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## **The eco-evolutionary landscape of power relationships between males and females**

**Authors:** Eve Davidian<sup>1,\*</sup>, Martin Surbeck<sup>2</sup>, Dieter Lukas<sup>3</sup>, Peter M. Kappeler<sup>4,5</sup>, Elise Huchard<sup>6,\*</sup>

<sup>1</sup> Ngorongoro Hyena Project, Ngorongoro Conservation Area, Arusha, Tanzania; email: [davidian.ceve@gmail.com](mailto:davidian.ceve@gmail.com); Twitter: @HyenaProject; ORCID: 0000-0001-9437-0720

<sup>2</sup> Harvard University, Department of Human Evolutionary Biology, Cambridge, MA, USA; email: [msurbeck@fas.harvard.edu](mailto:msurbeck@fas.harvard.edu); Twitter: @HarvardHEB; ORCID: 0000-0003-2910-2927

<sup>3</sup> Department of Human Behavior, Ecology and Culture, Max Planck Institute for Evolutionary Anthropology, Leipzig, Germany; email: [dieter\\_lukas@eva.mpg.de](mailto:dieter_lukas@eva.mpg.de); ORCID: 0000-0002-7141-3545

<sup>4</sup> German Primate Center, Behavioral Ecology Unit, Leibniz Institute of Primate Biology, Göttingen, Germany; email: [pkappel@gwdg.de](mailto:pkappel@gwdg.de) ; ORCID: 0000-0002-4801-487X

<sup>5</sup> University of Göttingen, Department of Sociobiology & Anthropology, Germany

<sup>6</sup> Anthropologie Évolutive, Institut des Sciences de l'Évolution de Montpellier (ISEM), France ; email : [elise.huchard@umontpellier.fr](mailto:elise.huchard@umontpellier.fr); ORCID: 0000-0002-6944-449X

\* These authors contributed equally

Corresponding authors: Davidian, E. ([davidian.ceve@gmail.com](mailto:davidian.ceve@gmail.com)), Huchard, E.

([elise.huchard@umontpellier.fr](mailto:elise.huchard@umontpellier.fr))

**Abstract.** In animal societies, control over resources and reproduction is often biased towards one sex. Yet, the ecological and evolutionary underpinnings of male-female power asymmetries remain poorly understood. We outline a comprehensive framework to quantify and predict the dynamics of male-female power relationships within and across mammalian species. We show that male-female power relationships are more nuanced and flexible than previously acknowledged. We then propose that enhanced reproductive control over when and with whom to mate predicts social empowerment across ecological and evolutionary contexts. The framework explains distinct pathways to sex-biased power: coercion and male-biased dimorphism constitute a co-evolutionary highway to male power, whereas female power emerges through multiple physiological, morphological, behavioural, and socio-ecological pathways.

**Keywords:** intersexual power inequality; sexual conflict; social dominance; sexual size dimorphism; reproductive control; social evolution

## **What is power?**

Power, defined as the ability to elicit particular behaviours in others [1,2], is a pervasive yet elusive characteristic of social relationships. Identifying the ultimate and proximate underpinnings of power can provide insights into the emergence, dynamics and consequences of social and gender inequalities in human and animal societies; topics of increasing societal significance [3–6]. Power can be divided into two main domains: ‘resource-holding’ and ‘decisional’ power [7]. Resource-holding power encapsulates the degree of control that one individual can exert over another when competing over resources and mating opportunities [1,2,8]. It emerges through three main non-mutually exclusive

mechanisms: (i) **coercion** (see Glossary), where asymmetries in coercive potential may for example arise from differences in intrinsic and social **attributes**, experience, or **motivation** [5], (ii) **deception**, whereby individuals manipulate the information available to others [9], and (iii) **trade**, whereby some individuals may have a bargaining advantage or ‘**leverage**’ because they possess valuable resources and services that can be exchanged (‘commodities’) but cannot be taken by force [1,8,10]. Decisional power, also termed ‘leadership’, commonly refers to the ability of an individual to influence the behaviour of others in ways that generate collective activities in various contexts, such as movement, foraging, hunting, and intergroup conflict [3,4,7]. Asymmetries in resource-holding power between individuals of the same and different sex are prevalent in nonhuman mammalian societies and presumably have a deeper and more direct impact on individual survival and reproduction than decisional power [6,11]. These asymmetries may thereby play a key role in shaping the **social organisation** and **mating system** of mammalian societies and will be the focus of this review.

## **A novel framework for studying power relationships between males and females**

Here we outline a comprehensive framework to describe, quantify and predict ecological and evolutionary variations in resource-holding power relationships between males and females (hereafter, “**intersexual power**”) within and across mammalian societies. We introduce three new principles for the framework. First, intersexual power arises from conflicts over **reproductive control** and social control (Box 1). Second, features of the mating system predict intersexual biases in reproductive control (Box 2). Third, enhanced reproductive control by one sex predicts its social empowerment.

## **Brief overview of traditional approaches to the study of intersexual power**

## *The historical hypotheses*

Early empirical and theoretical studies of intersexual power have largely focused on the evolutionary drivers of female-biased power [12]. In the centre of attention were a handful of species where females were considered ‘masculinised’ because they possess behavioural, physiological or morpho-anatomical traits that are more strongly expressed among males in most mammals, such as large body size, aggressiveness, levels of androgens, or erectile external genitalia – as in many lemurs [13–15], spotted hyenas (*Crocota crocuta*) [16], rock hyraxes (*Procavia capensis*) [17], and meerkats (*Suricata suricatta*) [18]. A predominant set of hypotheses has therefore emphasised the role of intrinsic attributes on female coercive potential [12,15,19]. They posited that females outcompete males when they match or surpass males in size, weaponry or fighting ability. Extensions of these hypotheses later integrated the role of behavioural and hormonal masculinisation in female empowerment [13,14,16,20]. A second set of hypotheses emphasised asymmetries in motivation that may arise from differences in life history between males and females. They posited that females are more prone to engage in and to win contests with males when they gain higher payoffs than males from accessing resource, e.g., in species where females may incur particularly high reproductive costs [21,22]. A third set of hypotheses highlighted asymmetries in trading ability between the sexes and the role of female leverage through mate choice, whereby females preferentially mate with submissive, deferent or non-coercive males [1,23,24].

## *The main conceptual and methodological limitations*

These hypotheses and our current understanding of intersexual power have suffered from long-standing stereotypical views of sex roles – as emphasized in pioneering work by women primatologists and anthropologists [e.g., 25,26] – and several other methodological and

conceptual limitations [12]. First, previous studies often equated power with coercive social control (also termed 'social dominance') [8,12] that is, control that is acquired and maintained forcefully over resources that are primarily related to an individual's self-maintenance and survival, such as food, shelter, shade or territory. These approaches thereby do not account for (i) alternative, non-coercive mechanisms [8,12] and (ii) the more cryptic, yet key contribution of reproductive control [2], and may not fully capture the subtlety and diversity of power relationships between males and females. Second, asymmetries in intersexual power and their causes have rarely been formally tested or quantified by systematically measuring the outcomes of intersexual conflicts. Instead, dominance relationships were, and still are, often studied separately for males and females or only for members of one sex. Species were often categorised as male- or female-dominated based on which sex the sexual dimorphism in size and weaponry was biased towards, or on which sex occupied the top positions in a group's **social hierarchy** [27,28]. Third, male-biased power has often been implicitly considered as the default state, whereas female-biased power has traditionally been viewed as anecdotal and emerging from lineage-specific oddities [16,20], such as the 'lemur syndrome' or the peculiar anatomy of female genitalia in spotted hyenas, moving the topic outside mainstream socio-ecology. These oversimplifications, and the lack of a standardised methodology to quantify intersexual power, have prevented the conceptual integration of intersexual power into the broader evolutionary theories of sexual conflict and mating systems as well as comparative analyses that could reveal general mechanisms driving biases in intersexual power.

