



Exaptation Traits for Megafaunal Mutualisms as a Factor in Plant Domestication

Robert N. Spengler^{1*}, Michael Petraglia^{1,2,3}, Patrick Roberts¹, Kseniia Ashastina¹, Logan Kistler², Natalie G. Mueller⁴ and Nicole Boivin^{1,2,3,5}

¹ Department of Archaeology, Max Planck Institute for the Science of Human History, Jena, Germany, ² Department of Anthropology, Smithsonian Institution, National Museum of Natural History, Washington, DC, United States, ³ School of Social Science, The University of Queensland, Brisbane, QLD, Australia, ⁴ Department of Archaeology, Washington University in St. Louis, St. Louis, MO, United States, ⁵ Department of Anthropology and Archaeology, University of Calgary, Calgary, AB, Canada

OPEN ACCESS

Edited by:

Nina Rønsted,
National Tropical Botanical Garden,
United States

Reviewed by:

Julia Heinen,
University of Copenhagen, Denmark
Jun Ying Lim,
Nanyang Technological
University, Singapore

*Correspondence:

Robert N. Spengler
spengler@shh.mpg.de

Specialty section:

This article was submitted to
Plant Systematics and Evolution,
a section of the journal
Frontiers in Plant Science

Received: 04 January 2021

Accepted: 25 February 2021

Published: 24 March 2021

Citation:

Spengler RN, Petraglia M, Roberts P, Ashastina K, Kistler L, Mueller NG and Boivin N (2021) Exaptation Traits for Megafaunal Mutualisms as a Factor in Plant Domestication. *Front. Plant Sci.* 12:649394. doi: 10.3389/fpls.2021.649394

Megafaunal extinctions are recurring events that cause evolutionary ripples, as cascades of secondary extinctions and shifting selective pressures reshape ecosystems. Megafaunal browsers and grazers are major ecosystem engineers, they: keep woody vegetation suppressed; are nitrogen cyclers; and serve as seed dispersers. Most angiosperms possess sets of physiological traits that allow for the fixation of mutualisms with megafauna; some of these traits appear to serve as exaptation (preadaptation) features for farming. As an easily recognized example, fleshy fruits are, an exaptation to agriculture, as they evolved to recruit a non-human disperser. We hypothesize that the traits of rapid annual growth, self-compatibility, heavy investment in reproduction, high plasticity (wide reaction norms), and rapid evolvability were part of an adaptive syndrome for megafaunal seed dispersal. We review the evolutionary importance that megafauna had for crop and weed progenitors and discuss possible ramifications of their extinction on: (1) seed dispersal; (2) population dynamics; and (3) habitat loss. Humans replaced some of the ecological services that had been lost as a result of late Quaternary extinctions and drove rapid evolutionary change resulting in domestication.

Keywords: ecosystem engineering, megafauna, crops, seed dispersal, endozoochory, domestication, origins of agriculture, exaptation

INTRODUCTION

All plants rely, to varying degrees, on ecosystem services performed by animals, including nitrogen transport, carbon respiration, topsoil reworking, removal of competitive vegetation, controlling of pests, decomposition of organic material, and dispersal of seeds and pollen. Many of these services, especially for seed and pollen dispersal, are so advantageous that plants evolve energetically costly traits to recruit animals into mutualistic relationships (Tiffney, 1981, 1984, 2004; Friis and Crepet, 1987; Wing and Tiffney, 1987; Meredith et al., 2011). Angiosperms are particularly adept at developing such mutualisms, and essentially all domesticated plants are angiosperms (Tiffney, 2004; Eriksson, 2008). The evolution of early domestication traits in plants is an example of the recruitment of humans for ecological services, notably seed dispersal (Spengler, 2020). Animals cyclically evolve into larger lifeforms, which fill a keystone ecological niche, but are highly

susceptible to population collapse and extinction (Ferretti, 2008; Froyd et al., 2014; Estes et al., 2016; Malhi et al., 2016). These repeated megafaunal (>40 kg, as a rather arbitrary cutoff commonly cited in the literature) extinctions and subsequent extinctions have affected angiosperm evolution. The late Pleistocene (126,000–12,000 years ago) megafaunal extinctions also applied different evolutionary pressures on the plants that relied on zoochory; in some cases, these adaptations involved the recruitment of new dispersers, notably humans. In this article, we suggest that some of the plant traits for long-distance endozoochoric dispersal, such as rapid annual growth, self-compatibility, heavy investment in reproduction, high plasticity, and rapid evolvability, provided exaptations for agriculture.

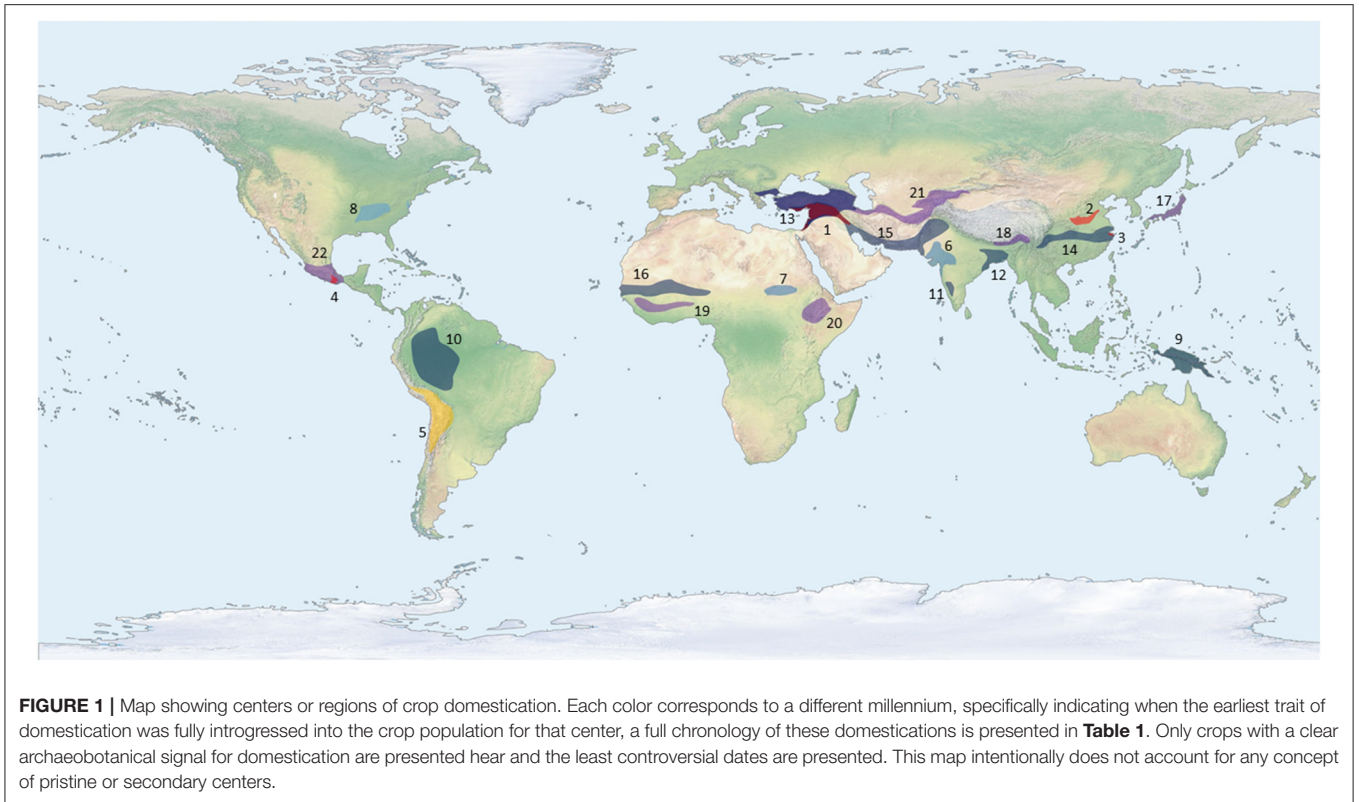
The two main categories of endozoochoric (dispersed through animal ingestion) plants include: (1) those that produce fleshy fruits (Janzen and Martin, 1982) and (2) annual herbaceous plants with small seeds (Janzen, 1984). Most of the first traits of domestication in plants during the early or mid-Holocene were associated with seed dispersal *via* one of these two categories (Rindos, 1984; Spengler, 2020; **Figure 1**; **Table 1**). Given that, these traits allowed for mutualisms in the wild, their importance under cultivation implies an exaptation. We define exaptation following Gould and Vrba's (1982:4) original usage, as: "features that now enhance fitness but were not built by natural selection for their current role." The use of exaptation, as opposed to preadaptation, avoids teleological pitfalls. We also recognize that an argument could be made for humans, as megafauna, being part of a broader adaptive guild, in which case the traits described here would not be exaptations, but simply adapted to earlier iterations of megafaunal mutualisms. Stepping away from the semantics or long-standing adaptationist debates, the key point remains pertinent, that many of the most important traits in plants for agriculture, evolved over the past 50 million years as parts of a series of systems for mutualisms with megafauna. Some of these physiological features, such as sugary fruits, are easy to visualize as exaptation traits allowing for agriculture. However, small-seeded annual crops have traits for survival in an anthropogenic ecosystem that were previously associated with non-human endozoochoric dispersal (Janzen, 1984; Kuznar, 1993; Spengler and Mueller, 2019). These traits include a lack of secondary defensive compounds, rapid annual growth, lack of defensive structures, dry small non-dehiscent fruits displayed on top of the plant, seeds smaller than 2.0 mm, hard protective seed coats, rapid evolvability, high developmental plasticity, and tolerance to trampling and readily disturbed environments. Domesticated plants with fleshy fruits, including many trees, cucurbits, and *Solanum* species, evolved larger fruits, higher sugar concentrations, thicker pericarp tissue, and often secondary traits, such as a reduction in alkaloids, protective structures, and increased size or architecture (Purugganan and Fuller, 2009; Fuller, 2018; Spengler, 2019). These suites of traits illustrate parallel evolutionary responses between: (1) the advantage of zoochory in the wild and (2) anthropogenic selective pressures under early cultivation.

Many crop progenitors are extinct or endangered today and in several cases, wild populations demonstrate low rates of gene flow, reflective of reduced ability to colonize and escape

ecological constraints. Likewise, changing herbivory pressures throughout the Holocene altered the distribution of suitable growing habitats and may have increased competition pressure (Spengler and Mueller, 2019). Often progenitors are restricted to marginal ecologies today, such as river floodplains (Hawkes, 1969; Rindos, 1984). The ecology of seasonally disturbed river edges mimics the disturbed habitats of agricultural fields, and both mimic the disturbance regimes of heavily grazed savannah. Riparian ecologies provide refuge to species with high dormancy rates and limited ability to disperse through space; some plants are able to continue in these settings longer than in non-disturbed locations, where they will be out competed. Exaptation traits from high-herbivory landscapes that increase adaptation to anthropogenic landscapes include a suite of endozoochoric features, rapid evolvability, and developmental plasticity. Links between megafaunal extinctions and the origins of agriculture have already been showcased in the Broad Spectrum Revolution Theory as a factor in human behavioral changes (Stiner, 2001; Kennett and Winterhalder, 2006); however, this theory states that the extinction of large game forced humans to focus more on seeds and less rewarding food sources. Rather than looking at possible economic pressures associated with the megafaunal extinctions, we look at changes in evolutionary pressures in plants and shifting ecosystem dynamics. We argue that the ecological ramifications of the late Quaternary extinctions set the stage for domestication in certain endozoochoric plants (Rindos, 1984; Purugganan and Fuller, 2009; Kistler et al., 2015; Spengler, 2019, 2020).

