

# THE QUARTERLY REVIEW *of* BIOLOGY



## A PARADIGM SHIFT, OR A PARADIGM ADJUSTMENT? THE EVOLUTION OF THE OLEACEAE MATING SYSTEM AS A SMALL-SCALE KUHNIAN CASE STUDY

ALEXANDRE FRANCO

*Univ. Lille, CNRS, UMR 8198—Evo-Eco-Paleo  
F-59000 Lille, France*

E-MAIL: AF.ALEXANDRE.FRANCO@GMAIL.COM

PIERRE SAUMITOU-LAPRADE

*Univ. Lille, CNRS, UMR 8198—Evo-Eco-Paleo  
F-59000 Lille, France*

E-MAIL: PIERRE.SAUMITOU-LAPRADE@UNIV-LILLE.FR

PHILIPPE VERNET

*Univ. Lille, CNRS, UMR 8198—Evo-Eco-Paleo  
F-59000 Lille, France*

E-MAIL: PHILIPPE.VERNET@UNIV-LILLE.FR

SYLVAIN BILLIARD

*Univ. Lille, CNRS, UMR 8198—Evo-Eco-Paleo  
F-59000 Lille, France*

E-MAIL: SYLVAIN.BILLIARD@UNIV-LILLE.FR

### KEYWORDS

evolutionary biology, plant evolution, Kuhn's evolutionary model, models, data,  
olive tree

*The Quarterly Review of Biology*, volume 98, number 2, June 2023.

© 2023 The University of Chicago. All rights reserved. Published by The University of Chicago Press. <https://doi.org/10.1086/725275>

## ABSTRACT

*Kuhn (1962) proposed an evolutionary model to explain how scientific knowledge is built, based on the concept of paradigm. Even though Kuhn's model is general, it has been applied to only a few topics in evolutionary biology, almost exclusively to broad-based paradigms. We analyze here, through the lens of Kuhn's theory, a small-scale paradigm change that occurred with the resolution of the controversy about the mating system of a Mediterranean shrub *Phillyrea angustifolia* (Oleaceae). We first summarize the different steps of the paradigm change and replace them in the more general context of sex ratio theory. Second, we show how the different steps of the paradigm change can be interpreted by Kuhnian concepts and tools. Finally, we discuss the actual and future status of the new paradigm.*

## INTRODUCTION

SINCE the publication of *The Structure of Scientific Revolutions* in 1962, Thomas Kuhn's theory of scientific research has been a cornerstone in the philosophy of science. His model seeks to identify global mechanisms underlying all scientific activities, at all scales, and aims at explaining the efficacy of scientific activity (Kuhn 1970). The Kuhnian model is an evolutionary theory (or "post-Darwinian Kantianism"; Kuhn 2000) as it is based on historical processes and because paradigm selection is analogous to natural selection. Kuhn's theory is also based on a realistic approach that humanizes scientific research, thus making it more plastic and less idealized. Even though the general applicability of his theory was debated very soon after the publication of *The Structure of Scientific Revolutions* (see Shapere 1964 for a critical statement about *incommensurability*, one of the fundamental concepts of the theory), Kuhn's model is certainly one of the most successful epistemological frameworks, especially among scientists themselves.

Biology is strikingly absent from *The Structure of Scientific Revolutions*, certainly because Kuhn was known to be more comfortable with the history of physics and chemistry. For more than 50 years, researchers have been trying to apply the Kuhnian model to different fields of biology, especially population biology, at different scales (Figure 1, see also the *epistemological metaparadigms* in Friedman 2002 for an alternative organization). With the adoption of Darwinian and Neo-Darwinian theory, high-level paradigms (Level 2 in Figure 1) received much attention (e.g., Ruse 1970, 2018; Jacob 1976; Greene 1981; Gayon 1992; Jablonka and Lamb 2005; Mo-

range 2017; Tanghe et al. 2021). Ruse (2018) in particular argued that Darwinism should not be considered as a paradigm but rather as an example of scientific consilience. Bertoldi (2018) further proposed that the Kuhnian category of paradigm does not allow us to accurately portray the specificities of Darwin's theory.

Besides the study of Darwinism, Kuhnian analyses of other population biology issues are scarce, but cover various subjects, including:

- The issue of the unit of selection in evolution (Level 3 in Figure 1; see Lloyd 2020 for a review; see Ruse 1987 for the particular case of sociobiology);
- The pace of evolution and the theory of the punctuated equilibria (Ruse 1989);
- Hoquet (2020) partly analyzed the status of Bateman's principle (1948; a Level 3 paradigm) especially focusing on the social components of the associated paradigm-based research; and
- Avise (2014) collected, reviewed, and evaluated the importance of a large number of paradigm changes in evolutionary genetics, from different levels (Levels 2 to 4 in Figure 1). However, its choices and evaluations of paradigm changes were mostly subjective, as confessed by the author himself. He did not precisely use the Kuhnian model to determine what could be considered as a paradigm, whether a scientific change should be considered as a paradigm shift, or how to evaluate the importance of a shift.

Our goal in this paper is to provide a new Kuhnian analysis of a low-level paradigm from population biology: the evolution of mating systems in plants applied to Oleaceae species, a paradigm embedded in sex ratio theory (Fisher 1930), one of the most celebrated in

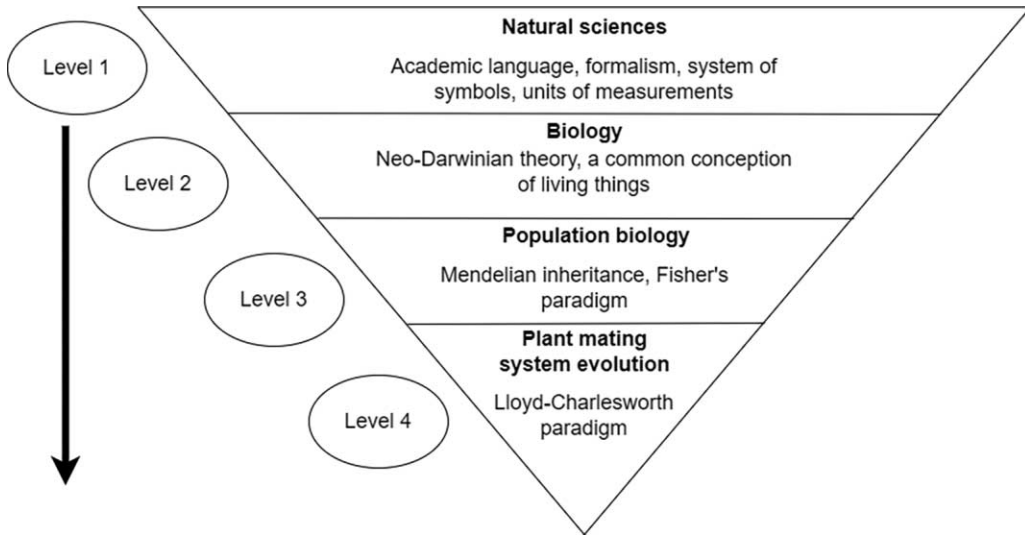


FIGURE 1. THE ENTANGLEMENT OF SCIENTIFIC FIELDS AND THEIR SHARED PARADIGMS APPLIED TO PLANT MATING SYSTEM EVOLUTION

This schematic representation shows that the field of plant mating system evolution and its paradigms (Level 4 paradigms) are nested within population biology (Level 3 paradigms), which is nested within biology (Level 2 paradigms) and natural sciences (Level 1). Note that the representation is not exhaustive as upper or lower levels could be added.

evolutionary theory (Edwards 1998). This paradigm has original specificities that make it an original and perfect candidate for an analysis of the evolution of a scientific paradigm in evolutionary biology: it is a small-scale and low-level paradigm (Level 4 in Figure 1); it is local in the sense that the involved scientific community and the associated literature is limited; and it is current since a crisis of this paradigm was only recently resolved (although the crisis is still marginally ongoing).

This paradigm crisis followed directly the basic idea of the sex ratio theory put forward by Fisher (1930): producing an offspring in sexual species necessarily involves the fusion of both male and female gametes. As a consequence, the sex ratio in a population should be balanced 1:1. Indeed, if one sex is rarer than the other, an individual that would produce more offspring of the former sex would have more grandchildren and thus would be favored by natural selection. The mating and sexual systems of the shrub *P. angustifolia* (Oleaceae) challenged the paradigm of plant mating system evolution as a subcase of the sex ratio theory. The controversy, which will be detailed below, was centered on the ob-

served frequency of males in *P. angustifolia* natural populations, which was much higher than expected (Pannell 2002): males were as frequent as hermaphrodites, which was not possible under the sex ratio theory in androdioecious populations (a population is androdioecious when hermaphrodites and male individuals co-occur). Hence, either the observations were wrong (i.e., hermaphrodites were in fact functionally females, which would give a 1:1 sex ratio in accordance with theoretical predictions) or the quantitative theoretical predictions from sex ratio theory were not correct because they were not adapted to the case of *P. angustifolia*. The controversy was resolved after the discovery of the link between sex determination and a diallelic self-incompatibility (DSI) system in this species (Saumitou-Laprade et al. 2010). From an epistemological point of view, this raises the question of whether this resolution constitutes a simple paradigm adjustment or a paradigm shift. Pannell and Korbecka (2010) claimed, without a proper analysis of the criteria proposed by Kuhn and others (e.g., Wray 2021), that only a paradigm adjustment was necessary (i.e., “normal science” is going on).