## **The fifty shades of intersexual power within and across species**

Recent empirical studies that quantified intersexual power in different socio-ecological contexts and in diverse mammalian taxa revealed that intersexual power relationships are much more nuanced and dynamic than previously thought, questioning the long-standing dichotomist views. First, intersexual power is not limited to strict male social dominance (as in Hamadryas baboons *Papio hamadryas* [29]) or strict female social dominance (as in ringed-tailed lemurs, *Lemur catta* [15]) but varies across species along a continuum, including more balanced male-female power – also termed ‘co-dominance’ or ‘egalitarianism’ – as in vervet monkeys (*Chlorocebus pygerythrus*) and meerkats (*Suricata suricatta*) [30–32]. Second, intersexual asymmetries in social dominance can exhibit flexibility within a species, as in rock hyraxes [17], European badgers (*Meles meles*) [33], and spotted hyenas [28]. These findings indicate that intersexual power relationships are not necessarily a fixed attribute of a species and are not invariably driven by any particular sex-specific trait. They also call for the broadening of the traditional framework and the study of intersexual power as a dynamic, emergent property of the socio-ecological environment experienced by individuals, groups, populations and species.

### **The distinct pathways to male and female reproductive control**

Reproductive control has been described as the ability to act in a situation in which conflict over reproduction exists [2,8]. In contexts of sexual conflict, fertilisable eggs can be viewed as a high-value commodity that is owned by females and desired by males [1,8,10]. To acquire this commodity, males try to strategically manipulate the reproductive opportunities and decisions of females, which, in turn, use strategies to resist, evade or discourage male manipulative attempts. This ‘male control-female resistance’ paradigm is at the heart of sexual conflict over copulation and fertilisation [34,35], which formalises the dynamic power

struggles between the sexes over reproductive control. Whether and to what extent reproductive control is biased between the sexes is largely predicted by key features of the mating system (Box 2 and Figure 1A).

#### *Male control pathway*

In many mammalian species, sexual dimorphism in size and weaponry is biased towards males. Striking examples include Hamadryas baboons and elephant seals (*Mirounga angustirostris*) where males can be twice as large and weigh four times as much as females, respectively [36,37]. Males often exploit their physical superiority by adopting coercive strategies such as forced copulation, sexual harassment, intimidation or infanticide, which directly increase their mating probability [9,38,39] (Box 2 and Figure 1A). Males may also form alliances, as in bottlenose dolphins (*Tursiops aduncus*), to overcome the difficulty of monopolising females in three-dimensional aquatic environments [40]. Other coercive strategies indirectly increase male mating probability and paternity certainty by preventing females from mating with rivals. For example, pre-copulatory mate-guarding allows males to monopolise access to a sexually receptive female by concurrently restraining female mate choice and aggressively deterring competitors [41,42]. In species with relatively low sexual dimorphism, such as in many rodents, males may deposit copulatory plugs in the reproductive tract of their mating partners as a non-aggressive form of mate-guarding [43]. They may also apply deceptive tactics, as in territorial ungulates, where males may falsely signal the presence of a predator to discourage females from leaving their territory [9].

#### *4.2. Female resistance pathways*



166 In contrast to males, pathways to female reproductive empowerment are manifold.  
167 Females, too, may employ coercion to retaliate against coercive males, either alone or as  
168 part of a coalition of females [38] but they may exhibit a wide variety of other behavioural,  
169 physiological, morpho-anatomical and life-history traits to resist male reproductive control  
170 (Figure 1A). Females may promote male-male competition by advertising their reproductive  
171 receptivity, which increases male costs of monopolisation [44]. They may also synchronise  
172 sexually receptive periods, which will reduce male incentive to monopolise a given fertile  
173 female, because many others are available [35]. Females may alternatively deceive males by  
174 displaying unreliable genital swellings or mating outside oestrus [45–47]; such a strategy  
175 discourages male attempts to monopolise sexually receptive females and probably played a  
176 pivotal role in the evolution of female-biased power in bonobos (*Pan paniscus*) (Box 3).  
177 Females may conversely shorten their receptivity period to a few hours per year, as in many  
178 lemurs [48,49], a strategy that makes the considerable energetic costs of resisting male  
179 monopolisation attempts more affordable [50]. They may also mate sneakily [51] or in  
180 locations promoting their reproductive control. For example, in fossas (*Cytoprocta ferox*), a  
181 non-arboreal carnivore, females mate in treetops, a location that facilitates their filtering of  
182 candidate males that queue on the ground, by letting preferred males climb and pushing the  
183 others back down [52]. To resist unwanted mating attempts, females may also hide their  
184 genitals, escape males [9,53], or associate with male allies for protection [54]. In banded  
185 mongooses (*Mungos mungo*), females incite intergroup conflicts to escape mate-guarding by  
186 males of their group and mate with extra-group males in the mist of the battle [55]. In some  
187 species, as in many ungulates and bottlenose dolphins, extreme female avoidance of male  
188 coercion leads to sexual segregation whereby members of each sex live in distinct  
189 (sub)groups for most of the year [56]. Anatomical alteration of the female reproductive tract

may further allow females to gain pre- or post-copulatory reproductive control by either requiring their cooperation for a mating to occur, as in the spotted hyena (Box 1), or by storing sperm from multiple mates and expressing post-copulatory ‘cryptic’ mate choice, as in many bat species [57].

## **How reproductive control promotes social control**

Here we propose that the degree of male and female reproductive control determines whether and how members of each sex can empower themselves socially, with respect to access to non-reproductive resources. We further illustrate how the mechanism by which power emerges may influence its durability.

### *Coercion: an evolutionary highway to male power*

Male coercive reproductive control is facilitated by large male-biased sexual dimorphism in size and weaponry, which is typical of **contest-based** mating systems, and includes most **polygynous** and some **polygynandrous** societies [37,44,58] (Figure 1). In these systems, males often extend their use of coercion to dominate females when competing over non-reproductive resources. Large males may further reinforce intersexual asymmetries in coercive potential and limit female empowerment by controlling their social environment and preventing them from recruiting social allies (Box 2). The tight association between the pervasive use of coercion by males and male-biased sexual dimorphism likely emerges from a co-evolutionary feedback with the mating system (Figure 2). When males gain reproductive payoffs from aggressively monopolising females against competitors, this will often (i) promote contest-based competition between males, (ii) subsequently drive the evolution of male-biased sexual dimorphism [36,37], which will (iii) promote male coercive

214 reproductive control over females, and (iv) drive male social empowerment and dominance  
215 over females [19,36,44]. This will (v) ultimately close the loop by strengthening male  
216 reproductive control over females [38,39,44,58]. When female resistance strategies fail to  
217 prevent or disrupt this potent self-reinforcing loop, it may catalyse the emergence and  
218 maintenance of male-biased power over evolutionary times. This likely explains why males  
219 often exert both high reproductive and social control over females in contest-based mating  
220 systems (Figure 1), and why contest-based systems are widespread among mammals. In  
221 addition, female strategies may sometimes reinforce rather than disrupt the loop favouring  
222 male power. For example, where males already have high reproductive control and females  
223 cannot mate promiscuously to dilute paternity, they may instead concentrate paternity to  
224 seek paternal protection in response to infanticidal threats [59]. Such paternity  
225 concentration strategies may contribute to locking females into male-dominated societies.