LOST SEED-DISPERSAL SERVICES

Mutualistic relationships for seed dispersal select for traits that allow seeds to move their offspring away from the parent plant, reducing parent-offspring and sibling-sibling competition (Howe and Miriti, 2000; Wang and Smith, 2002; Schupp et al., 2010; Tarszisz et al., 2018). Seed dispersal also reduces density-dependent mortality, a process linked to the Janzen-Connell Hypothesis (Nathan and Muller-Landau, 2000; Wang and Smith, 2002; Santamaría et al., 2007), interspecific competition (Garant et al., 2007), inbreeding depression (Nathan and Muller-Landau, 2000; Jara-Guerrero et al., 2018), and it promotes gene flow, diversification, adaptive evolution, and overall fitness (Eriksson, 2008). Seed dispersal allows for the colonization of new areas (Escribano-Avila et al., 2014), and dispersal by animals can increase the chance of seed deposition at ecologically favorable sites, what ecologists call directed dispersal (Eriksson, 2008). Nitrogen and water-absorbing organic material in dung also provides an ideal nursery for germinating seeds (Traveset and Verdú, 2002; Fuzessy et al., 2016). Studies show that megafaunal frugivores have a greater ability to disperse seeds and from a wider breadth of species than smaller animals (Peres and Van Roosmalen, 2002; Wotton and Kelly, 2011; Pires et al., 2018). Additionally, these large animals have longer gut passage times, allowing seeds to be moved over greater distances (Ruxton and Schaefer, 2012; Wotton and Kelly, 2012). Seed dispersal is an important evolutionary process in plants, as it ensures gene flow



and changes in allele frequencies (Mayr, 1963; Ellstrand, 2014). Therefore, it should not be surprising that so many angiosperms have evolved similar traits in parallel to recruit animal dispersers.

Small-Seeded Grains and Legumes

Among the ungulates, Artiodactyla (especially Ruminantia) evolved to digest plant cellulose in a four-chambered stomach, using fermentation and regurgitation with secondary mastication. The Perissodactyla retained a monogastric system, relying on intestines for digestion and are, therefore, unable to digest tough vegetative material and more often target sugary fruits. The success of the ruminant digestive system is evident in the sudden radiation and rapid dispersal of the clade across the Northern Hemisphere at about 53 million years ago. Seed dispersal of seeds <2 mm in diameter by Artiodactyla is well-documented (Nathan and Muller-Landau, 2000; O’Farrill et al., 2013) and can readily be observed through modern farm animals. Starting in the Eocene (56–33.9 million years ago), angiosperms began to evolve in response to heavy megafaunal herbivory pressure. Increased phytolith concentrations within and around plant cells appear to have coevolved with flat-crowned grinding teeth (Stebbins, 1981; Jacobs et al., 1999). The evolution of diminished lifecycles (annuals), rapid growth habits, and reduced floral and vegetative plant features, most pronounced in the florets of Poaceae or Cyperaceae, were likely responses to heavy herbivory (Vicentini et al., 2008). Likewise, the basal meristem of all grasses is presumed to be an adaptation for heavy herbivory and being consumed. Annuals invest more

energy into reproduction, express reduced rates of inbreeding depression, and are prone to self-compatibility (Friedman, 2020); scholars have suggested that these traits may be linked to colonization and long-distance seed dispersal (Stebbins, 1950; Baker, 1972, 1965). In this way, some of the widely shared traits that evolved in parallel along with annual lifecycles appear to be tied to endozoochory, as a seed-dispersal syndrome. The world’s first grasslands developed in the Northern Hemisphere in unison with herbivore ecosystem engineering, expressing mutualistic linkages between grazers and annuals (Jacobs et al., 1999). Following this, scholars have argued that herbivory with high-crowned dentition in grazing animals drove the expansion of grasslands and favored rapid-growing annual plants (Kaiser et al., 2013). Megafaunal extinctions of the Pleistocene appear to have led to denser vegetation cover and reduced diversity on some landscapes (Johnson, 2009; Bakker et al., 2016; Malhi et al., 2016) and forest loss on others (Escribano-Avila et al., 2014). Aggregate grazers can cause grasslands to grow faster and change community composition (Frank et al., 1998; Knapp et al., 1999; Geremia et al., 2019). While the fossil record appears to suggest the evolution of annuals prior to the Eocene, all of the most important clades of plants for agriculture evolved into their present forms during, arguably through coevolutionary linkages with megafauna, relatively recently in time.

Janzen (1984) first presented the theory of “Foliage as Fruit,” suggesting that these small annual plants evolved to be consumed by ruminant grazers as a way to disperse their seeds. Many of the plants in the grasslands of the early Holocene, when the

TABLE 1 | A list of centers or regions of domestication by millennium and the crops that evolved domestication traits there, linked numerically to **Figure 1**.

	Region/center	Crop vernacular	Crop binomial	Earliest
(1)	Southwest Asia	Einkorn, Emmer, Barley, Lentil, Pea, Chickpea, Bitter Vetch, Grass Pea, Fava Bean, and Flax ^a	<i>Triticum monococcum</i> ssp. <i>monococcum</i> ; <i>T. turgidum</i> ssp. <i>dicoccum</i> , <i>Hordeum vulgare</i> , <i>Lens culinaris</i> , <i>Pisum sativum</i> , <i>Cicer arietinum</i> , <i>Vicia ervilia</i> , <i>Lathyrus sativus</i> , <i>Vicia faba</i> , <i>Linum usitatissimum</i>	Eighth Mill BC
(2)	North China Plains	Broomcorn and Foxtail Millet	<i>Panicum miliaceum</i> , <i>Setaria italica</i>	Fifth Mill BC
(3)	Yangtze Basin	Melons, Rice, Peaches, Apricots	<i>Cucumis melo</i> , <i>O. sativa japonica</i> , <i>Prunus persica</i> <i>P. armeniaca</i>	Fifth Mill BC
(4)	Oxaca Valley	Maize, Beans, Squash, Pepper	<i>Zea mays</i> , <i>Cucurbita pepo</i> , <i>Capsicum annuum</i> , <i>Phaseolus vulgaris</i>	Fifth Mill BC
(5)	Central/South Andes	Beans, Quinoa, Amaranth	<i>Phaseolus vulgaris</i> , <i>Chenopodium quinoa</i> , <i>Amaranthus caudatus</i>	Fourth Mill BC
(6)	Savannahs of West India	Mung Beans, Little Millet, Sesame, Horse Gram, Kodo Millet	<i>Vigna radiata</i> , <i>Panicum sumatrense</i> , <i>Sesamum indicum</i> , <i>Macrotyloma uniflorum</i> , <i>Paspalum scrobiculatum</i>	Third Mill BC
(7)	East Africa Savannah	Sorghum, Bottle Gourds	<i>Sorghum bicolor</i> , <i>Lagenaria siceraria</i>	Third Mill BC
(8)	Eastern North American Plains	Sunflowers, Chenopods, Sumpweed, Erect Knotweed, Squash, Little Barley	<i>Helianthus annuus</i> , <i>Chenopodium berlandieri</i> , <i>Iva annua</i> , <i>Polygonum erectum</i> , <i>Cucurbita pepo</i> , <i>Hordeum pusillum</i>	Third Mill BC
(9)	Island Southeast Asia	Bananas	<i>Musa acuminata</i> , <i>M. balbisiana</i>	Second Mill BC
(10)	Lowlands South America	Squash, Chocolate, Guava, Peanut, Peach Palm	<i>Cucurbita moschata/maxima</i> , <i>Theobroma cacao</i> , <i>Psidium</i> sp., <i>Arachis hypogea</i> , <i>Bactris gasipaes</i>	Second Mill BC
(11)	South India	Millet and Mung Beans	<i>Brachiaria ramosa</i> , <i>Vigna radiata</i>	Second Mill BC
(12)	East Indian Plains	Cucurbits, Rice, Pigeon Peas	Cucurbits, <i>Cajanus cajan</i> , <i>O. sativa indica</i>	Second Mill BC
(13)	Eastern Mediterranean	Grapes, Olives, Oats, Spelt, Bread Wheat, Rye, Lupine, Common Vetch	<i>Vitis vinifera</i> , <i>Olea europaea</i> , <i>Avena sativa</i> , <i>Triticum aestivum</i> ssp. <i>spelta</i> , <i>T. aestivum</i> , <i>Secale cereal</i> , <i>Lupinus albus</i> , <i>Vicia sativa</i>	Second Mill BC
(14)	Central China	Peaches, Apricots, Persimmon, Cannabis, Soy Beans	<i>Prunus persica</i> , <i>P. armeniaca</i> , <i>Diospyros kaki</i> , <i>Cannabis sativa</i> , <i>Glycine max</i>	Second Mill BC
(15)	Southwest Asian Exchange	Jujube, Citrus, Date Palm	<i>Ziziphus jujube</i> , <i>Citrus</i> spp., <i>Phoenix dactylifera</i>	Second Mill BC
(16)	West Africa Sahel	Pearl Millet	<i>Pennisetum glaucum</i>	Second Mill BC
(17)	Japanese Islands	Adzuki, Soy Beans, Barnyard Millet	<i>Vigna angularis</i> , <i>Glycine max</i> , <i>Echinochloa esculenta</i>	First Mill BC
(18)	Southern Himalaya	Common and Tatar Buckwheat	<i>Fagopyrum esculentum</i> , <i>F. tataricum</i>	First Mill BC
(19)	West Africa	Fonio and Guinea Millet, African Rice, Cow Pea	<i>Digitaria exilis</i> , <i>Brachiaria deflexa</i> , <i>Oryza glaberrima</i> , <i>Vigna unguiculata</i>	First Mill BC
(20)	Ethiopian Plateau	Teff, Finger Millet, Peas, Oats	<i>Eragrostis tef</i> , <i>Eleusine coracana</i> , <i>Pisum abyssinicum</i> , <i>Avena abyssinica</i>	First Mill BC
(21)	Silk Road Mountain Belt	Apples, Walnuts, Pistachios, Almonds, Russian Olives	<i>Malus domestica/pumila</i> , <i>Juglans regia</i> , <i>Pistacia vera</i> , <i>Prunus dulcis</i> , <i>Elaeagnus angustifolia</i>	First Mill BC
(22)	Southern Meso-America	Lima Bean, Common Bean, Squash, Chili	<i>Phaseolus lunatus</i> , <i>Phaseolus vulgaris</i> , <i>Cucurbita pepo</i> , <i>Capsicum annuum</i>	First Mill BC

^aMuch of this information comes from Larson et al. (2014).

Note that we only present species with clear evidence for domestication, noting when the traits of domestication were significantly introgressed into the cultivated population. We have left out crops selected for vegetative parts, because currently there is a lack of reliable data on when they evolved domestication traits and there are few discussions of what the selective pressures would have been that drove domestication.

first seed foraging humans crossed the Northern Hemisphere, had specific adaptations to a heavy herbivory landscape. Notably, they display seeds above terminal foliage and lack secondary metabolites or physical (thorns/spikes) defenses, making them the ideal food for ungulate grazers. Their seeds have hard testae, they remain small enough to pass through a ruminant digestive system, and have high dormancy rates (Spengler and Mueller, 2019). Some of these phenotypic traits are the first traits to

change once humans take over as the main dispersers (Janzen, 1984; Kuznar, 1993; Spengler and Mueller, 2019). Many crop progenitors and ruderal or agricultural weeds express traits of endozoochoric dispersal and may have evolved on heavily grazed landscapes. The prominence of endozoochoric seeds in the stomach contents of frozen tundra Pleistocene grazers, studies of seed germination rates post-digestion in extant grazers, and rangeland composition studies all demonstrate the important