In this paper, we first detail the controversy about the mating system in *P. angustifolia*, especially highlighting the different crucial steps of the paradigm changes (Figure 2). Second, we study and analyze this controversy through the lens of Kuhn's theory. Third, we discuss the actual and future status of the paradigm changes regarding the case of *P. angustifolia*. In particular, we discuss whether the discovery of the diallelic self-incompatibility system in *P. angustifolia* constitutes an adjustment of the previous paradigm, as sug-

gested by Pannell and Korbecka (2010), or a paradigm shift. We also describe current and possible future resistances regarding research on the Oleaceae mating system evolution based on our understanding of Kuhn's theory. Overall, a Kuhnian analysis of this small-scale case study offers a unique opportunity to analyze how science works in action, study some phenomena that are rarely observed for high-level paradigms (e.g., scientists' conversion from the old to the new paradigm), and thoroughly analyze the roles

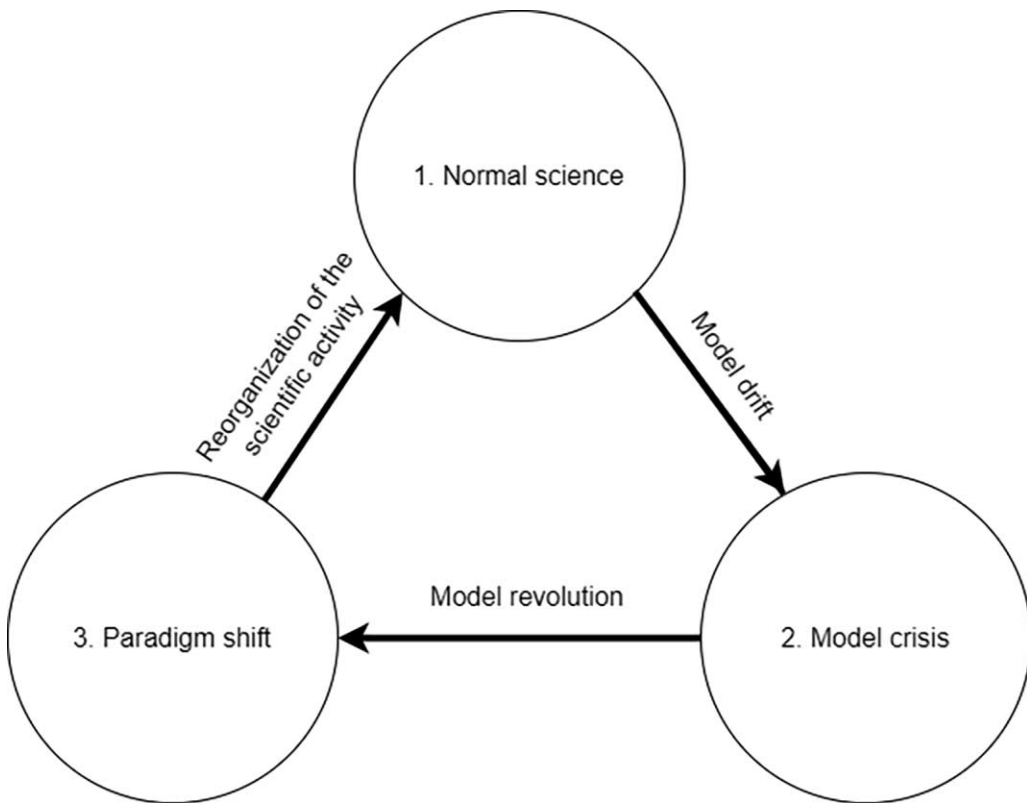


FIGURE 2. THE KUHN CYCLE OF SCIENTIFIC ACTIVITY

1. *Normal science*: the sex ratio and plant mating system evolution paradigm according to predictions by the Fisher-Lloyd-Charlesworth and Charlesworth paradigm. 2. *Model crisis*: the unexpected and unexplained high frequency of males in *P. angustifolia* populations was an anomaly, which led to a paradigm crisis with two opposing perceptions and interpretations of data: *P. angustifolia* was either indeed androdioecious or it was cryptically dioecious in agreement with the established Fisher-Lloyd-Charlesworth and Charlesworth paradigm. 3. *Paradigm shift*: the combination of experimental results and new models led to the acceptance that two new mechanisms should be considered to explain the high male frequencies in *P. angustifolia*: the diallelic self-incompatibility system and sexual distortion segregation. These discoveries were followed by new theoretical and empirical questions regarding the origin, maintenance, and coevolution of the two mechanisms, and the mating system status in other Oleaceae species.

played by the confrontation between models and data in a paradigm shift. It also raises the question whether the Kuhnian model applies to any paradigm level (as Kuhn suggests in the 1969 postscript to *The Structure of Scientific Revolutions*; see Kuhn 2012:168) or how important should a scientific change be in order to be considered as a paradigm shift.

THE *PHILLYREA ANGUSTIFOLIA*  
CONTROVERSY, ITS RESOLUTION,  
AND EXTENSIONS

NORMAL SCIENCE: SEX RATIO THEORY AND  
PLANT MATING EVOLUTION

Understanding the evolutionary mechanisms explaining why sex ratio should be balanced between males and females in dioecious species has a long history that dates back to Darwin (see Edwards 1998 for a review and references therein). In the first edition of *The Descent of Man, and Selection in Relation to Sex* (1871), Darwin proposed that selection should favor a 1:1 sex ratio between males and females. However, after the reports of exceptions to 1:1, he eventually admits that “the whole problem is so intricate that it is safer to leave its solution for the future” (Darwin 1874:260). It was almost 60 years later that Fisher (1930) proposed a simple explanation based on natural selection, as anticipated by Darwin (1871): since producing offspring by sexual reproduction necessarily needs male and female gametes, if the sex ratio is unbalanced, then one individual of the rare sex would have more descendants, on average, than an individual of the other sex. Hence, a mutation that would bias the sex ratio at birth in favor of the rare sex would be advantaged by natural selection because individuals bearing such a mutation will have more grandchildren on average. The evolutionary stable equilibrium sex ratio is thus expected to be 1:1.

Departures from the 1:1 prediction are observed in many different species. As a consequence, Fisher’s paradigm (1930) suffered many changes to explain unbalanced sex ratio. For instance, it has been necessary to distinguish between sex ratio at birth (i.e., the ratio of males versus females in offspring)

and operational sex ratio (the ratio of competing males versus females that can be fertilized) to explain why sex ratio is slightly male-biased in human populations (such a bias being compensated by a higher male death rate before sexual maturity; e.g., Ritchie and Roser 2019) or why sex ratio can be highly skewed when a population is structured (Hamilton 1967). Actually, sex ratio theory is regarded more generally as a sex allocation theory (Charnov 1982), i.e., the ratio of resources allocated to male versus female gametes by individuals taking into account life-history traits, environmental conditions, and interactions between individuals.

In sex ratio theory, plant mating systems evolution received particular attention, mainly because most plant species show hermaphroditic individuals. Hermaphroditism was a particular issue for the sex ratio paradigm because it makes possible: self-fertilization, and its counterpart strong inbreeding depression, which respectively affect male fitness because less ovules are available for siring and female fitness since inbred offspring are less fit; differential resource allocation between male and female gametes within individuals, which can lead to highly skewed pollen-ovule ratio or male versus female reproductive organs production; and a generalized definition of sex ratio as the relative frequency of a pair (or a triplet) of genders within a population: males, females, and/or hermaphrodites.

After the seminal work by Lewis (1941), new questions were raised in the 1970s to early 1980s about plant mating evolution considering those three consequences of hermaphroditism within sex ratio theory, especially regarding the expected frequency of unisexual individuals (males or females) when they co-occur with hermaphrodites in natural populations (populations with hermaphrodites and males, hermaphrodites and females, and hermaphrodites and both males and females are respectively called androdioecious, gynodioecious, and trioecious). Among others, Lewis (1941), Lloyd (1975), Charnov et al. (1976), and Charlesworth and Charlesworth (1981) provided theoretical quantitative predictions about the expected frequencies of unisexual individuals

in populations, and the conditions under which unisexuals are expected to be maintained with hermaphrodites. In other words: what is the evolutionarily stable sex ratio in androdioecious and gynodioecious populations?

As expected by quantitative models and confirmed by empirical observations, gynodioecy and androdioecy are not symmetrical mating systems (reviewed in Pannell 2002; Dufay and Billiard 2012). From a theoretical point of view, androdioecy is expected to be less stable (and thus rarer) than gynodioecy in plants. In addition, males are expected to be rare in androdioecious species, i.e., the sex ratio is expected to be highly skewed toward hermaphrodites. This is due to three main factors. First, Bateman's principle posits that male fitness is mostly limited by mating opportunities, whereas female fitness is mostly limited by resource acquisition (Bateman 1948). Second, since males can only have offspring by their male gametes, while hermaphrodites can provide offspring by both their male and female gametes, males should sire at least twice as many offspring as hermaphrodites to be maintained in a population (Lloyd 1975). This means that resource allocation to male gamete production (or efficiency) should be largely higher in males than in hermaphrodites. Third, because hermaphrodites can self-fertilize, it gives even less opportunities for males to sire available ovules, and it is easier for hermaphrodites to produce offspring through both their male and female gametes. Hence, as self-fertilization increases, it is expected that males are rarer and that androdioecy is less stable (Charlesworth and Charlesworth 1981).

As reviewed by Pannell (2002), empirical observations totally agree with theoretical predictions in flowering plants: androdioecious species are rare, males show low frequencies in natural androdioecious populations, and males show an obvious larger production of male gametes than hermaphrodites. Overall, it means that the sex ratio theory applied to plant mating systems perfectly fits observations. There was, however, one exception that challenged the theory for two decades, as acknowledged by Pannell (2002): the Mediterranean shrub *P. angustifolia* (Oleaceae).

#### CRISIS: SEX RATIO PARADIGM'S PREDICTIONS APPLIED TO *P. ANGUSTIFOLIA* POPULATIONS

In a nutshell, the sex ratio paradigm predicts that:

1. In dioecious populations, the male versus female sex ratio should be balanced 1:1, unless under particular situations; and
2. In androdioecious populations, the male versus hermaphrodite sex ratio should be highly skewed toward hermaphrodites for realistic values of male gamete production (or efficiency) by both males and hermaphrodites (in other words, if hermaphrodites effectively produce some male gametes, and if males produce a finite number of male gametes).

None of these conditions fit for in *P. angustifolia*. Gerber and Kieffer (1898) and Lepart and Dommée (1992) showed that three natural populations of this species in southern France were morphologically androdioecious, since they were composed of individuals showing flowers with only male organs (males) or with both male and female organs (hermaphrodites). They also showed that, in these three populations, hermaphrodites were male fertile when outcrossed onto another hermaphrodite. Unexpectedly for an androdioecious species (Pannell 2002), 45 populations from southern France, Portugal, and Spain showed that the distribution of the sex ratio within this species ranged between 0.34 to 0.77, with 0.49 on average (Strasberg 1988; Lepart and Dommée 1992; Pannell and Ojeda 2000; compiled in Husse et al. 2013). Even more surprising were the estimates of the male advantage in their production (or efficiency) of male gametes relatively to hermaphrodites: it was estimated to lie between 1 and 2 by Vassiliadis et al. (2000a) and Pannell and Korbecka (2010), a value largely incompatible with the observed sex ratio and the expectations from the sex ratio paradigm (Lloyd 1975). This generated a paradigm crisis.

From there, two research programs followed two different paths. One program hypothesized that, following the sex ratio paradigm, since the sex ratio in *P. angustifolia* is approximately balanced, then this species

should be dioecious and not androdioecious: hermaphrodites *morphologically* show both male and female reproductive organs, but they should be *functionally* females (Pannell 2002), a situation referred to as *cryptic dioecy*, in contradiction to Lepart and Dommée's (1992) conclusions. Roughly speaking, posing that the paradigm and its quantitative predictions are true, then observations should be wrong. Considering hermaphrodites were in fact females was the least costly explanation for maintaining the established paradigm. At the same time, using genetic markers and paternity analyses, Vassiliadis et al. (2002) assigned both parents based on genetic similarities between adult trees and seedlings. They detected that a substantial amount of seeds were effectively sired by hermaphrodites, and thus concluded that hermaphrodites were *functionally* and not only *morphologically* hermaphrodites, in agreement with Lepart and Dommée (1992).