226

#### 227 *Why the coercive pathway is not the females' way*

228 A similar coercive co-evolutionary pathway is unlikely to drive female social empowerment  
229 because mammalian species in which females concurrently exhibit contest-based intrasexual  
230 competition to monopolize access to multiple males and larger body sizes are currently  
231 unreported [60,61]. In some species, reproductive competition may be most intense among  
232 females; yet, contrary to what would be expected for this co-evolutionary pathway, these  
233 species either exhibit sexual monomorphism, as in the polyandrous moustached tamarins  
234 (*Saguinus mystax*) [58] and cooperatively-breeding meerkats [30] (Figure 1A), or male-biased  
235 size dimorphism as in Damaraland mole-rats (*Fukomis damarensis*) [61]. This apparent  
236 paradox probably reflects inherent differences in the life-history and modality of intrasexual  
237 competition in females and males [61]. Female mammals often compete using subtle forms

of coercion – e.g., using threats and agonistic signals – which do not select for increased body size and weaponry. There are at least three reasons for this: first, theory predicts avoidance of overt aggression for the sex that faces highest costs of offspring production [62]; second, the incentive for females to engage in physical contest may be low because sharing resources is often less costly for females than for males; third, mammalian females are usually philopatric and thus predominantly compete with close female kin, with whom they may avoid engaging in costly contests [61]. These insights emphasise key differences in the pathways to female and male power (Figure 2), in particular that a large body size and overt coercion are not pre-requisites for female empowerment.

#### *Female social empowerment from leverage based on sex*

When females retain some reproductive control – usually in species with moderate sexual size dimorphism as in **monogamous**, **polyandrous** and **scramble-based polygynandrous** species (Figure 1) – they can trade copulations for resources or services that males can provide, such as protection for themselves or their offspring against conspecifics or predators. Yet, such leverage is usually restricted to periods of female sexual receptivity and thereby only confer short-term social empowerment to females, as in some mouse lemurs (*Microcebus spp.*) where female social control over males is more pronounced during the breeding season [63,64]. Leverage-based power may therefore explain female social empowerment over males in species where males are non-permanent residents and join groups only during the mating season, as in rock hyraxes [17]. In species living in permanent groups where males and females maintain long-term, differentiated social relationships, females can extend their leverage beyond the receptive period. This strategy may durably promote cooperative behaviour or inhibit aggression from males through mating markets, as

in long-tailed macaques (*Macaca fascicularis*) [65] and Guinea baboons (*Papio papio*) [66]. Leverage can then represent a potent source of social control that may, even under male-biased dimorphism, allow females to manipulate the social rank of subordinate males, as in vervet monkeys [32], or to influence male social and competitive relationships, as in bonobos (Box 3). Similar to males, but through a different mechanism, increased social control by females may subsequently reinforce female reproductive control by facilitating their resistance to unwanted solicitations in a positive feedback loop (Figure 2).

#### *Female social empowerment from mate choice*

When female reproductive control enables them to exercise pre-copulatory mate choice, they may select male traits – e.g., social deference, cooperative personalities or a smaller body size – that may, over evolutionary time, increase female social control in a process described by the ‘docile male’ hypothesis [35]. In bonobos, the related ‘self-domestication’ hypothesis posits that selection for non-aggressive males, which may partly result from female choice, has contributed to the contrasts in morphology, physiology, behaviour and cognition between male bonobos and chimpanzees (*Pan troglodytes*) [24]. Empirical evidence of female mate choice for such male traits is scarce in mammals, however [12]. Female preferences for males with whom they are socially-bonded have been reported [67,68], but may reflect leverage rather than choice for male traits that are relevant to intersexual power. Alternatively, female mate choice can promote intersexual power asymmetries indirectly. For example, in spotted hyenas, female reproductive control and mate preferences drive male dispersal [69], which decreases the number of social allies that males can recruit and thus reduces male social control [28].

## **Ecological and evolutionary dynamics in intersexual power**

Power relationships between the sexes are closely intertwined with the social organisation and mating system of a species. Positive feedback loops between reproductive and social control suggest that dynamic changes in reproductive control and in the relative payoffs of male and female reproductive strategies are particularly likely to trigger shifts in intersexual power, across both ecological and evolutionary contexts.

### *Socio-ecological fluctuations in intersexual power*

Wherever power asymmetries between the sexes are moderate, fluctuations in the ecological or social environment – e.g., population density, food abundance, habitat structure, adult sex ratio or kin composition – may cause a change in the social and mating systems, which in turn may affect power relationships between males and females. For example, red foxes (*Vulpes vulpes*) changed their mating system from the typical contest-based polygyny to monogamy [70] following an abrupt decrease in the abundance and quality of prey species caused by an El Niño event, illustrating a shift from male-biased power to a more egalitarian system (Figure 1B). Ecological changes may also alter the relative payoffs of intersexual contests over non-reproductive resources, such as the relative value of one monopolisable food item for male and female reproductive success, therefore affecting the motivation of opponents and the likely outcome of their contest [21].

Fluctuations in the local mating market, for instance due to variation in the adult or operational sex ratio, may affect the extent of leverage possessed by females; e.g., the fewer females are available to mate, the more leverage they have over males [10,71]. This may explain why female power increases with the number of male group members in several primate species [27,31]. Conversely, where females have little reproductive control and low

leverage as in chimpanzees, sexual coercion by males may instead increase with the number of males in the community, as these dynamics reflect greater male-male competition, and associated male incentive to use coercive strategies [72,73].

#### *Evolutionary shifts in intersexual power*

Male-biased or female-biased power may be evolutionarily stable when reinforced by the positive feedback between reproductive control and social control (Figure 2). This framework predicts a relatively low stability, and thus the relative scarcity, of egalitarian societies where reproductive and/or social control is equally distributed between the sexes. It also predicts the potential for rapid directional evolution towards one or the other end of the intersexual power spectrum, following subtle changes in one keystone component of the system [74,75], which will affect the self-reinforcing evolutionary loop. Factors that reduce the extent of male reproductive control – and reciprocally increase that of females – may thus generate evolutionary shifts in intersexual power within or across species. For example, in the common brushtail possum (*Trichosurus vulpecula*), male sexual dimorphism and degree of polygyny vary in response to different degrees of breeding synchrony in females (influenced by food distribution and seasonality) [76]. Similarly, the emergence of reproductive synchrony in female Kinda baboons (*Papio kindae*) (Box 2) and of deceptive genital swellings in female bonobos (Box 3) have likely contributed to the abrupt shift from male-biased to female-biased power observed in the genera *Papio* and *Pan*, alongside a shift from contest-based to more scramble-based mating systems (Figure 1). Importantly, shifts in intersexual power will probably not occur in isolation but rather follow major transitions in the social organisation and mating system. While sex biases in reproductive control and

intersexual power can fluctuate across ecological contexts, broader and more impactful shifts may be expected at an evolutionary scale.

### **Concluding remarks**

We break here with the traditional, dichotomist and static view of intersexual power and offer a broader unifying framework that holds a central role for intersexual conflicts over reproductive control. By integrating concepts from theories of sexual conflict, sexual selection and social evolution, this new framework generates testable predictions regarding the ecological and evolutionary landscape of intersexual power within and across mammalian societies. The scientific investigation of intersexual power is a burgeoning and interdisciplinary research topic where much conceptual and empirical work remains to be done. We hope that this framework will be expanded and stimulate further studies (see Outstanding Questions), notably to develop standardised and widely applicable methods and transdisciplinary tools to quantify intersexual power in reproductive and social contexts. We also hope this work will encourage future studies to investigate how resource-holding power and decisional power (leadership) may interact and influence one another and ultimately develop a unifying framework for the study of intersexual power across domains, and across mammalian societies, including humans.