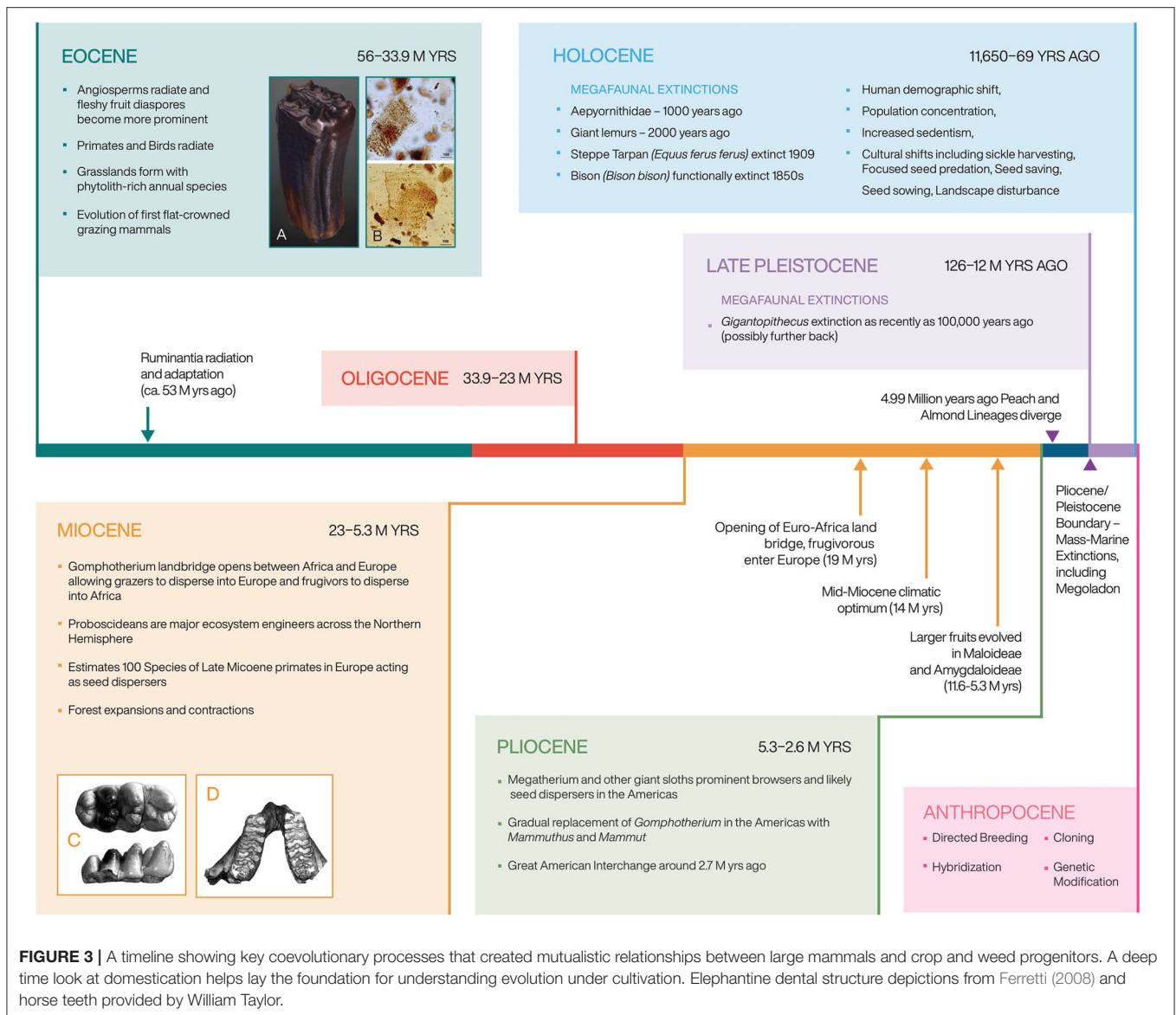
FIGURE 2 | From top left: (1) Photo of horses consuming wild apples in the Tien Shan apple forests of Kazakhstan, photo taken by Artur Stroscherer and Martin Stuchtey. (2) Yaks grazing in Tibet have significant ecological impacts, photo by lead author. (3) Giraffes near the Olduvai Camp in Tanzania browsing on Acacia trees, photo taken by Yiming Wang. (4) Bison in Missouri on a heavily grazed field of herbaceous annuals, most of which are well adapted to heavy herbivory and disturbed ecosystems, photo by lead author.

role that megafauna provided to many progenitors (Spengler and Mueller, 2019) (**Figure 2**).

Large Fleshy Fruiting Plants

Monogastric megafauna with browsing feeding behaviors readily disperse large seeds from within fleshy fruits. Extinction rates of Perissodactyla were much higher during the terminal Pleistocene. Despite high seed loss from mastication and digestion, the evolution of energetically costly sugary pericarp tissue and hard testae is testament to the adaptive advantage of endozoochory. Often the seeds within these fruits will not germinate without scarification or removal of the fruit coat; both are achieved through digestion (Traveset, 1998; Samuels and Levey, 2005; Robertson et al., 2006; Traveset et al., 2007). High dormancy rates in endozoochoric plants also prevent seed germination unless the seed has been consumed and dispersed (Baskin and Baskin, 2004; Tarszisz et al., 2018). In some cases, plants are able to maintain low rates of dispersal despite the loss of one or several dispersers (Guimarães et al., 2008; Jaroszewicz, 2013). Escribano-Avila et al. (2014) suggest that guilds of megafaunal browsers are responsible for woodland expansion in tropical South America. Jara-Guerrero et al. (2018) demonstrate that many large-fruited trees in Central and South America are still dispersed by the few remaining megafauna, but that the general ability for dispersal has been greatly reduced. Studies of isolated populations of *Euterpe edulis* in the Brazilian Atlantic forest, illustrate that the functional extinction of dispersers can drive astonishingly rapid genetic changes in fruit size (Galetti et al., 2013).

Their ability to consume large amounts of fruits with large seeds make large-bodied animals particularly effective seed dispersers (Brodie et al., 2009; Campos-Arceiz and Blake, 2011; Sekar et al., 2015). Most megafruits are too large to be effectively disperse by medium-sized animals, which often function as seed predators (Guimarães et al., 2008; Jansen et al., 2012; Pires et al., 2014). In this paper, we define megafruit as any fleshy fruit too large for an avian disperser; in the temperate zones this is usually >25 mm, as Corvids are the largest avian dispersers, but in tropical zones with parrots and their kin, a cutoff of >40 mm has been posed (Onstein et al., 2018). Seed dispersal studies of small or medium-sized mammals and megafruits rather consistently demonstrate seed predation or weak dispersal ability (Murphy, 2018). In tropical zones, the guild of megafaunal dispersers is not restricted to mammals, and during the past few million years has included large-bodied birds and reptiles, notably tortoises (Valido and Olesen, 2007; Falcón et al., 2020). Studies of megafruit dispersal by large-bodied animals have been conducted in ecosystems around the world, including savannas and drier forest patches (Campos-Arceiz et al., 2008; Sekar and Sukumar, 2013; Sridhara et al., 2016) and in tropical forests (Kitamura et al., 2007; Harich et al., 2016). Several recent studies have shown that the megafaunal extinctions caused a loss or complete termination of seed-dispersal processes in many arborescent plants (Guimarães et al., 2008; Galetti et al., 2017; McConkey et al., 2018; Onstein et al., 2018; Pires et al., 2018; Lim et al., 2020).



Onstein et al. (2018) link megafaunal extinctions to an evolutionary decrease in the size of the fruits produced across the Arecaceae family and high rates of species extinctions throughout the Holocene (Lim et al., 2020). Purugganan (2019) used date palms (*Phoenix* spp.) as a case study for domestication through hybridization, noting that isolated populations of palms were brought into contact by humans. Spengler (2019) suggested, using archaeobotanical data from across Europe, that the mid-Holocene distribution of wild apples was restricted to glacial refugia. Apples, as with many domesticated perennials, appear to have rapidly evolved through hybridization, as humans dispersed plants between distant populations (Hughes et al., 2007; Cornille et al., 2014). Large-fruited *Prunus* trees in Asia have restricted ranges and, in effect, the smaller the fruit, the wider the range today. Some of these species reproduce largely asexually despite heavy investment in a diaspore. The progenitor for the peach is

thought to have once being widely distributed across northern China, but is now presumed extinct (Kostina, 1971; Lu and Bartholomew, 2003; Weisskopf and Fuller, 2014). Yu et al. (2018) and Su et al. (2015) suggest that megafaunal primates may have been a key component in the evolution of large fleshy fruits during the late Miocene or Pliocene.

Another informative case study of the loss of a megafaunal seed disperser comes from the Madagascar mega-lemurs (e.g., *Megaladapis edwardsi*), which went extinct as recently as two millennia ago (Burney et al., 2004; Crowley et al., 2011; Federman et al., 2016). Two of these extinct species, *Archaeolemur majori* and *Pachylemur insignis*, have been described as seed dispersers (Godfrey et al., 2008), and several others are thought to have been largely frugivorous (Crowley et al., 2011). Crowley et al. (2011) provide a list of trees with fleshy fruits that were likely dispersed by these giant primates, which today are losing range and have no

ability to colonize new areas without human assistance (also see Rapanarivo, 1999). Some biologists have also noted that the most threatened large-fruited species have seeds that are too large for successful dispersal by extant lemurs, which have smaller bodies than their precursors (Godfrey et al., 2008; Federman et al., 2016). Genetic studies of some of the trees in these vanishing forests show that they express genetic differentiation and fragmentation into subpopulations (Voigt et al., 2009; Federman et al., 2016). The introduction of new invasive species and domesticated herbivores to Madagascar, including horses, may further serve as surrogate dispersers (Vavra et al., 2007). The observations made in Madagascar are not unique, the loss of frugivores has led to a corresponding decrease in large-fruited arborescent species around the world (Peres and Palacios, 2007; Stoner et al., 2007; Terborgh et al., 2008; Beaune et al., 2013; Tarszisz et al., 2018; Lim et al., 2020). Likewise, “rewilding” exercises, the practice of introducing animals with similar ecosystem functions as those lost during late Pleistocene or Holocene extinction events, serve as field studies to test theories relating to seed-dispersal mutualisms (Donlan, 2005; Griffiths et al., 2011). Many other studies have suggested population fragmentation, genetic heterogeneity, and extinction in plants that produce large fleshy fruits in response to megafaunal extinctions (Eriksson, 2008; Malhi et al., 2016; Galetti et al., 2017; Pires et al., 2018; van Zonneveld et al., 2018).

LOSS OF HERBIVORY AND DISTURBANCE REGIMES

Some paleoecologists have suggested that the extinction of herbivores caused shifts in vegetation cover and species composition, with some areas moving into secondary succession and dense forest cover, while other woody species lost their ability to colonize, causing forests to retreat (Johnson, 2009; Malhi et al., 2016), shifting vegetation densities and composition (Guimarães et al., 2008; Geremia et al., 2019). The reshaping of ecosystem structure shifted selection pressures and changed which plant species were dominant (Rule et al., 2012; Bakker et al., 2016; Malhi et al., 2016). The late Pleistocene megafaunal extinctions also may have caused evolutionary ripples in surviving animal populations, as predator pressures and competition changed (Estes et al., 2011). Scholars have also suggested that herbivore extinctions caused reductions in available nitrogen, calcium, and other soil nutrients (Doughty et al., 2013) and changed atmospheric composition (Doughty et al., 2010; Smith et al., 2010). Understanding the domestication of plants requires a better understanding of the shifting ecological pressures plant communities faced during the early Holocene.

Paleoecologists suggest that the extirpation of megafaunal grazers increased natural fire regimes and changed vegetation communities (Burney et al., 2003; Rule et al., 2012; Codding et al., 2014; Kaplan et al., 2016). As just one of many grazing and rangeland ecology studies, the results of years of fenced enclosure research of Yellowstone’s Northern Range (Council, 2002), ecologists state that elk (*Cervus canadensis*) cause extreme reductions in woody plant growth, suppressing the regeneration

of forests. Studies show that heavy bison (*Bison bison*) grazing on the North American grasslands dramatically alters vegetation communities and can replace the ecosystem services currently implemented by park service officials through controlled burning (Geremia et al., 2019). Bison and other megafaunal ruminants of the Pleistocene are described as creators and maintainers of grasslands or mosaic landscapes (Feranec et al., 2011; Gill et al., 2013; Malhi et al., 2016; Lundgren et al., 2018). Similar observations have been made regarding the loss of megafauna in Australia being linked to fire-prone landscapes before prehistoric peoples started partaking in controlled burning (Rule et al., 2012; Codding et al., 2014; Van Der Kaars et al., 2017). Many paleoecologists argue that the terminal Pleistocene extinctions in eastern North America increased woody vegetation and dead biomass buildup leading to increased magnitudes and rates of wild fires (Robinson et al., 2005; Gill et al., 2009; Gill, 2014). During the Holocene, in many parts of the world, humans became key ecosystem engineers, creating grasslands and mosaic landscapes through burning (Codding et al., 2014; Kaplan et al., 2016). Effectively, humans have replaced the ecosystem engineering roles of large mammals by clearing forests, burning vegetation, and turning over soil, constructing landscapes favorable to a certain community of plants (Kaplan et al., 2016).