Yet, doubts remained. Verdú et al. (2006) performed similar paternity analyses on another supposedly androdioecious species with a balanced sex ratio, hence with the same paradoxical observations as *P. angustifolia* (the ash tree *Fraxinus ornus*, also an Oleaceae). Even though Verdú et al. found that the proportion of seeds sired by hermaphrodites was substantial, they concluded that the "1:1 sex ratios of *F. ornus* populations indicate that the species is cryptically dioecious" (Verdú et al. 2006:2061). They further speculated that, since hermaphrodites were functional, a high *ovule discounting* was necessary, i.e., a very high mortality rate of seeds or offspring produced by a cross between hermaphrodites. Assuming ovule discounting is costly because there was no evidence of such a mechanism actually occurring, and because it would imply a very substantial mating cost, it is barely compatible with Darwinian selection, a higher level paradigm (Figure 1). By extrapolation, they also concluded that, most likely, a similar mechanism should occur in *P. angustifolia* and, therefore, that it also certainly was cryptically dioecious.

In another paper, Pannell and Ojeda (2000) proposed to explain the paradox of *P. angustifolia* by a phenomenon that could increase male advantage: males would tend

to flower more often than hermaphrodites and, consequently, would increase their opportunity to sire ovules produced by hermaphrodites. Higher flowering rates in males would also bias the estimation of the sex ratio toward males, suggesting that methodological limits could play a role in the paradox. However, Pannell and Ojeda (2000) did not provide quantitative predictions for the effect of differences in flowering rates on observed sex ratio. To what extent it might explain high male frequency in *P. angustifolia* populations was thus unclear. Overall, cryptic dioecy was the least costly explanation that makes the paradigm's predictions compatible with data.

A second research program took another path and challenged the sex ratio paradigm itself, especially the bases of Lloyd's model (1975). Fundamentally, the question remained: what factor could rebalance the male advantage between males and hermaphrodites: without making hermaphrodites cryptically females and such that the production (or efficiency) of male gametes by males are similar to that of hermaphrodites?

Vassiliadis et al. (2000a, 2002) observed that the siring success was homogeneous among male individuals, whereas it was highly variable among hermaphrodites. This observation led Vassiliadis et al. (2000b, 2002) to hypothesize that self-incompatibility could be a mechanism involved in androdioecy in *P. angustifolia*. Self-incompatibility (SI) is a very common mechanism in flowering plants that prevents mating between individuals belonging to the same self-incompatibility group. SI is a self-/nonself recognition system that makes mating possible between compatible genotypes only: individuals can cross only if they do not share the same self-incompatibility phenotype, in other words, if they do not belong to the same SI group (e.g., individuals from the  $G_a$  self-incompatibility group can only sire and be sired by individuals from the  $G_b$  SI group). A given individual expressing SI is thus necessarily unable to self-fertilize.

The idea of Vassiliadis et al. (2000b, 2002) relies on the following mechanism: if hermaphrodites express SI groups, while males do not or express a SI group unique to them, then males automatically have a fertilization

advantage relative to hermaphrodites. Indeed, since males could sire all hermaphrodites while hermaphrodites can only sire a part of all hermaphrodites in the population, pollen emitted by males would have a higher chance to successfully fertilize an ovule than pollen emitted by hermaphrodites. In addition, SI prevents self-fertilization that precludes a potential extra advantage for hermaphrodites (Charlesworth and Charlesworth 1981).

Vassiliadis et al. (2000b) showed in a model that their hypothesis could partly solve the paradox: males could indeed reach high frequency if the number of SI groups expressed by hermaphrodites was small. However, some information was still missing. First, there was no direct evidence of SI in *P. angustifolia*. Second, even if SI indeed exists in *P. angustifolia*, the number of SI groups in the hermaphrodites should be determined. Another difficulty thus arose: in the plant mating system literature, SI plant species typically show several dozen SI groups (Castric and Vekemans 2004), which made Vassiliadis et al.'s (2000b) predictions incompatible with data. Hence, the picture was still incomplete, and the paradox remained unsolved.

#### SOLVING THE PARADOX: DIALLELIC SELF-INCOMPATIBILITY AND SEGREGATION DISTORTION

The paradox was resolved in two steps. The first step was the discovery of a diallelic self-incompatibility system with only two homomorphic SI groups of hermaphrodites ( $G_a$  and  $G_b$ ), and no SI in males (Saumitou-Laprade et al. 2010). Such a homomorphic DSI system was unexpected for two reasons. First, because only heteromorphic SI systems with two SI groups were known in flowering plants (distyly or heterostyly). Second, because all known homomorphic SI in flowering are multiallelic. The discovery of DSI was also surprising because, with a single simple mechanism, males automatically compensate their fitness disadvantage compared to hermaphrodites: hermaphrodites can reproduce through their ovules and pollen, while males can only reproduce through their pollen, but hermaphrodites can only sire approximately

one-half of the hermaphrodites (the ones of the other SI group), while males can sire all hermaphrodites. Pannell and Korbecka (2010) and Husse et al. (2013) introduced DSI into Lloyd's (1975) model (or an equivalent population genetics model). They showed that indeed, DSI increases the expected frequency of males in populations. However, once again, not to the extent observed in natural populations.

The second step consisted of the observation of sex ratio distortion at birth, which was less surprising than DSI because segregation distortion was already observed multiple times in angiosperms. More precisely, it was observed that the inheritance of sexual phenotypes from parents to offspring did not follow Mendelian segregation rules for one particular crossing (Billiard et al. 2015): offspring from the mating between males and hermaphrodites from group  $G_b$  are all males. All other crosses with hermaphrodites from group  $G_a$  produced progeny with a proportion of hermaphrodites  $G_a$ ,  $G_b$ , and/or males as expected under Mendelian segregation where two unlinked loci controlled for DSI and sex determination. Billiard et al. (2015) introduced this distortion segregation into Husse et al.'s model (2013). They showed that the combination of DSI and distortion segregation gives expected male frequency compatible with observations in natural populations. This finally solved the paradox from the observation of a balanced sex ratio 1:1 in the androdioecious *P. angustifolia*.

Thanks to a combination of experiments in controlled conditions, theoretical quantitative modeling, and observations in natural populations, the research program that resolved the paradox in *P. angustifolia* allowed the discovery of a brand new biological phenomenon: homomorphic DSI in flowering plants. Analog mating systems were already known in fungi, ciliates, yeast, and green algae, but not in angiosperms (Billiard et al. 2011, 2012). This research program is an illustration of the scientific fecundity of confronting data with theory. The program also allowed the discovery of a clear association between two mechanisms controlling matings that are common in angiosperms, but



generally considered separately: sexual (males, females, or hermaphrodites) and self-incompatibility. Finally, this research program was pursued by the study of mating and sexual systems in other Oleaceae species. In all studied species, the existence of a DSI system was demonstrated either at the prezygotic stage by controlled stigma test or at the postzygotic stage by paternity analysis of seeds produced by controlled crosses and/or open pollination: the androdioecious manna ash *F. ornus*, which also shows high male frequencies (Vernet et al. 2016); the hermaphroditic olive tree *Olea europaea* (Saumitou-Laprade et al. 2017a; Besnard et al. 2020; Mariotti et al. 2021); the morphologically polygamous but functionally dioecious common ash *Fraxinus excelsior* (Saumitou-Laprade et al. 2018); and the hermaphroditic privet *Ligustrum vulgare* (De Cauwer et al. 2021). No other case of distortion segregation was yet detected. Hence, *P. angustifolia* is, on one hand, still a particular case because of this association between sex ratio distortion and DSI but, on the other, it is representative of the Oleaceae family since it shares DSI with all Oleaceae species studied so far.

REORGANIZATION OF SCIENTIFIC  
ACTIVITIES: WHAT DID THE RESOLUTION OF  
THE *P. ANGUSTIFOLIA* PARADOX CHANGE?

Solving the paradox needed two paradigm changes. First, incorporating that *P. angustifolia* population was structured both by sexes (males and hermaphrodites) and by SI groups for hermaphrodites ( $G_a$  and  $G_b$ ), and that mating relationships were asymmetrical (hermaphrodites can only sire one of the two SI groups, while males can sire both SI groups). Whether or not it was a big change to the sex ratio paradigm and Lloyd's (1975) model is partly a subjective matter. It was yet the first demonstration that the two factors responsible for the structure of mating patterns had to be considered altogether in plant mating evolution. In other words, at least in the Oleaceae, the evolution of sexes and self-incompatibility have to be considered jointly in order to understand mating system evolution. Previous works speculated on this joint evolution,

but in a different context, and without any direct demonstration: Ehlers and Schierup (2008) showed that a correlation between gynodioecy and SI could be possible (a theoretical prediction partly supported by data) and that heteromorphic SI (i.e., heterostyly) could evolve to dioecy, but without any direct evidence (Barrett 2019a).

Second, distortion segregation and biased sex ratio at birth were needed to fully explain the near-balanced sex ratio 1:1 in natural populations. In the case of gynodioecy, it was suspected very early on that genetic conflicts within individuals, between nuclear and cytoplasmic genomes, could result in the evolution, maintenance, and high frequency of females (Lewis 1941). It was further confirmed and demonstrated in many different species (Dufay and Billard 2012). In *P. angustifolia*, it was shown for the first time that genetic conflicts among nuclear genes resulted in the maintenance of high male frequency in an androdioecious species. This changes the nature of the models that could be used to study such a situation. Indeed, Lloyd's (1975) model is phenotypic, which means that the genetic architecture underlying the sex determination can be neglected, but only under the hypothesis that there are no such evolutionary conflicts at the level of the genes themselves. In this case, taking explicitly into account that genetic architecture is needed, as Billiard et al. (2015) did by modifying Husse et al.'s (2013) population genetics model. But here again, whether or not this should be considered as a big change in the paradigm is a matter of taste. Yet, it means that a whole category of models—phenotypic models—cannot be used, at least in the case of *P. angustifolia*.

Aside from the two previously exposed changes made to the paradigm to solve the *P. angustifolia* paradox, many new questions were raised. First, what are the consequences of the necessity to jointly consider the evolution of SI and sexes? To what extent can it be extrapolated to other plant families? How does it work at genomic, physiological, and morphological levels? Second, even though DSI was discovered in all of the other Oleaceae species that have been checked so far, all species had their own surprising specificities.