## Glossary

**Attribute:** trait possessed by an individual at a given time. Intrinsic attributes arise from an individual's physical, physiological or cognitive characteristics; social attributes arise from a relationship to others (e.g., social rank, kinship, coalitionary partners).

**Coercion:** strategy to influence the behaviour of others using some form of physical or psychological pressure, which often involves (the threat of) violence and may incur immediate, direct costs or delayed, indirect costs to the target.

**Contest-based:** mating system where males aggressively defend reproductive access to multiple females.

**Deception:** strategy to influence the behaviour of others that consists of withholding information or sharing inaccurate information to increase uncertainty and promote desired behaviour in other individuals.

**Intersexual power:** degree of control over resources and reproduction that members of one sex exert over members of the other sex. It shapes – and can thus be measured by – whether and how access to resources and reproduction is gained or retained by individuals of one sex.

**Leverage:** bargaining asymmetry in the control over the modality of an exchange that arises between trading individuals when one possesses a desirable commodity that cannot be taken by force by others (e.g., skills, information, and under certain conditions, fertilisable eggs).

**Mating system:** component of a social system that defines the modality of reproduction within a social unit; e.g., the average number of mates of males and females, the strategies applied to access mates, the timing and frequency of matings.

**Monogamy:** mating system where a single adult female and a single adult male mate mostly with each other during one or several mating season(s).

**Motivation:** incentive to engage in potentially costly behaviours or suboptimal trade to gain or retain access to a resource that is shaped by the value attributed to the resource, as determined by the individual state or life history.

**Polyandry:** mating system where individual females commonly mate with multiple males during a single mating season, whereas males do not mate with more than one female.

**Polygynandry:** mating system where males and females both are promiscuous and mate with multiple partners during a mating season.

**Polygyny:** mating system where a male can mate with multiple females but each female usually only mates with one male.

**Reproductive control:** extent to which an individual can influence the modality of its own reproduction and/or that of others (competitors and potential mates) in terms of the occurrence, timing and frequency of matings and the number and identity of mates.

**Scramble-based:** mating system where females cannot be monopolised by males and male-male competition is mostly non-coercive and post-copulatory.

**Social hierarchy:** ordinal ranking of individuals belonging to the same social group according to their relative power, typically resource-holding power as derived from the outcome of coercive interactions between pairs of individuals.

**Social organisation:** component of a species social system relating to the demographic composition of a social unit.

**Trade:** exchange of commodities between individuals. The modality and expected fitness payoff of the exchange are shaped by the local socio-ecological environment.

**Figure 1. Relationship between mating system, reproductive control and intersexual power. (A)** A selection of male and female traits proposed as criteria to quantify male-female biases in reproductive and social control in social mammals as illustrated by nine exemplary species. Colour tones indicate whether the trait promotes or reflects male control (blue), female control (red) or neither (yellow); “?” indicates either missing or non-consensual data in the current literature. Colour darkness reflects the strength of trait expression and, for overall scores of reproductive control and social control, the corresponding degree of intersexual bias. Scores of social control were derived from published indices of social dominance as quantified by the proportions of dyadic agonistic interactions won by males vs. females. **(B)** Putative position of the nine exemplary species (black icons) and their mating systems in relation to intersexual power measured along two axes – reproductive control and social control – as assessed by overall scores in panel A. Coloured areas delineate the proposed range of association between reproductive and social control for four mating systems; the gradient of contest- vs. scramble- based competition is depicted by a double arrow. Intersexual power follows a diagonal from species where power is strongly male-biased in both contexts (see Hamadryas baboons, *Papio hamadryas*) to strongly female-biased in both contexts (see ring-tailed lemurs, *Lemur catta*); mammalian societies where reproductive control is strongly female-biased and social control is strongly male-biased, or *vice versa*, are unlikely to emerge. Reference list: Hamadryas baboon [29,46]; chimpanzee (*Pan troglodytes*) and bonobo (*Pan paniscus*, see Box 3); Guinea baboon (*Papio papio*) [77] ; meerkat (*Suricata suricatta*) [30,78], moustached tamarin (*Saguinus mystax*) [79–83]; rock hyrax (*Procavia capensis*) [17,84]; spotted hyena (*Crocuta crocuta*) [16,28,85]; ring-tailed lemur [15,49]; cross-specific data [27,36,86].

424 **Figure 2. Eco-evolutionary pathways to male and female empowerment in mammals.**

425 Proposed framework to predict intersexual power asymmetries across mammalian societies.

426 There is interplay and feedback between the mating system and asymmetries in

427 reproductive and social control, and outlines the distinct pathways to male- and female-

428 biased power. Coercion and male-biased dimorphism constitute a co-evolutionary highway

429 to male power in species with contest-based polygyny and polygynandry (left side); Female

430 empowerment emerges through diverse physiological, morphological, behavioural, and

431 socio-ecological pathways (right side) in species with scramble-based polygynandrous,

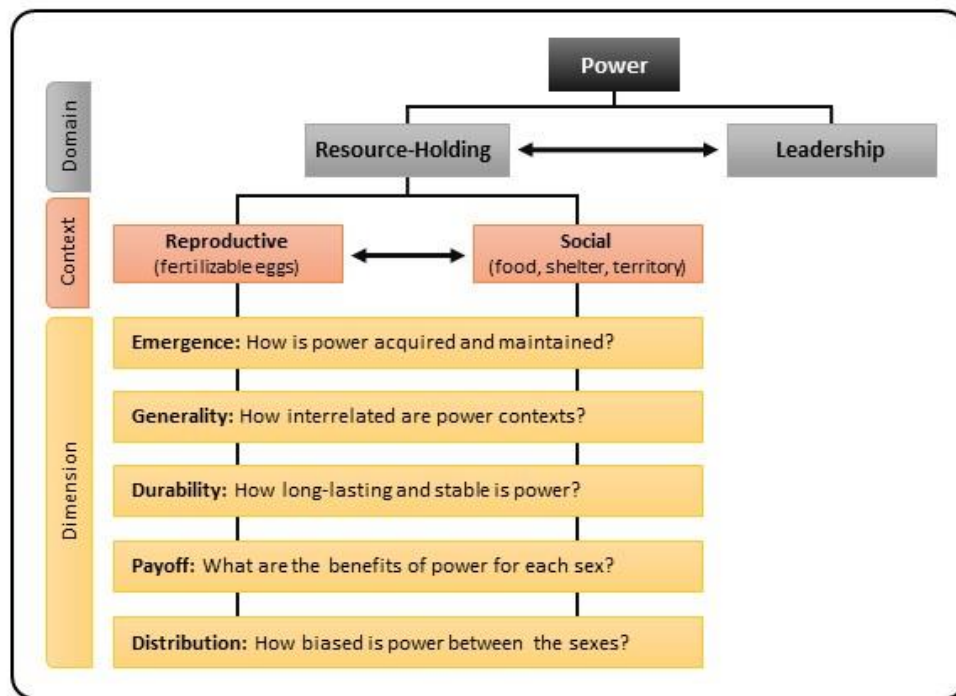
432 monogamous or polyandrous mating systems.