Many crop or weed progenitors favor anthropogenically disturbed soils around villages or in cultivated fields, the Dump Heap Hypothesis (Anderson, 1952; Sauer, 1952), paralleling the highly disturbed settings of a bison wallow or a mammoth run. Scholars have long recognized that progenitors of annual crops share similar characteristics with agricultural field weeds and likely evolved for similar disturbance habitats (Hawkes, 1969; Baker, 1974; Vigne, 2015). This suite of traits is variably referred to as weedy, anthropophilic, commensal, or prone to disturbed environments. Scholars have argued that the anthropophilic traits of certain plants brought them closer to human settlements or “dump heaps”—areas where humans cleared woody vegetation, removed slower growing herbaceous competition, increased soil nutrient levels, and eventually maintained these “camp followers” (Anderson, 1952). Anthropophilic species tend to: (1) express high rates of developmental plasticity; (2) be better adapted to disturbed or dynamic ecosystems; and (3) have greater genetic evolvability (Baker, 1974; Geiger et al., 2018). Interestingly, anthropophilic traits, such as rapid growth (at the expense of overall competitiveness with slower growing species), lack of secondary chemicals or defensive structures, high dormancy rates, high rates of developmental plasticity, greater adaptability, short generations, prolific generation production, and rapid rates of evolutionary change, are all traits that favor landscapes with heavy herbivory pressure.

Most ecologists agree that agricultural weeds and many progenitors have exaptation traits allowing for success under disturbance regimes (Adams and Baker, 1962; Baker, 1965, 1972; Clements et al., 2004). Roughly half of the weedy species of annuals are selfing, leading to populations that tend to be genetically homogenous within a population, but highly divergent between populations (Friedman, 2020)—laying the foundations for hybridization (Clements et al., 2004). Selfing also allows a colonizing seed or founder population to reproduce

even if it is not in pollinator range of a compatible mate, helping reestablish after disturbance (Stebbins, 1950; Baker, 1965, 1972). Weed ecologists have noted that selfing and high rates of dormancy often work together as mechanisms for dispersal, either through time on readily disturbed environments or across space as with long-distance colonization. Weedy annuals also tend to express remarkable capacities to tolerate treading, including by automobiles, suggesting adaptation to heavily trodden areas (Baker, 1974).

The mystery of this broad clade of weedy annuals is how they express such strong exaptation, given that the “level and frequency of fluctuations experienced in cropping systems is seldom encountered in more natural ecosystems” (Clements et al., 2004). Some ecologists have suggested that European weeds evolved on Pleistocene tundras, and gradually transitioned into agricultural fields (Salisbury, 1942; Godwin, 1960); other scholars have favored the riverbank or Floodplains Theory (Anderson, 1952; Sauer, 1952; Adams and Baker, 1962; Baker, 1965; Rindos, 1984). Given the prominence of seeds from weedy annuals in the stomachs of frozen megafauna in tundra ecosystems, it seems likely that weedy annuals evolved the suite of traits discussed above on heavily grazed fields with herbivores. These herbivore-heavy ecosystems would have dominated Pleistocene Europe and continued in many areas until the advent of farming, at which point endozoochoric annuals could have evolved into agricultural weeds, in a few cases they could have continued to evolve into field crops. The feedback-driven dynamics of human-weed ecology favor similar conditions as those seen on pre-Holocene landscapes, ecological conditions that today are limited to very small pockets outside anthropogenic setting, such as river floodplains or fire-prone landscapes.

The debate continues over what drove the high level of adaptability and invasiveness in weedy species, with two groups of thought: (1) those who argue that these plants primarily express a high degree of developmental plasticity (Neve et al., 2009); and (2) those who argue that they are capable of rapid genetic change (Baucom and Holt, 2009). Baker (1965) campaigned the view of adaptiveness through highly flexible developmental systems. He was also quick to note that many of these plants do evolve genetic responses rapidly and that plasticity itself is genetically programmed—i.e., not mutually exclusive viewpoints (Neve et al., 2009). Over the past few decades, with more genomic research on agricultural weed populations, scholars have increasingly recognized the remarkable ability for rapid evolutionary change in these plants. These genetic changes occur in a cultivated field on the scale of decades, and are increasingly being recognized as common ecological phenomena, notably in perturbed anthropogenic ecosystems (Thompson, 1999; Palumbi, 2001; Clements et al., 2004). The genetic basis of weedy and invasive traits may be tied into a genetically programmed exaptation, allowing both rapid evolution and developmental plasticity (Baucom and Holt, 2009; Stewart et al., 2009). We argue that, one plausible explanation for the evolution of these traits across a remarkably diverse guild of plants, representing dozens of unrelated families is the collective evolutionary pathway toward adaptation to disturbed landscapes, most notably heavily grazed fields.

PLANT DOMESTICATION

The earliest traits of domestication in plants are associated with changes in seed-dispersal mechanisms (Rindos, 1984); this is well-recognized in the switch from dispersal by brittle rachises in wild cereals to sickle harvesting and seed sowing by humans, causing an evolutionary shift to the tough-rachised allele in the cultivated population (Ladizinsky, 1979; Fuller and Allaby, 2009; Langlie et al., 2014; Larson et al., 2014; Li and Olsen, 2016; Jones et al., 2021). Most of the earliest plants to express evolutionary traits of domestication had seeds that were dispersed by animals in their wild forms (Spengler, 2020). These endozoochoric crop progenitors fall into two broad categories, plants with fleshy fruits and annual herbaceous plants with small seeds—in both cases, megafaunal mammals likely played a primary role as seed dispersers. In **Figure 1**, we present a revised version of the earliest centers of domestication map, whereas we do not separate primary or secondary centers, and instead, see domestication as the result of increasingly more intense interaction between humans and select plant species. Likewise, in **Table 1**, we present a summary of the earliest crops domesticated in each center, as illustrated by morphological changes seen in archaeobotanical remains (following data accumulated by Fuller et al., 2014; Larson et al., 2014; Spengler, 2020). Ultimately, agriculture is better thought of as a complex mutualism between humans and plants, whereas domestication is a set of evolutionary changes associated with that mutualism. Several factors influence the rates of these evolutionary changes, including the insularity of cultivated populations and the presence of exaptation traits.

Rindos (1984) posited that the reason the oak tree was never domesticated, despite intensive human harvesting of acorns on three continents over several millennia, is because of the continued successful dispersal by Sciuridae. The prominent seed dispersal rates and strong wind pollination result in a different population structure than most large fruit trees (Petit et al., 2003; Curtu et al., 2007). Extreme generation lengths may also have hindered domestication in *Quercus*, but many long-generation fruiting trees did evolve in response to human selective pressures during the mid- to late Holocene (Miller and Gross, 2011; Fuller, 2018). A similar argument could be made for avian dispersed fruits, which in most cases, maintained successful seed-dispersal regimes into the Holocene; tellingly, there are few examples of small berries evolving domestication traits prior to directed breeding and the intentional implementation of hybridization (except *Vitis vinifera*). Miller and Gross (2011) note that protracted models of domestication might be expected for arborescent crops, and Fuller (2018) argued for protracted models in some species, but this is not the case from even a cursory look at long-generation perennial crops. In addition to their long generations, clonal propagation further reduced the number of sexual cycles separating wild and domesticated forms (Zohary and Spiegel-Roy, 1975). It is clear that hybridization is a key factor in long-generation perennial evolution in the wild and under cultivation, especially for endozoochoric species. Self-incompatibility among long-generation perennials also facilitates greater rates of interspecific hybridization (Hamrick et al., 1992; Ellstrand et al., 1996; Petit et al., 2003; Ward et al., 2005; Curtu et al., 2007; Miller and Gross, 2011). Hamrick et al. (1992) claim

that among woody plants, endozoochoric species tend to have greater diversity within a species or a population; although, they do not consider whether extinction of dispersers increases or decreases diversity. The loss of megafaunal dispersers left many of the progenitors of modern crops with lower levels of gene flow, fragmentary and isolated populations, and heterogeneous population structures.

Domestication through hybridization is largely the result of genetically isolated populations being brought together as humans crossed their former seed-dispersal barriers. For example, roughly 3,000 years ago, as humans dispersed dates (*Phoenix dactylifera*) out of North Africa, they hybridized with *P. theophrasti*, a previously isolated wild relative (Purugganan and Fuller, 2009; Flowers et al., 2019). Isolated populations of phoenix palms existed across South and southwest Asia, notably in Yunnan, northern India, and the Arabian Peninsula during the mid-Holocene. Other South Asian fruits may have experienced similar range contractions when humans began to disperse their seeds, including *Ziziphus* spp., and *Citrus* spp., both of which were domesticated through hybridization for isolated wild populations. Similarly, in Central America, avocado (*Persea americana*), papaya (*Carica papaya*), and white sapote (*Casimiroa edulis*) were all megafaunal dispersed in their progenitor habitats, as were *Annona* spp., cashew (*Anacardium occidentale*), and guava (*Psidium guajava*) in South America.

Recent research into the domestication of peach palms (*Bactris gasipaes*) in South America has identified a heterogeneous mosaic of wild relatives, which appear to have played varying roles in the domestication process (Clement et al., 2017). Clement et al. (2017:149) note that “dispersal by humans is effective for crossing barriers” and shifting distributions of Late Glacial Maximum populations of the palms are a key aspect of domestication. *Musa* spp. provides another interesting case study of perennials evolving through human-driven barrier crossing. Most cultivated bananas today are in the Eumusa section, and are diploid or triploid hybrids, involving *Musa acuminata* × *M. balbisiana*. Genetic studies show that hybridizations between different *M. acuminata* subspecies occurred in island Southeast Asia and western Melanesia, and involved humans dispersing the plants between islands (Perrier et al., 2011). This hybridization led to parthenocarpy and eventually fed into a selection for sweeter fleshier fruits; illustrating, yet again, that the earliest traits of domestication are tied into seed dispersal (Heslop-Harrison and Schwarzacher, 2007). Further hybridization led to the tetraploid forms (Ude et al., 2002; Raboin et al., 2005).

North American wild cucurbits may have had a much wider range of distribution in the mid-Holocene than today (Fritz, 1999)—currently rare or extinct. There are at least five domesticated species, each brought under cultivation in different regions of the Americas and at different times (Nee, 1990). Kistler et al. (2015) used high-throughput sequencing to analyze complete placid genomes of 91 ancient domesticated, modern domesticated, and modern wild specimens. Additionally, they studied bitter taste receptors in extant megafaunal mammals and note that the toxic and bitter triterpenoid compounds in wild cucurbits would have been less likely to deter large mammals than small mammals. They argue that the extinction of large-bodied seed dispersers across the northern portion of the Americas

dramatically altered ecological pressures on wild populations and illustrated that population fragmentation was the direct result of the loss of megafaunal dispersers, and the domestication of all modern squashes is the result of humans taking on the role of surrogate megafaunal dispersers (Kistler et al., 2015).

Exaptation Traits Supporting Domestication

Scholars have argued that the “key to successful domestication of a crop lies... in its genetic endowment,” claiming some plants express genetic traits that allow for domestication (Murphy, 2007). Scholars have also pointed out that the small-seeded annual crop progenitors, including most agricultural weeds, were “specifically pre-adapted to growth and competition for resources in agriculture-like environments” (Vigueira et al., 2013). They further state that “species that are naturally adapted to disturbed habitats may be pre-adapted to the agricultural environment” (Vigueira et al., 2013). It is not a coincidence that only grasses and Core Eudicots have proven suitable for cultivation; furthermore, within this grouping, only a handful of families have been adopted as field crops, including Poaceae, Amaranthaceae, Fabaceae, Brassicaceae, Lamiaceae, Solanaceae, and Rosaceae. These families are all highly specious and, with the exception of the grasses, are all relatively recently evolved lineages. There is reason to argue that the speciation and diversification of these families occurred in part due to coevolutionary linkages with megafaunal mammals, mostly since the Eocene. We argue that many of the exaptation traits that allowed plants to become crops or weeds, evolved as adaptations to heavy herbivory, including rapid evolvability, developmental plasticity, and likelihood of change through hybridization. This argument would also imply that farming, as it exists today, could not have evolved on Earth during an earlier period (Figure 3).