*F. ornus* is androdioecious in some populations, but cryptically dioecious in others because one of the two SI groups was lost (Vernet et al. 2016). *F. ornus* populations also show high male frequencies, as high as in *P. angustifolia*, but so far there is no evidence of segregation distortion in this species. *L. vulgare* shows self-compatible hermaphrodites, but in a single direction: self-compatible hermaphrodites belong to the  $G_a$  SI group, they can self-fertilize, they can sire  $G_a$  and  $G_b$  SI hermaphrodites, and can be sired by  $G_b$  SI hermaphrodites, but cannot be sired by  $G_a$  SI hermaphrodites (De Cauwer et al. 2021). *F. excelsior* populations show a quantitative and continuous variation of the allocation to male and/or female reproductive organs, but structured into two SI groups: one group tends to contain hermaphrodites producing a large number of male flowers and a few hermaphroditic flowers, while the other group contains hermaphrodites producing a majority of female or hermaphroditic flowers (Saumitou-Laprade et al. 2018). *O. europaea* shows only hermaphroditic individuals, with no unisexuales and no self-compatible phenotype (Saumitou-Laprade et al. 2017a; Besnard et al. 2020; Mariotti et al. 2021), a situation that is expected to be theoretically unstable (Van de Paer et al. 2015). One can speculate that the existence of DSI opens evolutionary pathways that were not anticipated until now. In particular, it is not known so far which one of the situations encountered in the Oleaceae species are evolutionary stable or on their way to dioecy or other mating strategies.

Finally, maybe the most challenging questions are: How did DSI evolve first, and how is it maintained? Since the species at the roots of the Oleaceae phylogeny are heterostylous, and that all homostylous species derived after a polyploidization event, one can speculate that DSI is heterostyly's "ghost." However, it remains to be demonstrated. More problematic is the open question of the conditions of maintenance of only two SI groups. Indeed, SI always shows dozens of SI groups in all other SI angiosperm families. Indeed, a new SI group is necessarily advantaged, although it is rare because it has a higher number of mating

opportunities than frequent SI groups, which thus facilitates the emergence of new SI groups. Two SI groups is the most favorable situation where a new SI group can emerge. But, in the case of the Oleaceae, only two SI groups are observed, and this situation is certainly a dozen million years old (Vernet et al. 2016). Hence, solving the paradox in *P. angustifolia* finally opened new theoretical questions for which the actual sex ratio paradigm has no answer. As will be argued below on the bases of Kuhn's criteria, these new open questions justify on their own that resolving the *P. angustifolia* dilemma led to a paradigm shift and not simple paradigm adjustments.

#### ANALYSIS OF THE *P. ANGUSTIFOLIA* CONTROVERSY ACCORDING TO KUHN'S THEORY

Our first goal is to address whether or not the controversy about the mating system of *P. angustifolia* was indeed a *paradigm crisis*, in the sense of Kuhn. A paradigm crisis usually emerges from an anomaly. Before specifically analyzing the *P. angustifolia* controversy, we define the central concepts of Kuhn's theory: *paradigm*, *crisis*, and *anomaly*.

#### PARADIGM

A *paradigm* (later called a *disciplinary matrix* in Kuhn 1970) is a set of theories, rules, and tools adopted—often tacitly—and put into practice by a community of scientists. The term *disciplinary matrix* was introduced by Kuhn (1970) following a seminal paper written by Masterman (1970) where the 21 different definitions of *paradigm* in Kuhn (1962) were thoroughly analyzed. Although the term *disciplinary matrix* can be more clearly defined and appropriate for a Kuhnian analysis, we chose to keep the term *paradigm* in the rest of our analysis in order to be more in line with the existing literature on Kuhnian models.

Masterman (1970) identified three main categories under which *paradigm* can be understood: metaphysical, sociological, or construct paradigms (i.e., paradigms as an actual instrumentation). The sociological aspect of paradigms will be analyzed in the rest of this

section (the metaphysical and artifactual aspects will be analyzed in the section titled Models versus Data). Paradigms show two main features:

1. A greater explicative power than other scientific activities; and
2. A proposition of new problems or new approaches to extant problems.

When a paradigm change occurs, it is important that most of the progress previously acquired is conserved. In the case of the *P. angustifolia* controversy, the new paradigm should keep all of the progress achieved by the theories of Fisher (1930), Lloyd (1975), and Charlesworth and Charlesworth (1981). A paradigm shift is thus not a *tabula rasa*.

*Normal science*, also called *paradigm-based research*, is the application of a paradigm where a majority of researchers in a given scientific field attempt to solve particular problems (Figure 2). Normal science and paradigms are exchangeable. Hence, there are as many paradigms as normal science communities, and there are paradigms at all scales, as long as paradigms are required by a scientific community.

Kuhn (1962) exposed canonical paradigms such as Ptolemaic cosmology, Newtonian physics, phlogistics, Einstein's relativity, or quantum mechanics. These general (or high-level) paradigms are keystones to multiple disciplines (Figure 1). Local or small-scale paradigms also exist, such as the one applied to mating system evolution in *P. angustifolia*. However, paradigms at any scale obey the same mechanisms (Hacking 2012; Kuhn 2012). In his postscript, Kuhn (2012) indeed distinguished four levels of scientific communities where paradigms are applied (Figure 1):

1. All natural sciences, where the most general paradigms exist (e.g., units of measurement);
2. General fields, where global paradigms are adopted (e.g., Darwinian theory of evolution for biology);
3. Major subgroups such as ecology or molecular biology, where paradigms include elements of technology or theoretical tools (e.g., Lotka-Volterra and Wright-Fisher

models, the concept of fitness as a predictor of genetic changes, the genotype-to-phenotype unidirectional path, or Fisher's sex ratio paradigm); and

4. Specific research groups such as plant population biology, where local paradigms are adopted (e.g., some species are biological models such as *Arabidopsis thaliana*, the ovule/pollen ratio as predictors of mating systems in plants, or the sex ratio predicted in androdioecious plant populations after Lloyd 1975).

The concept of *paradigm* is thus adapted in the case of the androdioecious status of *P. angustifolia*, even though the associated scientific community interested in this question is small. Although Fisher's sex ratio theory might be considered as a Level 3 paradigm, what we call the Fisher-Lloyd-Charlesworth and Charlesworth paradigm is a Level 4 paradigm, since it was built to specifically explain plant mating system evolution (Figure 1). On a side note, this raises the question whether a Level 1, 2, or 3 paradigm could be refuted by an experiment that necessarily takes place within a Level 4 paradigm. As Duhem ([1906] 2016) pointed out, the different scales and the interdependence of paradigms challenge the notion of *experimentum crucis* and refutability.

#### ANOMALY

*Anomaly* refers to an object that does not fit into the paradigm and whose resolution is not possible within the paradigm it emerged from. In the evolutionary framework of Kuhnian models, anomalies are analogous to evolutionary pressures in ecology by negatively selecting a theoretical set's particular locus. Concerning *P. angustifolia*, the local paradigm that ruled normal science seemed to have been set since 1930 by Fisher and reinforced later on (Lloyd 1975; Charlesworth and Charlesworth 1981). Therefore, *P. angustifolia* is recognized as an anomaly since it does not match the paradigm's prediction (Pannell 2002). This characterization is amplified by the unusual features of *P. angustifolia*. In general, objects with rare characteristics tend to be naturally opposed to paradigms as the paradigm's goal is to be as

general as possible in a particular field. The recognition of an anomaly is chiefly a subjective and collective process that does not imply that what is identified as an anomaly actually is one (Kuhn 1970; Watkins 1970).

#### PARADIGM CRISIS

When an anomaly is identified and collectively accepted as such, a paradigm crisis usually occurs. A paradigm crisis is not a period of downfall for science, but instead leads to a proliferation of new propositions and discussions about the foundations of a paradigm. The start of the *P. angustifolia* crisis took place around 1992 (Lepart and Dommée 1992), while it ended around 2015 (Pannell and Korbecka 2010; Saumitou-Laprade et al. 2010; Billiard et al. 2015).

A paradigm crisis ineluctably leads to what Kuhn stated as *extraordinary science*, in opposition to *normal science*, where scientific activity and production are decreased, and where a scientific field becomes unstable. Extraordinary science naturally causes a division in the formerly solid adherence to a certain paradigm in a given group of scientists. The number of newly formed groups is very variable and sometimes depends on sociogeographic parameters or local scientific tradition. Sometimes there are as many scientific groups as laboratories dedicated to a discipline facing a crisis. Concerning *P. angustifolia*, two research groups were in opposition (see the section titled The *Phillyrea angustifolia* Controversy, Its Resolution, and Extensions for details): the first one, more in agreement with Lloyd's paradigm, assured that based on the validity of the theoretical framework, *P. angustifolia* had to be cryptically dioecious. The second one, based on laborious and clarifying experimentations, assured that *P. angustifolia* was androdioecious, yet without getting rid of the paradoxical situation and the anomaly.

#### ADOPTION OF A NEW PARADIGM

Because it is not a stable state of research, extraordinary science consequently leads to a conservation or a replacement of the formerly shared common paradigm. The con-

ditions for the identification of a paradigm change (Conant 1947; Kuhn 2012; Wray 2021) include:

1. The competing paradigm must resolve a primordial problem or an anomaly (e.g., the unexpected mating system in *P. angustifolia*);
2. The competing paradigm must conserve the majority of the previously acquired knowledge (e.g., DSI and distortion segregation were included into Fisher-Lloyd-Charlesworth and Charlesworth's models);
3. The competing paradigm must open new problematic fields (called *puzzles* by Kuhn) that nevertheless do not challenge the validity of the competing paradigm itself (e.g., the evolutionary origin and maintenance of DSI is still puzzling; see below); and
4. The competing paradigm must propose ideas contradictory to the former one (Conant 1947).

Criteria 2 and 4 seem contradictory at first sight: how can a paradigm shift conserve knowledge while contradicting it? This particularity is due to the evolutionary approach inherent to Kuhn's model. Anomalies are indeed submitted to a pressure analogous to natural selection in biological evolution, which plays on a particular locus of a genome. Anomalies can be considered as a particular locus of a paradigm that suffers epistemological pressure while leaving the rest of the paradigm relatively untouched. Criteria 2 and 4 are therefore compatible since what is contradicted and negatively selected in the old paradigm is only what the new paradigm resolves, while the main corpus of knowledge remains unthreatened.

#### INCOMMENSURABILITY

The competition between the former established paradigm and the propositions of a new paradigm is actually more complex because of what Kuhn called the *incommensurability* of paradigms (a central concept in Kuhn's theory strongly debated in philosophy of science; see Shapere 1964; Watkins 1970; Lakatos 1976; Ruse 1989). The general idea of incommensurability is that each scientific discourse is settled within a particular

theoretical framework that changes particular dimensions of the world. Four different types of incommensurabilities can be distinguished (Kuhn 1962, 2000; Devlin 2021; Wray 2021):

1. Ontological and lexical: paradigms create and carry a set of conceptual beings, while other types of beings cannot be conceptualized. In the *P. angustifolia* controversy, the same being could be at the same time dioecious for a given paradigm and androdioecious for the other. This dimension affects the way scientists see things, and the relationship they establish between these things.
2. Methodological: when beings differ, because of ontological incommensurability, the tools and methods to study them are also different. In the case of *P. angustifolia*, paternity analysis, diallelic crosses, or explicit genetic modeling were necessary to resolve the paradox in addition to Lloyd's (1975) phenotypic models.
3. The incommensurability between specialty communities: the communication and the research on a common ground are nearly impossible since the theoretical and functional frameworks of research groups are too distinct from one another. This was especially important when the controversy about the mating system in *P. angustifolia* was extended to the whole Oleaceae family, and especially in the case of olive oil (*O. europaea*) where one group was mostly constituted of theoreticians in evolutionary biology while the other groups came from agronomy and plant physiology academic backgrounds (see details below).
4. Dissociation: this occurs when writings from an earlier era are not comprehensible under our modern gaze. As the *P. angustifolia* controversy is recent and temporally narrow, this dimension of incommensurability is not relevant.

