentation. Second, helping does not appear to augment group size in sparrow-weavers, as helper numbers (whether female helpers, male helpers or both) do not positively predict either offspring survival to fledging or the breeding rate of dominant females. Indeed, it seems unlikely that helping yields an appreciable direct benefit of any kind in sparrow-weaver societies (e.g., via other mechanisms such as "pay to stay" or signalling "quality"; Clarke 1989; Gaston 1978; Reyer 1986; Zöttl et al. 2013), as subordinates rarely contribute to cooperative care at all following dispersal from their family groups (Harrison *et al.* 2013a; Walker *et al.* 2021). Together, our findings suggest that the Philopatry hypothesis cannot readily account for the evolution of female-biased natal cooperation in this species.

The Dispersal trade-off hypothesis proposes that the more dispersive sex contributes less to natal cooperation because all individuals face a trade-off between activities that promote dispersal (such as prospecting for dispersal opportunities) and natal cooperation (Williams & Hale 2007; Young *et al.* 2005). It thus envisages that sex differences in cooperation arise not from a sex difference in the direct benefits of cooperation but from a trade-off between cooperation and other fitness-relevant traits in which sex differences exist. Our findings support the key prediction of this hypothesis, that male-biased dispersal should be accompanied by female-biased natal cooperation, and provide evidence in support of the

underlying mechanisms that it envisages. First, our automated radio-tracking study revealed that natal subordinates of both sexes prospect, but males (the more dispersive sex) prospect at significantly higher rates than females, while making forays of comparable duration and distance. Our direct observations of prospectors visiting our study groups also reflect this male-bias in prospecting incidence. Subordinate males may benefit more from prospecting than females because they are more likely than females to disperse in all age classes, and because they disperse further on average than females between their natal and breeding sites (Harrison *et al.* 2014). While forays in some species function in extra-group mating (Double & Cockburn 2000; Peters 2002; Young *et al.* 2007), the forays of subordinate sparrow-weavers do not appear to serve this purpose (as both within- and extra-group paternity are monopolised by dominant males (Harrison *et al.* 2013b, a). That this male bias in prospecting was apparent during nestling provisioning periods highlights its potential to drive sex differences in cooperative care. That male-biased prospecting was also apparent *outside* nestling provisioning periods is important too, as this suggests that the sex difference in prospecting is unlikely to be a *product* of the sex difference in natal cooperation (i.e., a reversal of the causal direction suggested above), as it also occurs when subordinates are not engaged in cooperative care.

Our automated radio-tracking study also revealed support for the second key assumption of the Dispersal trade-off hypothesis: that investments in prospecting trade off against investments in cooperative care. Individuals that prospected at higher rates during nestling provisioning periods contributed significantly less to cooperative care. Consistent with the sexes facing a similar trade-off between prospecting and cooperation, and thus differing in their mean levels of cooperative care because they differ in their mean prospecting rates (the rationale of the Dispersal trade-off hypothesis), we found no evidence that this apparent trade-off regression line differed in either intercept or slope between males and females, though our sample size for this comparison was small. As prospecting and cooperative care are both likely to entail energetic costs that may compromise body condition (Heinsohn & Legge 1999; Kingma *et al.* 2016b; Young *et al.* 2005), the observed negative association between prospecting and cooperation could reflect a simple resource allocation trade-off between the two traits (potentially mediated by endocrine changes associated with prospecting (Peters 2002; Young *et al.* 2005). While these forays could well entail intense energetic expenditures, the total energetic costs entailed in active prospecting and any recovery from it might not be terribly high, as the detected forays occurred on average at modest daily rates (though our analyses could underestimate this) and were often short in duration. However, prospecting is also likely to entail significant risks, as prospectors are frequently attacked when visiting other groups (Kingma *et al.* 2016b, a; Williams & Hale 2007; Young *et al.* 2005) and could face elevated predation risk while moving alone (Maag *et al.* 2022).

As such, regular prospectors may benefit more than others from maintaining a competitive phenotype (to mitigate the risk of injury and enhance dispersal success), which could itself trade-off against investments in costly cooperation, compounding the effects of any energetic and/or time costs entailed in the conduct of forays. As both of these trade-off scenarios are consistent with the rationale of the Dispersal trade-off hypothesis, this hypothesis seems well placed to explain the evolution of female-biased natal cooperation in this species: males contribute less to natal cooperation because male-biased dispersal leaves them prospecting more, and all individuals face a trade-off between prospecting and natal cooperation.

Our study has tested both the predictions and assumptions of the Philopatry and Dispersal trade-off hypotheses for the evolution of sex differences in cooperation. Our findings suggest that the mechanisms envisaged in the Dispersal trade-off hypothesis are better placed to explain the evolution of female-biased natal helping in sparrow-weaver societies than those envisaged in the Philopatry hypothesis. This is important because the Philopatry hypothesis is commonly invoked as the likely driver of the association between sex differences in philopatry and cooperation across taxa (e.g., Clutton-Brock et al. 2002; Downing et al. 2018; Fenner et al. 2023), but the Dispersal trade-off hypothesis also predicts this pattern (Fenner *et al.* 2023; Williams & Hale 2007; Young *et al.* 2005) and could apply more widely across taxa as it does not require that helping yields a direct fitness benefit contingent upon remaining in the natal group (a scenario whose generality is unclear). The mechanisms envisaged in the Philopatry hypothesis could well apply in some species, such as those in which helping augments group size and in which group augmentation has the potential to yield a downstream direct benefit to helpers (e.g., Florida scrub jays, *Aphelocoma coerulescens* [Woolfenden & Fitzpatrick 1978] and meerkats [Clutton-Brock et al. 2002]). Indeed, they may act in concert with those envisaged in the Dispersal trade-off hypothesis to select for sex differences in helping in such species (e.g., both Florida scrub jays and meerkats also show sex differences in helper prospecting; Woolfenden & Fitzpatrick 1978; Young et al. 2005, 2007). But our findings highlight the need for caution when extrapolating the Philopatry hypothesis to species in which helping may not yield direct fitness benefits or in which mechanisms for the differential accrual of downstream direct benefits by the more philopatric sex have yet to be identified. On balance, it seems likely that the mechanisms envisaged in the Philopatry and Dispersal trade-off hypotheses have both played a role in the evolution of sex differences in natal cooperation, but their relative importance may vary markedly among species.

Our findings have wider implications too for the role that direct benefits may have played in the evolution of cooperation. Recent comparative studies have highlighted that the probability that helpers will breed within

their natal group predicts helper contributions to cooperative care across taxa (Downing *et al.* 2018; Kingma 2017), and it has been suggested that these findings constitute rare evidence that direct fitness benefits of helping have played a widespread role in the evolution of helping, alongside kin selection (Downing *et al.* 2018; Fenner *et al.* 2023; Kingma 2017). However, our findings highlight that such associations could also arise in the absence of direct benefits of helping, via trade-offs between dispersal and cooperation: helpers with higher probabilities of natal breeding might generally contribute more to natal cooperation across taxa because they need not invest as much in preparations for dispersal. Thus, while direct benefits of helping may indeed be important in some contexts (e.g., Clutton-Brock et al. 2002; Reyer 1986; Zöttl et al. 2013), how widespread a role they have played in the evolution of helping behaviour across taxa would seem to remain an open question.

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