Each interglacial forced the ancestral populations of domesticated plants to migrate across an entire continent (Clark, 1998). The remnant fragmentary populations of large-fruited trees across: (1) North America, *Maclura pomifera*, *Asimina triloba*, and *Prunus Americana*; (2) Europe, *Sorbus domestica*, *Mespilus germanica*, *Pyrus pyraeaster*, and *Malus sylvestris*; (3) Asia, *Prunus mira*, *P. davidiana*, and *Diospyros kaki*; and (4) South America, *Annona cherimola*, *A. muricataspeak*, and *Eugenia stipitata*, speak to the ecological effects of the large mammalian extinctions (Janzen and Martin, 1982). Genetically isolated refugial populations often hybridize when brought into contact, expressing heterosis and in some cases polyploidy (Faurby and Svenning, 2015). There were rapid changes in allele frequencies in these populations once humans dispersed them across the melted glacial barriers (or between oases, islands, or isolated forest patches) during the mid-Holocene. Domestication occurs quicker on mosaic or fragmentary landscapes with lower levels of gene flow, following the basic principles of Island Biogeography (Rindos, 1984).

The idea that domestication in certain species arose from hybridization of glacial refugial or isolated forest populations has been suggested for several crops, including: apples (*Malus domestica/pumila*) (Cornille et al., 2014); bananas (polyploidy) (Heslop-Harrison and Schwarzacher, 2007); dates (*Phoenix dactylifera* × *P. theophrasti*; Flowers et al., 2019), (*P. dactylifera* ×

P. canariensis; (González-Pérez et al., 2004); cannabis (*Cannabis sativa*) (Clarke and Merlin, 2013; Small, 2015); citrus (Moore, 2001; Wu et al., 2014); cucurbits (Kistler et al., 2015); walnuts (*Juglans regia*) (Pollegioni et al., 2015); and wheat (polyploidy) (Marcussen et al., 2014). Hybridization can also introgress favorable traits into crops, such as non-shattering into *indica* rice (Gross and Zhao, 2014) or frost tolerance into Himalayan barley (Zeng et al., 2018), allowing for the expansion of crops into regions with novel ecological constraints (Pfennig et al., 2016; Janzen et al., 2019). Scholars have long recognized the important role of hybridization in plant evolution and domestication more specifically (Anderson, 1949; Stebbins, 1950; Anderson and Stebbins, 1954; Ellstrand et al., 1996; Arnold, 1997; Rieseberg and Carney, 1998; Hughes et al., 2007). The prominence of interspecific hybridization in plants often drives adaptive evolution, a process well-documented in early agricultural fields (Purugganan and Fuller, 2009; Hancock, 2012; Arnold, 2014). Perennials are disproportionately more likely to double their genome upon hybridization, due to the increased number of replication errors possible throughout one individual's lifetime (Friedman, 2020). Miller and Gross (2011) note that among fruit trees, most crops have multiple domestication events, often through hybridization, and in many cases resulting in multiple domesticated species. They single out the genera of *Annona*, *Artocarpus*, *Citrus*, *Diospyros*, and *Prunus*, as having multiple domesticated species and hybrid complexes. The switch from unisex to bisexual flowers is possibly linked to limited mate selection and self-incompatibility, especially on clonally propagated orchards (Miller and Gross, 2011). Ultimately, the isolated nature of many of the crop-progenitor populations facilitated hybridization and the expression of heterosis or polyploidy once humans began dispersing them.

ANTHROPOGENIC ECOSYSTEM SERVICES

Megafaunal extinctions caused range reductions, genetic heterogeneity, and extinction of many crop progenitors. The loss of ecosystem services performed by megafaunal grazers and browsers set the stage for plant domestication in the early and mid-Holocene. It is also interesting to note that many domesticated animals represent megafauna that were approaching extinction when they developed mutualistic relationships with humans. Likewise, many field weeds and anthropophilic plants have traits that make them prone to disturbed environments, which include areas around human settlement and heavily grazed fields with large herd animals (Hawkes, 1969). The earliest traits of domestication in many crops are associated with endozoochoric dispersal (Janzen, 1984; Kuznar, 1993; Purugganan and Fuller, 2009; Spengler and Mueller, 2019). Ultimately, humans have served as the greatest "rewilding" force for the megafaunal anachronisms, simultaneously: (1) dispersing seeds of plants that had lost part or all of their ability for dispersal; (2) clearing and disturbing soils creating conditions favored by species that evolved on

heavy herbivory landscapes; (3) maintaining fruiting tree species; and (4) causing hybridization between previously isolated plant lineages. When considering plant an animal domestication, it is essential to explore the evolutionary legacy that led up to the mutualistic relationships that humans engage in today.

The reframing of plant domestication as a set of traits evolving to support an mutualistic relationship is a growing perspective in the literature. As scholars, reframe the discussion, they can stop searching for rational drivers of human decision making, such as climate change or population pressure, and instead focus on the ecology and evolutionary history of the plants involved. Future discussions of these dispersal processes in the progenitors of modern crops would better be served by a more detailed understanding of the megafaunal mammals that existed in the regions where plants evolved domestication traits. Likewise, a clearer understanding of the digestion and preferences in food choices among extant representatives of these mammalian lineages would help pinpoint specific species or guilds linked in coevolutionary relationships with the respective plant clades. New applications of aDNA can better link ancient dispersers and their plants, as well as identifying the population legacies of disperser extinctions (e.g., Kistler et al., 2015). Future studies of plant domestication would greatly benefit from collaborations between ecologists, paleontologists, and archaeologists. Traditionally, these are disciplines that have not mingled and we would argue that the lack of crosspollination of ideas has hampered the development of domestication research. The traits of domestication seen in the archaeological record parallel evolutionary processes in the wild; therefore, a better understanding of the archaeological record could come from discussions with evolutionary ecologists. Likewise, many traits in plants that favored farming originally evolved on now extinct landscapes; therefore, the missing variables for understanding early plant domestication may come from restored megafaunal landscapes, such as the bison preserves of the American Midwest, Białowiecki National Park in Poland, or Pleistocene Park in Sakha Republic, Russia (e.g., Mueller et al., 2020).

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS

All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

FUNDING

Research funds and support were provided by the Max Planck Institute for the Science of Human History and the European Research Council, grant number 851102, Fruits of Eurasia: Domestication and Dispersal (FEDD).

REFERENCES

- Adams, C. D., and Baker, H. G. (1962). "Weeds of cultivation and grazing lands," in *Agriculture and Land Use in Ghana*, ed G. B. Wills (London: Oxford University Press), 402–415.
- Anderson, E. (1949). *Introgressive Hybridization*. New York, NY: John Wiley.
- Anderson, E. (1952). *Plants, Man and Life*. Berkeley, CA: University of California Press.
- Anderson, E., and Stebbins, G. L. (1954). Hybridization as an evolutionary stimulus. *Evolution* 8, 378–388. doi: 10.1111/j.1558-5646.1954.tb01504.x
- Arnold, M. L. (1997). *Natural Hybridization and Evolution*. Oxford: Oxford University Press.
- Arnold, M. L. (2014). Natural hybridization and the evolution of domesticated, pest and disease organisms. *Mol. Ecol.* 13, 997–1007. doi: 10.1111/j.1365-294X.2004.02145.x
- Baker, H. G. (1965). "Characteristics and modes of origin of weeds," in *The Genetics of Colonizing Species*, eds H. G. Baker and G. L. Stebbins (New York, NY: Academic Press), 147–172.
- Baker, H. G. (1972). Human influences on plant evolution. *Econ. Bot.* 26, 32–43. doi: 10.1007/BF02862259
- Baker, H. G. (1974). The evolution of weeds. *Annu. Rev. Ecol. Syst.* 5, 1–24. doi: 10.1146/annurev.es.05.110174.000245
- Bakker, E. S., Gill, J. L., Johnson, C. N., Vera, F. W., Sandom, C. J., Asner, G. P., et al. (2016). Combining paleo-data and modern enclosure experiments to assess the impact of megafauna extinctions on woody vegetation. *PNAS* 113, 847–855. doi: 10.1073/pnas.1502545112
- Baskin, J. M., and Baskin, C. C. (2004). A classification system for seed dormancy. *Seed Sci. Res.* 14, 1–16. doi: 10.1079/SSR2003150
- Baucom, R. S., and Holt, J. S. (2009). Weeds of agricultural importance: bridging the gap between evolutionary ecology and crop and weed science. *New Phytol.* 184, 741–743. doi: 10.1111/j.1469-8137.2009.03077.x
- Beaune, D., Fruth, B., Bollache, L., Hohmann, G., and Bretagnolle, F. (2013). Doom of the elephant-dependent trees in a Congo tropical forest. *For. Ecol. Manag.* 295, 109–117. doi: 10.1016/j.foreco.2012.12.041
- Brodie, J. F., Helmy, O. E., Brockelman, W. Y., and Maron, J. L. (2009). Functional differences within a guild of tropical mammalian frugivores. *Ecology* 90, 688–698. doi: 10.1890/08-0111.1
- Burney, D. A., Burney, L. P., Godfrey, L. R., Jungers, W. L., Goodman, S. M., Wright, H. T., et al. (2004). A chronology for late prehistoric Madagascar. *J. Hum. Evol.* 47, 25–63. doi: 10.1016/j.jhevol.2004.05.005
- Burney, D. A., Robinson, G. S., and Burney, L. P. (2003). Sporormiella and the late Holocene extinctions in Madagascar. *PNAS* 100, 10800–10805. doi: 10.1073/pnas.1534700100
- Campos-Arceiz, A., and Blake, S. (2011). Megagardeners of the forest—the role of elephants in seed dispersal. *Acta Oecol.* 37, 542–553. doi: 10.1016/j.actao.2011.01.014
- Campos-Arceiz, A., Larrinaga, A. R., Weerasinghe, U. R., Takatsuki, S., Pastorini, J., Leimgruber, P., et al. (2008). Behavior rather than diet mediates seasonal differences in seed dispersal by Asian elephants. *Ecology* 89, 2684–2691. doi: 10.1890/07-1573.1
- Clark, J. S. (1998). Why trees migrate so fast: confronting theory with dispersal biology and the paleorecord. *Am. Nat.* 152, 204–224. doi: 10.1086/286162
- Clarke, R. C., and Merlin, M. D. (2013). *Cannabis: Evolution and Ethnobotany*. Berkeley, CA: University of California Press.
- Clement, C. R., de Cristo-Araújo, M., Coppens d'Eeckenbrugge, G., de Reis, V. M., Lehnbech, R., and Picanço-Rodrigues, D. (2017). Origin and dispersal of domesticated peach palm. *Front. Ecol. Evol.* 5:148. doi: 10.3389/fevo.2017.00148
- Clements, D. R., DiTommaso, A., Jordan, N., Booth, B. D., Cardina, J., Doohan, D., et al. (2004). Adaptability of plants invading North American cropland. *Agric. Ecosyst. Environ.* 104, 379–398. doi: 10.1016/j.agee.2004.03.003
- Codding, B. F., Bird, R. B., Kauhane, P. G., and Bird, D. W. (2014). Conservation or co-evolution? Intermediate levels of aboriginal burning and hunting have positive effects on kangaroo populations in Western Australia. *Hum. Ecol.* 42, 659–669. doi: 10.1007/s10745-014-9682-4
- Cornille, A., Giraud, T., Smulders, M. J., Roldán-Ruiz, I., and Gladieux, P. (2014). The domestication and evolutionary ecology of apples. *Trends Genet.* 30, 57–65. doi: 10.1016/j.tig.2013.10.002
- Council, N. R. (2002). *Ecological Dynamics on Yellowstone's Northern Range*. Washington, DC: National Academies Press.
- Crowley, B. E., Godfrey, L. R., and Irwin, M. T. (2011). A glance to the past: subfossils, stable isotopes, seed dispersal, and lemur species loss in southern Madagascar. *Am. J. Primatol.* 73, 25–37. doi: 10.1002/ajp.20817
- Curtu, A. L., Gailing, O., and Finkeldey, R. (2007). Evidence for hybridization and introgression within a species-rich oak (*Quercus* spp.) community. *BMC Evol. Biol.* 7, 1–15. doi: 10.1186/1471-2148-7-218
- Donlan, J. (2005). Re-wilding North America. *Nature* 436, 913–914. doi: 10.1038/436913a
- Doughty, C. E., Wolf, A., and Field, C. B. (2010). Biophysical feedbacks between the Pleistocene megafauna extinction and climate: The first human-induced global warming? *Geophys. Res. Lett.* 37, 1–5. doi: 10.1029/2010GL043985
- Doughty, C. E., Wolf, A., and Malhi, Y. (2013). The legacy of the Pleistocene megafauna extinctions on nutrient availability in Amazonia. *Nat. Geosci.* 6, 761–764. doi: 10.1038/ngeo1895
- Ellstrand, N. C. (2014). Is gene flow the most important evolutionary force in plants? *Am. J. Bot.* 101, 737–753. doi: 10.3732/ajb.1400024
- Ellstrand, N. C., Whitkus, R., and Rieseberg, L. H. (1996). Distribution of spontaneous plant hybrids. *PNAS* 93, 5090–5093. doi: 10.1073/pnas.93.10.5090
- Eriksson, O. (2008). Evolution of seed size and biotic seed dispersal in angiosperms: paleoecological and neoecological evidence. *Int. J. Plant Sci.* 169, 863–870. doi: 10.1086/589888
- Escribano-Avila, G., Calviño-Cancela, M., Pias, B., Virgos, E., Valladares, F., and Escudero, A. (2014). Diverse guilds provide complementary dispersal services in a woodland expansion process after land abandonment. *J. Appl. Ecol.* 51, 1701–1711. doi: 10.1111/1365-2664.12340
- Estes, J. A., Heithaus, M., McCauley, D. J., Rasher, D. B., and Worm, B. (2016). Megafaunal impacts on structure and function of ocean ecosystems. *Annu. Rev. Environ. Resour.* 41, 83–116. doi: 10.1146/annurev-environ-110615-085622
- Estes, J. A., Terborgh, J., Brashares, J. S., Power, M. E., Berger, J., Bond, W. J., et al. (2011). Trophic downgrading of planet Earth. *Science* 333, 301–306. doi: 10.1126/science.1205106
- Falcón, W., Moll, D., and Hansen, D. M. (2020). Frugivory and seed dispersal by chelonians: a review and synthesis. *Biol. Rev.* 95, 142–166. doi: 10.1111/brv.12558
- Faurby, S., and Svenning, J. C. (2015). Historic and prehistoric human-driven extinctions have reshaped global mammal diversity patterns. *Divers. Distrib.* 21, 1155–1166. doi: 10.1111/ddi.12369
- Federman, S., Dornburg, A., Daly, D. C., Downie, A., Perry, G. H., Yoder, A. D., et al. (2016). Implications of Lemuriform extinctions for the Malagasy flora. *PNAS* 113, 5041–5046. doi: 10.1073/pnas.1523825113
- Feranec, R. S., Miller, N. G., Lothrop, J. C., and Graham, R. W. (2011). The Sporormiella proxy and end-Pleistocene megafaunal extinction: a perspective. *Quat. Int.* 245, 333–338. doi: 10.1016/j.quaint.2011.06.004
- Ferretti, M. P. (2008). Miocene proboscideans from Italy: African elements and palaeogeographic implications. *Geol. East Libya* 3, 325–334.
- Flowers, J. M., Hazzouri, K. M., Gros-Balthazard, M., Mo, Z., Koutroumpa, K., Perrakis, A., et al. (2019). Cross-species hybridization and the origin of North African date palms. *PNAS* 116, 1651–1658. doi: 10.1073/pnas.1817453116
- Frank, D. A., McNaughton, S. J., and Tracy, B. F. (1998). The ecology of the earth's grazing ecosystems. *Bioscience* 48, 513–521. doi: 10.2307/1313313
- Friedman, J. (2020). The evolution of annual and perennial plant life histories: Ecological correlates and genetic mechanisms. *Annu. Rev. Ecol. Evol. Syst.* 51, 461–481. doi: 10.1146/annurev-ecolsys-110218-024638
- Friis, E., and Crepet, W. (1987). "Time of appearance of floral features," in *The Origins of Angiosperms and Their Biological Consequences*, eds E. M. Friis, W. G. Chaloner, and P. R. Crane (Cambridge, MA: Cambridge University Press), 259–304.
- Fritz, G. J. (1999). Gender and the early cultivation of gourds in eastern North America. *Am. Antiq.* 64, 417–430. doi: 10.2307/2694142
- Froyd, C. A., Coffey, E. E., van der Knaap, W. O., van Leeuwen, J. F., Tye, A., and Willis, K. J. (2014). The ecological consequences of megafaunal loss: giant tortoises and wetland biodiversity. *Ecol. Lett.* 17, 144–154. doi: 10.1111/ele.12203
- Fuller, D. Q. (2018). Long and attenuated: comparative trends in the domestication of tree fruits. *Veg. Hist. Archaeobot.* 27, 165–176. doi: 10.1007/s00334-017-0659-2