The problematic aspect of incommensurability comes from the idea that two scientists in two different theoretical frameworks live in two distinct worlds while still looking at the same reality. At best, the different frameworks partially overlap, but they can be mutually exclusive. However, incommensurability should not be confused with incommunicability. Two scientists in distinct

theoretical frameworks can discuss together, but it requires a translation from one lexicon to the other. It even requires a transformation since both of the two participants have to try to see the world as the other sees it (Kuhn 1970:277). This transformation is made possible in Kuhn's theory by its idealistic framework (see the section titled Models versus Data below for details and a discussion of that point in the context of the *P. angustifolia* controversy). On a side note, translation and transformation make *Gestalt* switch possible for philosophers, thus allowing them to roam and temporarily settle within two concurrent paradigms.

Overall, the paradigm change that occurred in *P. angustifolia* can be compared with the discovery of oxygen at the end of the 18th century (Kuhn 1970). In 1774, Joseph Priestley identified the gas collected from heating red oxide of mercury as dephlogisticated common air. In 1775, Lavoisier identified the same gas as unaltered air and concluded that it was a constituent of the atmosphere. Perceptions and interpretations of the same object by Priestley and Lavoisier were incommensurable: they were two different paradigms and two distinct scientific "worlds" (see also Ruse 1989). A similar situation occurred in the case of *P. angustifolia*, based on the opposition of different perceptions and interpretations of the same observations: one research group considered *P. angustifolia* a cryptically dioecious species *because* it showed a 1:1 sex ratio; the other considered *P. angustifolia* androdioecious *because* hermaphrodites had a siring success similar to that of males.

#### INTERNAL RESILIENCE AND EXTERNAL RESISTANCE

The incommensurability of paradigms prevents the adoption of a new paradigm solely based on rational arguments and mathematical reasoning. The stability of a paradigm actually depends on two variables:

1. Internal resilience (coined by us as a synthesis of Kuhn's main ideas): e.g., the paradigm general coherence, its ability to easily incorporate new phenomena, and

the amount of theories whose existence depend on it since all paradigms are entangled vertically and horizontally (Figure 1); and

2. External resistance to the crisis: e.g., to what degree are researchers attached to the paradigm, its place in the learning process of new students, its historical background, and its prestige.

External resistance mechanisms inevitably lead to a form of “scientific crystallization.” When a set of theories are clearly adopted as paradigmatic, they are immediately endowed with a high resistance to anomalous data. Such resistance has multiple possible external sources, since the theoretical core of the paradigms are not drastically changed in the process of adoption. Hoquet (2020), for example, showed that resistance in favor of Bateman’s principle was mostly due to an excessive amount of citations of Bateman’s seminal paper (1948). Scientific crystallization is not to be understood as a pejorative term in this context: it is the tendency to give more confidence and value to the paradigm in times of crisis than in times of normal science. Even though crystallization might lead to regrettable outcomes (e.g., the fervent adherence to a recently abandoned paradigm), it is a vital mechanism to conserve the stability of normal science. Even though scientific crystallization has probably been encountered in the case of *P. angustifolia*, the new paradigm (i.e., the existence of diallelic self-incompatibility and distortion segregation bias in *P. angustifolia*) is now largely adopted by the community (Pannell and Korbecka 2010; Barrett 2019b), even though some resistance is still encountered (e.g., Breton et al. 2021; see below).

The resistances encountered by the new paradigm in the *P. angustifolia* controversy were certainly mainly due to an application of the principle of parsimony to the designated models, where the costs of hypotheses and mechanisms involved are evaluated and balanced. Three elements were differently evaluated by the two competing research groups: changing versus maintaining the Fisher-Lloyd-Charlesworth and Charlesworth’s models and predictions; accepting versus rejecting the observations that hermaphro-

dites were functionally males and females in *P. angustifolia*; and introducing self-incompatibility versus ovule discounting as a new mechanism to reconcile high male frequencies in natural populations and models’ predictions.

The two competing research groups reached two opposite conclusions:

1. Either Fisher-Lloyd-Charlesworth and Charlesworth’s models are too costly to be dismissed and high male frequencies in natural populations of *P. angustifolia* are likely explained by imprecisions or misinterpretations of the experimental observations. As a consequence, according to the established paradigm, hermaphrodites in the populations of *P. angustifolia* are females in a cryptically dioecious system. Ovule discounting was hypothesized as a heavy mechanism to reconcile data and predictions.
2. Or rejecting observations is too costly and high male frequencies in populations of *P. angustifolia* are likely explained by the incompleteness of the paradigm. As a consequence, hermaphrodites are functional and the paradigm has to be changed. According to the observations, heavy adjustments (i.e., DSI and genetic conflicts with distortion segregation bias) had to be included in the established paradigm.

Even though a heavy new mechanism had to be considered in both propositions (ovule discounting versus DSI and distortion segregation), the principle of parsimony indicates that proposition 2 was heavier than proposition 1 because of the cost of rejecting the established paradigm in proposition 2.

According to Kuhn’s theory, it is always possible to challenge the status and the validity of an observed anomaly. It is only when the anomaly is recurrent in time and in different experimental protocols that the parsimony of the models has to be questioned, and that more costly mechanisms must be considered. In the case of *P. angustifolia*, it appears that the anomaly and paradigm crisis needed paradigm changes by the costly introduction into the paradigm of new surprising mechanisms: DSI (an unexpected homomorphic SI system) and distortion segregation

TABLE 1

*Proposition of necessary and sufficient criteria to identify a paradigm shift over a paradigm adjustment (after Kuhn 1962, 2012; Wray 2021)*

	Paradigm adjustment	Paradigm shift
Conservation of the previously acquired knowledge	X	X
Resolution of an anomaly or of a fundamental problem	X	X
Opening of a new problematic field		X
Conant's criteria: the new paradigm partly or totally <b>contradicts</b> the former one		X

(unexpected conflicts between *nuclear* genes involved in sex and SI determination).

THE PRESENT AND FUTURE STATUS OF THE PARADIGM CHANGE IN PLANT MATING SYSTEM EVOLUTION

IS THE DISCOVERY OF DSI IN *P. ANGUSTIFOLIA* A PARADIGM SHIFT OR A PARADIGM ADJUSTMENT?

The identification of a paradigm shift requires four conditions (Kuhn 1962; Wray 2021): (i) the conservation of the previously acquired knowledge; (ii) the resolution of an anomaly; (iii) the opening of a new problematic field, since new paradigms are always formed around a set of problems; and (iv) the new paradigm partly contradicts the former one.

We propose that, although conditions (i), (ii), (iii), and (iv) are necessary to identify a paradigm shift, only conditions (i) and (ii) are sufficient to identify a paradigm adjustment. Conditions (iii) and (iv) are determinative, and have to be fulfilled to conclude the identification of a paradigm shift (Table 1). To summarize our definitions: it is a paradigm *adjustment* if only conditions (i) and (ii) are fulfilled, or a paradigm *shift* if conditions (iii) and (iv) also are. The choice of these two necessary and sufficient criteria can be discussed, but nonetheless offers a framework under which paradigm dynamics can be subsumed while being directly extracted from both Kuhn and Conant's works. As Shapere stated: "where do we draw the line between different paradigms and different articulations of the same paradigm?" (Shapere 1964:387). The problem of the identification of necessary and sufficient criteria to characterize a paradigm shift has

always been present in Kuhn's theory. Nonetheless, it does not mean that it is impossible to find efficient criteria to operate an analysis.

Regarding *P. angustifolia*, condition (i) is fulfilled because the seminal paradigms from sex ratio theory and plant mating system evolution by Fisher-Lloyd-Charlesworth and Charlesworth have been conserved. Condition (ii) is also fulfilled because the anomaly that initiated the crisis about the status of the mating system of *P. angustifolia* has been solved. However, the discovery of DSI in *P. angustifolia* was not the end of the story since DSI became an object of study on its own. Indeed, DSI generated new problems not yet explained by the new paradigm: the evolutionary origins and maintenance of DSI in the whole Oleaceae family and beyond, and the possible consequence that the evolution of sexes, self-incompatibility, and genetic conflicts are generally linked (Billiard et al. 2011; Barrett 2019a,b). Hence, condition (iii) is also fulfilled. Condition (iv) is also fulfilled since we showed that the two propositions "*P. angustifolia* is dioecious" and "*P. angustifolia* is androdioecious" are contradictory. Finally, we conclude that the discovery of DSI in *P. angustifolia* should be considered a paradigm shift, and not only as a paradigm adjustment, contrary to Pannell and Korbecka's (2010) analysis.

Moreover, in his 1970 contribution, Kuhn mentions another criterion that might be called the *consequential* criterion: "The gist of the problem is that to answer the question 'normal or revolutionary?' one must first ask, 'for whom?'" (Kuhn 1970:252). This criterion tries to take account of the *locality* of shifts. It states that in order to call something revolutionary—e.g., to identify a paradigm shift—we have to consider how the practice of

science has been affected relatively to each scientific group. In the case of *P. angustifolia*, although the discovery of DSI did not change anything to the way most biologists of evolution practice science, it drastically changed the way groups dedicated to plant mating systems pursued their research. As Kuhn stated it: “Many episodes will then be revolutionary for no communities, many others for only a single small group, still others for several communities together, a few for all of science” (Kuhn 1970:253).

#### A VARIETY OF RESISTANCES TO THE PARADIGM SHIFT

Resistance to changes is inherent to paradigms as Kuhn’s theory suggests. We presented one such resistance from a single research program, which was resolved between 2010 and 2015. Our demonstration focused on the opposition we thought was the most important, i.e., cryptic dioecy versus androdioecy. However, other resistances to the paradigm can be identified. We present here three categories of opposition, playing at different times. First, a resistance that appeared instantaneously after the paradigm shift, but which has known no posterior developments. Second, a resistance that runs over several years and is still ongoing because one research program only partly accepts the DSI paradigm shift in the olive tree. Third, since new resistances necessarily emerge once a paradigm shift is adopted, we propose one such possible new resistance that can slow down future discoveries in the Oleaceae.