**Box 1. The duality of intersexual power: reproductive and social control.**

Power can be divided into two main domains, namely resource-holding power and decisional power (or ‘leadership’) [6,7]. We propose that resource-holding power arises in two main contexts – namely, reproductive and social contexts – defined according to the nature of the resource the two sexes compete over [8] (Figure I). Spotted hyenas (*Crocuta crocuta*) nicely illustrate the duality and context-dependence of intersexual power. In this species, females have complete reproductive control owing to the peculiar anatomy of their external genitalia. Females lack a vaginal opening (because their labia are fused) and instead possess an elongated and erectile clitoris through which they copulate, urinate and give birth [16]. A direct consequence is that copulation requires the full cooperation of the female and that females can actively choose when and with whom they mate [85]. In contrast, social control emerges from asymmetries in the number of recruitable social allies. The extent of intersexual biases in social control may fluctuate between strictly female-biased power structures and balanced social power between males and females, depending on the kin and demographic structure of the groups [28]. Building on previous conceptual categorisation of dominance [8], power [87] and leadership [11], we further propose that resource-holding power can be described following five dimensions: distribution (extent of asymmetries between the sexes), emergence (whether acquired *via* coercion, deception or trade), durability (stability versus lability), payoffs (sex differences in the fitness benefits to win intersexual contests) and generality (context-dependence or consistency across contexts) (Figure I).

**Figure I. Domains, contexts and dimensions of intersexual power.** The proposed framework captures the multifaceted nature and variability of intersexual power in animal societies. Bi-

directional arrows indicate the inter-dependence and potential feedback among domains and contexts of power. We here focus on the categorisation of power that emerges from the resource-holding domain.



## Box 2. Covariation between male-female asymmetries in reproductive control and mating systems.

Asymmetries in reproductive control between males and females may vary in predictable ways across mating systems. We illustrate such covariation along a gradient across the six baboon species, before generalising (Figure 1). At one end, Hamadryas baboons (*Papio hamadryas*) exhibit contest-based **polygyny** with full male reproductive control; males are much larger than females and maintain exclusive access to multiple females by forcibly herding females away from their natal group, thereby breaking-up their social ties and support. Leaders of such one-male units use various coercive strategies including infanticide

after take-overs, and neck-biting when females leave their vicinity [29]. Next, chacma baboons (*P. ursinus*) exhibit predominantly **contest-based polygynandry**. They live in large multimale-multifemale groups with high male reproductive skew – i.e., where access to mates is monopolised by a few males – and intense sexual coercion, including mate-guarding, intimidation and infanticide [88]. Unlike Hamadryas females, chacma females mate promiscuously before ovulation and remain in their natal group; they can form alliances with kin and adult males to seek protection against other coercive males [54]. Next are yellow (*P. cynocephalus*), olive (*P. anubis*) and then Guinea baboons (*P. papio*), with a progressive decline in male reproductive skew, sexual dimorphism and coercion, and a concomitant increase in the degree of female reproductive control. Females of these species may shorten or evade mate-guarding bouts by using fertility signals to exacerbate costly male-male competition [44], by resisting mate-guarding attempts or by mating sneakily with other males [89]. In Guinea baboons, males maintain long-term social and sexual bonds with maximum two females, females can freely disperse to other social units and infanticide is unreported [77]. Finally, in Kinda baboons (*P. kindae*), females mate promiscuously even in the ovulatory period, male reproductive skew is low and there is no evidence of infanticide [90]. Unlike other species, Kinda baboons breed seasonally and live in very large groups (>200 individuals). These features likely restrict male reproductive control and triggered an evolutionary shift from contest-based to **scramble-based** polygynandry – where male-male competition primarily occurs *via* sperm competition – as suggested by their low sexual dimorphism, relatively large testes and the fact that males queue rather than fight over rank [90]. This gradient illustrates general links between the mating system, reproductive control and intersexual power (Figure 1): as female reproductive control grows, contest-based mating systems are progressively replaced by scramble-based systems, associated with more balanced intersexual power.

These changes in reproductive control likely emerge from the combined effect of passive ecological and demographic processes (e.g., increased group size) and of strategies used by females to resist male monopolization attempts. Where males retain partial reproductive control, but where reproductive competition between females is too high to allow group-living, **monogamy** may initially evolve as a form of permanent mate-guarding [91], decreasing subsequent sexual conflict and paving the way to more equal reproductive control between the sexes and less-coercive mechanisms of intersexual power.

**Box 3. Contrasts in female reproductive control and power in our closest living relatives.**

Chimpanzees (*Pan troglodytes*) and bonobos (*Pan paniscus*) live in multimale-multifemale groups in equatorial Africa. In both species, males typically remain in their natal group and females disperse [92]. Their common ancestor likely exhibited male-biased power, as all Great Apes except bonobos [92].

In chimpanzees, adult males dominate adult females and have priority over access to high-value resources like meat [92]. In bonobos, females win most dyadic conflicts against males and enjoy privileged access to meat [23,92,93]. Females in both species exhibit genital swellings when sexually receptive, but bonobo females exhibit more and longer swollen periods during an interbirth interval [45]. Consequently, the swellings of female bonobos advertise fertility less reliably than those of female chimpanzees [45,94]. Chimpanzee males can commit infanticide, are frequently coercive towards receptive females [95], and alpha males have the highest reproductive success, because they can effectively mate-guard females throughout their ovulatory period [94,96]. In contrast, bonobo males are not infanticidal nor sexually coercive, they fight less over females, and the high male reproductive skew likely reflects differences in maternal support [97,98].



Following our framework, the initial step towards loss of male-biased power may be the confusion of ovulation, preventing male bonobos from monopolising receptive females and concurrently increasing female reproductive control. This probably reduced the payoff and incentive of males to fight over females, leading to a shift towards scramble-based polygynandry (Figure 1) – as suggested by the relatively larger testes and decreased sexual dimorphism in bonobos compared to chimpanzees [92]. Increased female promiscuity likely further led to the disappearance of male infanticide and sexual coercion. In line with this, male chimpanzees seem more coercive in populations where female fertility signals are shortest (presumably most reliable) [99]. Female-biased control over resources probably evolved secondarily in relation to their ability to use leverage and exercise mate choice [23,24]. In contrast to males, female bonobos maintain strong relationships, which may be favoured by prolonged sexual receptivity [100,101]; they can form coalitions against males and interfere in male-male competition to support their sons [92,93,97]. Such male-female differences in cooperation likely reinforced female reproductive and social control [92,93,100] rather than constituted the initial step towards female empowerment. Overall, current evidence is compatible with the scenario that strong contrasts in mating system, intersexual power and social behaviour in chimpanzees and bonobos emerged from subtle changes in female reproductive control.