- Fuller, D. Q., and Allaby, R. (2009). "Seed dispersal and crop domestication: shattering, germination and seasonality in evolution under cultivation," in *Annual Plant Reviews*, ed L. Østergaard (Oxford: Wiley-Blackwell), 238–295. doi: 10.1002/9781444314557.ch7
- Fuller, D. Q., Denham, T., Arroyo-Kalin, M., Lucas, L., Stevens, C. J., Qin, L., et al. (2014). Convergent evolution and parallelism in plant domestication revealed by an expanding archaeological record. *PNAS* 111, 6147–6152. doi: 10.1073/pnas.1308937110
- Fuzessy, L. F., Cornelissen, T. G., Janson, C., and Silveira, F. A. (2016). How do primates affect seed germination? A meta-analysis of gut passage effects on neotropical plants. *Oikos* 125, 1069–1080. doi: 10.1111/oik.02986
- Galetti, M., Guevara, R., Córtes, M. C., Fadini, R., Von Matter, S., Leite, A. B., et al. (2013). Functional extinction of birds drives rapid evolutionary changes in seed size. *Science* 340, 1086–1090. doi: 10.1126/science.1233774
- Galetti, M., Moleón, M., Jordano, P., Pires, M. M., Guimaraes, P. R. Jr., Pape, T., et al. (2017). Ecological and evolutionary legacy of megafauna extinctions. *Biol. Rev.* 93, 845–862. doi: 10.1111/brv.12374
- Garant, D., Forde, S. E., and Hendry, A. P. (2007). The multifarious effects of dispersal and gene flow on contemporary adaptation. *Functional Ecol.* 21, 434–443. doi: 10.1111/j.1365-2435.2006.01228.x
- Geiger, M., Sánchez-Villagra, M. R., and Lindholm, A. K. (2018). A longitudinal study of phenotypic changes in early domestication of house mice. *R. Soc. Open Sci.* 5:172099. doi: 10.1098/rsos.172099
- Geremia, C., Merkle, J. A., Eacker, D. R., Wallen, R. L., White, P., Hebblewhite, M., et al. (2019). Migrating bison engineer the green wave. *PNAS* 116, 25707–25713. doi: 10.1073/pnas.1913783116
- Gill, J. L. (2014). Ecological impacts of the late Quaternary megaherbivore extinctions. *New Phytol.* 201, 1163–1169. doi: 10.1111/nph.12576
- Gill, J. L., McLauchlan, K. K., Skibbe, A. M., Goring, S., Zirbel, C. R., and Williams, J. W. (2013). Linking abundances of the dung fungus *Sporormiella* to the density of bison: implications for assessing grazing by megaherbivores in palaeorecords. *J. Ecol.* 101, 1125–1136. doi: 10.1111/1365-2745.12130
- Gill, J. L., Williams, J. W., Jackson, S. T., Lininger, K. B., and Robinson, G. S. (2009). Pleistocene megafaunal collapse, novel plant communities, and enhanced fire regimes in North America. *Science* 326, 1100–1103. doi: 10.1126/science.1179504
- Godfrey, L. R., Jungers, W. L., Schwartz, G. T., and Irwin, M. T. (2008). "Ghosts and orphans," in *Elwyn Simons: A search for Origins*, eds J. G. Fleagle and C. C. Gilbert (New York, NY: Springer), 361–395. doi: 10.1007/978-0-387-73896-3_24
- Godwin, H. (1960). "The history of weeds in Britain," in *The Biology of Weeds*, ed J. L. Harper (Oxford: Blackwell), 1–10.
- González-Pérez, M., Caujapé-Castells, J., and Sosa, P. (2004). Molecular evidence of hybridisation between the endemic *Phoenix canariensis* and the widespread *P. dactylifera* with Random Amplified Polymorphic DNA (RAPD) markers. *Plant Syst. Evol.* 247, 165–175. doi: 10.1007/s00606-004-0166-7
- Griffiths, C. J., Hansen, D. M., Jones, C. G., Züel, N., and Harris, S. (2011). Resurrecting extinct interactions with extant substitutes. *Curr. Biol.* 21, 762–765. doi: 10.1016/j.cub.2011.03.042
- Gross, B. L., and Zhao, Z. (2014). Archaeological and genetic insights into the origins of domesticated rice. *PNAS* 111, 6190–6197. doi: 10.1073/pnas.1308942110
- Guimarães, P. R. Jr., Galetti, M., and Jordano, P. (2008). Seed dispersal anachronisms: rethinking the fruits extinct megafauna ate. *PLoS ONE* 3:e1745. doi: 10.1371/journal.pone.0001745
- Hamrick, J. L., Godt, M. J. W., and Sherman-Broyles, S. L. (1992). Factors influencing levels of genetic diversity in woody plant species. *New Forest.* 6, 95–124. doi: 10.1007/978-94-011-2815-5_7
- Hancock, J. F. (2012). *Plant Evolution and the Origin of Crop Species*. Wallingford, CT: CABI. doi: 10.1079/9781845938017.0000
- Harich, F. K., Treydte, A. C., Ogutu, J. O., Roberts, J. E., Savini, C., Bauer, J. M., et al. (2016). Seed dispersal potential of Asian elephants. *Acta Oecol.* 77, 144–151. doi: 10.1016/j.actao.2016.10.005
- Hawkes, J. G. (1969). "The ecological background of plant domestication," in *The Domestication and Exploitation of Plants and Animals*, eds P. Ucko and G. W. Dimbleby (Chicago, IL: Aldine), 17–29.
- Heslop-Harrison, J. S., and Schwarzhacher, T. (2007). Domestication, genomics and the future for banana. *Ann. Bot.* 100, 1073–1084. doi: 10.1093/aob/mcm191
- Howe, H. F., and Miriti, M. N. (2000). No question: seed dispersal matters. *Trends Ecol. Evol.* 15, 434–436. doi: 10.1016/S0169-5347(00)01965-0
- Hughes, C. E., Govindarajulu, R., Robertson, A., Filer, D. L., Harris, S. A., and Bailey, C. D. (2007). Serendipitous backyard hybridization and the origin of crops. *PNAS* 104, 14389–14394. doi: 10.1073/pnas.0702193104
- Jacobs, B. F., Kingston, J. D., and Jacobs, L. L. (1999). The origin of grass-dominated ecosystems. *Ann. Missouri Bot. Gard.* 86, 590–643. doi: 10.2307/2666186
- Jansen, P. A., Hirsch, B. T., Emsens, W.-J., Zamora-Gutierrez, V., Wikelski, M., and Kays, R. (2012). Thieving rodents as substitute dispersers of megafaunal seeds. *PNAS* 109, 12610–12615. doi: 10.1073/pnas.1205184109
- Janzen, D. H. (1984). Dispersal of small seeds by big herbivores: foliage is the fruit. *Am. Nat.* 123, 338–353. doi: 10.1086/284208
- Janzen, D. H., and Martin, P. S. (1982). Neotropical anachronisms: the fruits the gomphotheres ate. *Science* 215, 19–27. doi: 10.1126/science.215.4528.19
- Janzen, G. M., Wang, L., and Hufford, M. B. (2019). The extent of adaptive wild introgression in crops. *New Phytol.* 221, 1279–1288. doi: 10.1111/nph.15457
- Jara-Guerrero, A., Escribano-Avila, G., Espinosa, C. I., De la Cruz, M., and Méndez, M. (2018). White-tailed deer as the last megafauna dispersing seeds in neotropical dry forests: the role of fruit and seed traits. *Biotropica* 50, 169–177. doi: 10.1111/btp.12507
- Jaroszewicz, B. (2013). Endozoochory by European bison influences the build-up of the soil seed bank in subcontinental coniferous forest. *Eur. J. Forest Res.* 132, 445–452. doi: 10.1007/s10342-013-0683-4
- Johnson, C. N. (2009). Ecological consequences of Late Quaternary extinctions of megafauna. *P. Roy. Soc. B Biol. Sci.* 276, 2509–2519. doi: 10.1098/rspb.2008.1921
- Jones, G., Kluyver, T., Preece, C., Swarbrick, J., Forster, E., Wallace, M., et al. (2021). The origins of agriculture: Intentions and consequences. *J. Archaeol. Sci.* 125:105290. doi: 10.1016/j.jas.2020.105290
- Kaiser, T. M., Müller, D. W., Fortelius, M., Schulz, E., Codron, D., and Clauss, M. (2013). Hypsodonty and tooth facet development in relation to diet and habitat in herbivorous ungulates: implications for understanding tooth wear. *Mammal Rev.* 43, 34–46. doi: 10.1111/j.1365-2907.2011.00203.x
- Kaplan, J. O., Pfeiffer, M., Kolen, J. C., and Davis, B. A. (2016). Large scale anthropogenic reduction of forest cover in Last Glacial Maximum Europe. *PLoS ONE* 11:e0166726. doi: 10.1371/journal.pone.0166726
- Kennett, D. J., and Winterhalder, B. (2006). *Behavioral Ecology and the Transition to Agriculture*. Berkeley, CA: University of California Press.
- Kistler, L., Newsom, L. A., Ryan, T. M., Clarke, A. C., Smith, B. D., and Perry, G. H. (2015). Gourds and squashes (*Cucurbita* spp.) adapted to megafaunal extinction and ecological anachronism through domestication. *PNAS* 112, 15107–15112. doi: 10.1073/pnas.1516109112
- Kitamura, S., Yumoto, T., Poonswad, P., and Wohandee, P. (2007). Frugivory and seed dispersal by Asian elephants, *Elephas maximus*, in a moist evergreen forest of Thailand. *J. Trop. Ecol.* 23, 373–376. doi: 10.1017/S0266467407004026
- Knapp, A. K., Blair, J. M., Briggs, J. M., Collins, S. L., Hartnett, D. C., Johnson, L. C., et al. (1999). The keystone role of bison in North American tallgrass prairie: bison increase habitat heterogeneity and alter a broad array of plant, community, and ecosystem processes. *Bioscience* 49, 39–50. doi: 10.2307/1313492
- Kostina, K. (1971). "Genera armeniaca, persica," in *Flora of the USSR, Vol. X.*, ed V. L. Kamorov (Jerusalem: Israel Program of Scientific Translations), 434–448.
- Kuznar, L. A. (1993). Mutualism between *Chenopodium*, herd animals, and herders in the south Central Andes. *Mountain Res. Dev.* 3, 257–265. doi: 10.2307/3673655
- Ladizinsky, G. (1979). Seed dispersal in relation to the domestication of Middle East legumes. *Econ. Bot.* 33, 284–289. doi: 10.1007/BF02858256
- Langlie, B. S., Mueller, N. G., Spengler, R. N., and Fritz, G. J. (2014). Agricultural origins from the ground up: archaeological approaches to plant domestication. *Am. J. Bot.* 101, 1601–1617. doi: 10.3732/ajb.1400145
- Larson, G., Piperno, D. R., Allaby, R. G., Purugganan, M. D., Andersson, L., Arroyo-Kalin, M., et al. (2014). Current perspectives and the future of domestication studies. *PNAS* 111, 6139–6146. doi: 10.1073/pnas.1323964111
- Li, L.-F., and Olsen, K. (2016). To have and to hold: selection for seed and fruit retention during crop domestication. *Curr. Top. Dev. Biol.* 119, 63–109. doi: 10.1016/b978-0-12-081202-0.0002
- Lim, J. Y., Svenning, J.-C., Göddel, B., Faurby, S., and Kissling, W. D. (2020). Frugivore-fruit size relationships between palms and mammals