#### Cryptic Distyly

A new resistance instantaneously appeared after DSI was adopted. Pannell and Korbecka concluded that the discovery of DSI was indeed a paradigm change but, at the same time, that “*P. angustifolia* is not cryptically dioecious, but displays something akin to cryptic ‘distyly’” (Pannell and Korbecka 2010:R483). Their conclusion was, however, ambiguous. It would mean that nothing was new under the sun as distyly is common in angiosperms. In other words, if *P. angustifolia* was indeed cryptically disty-

lous, then the discovery of DSI is finally not a paradigm change. Calling for cryptic distyly can thus be interpreted as a resistance to the paradigm change. However, resisting the paradigm change with cryptic distyly raises unnecessary difficulties. Distyly is by definition heteromorphic: these populations show individuals with either long or short styles that coevolved with their animal pollinators. How can heteromorphism be cryptic? What about Oleaceae species not pollinated by animals? Most importantly, adopting cryptic distyly in *P. angustifolia* would have oriented future research about the evolution of the mating system in Oleaceae to a single question: How has heteromorphic distyly been lost while SI is conserved? On the contrary, adopting DSI makes the latter question only a possible evolutionary scenario among others, which are still to be investigated. Adopting the DSI paradigm also raises a general question about the relationships between homomorphic self-incompatibility, heterostyly, and sexes (Barrett 2019a,b).

#### Self-Incompatibility in the Olive Tree

Determining the mating system of the olive tree (*O. europaea*) is a long-standing issue that involved two isolated communities of researchers, with their own paradigms: a community mostly involved in addressing agricultural and physiological issues (e.g., Wu et al. 2002; Breton and Bervillé 2012) and the other community mostly interested in the evolution of plant mating systems (e.g., Saumitou-Laprade et al. 2017a). Observations regarding the mating system in *O. europaea* appeared complex and confusing: the species seemed at the same time self-incompatible, with some cultivars capable of recurrent self-fertilization under some crossing conditions, while other cultivars were only partially self-incompatible. This led a research group to propose a complex organization of the SI system with dozens of *S*-alleles, expressed at the sporophytic level, with complex dominance relationships between *S*-alleles (Breton et al. 2014, 2016). The discovery of DSI dismissed such a complex SI system (Saumitou-Laprade et al. 2017a,b; Besnard et al. 2020; Mariotti et al.



2021). Interestingly, the paradigm change about the mating system in *P. angustifolia* in the community of evolutionary biologists rapidly affected the community interested in *O. europaea*, as shown by the sudden cross-referencing literature between both communities after Saumitou-Laprade et al. (2017a). As predicted by Kuhn's theory, and especially by the existence of incommensurability, there was, and there still is, an ongoing resistance against the DSI paradigm as a relevant mating system in the olive tree (Breton et al. 2017, 2021; Farinelli et al. 2018). This raises the question about the frontiers of a community for a given paradigm: a paradigm change can incidentally affect an independent community, which can become an important source of resistance to the paradigm change. This suggests that the frontiers of paradigms and their communities are not impermeable, leading to incommensurabilities between specialty groups.

The ongoing resistance about DSI in the olive tree can be explained by two different mechanisms. First, due to sociological reasons. The concurrent research program is led by scholars involved in the study of the olive tree for dozens of years, especially for industrial and agricultural purposes. Second, for some scholars, the mating patterns observed in the olive tree are still unexplained by DSI (Farinelli et al. 2018; Breton et al. 2021). These authors concluded that the crossing patterns observed in the olive tree are, on one hand, too complex to be compatible with DSI and, on the other, DSI cannot explain that self-fertilization is recurrently reported in some cultivars. However, it has been shown that self-compatible genotypes can stably coexist with DSI, both theoretically (Van de Paer et al. 2015) and empirically in *L. vulgare* populations (De Cauwer et al. 2021), which makes the existence of self-compatibility in the olive tree perfectly compatible with the DSI paradigm. Yet, the situation in the olive tree is certainly different than in *L. vulgare*: a self-compatible genotype has been identified in the latter, while self-compatibility seems not to be associated to a particular genotype in the former, suggesting that self-compatibility might be a plastic response in the olive tree.

The disagreement between the DSI paradigm's predictions and mating patterns reported in the olive tree can be explained by observation errors in experiments, with at least two different origins. The cultivars used in the controlled crossing experiments could be composed by more than a single genotype as putatively supposed (Mariotti et al. 2021). If a cultivar indeed contains a single genotype, crosses between different individuals from the same cultivar would not be possible under DSI. Hence, a compatible cross within a cultivar might indeed be evidence against DSI. However, this conclusion can be reached only if the genetic composition of the cultivar is verified and controlled, which needs a particular experimental design. Without such a perfectly controlled experimental design, where cultivars are indubiously associated to a single genotype, crosses between and within cultivars can be found compatible or incompatible because apples would be compared to oranges (Bergelson et al. 2016; Saumitou-Laprade et al. 2017b).

Another possible origin of experimental errors could come from crosses not controlled enough. *O. europaea* is wind pollinated, which means that the air surrounding trees can be saturated in pollen, especially by long-distance pollen. If inflorescences are not bagged early enough, flowers might have received pollen from many different trees, especially by compatible trees, which can lead experimenters to false interpretations. Overall, the adoption of the new DSI paradigm leads to casting doubts on the validity of empirical observations, and legitimizes requests to verify the robustness of the data reports (e.g., validation of crosses by paternity analysis; Saumitou-Laprade et al. 2017a,b; Mariotti et al. 2021). Ironically, the validity of the observations was one of the arguments against the interpretation of *P. angustifolia* as an androdioecious species. This shows that even though paradigms change, the nature of resistances do not.

#### New Paradigm, New Resistance: A Never-Ending Rise of Experimental Standards?

Once a new paradigm is adopted, scientific activities are reorganized and science returns

to its normal state (Figure 2), which involves new resistances to paradigm changes. Kuhn's theory thus cannot only help in understanding a paradigm change, but also help in anticipating and identifying new established resistances. The ongoing debate about the mating system in *O. europaea* illustrates such an issue.

The large adoption of the DSI paradigm is accompanied in particular by the prediction that by default an Oleaceae species should show two SI groups, possibly in association with an additional mating phenotype such as males, as in *P. angustifolia*, or self-compatible individuals, as in *L. vulgare*. Yet, the existence of DSI has been demonstrated in a few species only. It is always possible, in principle, to find a species with a different mating system; for example, with more than two SI groups. Since DSI is now the default situation, a research program that would aim to demonstrate, for example, the existence of at least a third SI group of hermaphrodites, would be faced with a high resistance: only high-standard experimental results would convince the new paradigm defenders that DSI is not general as actually thought. What are these high-standard experiments? The most important requirements would certainly be high-standard controlled crosses: on a sufficiently large number of different individuals, with all needed treatments (two positive controls showing that the stigma of the pollen receiver and the pollen of the pollen donor are functional, and two negative controls showing that self-pollen or crossing between individuals of a given group give no fertilization), such that crosses are indubiously controlled enough (pollen contamination should be excluded), definitely assessed by paternity analyses.

Such high-standard experimental conditions have been progressively built to defend DSI as a factor explaining the situation encountered in *P. angustifolia*. Now that the new paradigm has been adopted, these experimental conditions have become the new experimental standards. We can thus speculate that demonstrating an exception to DSI would need to push even further these experimental standards in order to convince the DSI defenders.

#### MODELS VERSUS DATA

The predominant role of models over data in the discussions around *P. angustifolia* is striking. To a certain degree, Fisher's (1930) sex ratio paradigm was the main element of verification since data were faced with its predictions since it led one research group to cast doubts on the validity of observations. Why were models so predominant over data in the case of *P. angustifolia*? Despite such a predominance, why could the paradigm shift occur against models?

Experiments are often naively considered as testing a theory's predictions. However, in the *P. angustifolia* controversy, the reverse occurred since experiments were considered dubious because they were not in agreement with theoretical predictions: the predictions from the Fisher-Lloyd-Charlesworth and Charlesworth's models were used to challenge data interpretations. This turnaround could be due to the nature of experimentation in macrobiology. In physics, for example, the canonical method is relatively adequate since it is easier to perform controlled experiments with isolated objects. In macrobiology, experimental conditions are less easily controlled. In particular, performing experiments of macrobiological close and isolated systems is in practice almost impossible. Models and data are in a balance where the decrease of trust in the validity of experimental conditions increases the trust in models. This raises the general issue about the relationship and agreement between models and data: How can we know whether or not ingredients of models are coherent with observations from the physical world?

The concept of "world" proposed by Kuhn partly addresses this question. Each paradigm adopted by a scientific group corresponds to different scientific worlds, where what is observed is directly influenced by its own set of theories (e.g., Priestley and Lavoisier about oxygen). However, different worlds are not to be conflated with different interpretations—see Fodor (1984, 1988) and Devlin (2021) for the distinction between *weak* and *strong* theory dependence. Lavoisier did not consider the gas extracted from heating red mercury oxide *as pure air, it was*

air. In the case of *P. angustifolia*, one group saw an androdioecious species, the other one saw a cryptically dioecious species. The paradigm thus provides an *a priori* signification to data themselves, which can consequently explain the predominant role of models over experiments (Figure 3).

This incommensurability concerning perception and interpretation is clearly stated by Kuhn (1970:276). The ontological baggage that paradigms carry is inseparably linked to Kuhn’s theory of language. One of the main concerns of his work is indeed to determine the specificity of the scientific languages, and how the terms developed by these attach to nature. The idea of world in Kuhn’s work consequently contains a theory of reference, which permits establishing a relation between a supposed external world, and a lexicon that we attach to it. This point is illustrated in the “double coinage” metaphor (Kuhn 2000:29). Actually, the familiarity between Kuhn and idealist philosophy is something that has already been underlined by Ruse (1970) and Wray (2021).

All of the above leads to questioning the very nature of data, in particular in the case of population biology and evolutionary biology, as illustrated here by the *P. angustifolia* controversy. A long tradition of philosophers, starting from Kant and his *Critique of Pure Reason*

([1787] 2021), followed by Schopenhauer ([1813] 1997, [1844] 2009), Duhem ([1906] 2016), Bachelard (1938), Kuhn (1970), and Hacking (2000), among others, suggested that the notion of “raw data” was dubious. Supposing that knowledge can be built from raw data means that it is possible to extract pure elements of intelligible reality that can be included in theoretical models afterwards. Instead, a less radical approach to experimentation seems to be more adequate: models predetermine experimentations and data are elements of theories that we inject into the empirical reality. This has two implications. First, the interpretation of data varies according to the paradigm. Second, the very nature of data itself is also determined by the paradigm (Ruse 1970; Kuhn 2012). This can explain the origin of the incommensurability of the paradigms in the case of *P. angustifolia* that led to the paradigm crisis.

This does not, however, lead to a total relativism where “everyone has its own equally viable view of the world.” Scientific discourse is mainly descriptive, and description is principally a matter of abstract properties applied to concrete or abstract objects. Proposing that scientific change is not only a matter of logical reasoning does not mean that there is no argument to persuade a colleague to convert to another paradigm (Kuhn 2012).