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546 **Reference list:**

- 547 1 Lewis, R.J. (2002) Beyond dominance: the importance of leverage. *Q. Rev. Biol.* 77, 149–164  
548 2 Beekman, M. *et al.* (2003) Reproductive conflicts in social animals: who has power? *Trends Ecol.*  
549 *Evol.* 18, 277–282  
550 3 Smith, J.E. *et al.* (2021) An evolutionary explanation for the female leadership paradox. *Front.*  
551 *Ecol. Evol.* 9, 468  
552 4 Garfield, Z.H. *et al.* (2020) Universal and variable leadership dimensions across human societies.  
553 *Evol. Hum. Behav.* 41, 397–414  
554 5 Tibbetts, E.A. *et al.* (2022) The establishment and maintenance of dominance hierarchies.  
555 *Philos. Trans. R. Soc. B Biol. Sci.* 377, 20200450  
556 6 Cheng, J.T. and Tracy, J.L. (2020) Why social status is essential (but sometimes insufficient) for  
557 leadership. *Trends Cogn. Sci.* 24, 261–263  
558 7 Van Vugt, M. and Smith, J.E. (2019) A dual model of leadership and hierarchy: evolutionary  
559 synthesis. *Trends Cogn. Sci.* 23, 952–967  
560 8 Hand, J.L. (1986) Resolution of social conflicts: dominance, egalitarianism, spheres of  
561 dominance, and game theory. *Q. Rev. Biol.* 61, 201–220  
562 9 Bro-Jørgensen, J. (2011) Intra- and intersexual conflicts and cooperation in the evolution of  
563 mating strategies: lessons learnt from ungulates. *Evol. Biol.* 38, 28–41  
564 10 Noë, R. (2017) Local mating markets in humans and non-human animals. *Behav. Ecol. Sociobiol.*  
565 71, 148  
566 11 Smith, J.E. *et al.* (2016) Leadership in mammalian societies: emergence, distribution, power,  
567 and payoff. *Trends Ecol. Evol.* 31, 54–66  
568 12 Lewis, R.J. (2018) Female power in primates and the phenomenon of female dominance. *Annu.*  
569 *Rev. Anthropol.* 47, 533–551  
570 13 Petty, J.M.A. and Drea, C.M. (2015) Female rule in lemurs is ancestral and hormonally  
571 mediated. *Sci. Rep.* 5, 1–5  
572 14 Grebe, N.M. *et al.* (2019) Organizational and activational androgens, lemur social play, and the  
573 ontogeny of female dominance. *Horm. Behav.* 115, 104554  
574 15 Jolly, A. (1984) The puzzle of female feeding priority. In *Female primates; studies by women*  
575 *primatologists*. Alan R. Liss.pp. 197–216, Small, M.  
576 16 McCormick, S.K. *et al.* (2021) Sex differences in spotted hyenas. *Cold Spring Harb. Perspect.*  
577 *Biol.* DOI: 10.1101/cshperspect.a039180  
578 17 Koren, L. *et al.* (2006) Elevated testosterone levels and social ranks in female rock hyrax. *Horm.*  
579 *Behav.* 49, 470–477  
580 18 Davies, C.S. *et al.* (2016) Exceptional endocrine profiles characterise the meerkat: sex, status,  
581 and reproductive patterns. *Sci. Rep.* 6, 35492  
582 19 Kappeler, P.M. (1993) Female dominance in primates and other mammals. *Perspect. Ethol.* 10,  
583 143–158  
584 20 Kappeler, P.M. and Fichtel, C. (2015) Eco-evo-devo of the lemur syndrome: did adaptive  
585 behavioral plasticity get canalized in a large primate radiation? *Front. Zool.* 12 Suppl 1, S15  
586 21 Dunham, A.E. (2008) Battle of the sexes: cost asymmetry explains female dominance in lemurs.  
587 *Animal Behaviour*, 1435–39  
588 22 Young, A.L. *et al.* (1990) Female dominance and maternal investment in strepsirrhine primates.  
589 *Am. Nat.* 135, 473–488

- 590 23 Surbeck, M. and Hohmann, G. (2013) Intersexual dominance relationships and the influence of  
591 leverage on the outcome of conflicts in wild bonobos (*Pan paniscus*). *Behav. Ecol. Sociobiol.* 67,  
592 1767–1780
- 593 24 Hare, B. *et al.* (2012) The self-domestication hypothesis: evolution of bonobo psychology is due  
594 to selection against aggression. *Anim. Behav.* 83, 573–585
- 595 25 Haraway, D.J. (2013) *Primate visions: gender, race, and nature in the world of modern science*,  
596 Routledge.
- 597 26 Strum, S.C. and Fedigan, L.M. (2000) *Primate encounters: models of science, gender, and*  
598 *society*, University of Chicago Press.
- 599 27 Hemelrijk, C.K. *et al.* (2008) Female dominance over males in primates: self-organisation and  
600 sexual dimorphism. *PLoS ONE* 3, e2678
- 601 28 Vulliamy, C. *et al.* (2019) Social support drives female dominance in the spotted hyaena. *Nat.*  
602 *Ecol. Evol.* 3, 71–76
- 603 29 Swedell, L. *et al.* (2014) Sexual conflict in a polygynous primate: costs and benefits of a male-  
604 imposed mating system. *Behav. Ecol. Sociobiol.* 68, 263–273
- 605 30 Clutton-Brock, T.H. *et al.* (2006) Intrasexual competition and sexual selection in cooperative  
606 mammals. *Nature* 444, 1065–1068
- 607 31 Hemelrijk, C.K. *et al.* (2020) Dynamics of intersexual dominance and adult sex-ratio in wild  
608 vervet monkeys. *Front. Psychol.* 11,
- 609 32 Young, C. *et al.* (2017) Formidable females and the power trajectories of socially integrated  
610 male vervet monkeys. *Anim. Behav.* 125, 61–67
- 611 33 Hewitt, S.E. *et al.* (2009) Context-dependent linear dominance hierarchies in social groups of  
612 European badgers, *Meles meles*. *Anim. Behav.* 77, 161–169
- 613 34 Parker, G.A. (2006) Sexual conflict over mating and fertilization: an overview. *Philos. Trans. R.*  
614 *Soc. B Biol. Sci.* 361, 235–259
- 615 35 Gowaty, P.A. (2004) Sex roles, contests for the control of reproduction, and sexual selection. In  
616 *Sexual selection in primates: New and comparative perspectives* Cambridge University Press. pp.  
617 37–54, Kappeler PM, van Schaik CL
- 618 36 Cassini, M.H. (2020) Sexual size dimorphism and sexual selection in primates. *Mammal Rev.* 50,  
619 231–239
- 620 37 Weckerly, F.W. (1998) Sexual-size dimorphism: influence of mass and mating systems in the  
621 most dimorphic mammals. *J. Mammal.* 79, 33–52
- 622 38 Smuts, B.B. and Smuts, R.W. (1993) Male aggression and sexual coercion of females in  
623 nonhuman primates and other mammals: evidence and theoretical implications. *Adv. Study*  
624 *Behav.* 22, 1–63
- 625 39 Clutton-Brock, T.H. and Parker, G.A. (1995) Sexual coercion in animal societies. *Anim. Behav.*  
626 49, 1345–1365
- 627 40 Wiszniewski, J. *et al.* (2012) Male reproductive success increases with alliance size in Indo-  
628 Pacific bottlenose dolphins (*Tursiops aduncus*). *J. Anim. Ecol.* 81, 423–431
- 629 41 Setchell, J.M. *et al.* (2005) Mate guarding and paternity in mandrills: factors influencing alpha  
630 male monopoly. *Anim. Behav.* 70, 1105–1120
- 631 42 Mass, V. *et al.* (2009) Mate-guarding as a male reproductive tactic in *Propithecus verreauxi*. *Int.*  
632 *J. Primatol.* 30, 389–409
- 633 43 Stockley, P. *et al.* (2020) Revealing mechanisms of mating plug function under sexual selection.  
634 *Proc. Natl. Acad. Sci.* 117, 27465–27473
- 635 44 Clarke, P. *et al.* (2009) Infanticide, paternity allocation, and the role of coercion. In *Sexual*  
636 *coercion in primates and humans: an evolutionary perspective on male aggression against*  
637 *females* pp. 42–80, Harvard University Press
- 638 45 Douglas, P.H. *et al.* (2016) Mixed messages: wild female bonobos show high variability in the  
639 timing of ovulation in relation to sexual swelling patterns. *BMC Evol. Biol.* 16, 140