- reveal past and future defaunation impacts. *Nat. Commun.* 11:4904. doi: 10.1038/s41467-020-18530-5
- Lu, L., and Bartholomew, B. (2003). "Amygdalus L," in *Flora of China*, eds Z. Y. Wu and P. H. Raven (St Louis: Missouri Botanical Garden Press), 391–395.
- Lundgren, E. J., Ramp, D., Ripple, W. J., and Wallach, A. D. (2018). Introduced megafauna are rewilding the Anthropocene. *Ecography* 41, 857–866. doi: 10.1111/ecog.03430
- Malhi, Y., Doughty, C. E., Galetti, M., Smith, F. A., Svenning, J.-C., and Terborgh, J. W. (2016). Megafauna and ecosystem function from the pleistocene to the anthropocene. *PNAS* 113, 838–846. doi: 10.1073/pnas.1502540113
- Marcussen, T., Sandve, S. R., Heier, L., Spannagl, M., Pfeifer, M., Jakobsen, K. S., et al. (2014). Ancient hybridizations among the ancestral genomes of bread wheat. *Science* 345:1250092. doi: 10.1126/science.1250092
- Mayr, E. (1963). *Animal Species and Evolution*. Cambridge, MA: Harvard University Press.
- McConkey, K. R., Nathalang, A., Brockelman, W. Y., Saralamba, C., Santon, J., Matmoon, U., et al. (2018). Different megafauna vary in their seed dispersal effectiveness of the megafaunal fruit *Platymitra macrocarpa* (Annonaceae). *PLoS ONE* 13:e0198960. doi: 10.1371/journal.pone.0198960
- Meredith, R. W., Janečka, J. E., Gatesy, J., Ryder, O. A., Fisher, C. A., Teeling, E. C., et al. (2011). Impacts of the cretaceous terrestrial revolution and KPg extinction on mammal diversification. *Science* 334, 521–524. doi: 10.1126/science.1211028
- Miller, A. J., and Gross, B. L. (2011). From forest to field: perennial fruit crop domestication. *Am. J. Bot.* 98, 1389–1414. doi: 10.3732/ajb.1000522
- Moore, G. A. (2001). Oranges and lemons: clues to the taxonomy of Citrus from molecular markers. *Trends Genet.* 17, 536–540. doi: 10.1016/S0168-9525(01)02442-8
- Mueller, N. G., Spengler, R. N., Glenn, A., and Lama, K. (2020). Bison, anthropogenic fire, and the origins of agriculture in eastern North America. *Anthropocene Review*. doi: 10.1177/2053019620961119
- Murphy, D. J. (2007). *People, Plants and Genes: The Story of Crops and Humanity*. Oxford: Oxford University Press.
- Murphy, S. (2018). Seed dispersal in osage orange (*Maclura pomifera*) by squirrels (*Sciurus* spp.). *Am. Midl. Nat.* 180, 312–317. doi: 10.1674/0003-0031-180.2.312
- Nathan, R., and Muller-Landau, H. C. (2000). Spatial patterns of seed dispersal, their determinants and consequences for recruitment. *Trends Ecol. Evol.* 15, 278–285. doi: 10.1016/S0169-5347(00)01874-7
- Nee, M. (1990). The domestication of Cucurbita (Cucurbitaceae). *Econ. Bot.* 44:56. doi: 10.1007/BF02860475
- Neve, P., Vila-Aiub, M., and Roux, F. (2009). Evolutionary-thinking in agricultural weed management. *New Phytol.* 184, 783–793. doi: 10.1111/j.1469-8137.2009.03034.x
- O'Farrill, G., Galetti, M., and Campos-Arceiz, A. (2013). Frugivory and seed dispersal by tapirs: an insight on their ecological role. *Integr. Zool.* 8, 4–17. doi: 10.1111/j.1749-4877.2012.00316.x
- Onstein, R. E., Baker, W. J., Couvreur, T. L., Faurby, S., Herrera-Alsina, L., Svenning, J.-C., et al. (2018). To adapt or go extinct? The fate of megafaunal palm fruits under past global change. *P. Roy. Soc. B Biol. Sci.* 285:20180882. doi: 10.1098/rspb.2018.0882
- Palumbi, S. R. (2001). Humans as the world's greatest evolutionary force. *Science* 293, 1786–1790. doi: 10.1126/science.293.5536.1786
- Peres, C. A., and Palacios, E. (2007). Basin-wide effects of game harvest on vertebrate population densities in Amazonian forests: implications for animal-mediated seed dispersal. *Biotropica* 39, 304–315. doi: 10.1111/j.1744-7429.2007.00272.x
- Peres, C. A., and Van Roosmalen, M. (2002). "Primate frugivory in two species-rich neotropical forests: implications for the demography of large-seeded plants in overhunted areas," in *Seed Dispersal and Frugivory: Ecology, Evolution and Conservation*, eds D. J. Levey, W. Silva, and M. Galetti (New York, NY: CABI), 407–422. doi: 10.1079/9780851995250.0407
- Perrier, X., De Langhe, E., Donohue, M., Lentfer, C., Vrydaghs, L., Bakry, F., et al. (2011). Multidisciplinary perspectives on banana (*Musa* spp.) domestication. *PNAS* 108, 11311–11318. doi: 10.1073/pnas.1102001108
- Petit, R. J., Bodénès, C., Ducouso, A., Roussel, G., and Kremer, A. (2003). Hybridization as a mechanism of invasion in oaks. *New Phytol.* 161, 151–164. doi: 10.1046/j.1469-8137.2003.00944.x
- Pfennig, K. S., Kelly, A. L., and Pierce, A. A. (2016). Hybridization as a facilitator of species range expansion. *P. Roy. Soc. B Biol. Sci.* 283:20161329. doi: 10.1098/rspb.2016.1329
- Pires, M. M., Galetti, M., Donatti, C. I., Pizo, M. A., Dirzo, R., and Guimarães, P. R. (2014). Reconstructing past ecological networks: the reconfiguration of seed-dispersal interactions after megafaunal extinction. *Oecologia* 175, 1247–1256. doi: 10.1007/s00442-014-2971-1
- Pires, M. M., Guimarães, P. R., Galetti, M., and Jordano, P. (2018). Pleistocene megafaunal extinctions and the functional loss of long-distance seed-dispersal services. *Ecography* 41, 153–163. doi: 10.1111/ecog.03163
- Pollegioni, P., Woeste, K. E., Chiochini, F., Del Lungo, S., Olimpieri, I., Tortolano, V., et al. (2015). Ancient humans influenced the current spatial genetic structure of common Walnut populations in Asia. *PLoS ONE* 10:e0135980. doi: 10.1371/journal.pone.0135980
- Purugganan, M. D. (2019). Evolutionary insights into the nature of plant domestication. *Curr. Biol.* 29, R705–R714. doi: 10.1016/j.cub.2019.05.053
- Purugganan, M. D., and Fuller, D. Q. (2009). The nature of selection during plant domestication. *Nature* 457, 843–848. doi: 10.1038/nature07895
- Raboin, L.-M., Carreel, F., Noyer, J.-L., Baurens, F.-C., Horry, J.-P., Bakry, F., et al. (2005). Diploid ancestors of triploid export banana cultivars: molecular identification of 2n restitution gamete donors and n gamete donors. *Mol. Breed.* 16, 333–341. doi: 10.1007/s11032-005-2452-7
- Rapanarivo, S. H. J. V. (1999). *Pachypodium (Apocynaceae): Taxonomy, Habitats and Cultivation*. Rotterdam: Brookfield.
- Rieseberg, L., and Carney, S. (1998). Plant hybridization. *New Phytol.* 140, 599–624. doi: 10.1046/j.1469-8137.1998.00315.x
- Rindos, D. (1984). *The Origins of Agriculture. An Evolutionary Perspective*. London: Academic Press.
- Robertson, A., Trass, A., Ladley, J., and Kelly, D. (2006). Assessing the benefits of frugivory for seed germination: the importance of the deinhibition effect. *Funct. Ecol.* 20, 58–66. doi: 10.1111/j.1365-2435.2005.01057.x
- Robinson, G. S., Pigott Burney, L., and Burney, D. A. (2005). Landscape paleoecology and megafaunal extinction in southeastern New York State. *Ecol. Monogr.* 75, 295–315. doi: 10.1890/03-4064
- Rule, S., Brook, B. W., Haberle, S. G., Turney, C. S., Kershaw, A. P., and Johnson, C. N. (2012). The aftermath of megafaunal extinction: ecosystem transformation in Pleistocene Australia. *Science* 335, 1483–1486. doi: 10.1126/science.1214261
- Ruxton, G. D., and Schaefer, H. M. (2012). The conservation physiology of seed dispersal. *Philos. Trans. R. Soc. B.* 367, 1708–1718. doi: 10.1098/rstb.2012.0001
- Salisbury, E. J. (1942). *The Reproductive Capacity of Plants. Studies in Quantitative Biology*. London: Bell.
- Samuels, I., and Levey, D. (2005). Effects of gut passage on seed germination: do experiments answer the questions they ask? *Funct. Ecol.* 19, 365–368. doi: 10.1111/j.1365-2435.2005.00973.x
- Santamaría, L., Rodríguez-Pérez, J., Larrinaga, A. R., and Pias, B. (2007). Predicting spatial patterns of plant recruitment using animal-displacement kernels. *PLoS ONE* 2:e1008. doi: 10.1371/journal.pone.0001008
- Sauer, C. (1952). *Seeds, Spades, Hearths and Herds: The Domestication of Animals and Foodstuffs*. Dordrecht: MIT Press.
- Schupp, E. W., Jordano, P., and Gómez, J. M. (2010). Seed dispersal effectiveness revisited: a conceptual review. *New Phytol.* 188, 333–353. doi: 10.1111/j.1469-8137.2010.03402.x
- Sekar, N., Lee, C.-L., and Sukumar, R. (2015). In the elephant's seed shadow: the prospects of domestic bovids as replacement dispersers of three tropical Asian trees. *Ecology* 96, 2093–2105. doi: 10.1890/14-1543.1
- Sekar, N., and Sukumar, R. (2013). Waiting for Gajah: an elephant mutualist's contingency plan for an endangered megafaunal disperser. *J. Ecol.* 101, 1379–1388. doi: 10.1111/1365-2745.12157
- Small, E. (2015). Evolution and Classification of *Cannabis sativa* (marijuana, hemp) in relation to human utilization. *Bot. Rev.* 81, 189–294. doi: 10.1007/s12229-015-9157-3
- Smith, F. A., Elliott, S. M., and Lyons, S. K. (2010). Methane emissions from extinct megafauna. *Nat. Geosci.* 3, 374–375. doi: 10.1038/ngeo877
- Spengler, R., and Mueller, N. (2019). The unexplored role of endozoochoric seed dispersal in the domestication of grain crops. *Nat. Plants* 5, 656–662. doi: 10.1038/s41477-019-0470-4