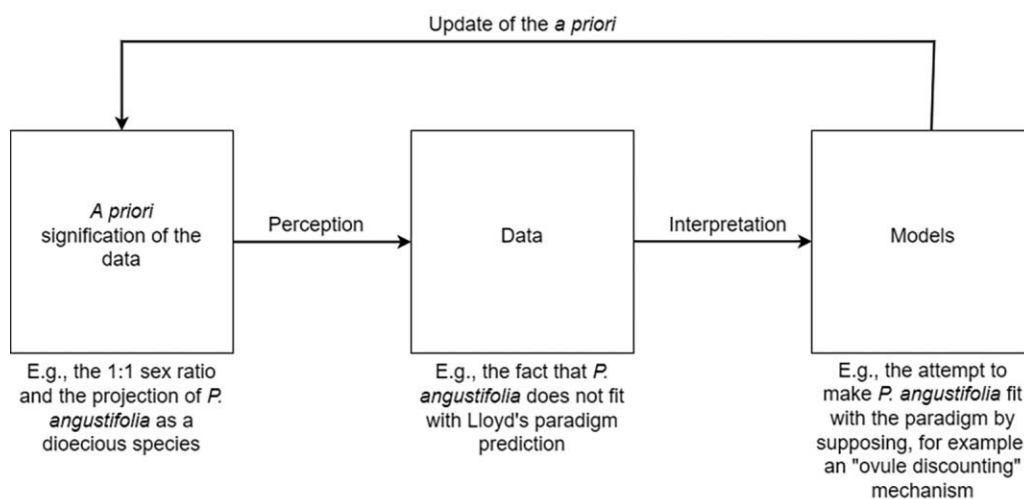


FIGURE 3. INTERACTIONS BETWEEN DATA AND MODELS

As previously mentioned, internal criteria can be found, such as the amount of theories that rely on the existence of a given paradigm. This internal dimension is also why Kuhn should not be misidentified as a relativist but rather as a functionalist. His concern was to determine *why* science works, and fruitfulness—a criteria that Conant (1947) also proposed—is maybe the internal golden standard of Kuhn's theory.

The notion of world also raises a concern that comes from Kuhnian models. If paradigms are two distinct worlds where scientists evolve, then conversions from one paradigm to another should not be possible because it would break the perceptual frameworks in which these individuals have been settled. In most of the examples used by Kuhn, namely the controversy between Lavoisier and Priestley, the tenants of the former paradigm stayed attached to it until they died. This leads to the idea that conversion between paradigms should not be possible, reinforcing the idea that a paradigm change really occurs once the tenants of the former one die. However, in the case of *P. angustifolia*, we can observe that these conversions have occurred, some tenants of the former paradigm have “changed their worlds” to the new one. Is this a refutation of the notion of incommensurability or a refutation of the identification of DSI as a change of paradigm? It is neither.

First, we have to mention that Kuhn was perfectly aware of the fact that conversions can occur: “Note, however, that the possibility of translation does not make the term ‘conversion’ inappropriate. In the absence of a neutral language, the choice of a new theory is a decision to adopt a different native language and to deploy it in a correspondingly different world” (Kuhn 1970:277).

Kuhn did not, however, specify that conversion is chiefly an effect of scale. Since each new paradigm imposes a distinct world from the previous one, and that some paradigms are broader than others (Figure 1), we can suppose that conversions are seldom or absent at Level 2, while being much more common at Levels 3 and 4. Indeed, the broader a paradigm is, the more radically distinct the world it proposes is from the one it

replaces, making conversions impossible for the mind of a single scientist. In lower scales, however, the change of a paradigm only affects particular *loci* of reality. This finding is notably one of the main reasons why the Kuhnian approach should be applied more often to low-level paradigms.

#### UNEXPECTED ORIGINS AND CONSEQUENCES OF PARADIGM SHIFTS

The paradigm shift generated by the resolution of the *P. angustifolia* controversy also illustrates how the scientific analysis of an object with high heuristic assets can have unpredictable economic impacts. This shows that the putative economic value of an object is certainly not a sufficient *criterion* to decide its scientific value. The economic value of *P. angustifolia* is not intrinsic, but especially depends on its phylogenetic relationship with the olive tree. As many other species or specific problems could be studied in close relationships with the olive tree, the economic value of *P. angustifolia* could not be predicted. Even though the origin of the paradigm change was certainly independent of any economic considerations in the case of *P. angustifolia*, this case study suggests that paradigm changes can lead to discoveries with potentially important economic consequences. The demonstration of DSI in olive trees (Saumitou-Laprade et al. 2017a), as expected under the new paradigm, initiated a reflection about how olive tree orchards should be organized, composed, and managed in order to improve and better control the quantity and quality of the production of olive fruits.

#### ACKNOWLEDGMENTS

We thank Lucien Platon for his useful advice and the continual review of our work and Cyprien Cocquyt for his precious reading advice and guidance through the preparation process of this article. We also thank Stefaan Blancke, Koen Tanghe, Pierre Gérard, and Gauvain Leconte-Chevillard for precious discussions and advice. We warmly thank Thibault Masset, Jacques Capelle, and l'Atelier Critique without whom this work would not have existed. We thank Michael Ruse for providing advice and kindly gifting a book we had difficulty

finding. Sylvain Billiard does not thank the administration team of the University of Lille in charge of the man-

agement of students' internships with whom this work would not have existed.

## REFERENCES

- Avisé J. C. 2014. *Conceptual Breakthroughs in Evolutionary Genetics: A Brief History of Shifting Paradigms*. Amsterdam (The Netherlands): Academic Press.
- Bachelard G. 1938. *La formation de l'esprit scientifique: contribution à une psychanalyse de la connaissance objective*. Paris (France): J. Vrin.
- Barrett S. C. H. 2019a. "A most complex marriage arrangement": recent advances on heterostyly and unresolved questions. *New Phytologist* 224:1051–1067.
- Barrett S. C. H. 2019b. Spencer C. H. Barrett. *New Phytologist* 224:1048–1050.
- Bateman A. J. 1948. Intra-sexual selection in *Drosophila*. *Heredity* 2:349–368.
- Bergelson J., Buckler E. S., Ecker J. R., Nordborg M., Weigel D. 2016. A proposal regarding best practices for validating the identity of genetic stocks and the effects of genetic variants. *Plant Cell* 28:606–609.
- Bertoldi N. 2018. Existe-t-il un paradigme darwinien? Pour une ontologie historique de la théorie de l'évolution. *Lato Sensu: Revue De La Société De Philosophie Des Sciences* 5:50–60.
- Besnard G., Cheptou P.-O., Debbaoui M., Lafont P., Huguény B., Dupin J., Baali-Cherif D. 2020. Paternity tests support a diallelic self-incompatibility system in a wild olive (*Olea europaea* subsp. *laperrinei*, Oleaceae). *Ecology and Evolution* 10:1876–1888.
- Billiard S., López-Villavicencio M., Devier B., Hood M. E., Fairhead C., Giraud T. 2011. Having sex, yes, but with whom? Inferences from fungi on the evolution of anisogamy and mating types. *Biological Reviews* 86:421–442.
- Billiard S., López-Villavicencio M., Hood M. E., Giraud T. 2012. Sex, outcrossing and mating types: unsolved questions in fungi and beyond. *Journal of Evolutionary Biology* 25:1020–1038.
- Billiard S., Husse L., Lepercq P., Godé C., Bourceaux A., Lepart J., Vernet P., Saumitou-Laprade P. 2015. Selfish male-determining element favors the transition from hermaphroditism to androdioecy. *Evolution* 69:683–693.
- Breton C. M., Bervillé A. 2012. New hypothesis elucidates self-incompatibility in the olive tree regarding S-alleles dominance relationships as in the sporophytic model. *Comptes Rendus Biologies* 335:563–572.
- Breton C. M., Farinelli D., Shafiq S., Heslop-Harrison J. S., Sedgley M., Bervillé A. J. 2014. The self-incompatibility mating system of the olive (*Olea europaea* L.) functions with dominance between S-alleles. *Tree Genetics & Genomes* 10:1055–1067.
- Breton C. M., Farinelli D., Koubouris G., Bervillé A. 2016. A model based on S-allele dominance relationships to explain pseudo self-fertility of varieties in the olive tree. *Euphytica* 210:105–117.
- Breton C. M., Koubouris G., Villemur P., Bervillé A. J. 2017. "Comment on Saumitou et al. 2017): Elucidation of the genetic architecture of self-incompatibility in olive: evolutionary consequences and perspectives for orchard management." *Evolutionary Applications* 10:855–859.
- Breton C. M., Farinelli D., Koubouris G., Famiani F., Raymond M., Bervillé A. 2021. A dual-successive-screen model at pollen/stigma and pollen tube/ovary explaining paradoxical self-incompatibility diagnosis in the olive tree—an interpretative update of the literature. *Plants* 10:1938.
- Castric V., Vekemans X. 2004. Plant self-incompatibility in natural populations: a critical assessment of recent theoretical and empirical advances. *Molecular Ecology* 13:2873–2889.
- Charlesworth D., Charlesworth B. 1981. Allocation of resources to male and female functions in hermaphrodites. *Biological Journal of the Linnean Society* 15:57–74.
- Charnov E. L. 1982. *The Theory of Sex Allocation*. Princeton (New Jersey): Princeton University Press.
- Charnov E. L., Maynard Smith J., Bull J. J. 1976. Why be an hermaphrodite? *Nature* 263:125–126.
- Conant J. B. 1947. *On Understanding Science: An Historical Approach*. New Haven (Connecticut): Yale University Press.
- Darwin C. 1871. *The Descent of Man, and Selection in Relation to Sex*. London (United Kingdom): John Murray.
- Darwin C. 1874. *The Descent of Man, and Selection in Relation to Sex*. Second Edition. London (United Kingdom): John Murray.
- De Cauwer L., Vernet P., Billiard S., Godé C., Bourceaux A., Ponitzki C., Saumitou-Laprade P. 2021. Widespread coexistence of self-compatible and self-incompatible phenotypes in a diallelic self-incompatibility system in *Ligustrum vulgare* (Oleaceae). *Heredity* 127:384–392.
- Devlin W. J. 2021. Kuhn and the varieties of incommensurability. Pages 105–124 in *Interpreting Kuhn: Critical Essays*, edited by K. B. Wray. Cambridge (United Kingdom): Cambridge University Press.
- Dufay M., Billard E. 2012. How much better are females? The occurrence of female advantage, its proximal causes and its variation within and among gynodioecious species. *Annals of Botany* 109:505–519.
- Duhem P. 1906. *La théorie physique. Son objet, sa structure*. [2016 reprint.] Lyon (France): ENS Éditions.