- 640 46 Zinner, D. and Deschner, T. (2000) Sexual swellings in female hamadryas baboons after male  
641 take-overs: “deceptive” swellings as a possible female counter-strategy against infanticide. *Am.*  
642 *J. Primatol.* 52, 157–168
- 643 47 Andelman, S.J. (1987) Evolution of concealed ovulation in vervet monkeys (*Cercopithecus*  
644 *aethiops*). *Am. Nat.* 129, 785–799
- 645 48 Eberle, M. and Kappeler, P.M. (2004) Selected polyandry: female choice and inter-sexual  
646 conflict in a small nocturnal solitary primate (*Microcebus murinus*). *Behav. Ecol. Sociobiol.* 57,  
647 91–100
- 648 49 Parga, J.A. (2006) Sexual selection in the ring-tailed lemur (*Lemur catta*): female choice, male  
649 mating strategies, and male mating success in a female dominant primate. , University of Texas
- 650 50 Huchard, E. *et al.* (2012) Convenience polyandry or convenience polygyny? Costly sex under  
651 female control in a promiscuous primate. *Proc. R. Soc. B Biol. Sci.* 279, 1371–1379
- 652 51 le Roux, A. *et al.* (2013) Evidence for tactical concealment in a wild primate. *Nat. Commun.* 4,  
653 1462
- 654 52 Lührs, M.-L. and Kappeler, P.M. (2014) Polyandrous mating in treetops: how male competition  
655 and female choice interact to determine an unusual carnivore mating system. *Behav. Ecol.*  
656 *Sociobiol.* 68, 879–889
- 657 53 Keddy, A.C. (1986) Female mate choice in vervet monkeys (*Cercopithecus aethiops sabaesus*). ,  
658 *American Journal of Primatology*, 125–134
- 659 54 Palombit, R. (2012) Infanticide: male strategies and female counter-strategies. In *The Evolution*  
660 *of Primate Societies* University of Chicago Press.pp. 432–468, Mitani JC, Call J, Kappeler PM,  
661 Palombit RA, Silk JB
- 662 55 Johnstone, R.A. *et al.* (2020) Exploitative leaders incite intergroup warfare in a social mammal.  
663 *Proc. Natl. Acad. Sci.* 117, 29759–29766
- 664 56 Galezo, A.A. *et al.* (2018) Sexual segregation in Indo-Pacific bottlenose dolphins is driven by  
665 female avoidance of males. *Behav. Ecol.* 29, 377–386
- 666 57 Archer, C.R. *et al.* (2017) Sperm competition and cryptic female choice in bats. In *Reproductive*  
667 *Biology, Physiology and Biochemistry of Male Bats* Edith Arenas-Ríos., 1pp. 119–147, Bentham  
668 Science Publishers
- 669 58 Cassini, M.H. (2020) A mixed model of the evolution of polygyny and sexual size dimorphism in  
670 mammals. *Mammal Rev.* 50, 112–120
- 671 59 Swedell, L. and Saunders, J. (2006) Infant mortality, paternity certainty, and female  
672 reproductive strategies in Hamadryas baboons. In *Reproduction and fitness in baboons:*  
673 *behavioral, ecological, and life history perspectives* (Swedell, L. and Leigh, S. R., eds), pp. 19–51,  
674 Springer US
- 675 60 Ralls, K. (1976) Mammals in which females are larger than males. *Q. Rev. Biol.* 51, 245–276
- 676 61 Young, A.J. and Bennett, N.C. (2013) Intra-sexual selection in cooperative mammals and birds:  
677 why are females not bigger and better armed? *Philos. Trans. R. Soc. B Biol. Sci.* 368, 20130075
- 678 62 Cant, M.A. and Young, A.J. (2013) Resolving social conflict among females without overt  
679 aggression. *Philos. Trans. R. Soc. B Biol. Sci.* 368, 20130076
- 680 63 Evasoa, M.R. *et al.* (2019) Sources of variation in social tolerance in mouse lemurs (*Microcebus*  
681 *spp.*). *BMC Ecol.* 19,
- 682 64 Hohenbrink, S. *et al.* (2015) Need for speed: sexual maturation precedes social maturation in  
683 gray mouse lemurs. *Am. J. Primatol.* 77, 1049–1059
- 684 65 Gumert, M.D. (2007) Payment for sex in a macaque mating market. *Anim. Behav.* 74, 1655–  
685 1667
- 686 66 Goffe, A. and Fischer, J. (2016) Meat sharing between male and female Guinea baboons (*Papio*  
687 *papio*). *Primate Biol.* 3, 1–8
- 688 67 Szykman, M. *et al.* (2001) Association patterns among male and female spotted hyenas  
689 (*Crocuta crocuta*) reflect male mate choice. *Behavioral Ecology and Sociobiology*, 231–238
- 690 68 Smuts, B.B. (1985) *Sex and friendship in baboons*, Routledge.

- 69 Höner, O.P. *et al.* (2007) Female mate-choice drives the evolution of male-biased dispersal in a social mammal. *Nature* 448, 798–801
- 70 Zabel, C.J. and Taggart, S., J. (1989) Shift in red fox, *Vulpes vulpes*, mating system associated with El Niño in the Bering Sea. *Anim. Behav.* 38, 830–838
- 71 Kappeler, P.M. (2017) Sex roles and adult sex ratios: insights from mammalian biology and consequences for primate behaviour. *Philos. Trans. R. Soc. B Biol. Sci.* 372, 20160321
- 72 Muller, M.N. *et al.* (2011) Sexual coercion by male chimpanzees shows that female choice may be more apparent than real. *Behav. Ecol. Sociobiol.* 65, 921–933
- 73 Watts, D.P. (2022) Male chimpanzee sexual coercion and mating success at Ngogo. *Am. J. Primatol.* 84, e23361
- 74 Alonzo, S.H. (2010) Social and coevolutionary feedbacks between mating and parental investment. *Trends Ecol. Evol.* 25, 99–108
- 75 Crespi, B.J. (2004) Vicious circles: positive feedback in major evolutionary and ecological transitions. *Trends Ecol. Evol.* 19, 627–633
- 76 Isaac, J.L. and Johnson, C.N. (2003) Sexual dimorphism and synchrony of breeding: variation in polygyny potential among populations in the common brushtail possum, *Trichosurus vulpecula*. *Behav. Ecol.* 14, 818–822
- 77 Fischer, J. *et al.* (2017) Charting the neglected West: the social system of Guinea baboons. *Am. J. Phys. Anthropol.* 162 Suppl 63, 15–31
- 78 Jordan, N.R. *et al.* (2007) Latrine distribution and patterns of use by wild meerkats: implications for territory and mate defence. *Anim. Behav.* 73, 613–622
- 79 Huck, M. *et al.* (2004) Proximate mechanisms of reproductive monopolization in male moustached tamarins (*Saguinus mystax*). *Am. J. Primatol.* 64, 39–56
- 80 Huck, M. *et al.* (2005) Paternity and kinship patterns in polyandrous moustached tamarins (*Saguinus mystax*). *Am. J. Phys. Anthropol.* 127, 449–464
- 81 Heymann, E.W. (1998) Sex differences in olfactory communication in a primate, the moustached tamarin, *Saguinus mystax* (Callitrichinae). *Behav. Ecol. Sociobiol.* 43, 37–45
- 82 Heymann, E.W. (1996) Social behavior of wild moustached tamarins, *Saguinus mystax*, at the Estación Biológica Quebrada Blanco, Peruvian Amazonia. *Am. J. Primatol.* 38, 101–113
- 83 Bicca-Marques, J.C. (2003) Sexual selection and the evolution of foraging behavior in male and female tamarins and marmosets. In *Sexual selection and reproductive competition in primates: New perspectives and directions* American Society of Primatologists, Norman.pp. 455–475
- 84 Bar-Ziv, E. *et al.* (2016) Individual, social, and sexual niche traits affect copulation success in a polygynandrous mating system. *Behav. Ecol. Sociobiol.* 70, 901–912
- 85 East, M.L. *et al.* (2003) Sexual conflicts in spotted hyenas: male and female mating tactics and their reproductive outcome with respect to age, social status and tenure. *Proc. R. Soc. Lond. B Biol. Sci.* 270, 1247–1254
- 86 Lukas, D. and Huchard, E. (2014) The evolution of infanticide by males in mammalian societies. *Science* 346, 841–844
- 87 Lewis, R.J. (2020) Female power: a new framework for understanding “female dominance” in Lemurs. *Folia Primatol. (Basel)* 91, 48–68
- 88 Baniel, A. *et al.* (2017) Male violence and sexual intimidation in a wild primate society. *Curr. Biol.* 27, 2163–2168.e3
- 89 Bercovitch, F.B. (1995) Female cooperation, consortship maintenance, and male mating success in savanna baboons. *Anim. Behav.* 50, 137–149
- 90 Petersdorf, M. *et al.* (2019) Sexual selection in the Kinda baboon. *J. Hum. Evol.* 135, 102635
- 91 Lukas, D. and Clutton-Brock, T.H. (2013) The evolution of social monogamy in mammals. *Science* 341, 526–530
- 92 Parish, A.R. *et al.* (2000) The other “closest living relative”: how bonobos (*Pan paniscus*) challenge traditional assumptions about females, dominance, intra- and intersexual interactions, and Hominid evolution. *Ann. N. Y. Acad. Sci.* 907, 97–113