- Spengler, R. N. (2019). Origins of the Apple: the role of megafaunal mutualism in the domestication of malus and rosaceous trees. *Front. Plant Sci.* 10:617. doi: 10.3389/fpls.2019.00617
- Spengler, R. N. (2020). Anthropogenic seed dispersal: rethinking the origins of plant domestication. *Trends Plant Sci.* 25, 340–348. doi: 10.1016/j.tplants.2020.01.005
- Sridhara, S., McConkey, K., Prasad, S., and Corlett, R. T. (2016). “Frugivory and seed dispersal by large herbivores of Asia,” in *The Ecology of Large Herbivores in South and Southeast Asia*, eds F. S. Ahrestani and M. Sankaran (Dordrecht: Springer), 121–150. doi: 10.1007/978-94-017-7570-0_5
- Stebbins, G. L. (1981). Coevolution of grasses and herbivores. *Ann. Missouri Botanical Garden* 68, 75–86. doi: 10.2307/2398811
- Stebbins, J. L. (1950). *Variation and Evolution in Plants*. New York, NY: Columbia University Press. doi: 10.7312/steb94536
- Stewart, C. N., Tranel, P. J., Horvath, D. P., Anderson, J. V., Rieseberg, L. H., Westwood, J. H., et al. (2009). Evolution of weediness and invasiveness: charting the course for weed genomics. *Weed Sci.* 57, 451–462. doi: 10.1614/WS-09-011.1
- Stiner, M. C. (2001). Thirty years on the “Broad Spectrum Revolution” and paleolithic demography. *PNAS* 98, 6993–6996. doi: 10.1073/pnas.121176198
- Stoner, K. E., Riba-Hernández, P., Vulinec, K., and Lambert, J. E. (2007). The role of mammals in creating and modifying seedshadows in tropical forests and some possible consequences of their elimination. *Biotropica* 39, 316–327. doi: 10.1111/j.1744-7429.2007.00292.x
- Su, T., Wilf, P., Huang, Y., Zhang, S., and Zhou, Z. (2015). Peaches preceded humans: fossil evidence from SW China. *Sci. Rep.* 5, 1–7. doi: 10.1038/srep16794
- Tarsizis, E., Tomlinson, S., Harrison, M. E., Morrogh-Bernard, H. C., and Munn, A. J. (2018). Gardeners of the forest: effects of seed handling and ingestion by orangutans on germination success of peat forest plants. *Biol. J. Linn. Soc.* 123, 125–134. doi: 10.1093/biolinnean/blx133
- Terborgh, J., Nuñez-Iturri, G., Pitman, N. C., Valverde, F. H. C., Alvarez, P., Swamy, V., et al. (2008). Tree recruitment in an empty forest. *Ecology* 89, 1757–1768. doi: 10.1890/07-0479.1
- Thompson, J. N. (1999). The evolution of species interactions. *Science* 284, 2116–2118. doi: 10.1126/science.284.5423.2116
- Tiffney, B. H. (1981). “Diversity and major events in the evolution of land plants,” in *Paleobotany, Paleocology, and Evolution*, eds K. J. Niklas (New York, NY: Praeger), 193–230.
- Tiffney, B. H. (1984). Seed size, dispersal syndromes, and the rise of the angiosperms: evidence and hypothesis. *Ann. Missouri Bot. Gard.* 71, 551–576. doi: 10.2307/2399037
- Tiffney, B. H. (2004). Vertebrate dispersal of seed plants through time. *Annu. Rev. Ecol. Syst.* 35, 1–29. doi: 10.1146/annurev.ecolsys.34.011802.132535
- Traveset, A. (1998). Effect of seed passage through vertebrate frugivores’ guts on germination: a review. *Perspect. Plant Ecol. Evol. Syst.* 1, 151–190. doi: 10.1078/1433-8319-00057
- Traveset, A., Robertson, A., and Rodríguez-Pérez, J. (2007). A review on the role of endozoochory in seed germination,” in *Seed Dispersal: Theory and Its Application in a Changing World*, eds A. J. Dennis, R. J. Green, and E. W. Schupp (Wallingford, CT: CABI), 78–103. doi: 10.1079/978184593165.0.0078
- Traveset, A., and Verdú, M. (2002). “A meta-analysis of the effect of gut treatment on seed germination,” in *Seed Dispersal and Frugivory: Ecology, Evolution and Conservation*, eds D. J. Levey and M. Galetti (Wallingford, CT: CABI), 339–350. doi: 10.1079/9780851995250.0339
- Ude, G., Pillay, M., Nwakanma, D., and Tenkouano, A. (2002). Analysis of genetic diversity and sectional relationships in *Musa* using AFLP markers. *Theor. Appl. Genet.* 104, 1239–1245. doi: 10.1007/s00122-001-0802-3
- Valido, A., and Olesen, J. M. (2007). “The importance of lizards as frugivores and seed dispersers,” in *Seed Dispersal: Theory and Its Application in a Changing World*, eds A. J. Dennis, R. J. Green, and E. W. Schupp (Wallingford, CT: CABI), 124–147. doi: 10.1079/9781845931650.0124
- Van Der Kaars, S., Miller, G. H., Turney, C. S., Cook, E. J., Nürnberg, D., Schönfeld, J., et al. (2017). Humans rather than climate the primary cause of Pleistocene megafaunal extinction in Australia. *Nat. Commun.* 8, 1–7. doi: 10.1038/ncomms14142
- van Zonneveld, M., Larranaga, N., Blonder, B., Coradin, L., Hormaza, J. I., and Hunter, D. (2018). Human diets drive range expansion of megafauna-dispersed fruit species. in *Proceedings of the National Academy of Sciences*, 3326–3331. doi: 10.1073/pnas.1718045115
- Vavra, M., Parks, C. G., and Wisdom, M. J. (2007). Biodiversity, exotic plant species, and herbivory: the good, the bad, and the ungulate. *For. Ecol. Manage.* 246, 66–72. doi: 10.1016/j.foreco.2007.03.051
- Vicentini, A., Barber, J. C., Aliscioni, S. S., Giussani, L. M., and Kellogg, E. A. (2008). The age of the grasses and clusters of origins of C4 photosynthesis. *Glob. Chang. Biol.* 14, 2963–2977. doi: 10.1111/j.1365-2486.2008.01688.x
- Vigne, J.-D. (2015). Early domestication and farming: what should we know or do for a better understanding? *Anthropozoologica* 50, 123–150. doi: 10.5252/az2015n2a5
- Vigueira, C., Olsen, K., and Caicedo, A. (2013). The red queen in the corn: agricultural weeds as models of rapid adaptive evolution. *Heredity* 110, 303–311. doi: 10.1038/hdy.2012.104
- Voigt, F. A., Arafeh, R., Farwig, N., Griebeler, E. M., and Böhning-Gaese, K. (2009). Linking seed dispersal and genetic structure of trees: a biogeographical approach. *J. Biogeogr.* 36, 242–254. doi: 10.1111/j.1365-2699.2008.02002.x
- Wang, B. C., and Smith, T. B. (2002). Closing the seed dispersal loop. *Trends Ecol. Evol.* 17, 379–386. doi: 10.1016/S0169-5347(02)02541-7
- Ward, M., Dick, C. W., Gribel, R., and Lowe, A. J. (2005). To self, or not to self... a review of outcrossing and pollen-mediated gene flow in neotropical trees. *Heredity* 95, 246–254. doi: 10.1038/sj.hdy.6800712
- Weisskopf, A., and Fuller, D. Q. (2014). “Peach: origins and development,” in *Encyclopedia of Global Archaeology*, ed C. Smith (New York, NY: Springer), 5840–5842. doi: 10.1007/978-1-4419-0465-2_2311
- Wing, S., and Tiffney, B. (1987). “Interactions of angiosperms and herbivorous tetrapods through time,” in *The Origins of Angiosperms and Their Biological Consequences*, eds E. M. Friis, W. G. Chaloner, and P. R. Crane (Cambridge, MA: Cambridge University Press), 203–224.
- Wotton, D. M., and Kelly, D. (2011). Frugivore loss limits recruitment of large-seeded trees. *P. Roy. Soc. B Biol. Sci.* 278, 3345–3354. doi: 10.1098/rspb.2011.0185
- Wotton, D. M., and Kelly, D. (2012). Do larger frugivores move seeds further? Body size, seed dispersal distance, and a case study of a large, sedentary pigeon. *J. Biogeogr.* 39, 1973–1983. doi: 10.1111/jbi.12000
- Wu, G. A., Prochnik, S., Jenkins, J., Salse, J., Hellsten, U., Murat, F., et al. (2014). Sequencing of diverse mandarin, pummelo and orange genomes reveals complex history of admixture during citrus domestication. *Nat. Biotechnol.* 32, 656–662. doi: 10.1038/nbt.2906
- Yu, Y., Fu, J., Xu, Y., Zhang, J., Ren, F., Zhao, H., et al. (2018). Genome re-sequencing reveals the evolutionary history of peach fruit edibility. *Nat. Commun.* 9, 1–13. doi: 10.1038/s41467-018-07744-3
- Zeng, X., Guo, Y., Xu, Q., Mascher, M., Guo, G., Li, S., et al. (2018). Origin and evolution of qingke barley in Tibet. *Nat. Commun.* 9, 1–11. doi: 10.1038/s41467-018-07920-5
- Zohary, D., and Spiegel-Roy, P. (1975). Beginnings of fruit growing in the old world. *Science* 187, 319–327. doi: 10.1126/science.187.4174.319

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2021 Spengler, Petraglia, Roberts, Ashastina, Kistler, Mueller and Boivin. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.