- Edwards A. W. F. 1998. Natural selection and the sex ratio: Fisher's sources. *American Naturalist* 151:564–569.
- Ehlers B. K., Schierup M. H. 2008. When gametophytic self-incompatibility meets gynodioecy. *Genetics Research* 90:27–35.
- Farinelli D., Breton C., Koubouris G., Famiani F., Villemur P., Bervillé A. 2018. Reply to Saumitou-Laprade et al. (2017). "Controlling for genetic identity of varieties, pollen contamination and stigma receptivity is essential to characterize the self-incompatibility system of *Olea europaea* L." *Evolutionary Applications* 11:1465–1470.
- Fisher R. A. 1930. *The Genetical Theory of Natural Selection*. Oxford (United Kingdom): Clarendon Press.
- Fodor J. 1984. Observation reconsidered. *Philosophy of Science* 51:23–43.
- Fodor J. A. 1988. A reply to Churchland's "Perceptual plasticity and theoretical neutrality." *Philosophy of Science* 55:188–198.
- Friedman M. 2002. Kant, Kuhn, and the rationality of science. *Philosophy of Science* 69:171–190.
- Gayon J. 1992. *Darwin et l'Après Darwin: une Histoire de l'hypothèse de sélection naturelle*. Paris (France): Éditions Kimé.
- Gerber C., Kieffer H. 1898. Androdioécie du *Phillyrea angustifolia* L. et essai sur la filiation des *Phillyrea* de la région méditerranéenne. *Bulletin Scientifique Français et Belge* 31:236–244.
- Greene J. C. 1981. The Kuhnian paradigm and the Darwinian revolution in natural history. Pages 30–59 in *Science, Ideology, and World View: Essays in the History of Evolutionary Ideas*, by J. C. Greene. Berkeley (California): University of California Press.
- Hacking I. 2000. *The Social Construction of What?* Revised Edition. Cambridge (Massachusetts): Harvard University Press.
- Hacking I. 2012. Introductory essay. Pages vii–xxxviii in *The Structure of Scientific Revolutions: 50th Anniversary Edition*, by T. S. Kuhn. Chicago (Illinois): University of Chicago Press.
- Hamilton W. D. 1967. Extraordinary sex ratios: a sex-ratio theory for sex linkage and inbreeding has new implications in cytogenetics and entomology. *Science* 156:477–488.
- Hoquet T. 2020. Bateman (1948): rise and fall of a paradigm? *Animal Behaviour* 164:223–231.
- Husse L., Billiard S., Lepart J., Vernet P., Saumitou-Laprade P. 2013. A one-locus model of androdioecy with two homomorphic self-incompatibility groups: expected vs. observed male frequencies. *Journal of Evolutionary Biology* 26:1269–1280.
- Jablonka E., Lamb M. J. 2005. *Evolution in Four Dimensions: Genetic, Epigenetic, Behavioral, and Symbolic Variations in the History of Life*. Cambridge (Massachusetts): MIT Press.
- Jacob F. 1976. *La Logique du vivant: Une histoire de l'hérédité*. Paris (France): Gallimard.
- Kant E. 1787. *Critique de la raison pure (Philosophie)*. [2021 reprint.] Paris (France): Flammarion.
- Kuhn T. S. 1962. *The Structure of Scientific Revolutions*. Chicago (Illinois): University of Chicago Press.
- Kuhn T. S. 1970. Reflections on my critics. Pages 231–278 in *Criticism and the Growth of Knowledge*, edited by I. Lakatos and A. Musgrave. Cambridge (United Kingdom): Cambridge University Press.
- Kuhn T. S. 2000. *The Road Since Structure: Philosophical Essays, 1970–1993, with an Autobiographical Interview*, edited by J. Conant and J. Haugeland. Chicago (Illinois): University of Chicago Press.
- Kuhn T. S. 2012. *The Structure of Scientific Revolutions: 50th Anniversary Edition*. Chicago (Illinois): University of Chicago Press.
- Lakatos I. 1976. Falsification and the methodology of scientific research programmes. Pages 205–259 in *Can Theories be Refuted?*, edited by S. G. Harding. Dordrecht (The Netherlands): Springer.
- Lepart J., Dommée B. 1992. Is *Phillyrea angustifolia* L. (Oleaceae) an androdioecious species? *Botanical Journal of the Linnean Society* 108:375–387.
- Lewis D. 1941. Male sterility in natural populations of hermaphrodite plants: the equilibrium between females and hermaphrodites to be expected with different types of inheritance. *New Phytologist* 40:56–63.
- Lloyd D. G. 1975. The maintenance of gynodioecy and androdioecy in angiosperms. *Genetica* 45:325–339.
- Lloyd E. 2020. Units and levels of selection. In *The Stanford Encyclopedia of Philosophy* (Spring 2020 Edition), edited by E. N. Zalta. <https://plato.stanford.edu/archives/spr2020/entries/selection-units>.
- Mariotti R., Pandolfi S., De Cauwer I., Saumitou-Laprade P., Vernet P., Rossi M., Baglivo F., Baldoni L., Mousavi S. 2021. Diallelic self-incompatibility is the main determinant of fertilization patterns in olive orchards. *Evolutionary Applications* 14:983–995.
- Masterman M. 1970. The nature of a paradigm. Pages 59–90 in *Criticism and the Growth of Knowledge*, edited by I. Lakatos and A. Musgrave. Cambridge (United Kingdom): Cambridge University Press.
- Morange M. 2017. *Une histoire de la biologie*. Paris (France): Éditions Points.
- Pannell J. R. 2002. The evolution and maintenance of androdioecy. *Annual Review of Ecology and Systematics* 33:397–425.
- Pannell J. R., Korbecka G. 2010. Mating-system evolution: rise of the irresistible males. *Current Biology* 20:R482–R484.
- Pannell J. R., Ojeda F. 2000. Patterns of flowering and sex-ratio variation in the Mediterranean shrub *Phillyrea angustifolia* (Oleaceae): implications for the maintenance of males with hermaphrodites. *Ecology Letters* 3:495–502.

- Ritchie H., Roser M. 2019. Gender ratio. Oxford (United Kingdom): OurWorldInData.org. <https://ourworldindata.org/gender-ratio>.
- Ruse M. E. 1970. The revolution in biology. *Theoria* 36:1–22.
- Ruse M. E. 1987. Is sociobiology a new paradigm? *Philosophy of Science* 54:98–104.
- Ruse M. E. 1989. Is the theory of punctuated equilibria a new paradigm? Pages 117–144 in *The Darwinian Paradigm: Essays on Its History, Philosophy, and Religious Implications*, by M. E. Ruse. London (United Kingdom): Routledge.
- Ruse, M. 2018. The Darwinian revolution: was it a Kuhnian revolution? Pages 195–218 in *Philosophie, Histoire, Biologie: Mélanges offerts à Jean Gayon*, edited by F. Merlin and P. Huneman. Paris (France): Éditions Matériologiques.
- Saumitou-Laprade P., Vernet P., Vassiliadis C., Hoareau Y., de Magny G., Dommée B., Lepart J. 2010. A self-incompatibility system explains high male frequencies in an androdioecious plant. *Science* 327:1648–1650.
- Saumitou-Laprade P., Vernet P., Vekemans X., Billiard S., Gallina S., Essalouh L., Mhaïs A., Moukhlil A., El Bakkali A., Barcaccia G., Alagna F., Mariotti R., Cultrera N. G. M., Pandolfi S., Rossi M., Khadari B., Baldoni L. 2017a. Elucidation of the genetic architecture of self-incompatibility in olive: evolutionary consequences and perspectives for orchard management. *Evolutionary Applications* 10:867–880.
- Saumitou-Laprade P., Vernet P., Vekemans X., Castric V., Barcaccia G., Khadari B., Baldoni L. 2017b. Controlling for genetic identity of varieties, pollen contamination and stigma receptivity is essential to characterize the self-incompatibility system of *Olea europaea* L. *Evolutionary Applications* 10:860–866.
- Saumitou-Laprade P., Vernet P., Dowkiw A., Bertrand S., Billiard S., Albert B., Gouyon P.-H., Dufay M. 2018. Polygamy or subdioecy? The key impact of diallelic self-incompatibility on the sexual system in *Fraxinus excelsior* (Oleaceae). *Proceedings of the Royal Society of London B: Biological Sciences* 285:20180004.
- Schopenhauer A. 1813. *De la quadruple racine du principe de raison suffisante*. [1997 reprint.] Paris (France): J. Vrin.
- Schopenhauer A. 1844. *Monde comme volonté et comme représentation*. [2009 reprint.] Paris (France): Gallimard Education.
- Shapere D. 1964. The structure of scientific revolutions. *Philosophical Review* 73:383–394.
- Strasberg D. 1988. Androdioécie et dynamique de populations chez *Phillyrea angustifolia* en Camargue. PhD diss., Université des Sciences et Technologies de Lille.
- Tanghe K. B., Pauwels L., De Tiège A., Braeckman J. 2021. Interpreting the history of evolutionary biology through a Kuhnian prism: sense or nonsense? *Perspectives on Science* 29:1–35.
- Van de Paer C., Saumitou-Laprade P., Vernet P., Billiard S. 2015. The joint evolution and maintenance of self-incompatibility with gynodioecy or androdioecy. *Journal of Theoretical Biology* 371:90–101.
- Vassiliadis C., Lepart J., Saumitou-Laprade P., Vernet P. 2000a. Self-incompatibility and male fertilization success in *Phillyrea angustifolia* (Oleaceae). *International Journal of Plant Sciences* 161:393–402.
- Vassiliadis C., Valero M., Saumitou-Laprade P., Godelle B. 2000b. A model for the evolution of high frequencies of males in an androdioecious plant based on a cross-compatibility advantage of males. *Heredity* 85:413–422.
- Vassiliadis C., Saumitou-Laprade P., Lepart J., Viard F. 2002. High male reproductive success of hermaphrodites in the androdioecious *Phillyrea angustifolia*. *Evolution* 56:1362–1373.
- Verdú M., González-Martínez S. C., Montilla A., Mateu I., Pannell J. R. 2006. Ovule discounting in an outcrossing, cryptically dioecious tree. *Evolution* 60:2056–2063.
- Vernet P., Lepercq P., Billiard S., Bourceaux A., Lepart J., Dommée B., Saumitou-Laprade P. 2016. Evidence for the long-term maintenance of a rare self-incompatibility system in Oleaceae. *New Phytologist* 210:1408–1417.
- Watkins J. W. N. 1970. Against “normal science.” Pages 25–38 in *Criticism and the Growth of Knowledge*, edited by I. Lakatos and A. Musgrave. Cambridge (United Kingdom): Cambridge University Press.
- Wray B. K. 2021. *Kuhn's Intellectual Path: Charting The Structure of Scientific Revolutions*. Cambridge (United Kingdom): Cambridge University Press.
- Wu S.-B., Collins G., Sedgley M. 2002. Sexual compatibility within and between olive cultivars. *Journal of Horticultural Sciences and Biotechnology* 77:665–673.

Associate Editor: Michael Ruse  
 Handling Editor: Liliana M. Dávalos