742 93 Furuichi, T. (2011) Female contributions to the peaceful nature of bonobo society. *Evol.*  
743 *Anthropol.* 20, 131–142

744 94 Deschner, T. *et al.* (2004) Female sexual swelling size, timing of ovulation, and male behavior in  
745 wild West African chimpanzees. *Horm. Behav.* 46, 204–215

746 95 Muller, M.N. *et al.* (2007) Male coercion and the costs of promiscuous mating for female  
747 chimpanzees. *Proc. R. Soc. B Biol. Sci.* 274, 1009–1014

748 96 Bray, J. *et al.* (2016) Incomplete control and concessions explain mating skew in male  
749 chimpanzees. *Proc. R. Soc. B Biol. Sci.* 283, 20162071

750 97 Surbeck, M. *et al.* (2019) Males with a mother living in their group have higher paternity  
751 success in bonobos but not chimpanzees. *Curr. Biol.* 29, R354–R355

752 98 Surbeck, M. *et al.* (2017) Male reproductive skew is higher in bonobos than chimpanzees. *Curr.*  
753 *Biol. CB* 27, R640–R641

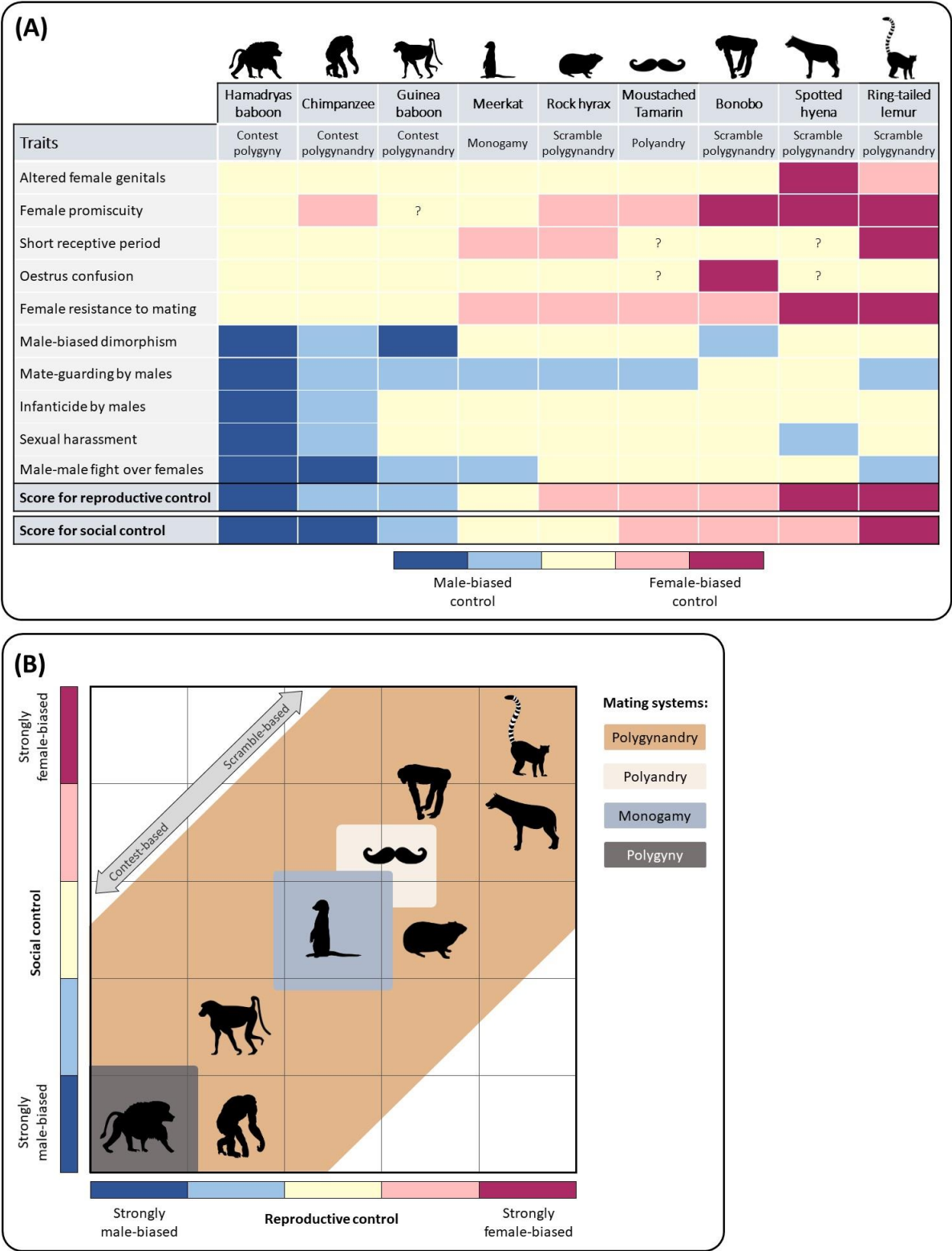
754 99 Wrangham, R.W. (2002) The cost of sexual attraction: is there a trade-off in female Pan  
755 between sex appeal and received coercion? In *Behavioural diversity in chimpanzees and*  
756 *bonobos* C. Boesch, G. Hohmann, L.F. Marchant. pp. 204–215

757 100 Surbeck, M. *et al.* (2021) Attractiveness of female sexual signaling predicts differences in female  
758 grouping patterns between bonobos and chimpanzees. *Commun. Biol.* 4, 1–11

759 101 Moscovice, L.R. *et al.* (2019) The cooperative sex: Sexual interactions among female bonobos  
760 are linked to increases in oxytocin, proximity and coalitions. *Horm. Behav.* 116, 104581

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762 Figure 1



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