

This is a repository copy of *Still armed after domestication? Impacts of domestication and agronomic selection on silicon defences in cereals*.

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/129528/>

Version: Published Version

Article:

Simpson, Kimberley J., Wade, Ruth Nicola, Rees, Mark et al. (2 more authors) (2017) Still armed after domestication? Impacts of domestication and agronomic selection on silicon defences in cereals. *Functional Ecology*. pp. 2108-2117. ISSN 0269-8463

<https://doi.org/10.1111/1365-2435.12935>

Reuse

This article is distributed under the terms of the Creative Commons Attribution (CC BY) licence. This licence allows you to distribute, remix, tweak, and build upon the work, even commercially, as long as you credit the authors for the original work. More information and the full terms of the licence here:

<https://creativecommons.org/licenses/>

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.

Still armed after domestication? Impacts of domestication and agronomic selection on silicon defences in cereals

Kimberley J. Simpson¹ | Ruth N. Wade²  | Mark Rees¹ | Colin P. Osborne¹ | Sue E. Hartley² 

¹Department of Animal and Plant Sciences, University of Sheffield, Sheffield, UK

²Department of Biology, University of York, York, UK

Correspondence

Sue E. Hartley
Email: sue.hartley@york.ac.uk

Funding information

Natural Environment Research Council,
Grant/Award Number: NE/H022716/1 and
NE/F003137/1

Handling Editor: Sergio Rasman

Abstract

1. Plant phenotypes reflect trade-offs between competing resource-intensive physiological processes. A shift in resource allocation, away from anti-herbivore defences and towards growth and reproduction, is predicted through plant domestication, such that crops are faster growing and higher yielding than their wild ancestors. These changes are hypothesized to have come at the cost of defence investment, leaving crops “disarmed by domestication”. Silicon is the principal anti-herbivore defence in grasses, including many of our most important staple cereal crops, but the impact of domestication on silicon-based defences is unknown.
2. We measured the effects of both domestication and modern agronomic selection on growth rate and a suite of anti-herbivore defences, specifically leaf toughness, silicon and phenolic concentrations. Our comparison of wild, landrace and modern cultivated cereals spanned multiple cereal species, including wheat, barley and maize, sampling eight independent domestication events and five examples of modern agronomic selection.
3. Leaf silicon concentration showed a small, but significant, 10% reduction through domestication, but there was no effect of modern agronomic selection, and phenolic concentration was not affected by either factor. Silicon concentration correlated positively with leaf tensile strength, but negatively with foliar phenolic concentrations, suggesting a trade-off between chemical and physical defences. Size-standardized growth rate was independent of domestication status, and did not trade-off with silicon or phenolic defences. However, modelling showed that relative growth rate slowed more with increasing size in plants with higher silicon levels, so that they reached a smaller final size, implying a cost of silicon-based defence. We found the opposite pattern for phenolic-based defence, with increasing phenolic concentrations associated with a greater plant size at maturity, and faster maximum relative growth rates.
4. Silicon-based defences have been reduced in cereals through domestication, consistent with our predicted costs of these defences to growth. However, modern

Paper previously published as Standard Paper.

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

© 2017 The Authors. *Functional Ecology* published by John Wiley & Sons Ltd on behalf of British Ecological Society.

agronomic selection has not influenced silicon defences in cereal crops and the small decrease in silicon concentration associated with domestication is unlikely to have a major effect on the ability of cereals to withstand a range of abiotic and biotic stresses. These findings have broad implications for crop protection and our understanding of plant trade-offs.

KEYWORDS

cereals, crop protection, growth-defence trade-offs, herbivory, landraces, modern cultivars, phenolics, silica, wild ancestors

1 | INTRODUCTION

Trade-offs between competing, energetically demanding traits are fundamental in shaping an organism's phenotype. The substantial phenotypic changes associated with plant domestication (Hammer, 1984; Harlan, 1971; Meyer, Duval, & Jensen, 2012) are thought to have resulted, in part, from changes in resource allocation between such traits, rather than an increase in productivity (Evans, 1993). Strong directional selection by humans for traits of agronomic importance (Gepts, 2004), such as enhanced seed production and growth rate, is predicted to have driven a re-partitioning of resources in domesticated landraces compared with their wild relatives. Modern crop cultivars, which have been shaped by recent agronomic selection and breeding (Lindig-Cisneros, Dirzo, & Espinosa-Garcia, 2002), are expected to be the product of an even greater reallocation of resources than domestication alone.

Life history theory predicts that a consequence of such a shift in resources through domestication and agronomic selection may be a reduction in other traits not of agronomic interest to humans (Evans, 1993; Milla, Osborne, Turcotte, & Violle, 2015; Rosenthal & Dirzo, 1997). In particular, plant anti-herbivore defences may be metabolically costly and thus trade-off with traits such as growth (Coley, Bryant, & Chapin, 1985; Herms & Mattson, 1992). Evidence from between-species comparisons shows that growth-defence trade-offs are common and widespread in natural ecosystems (Kneitel & Chase, 2004; Lind et al., 2013), although they are not universally demonstrated (e.g. Siemens, Garner, Mitchell-Olds, & Callaway, 2002).

In crop systems, a strong directional selection for faster growth is predicted through domestication, so that a reduction in resources available for defence investment is expected to have occurred concurrently (Massei & Hartley, 2000). In general, domestication has led to a reduction in defences in many crops (Bellota, Medina, & Bernal, 2013; Chen, Gols, & Benrey, 2015; Massei & Hartley, 2000; Mayrose, Kane, Mayrose, Dlugosch, & Rieseberg, 2011; Meyer et al., 2012; Mondolot et al., 2008; Rodriguez-Saona et al., 2011; Rosenthal & Dirzo, 1997). However, the increase in growth rate through domestication and agronomic selection assumed to be driving this reduction has not been found in several studies (Cook & Evans, 1983; Evans, 1993; Preece et al., 2017; Welter, 2000), suggesting that changes in defence investment through domestication may not result from a trade-off with

growth rate. Additionally, some studies have not found a defence reduction through domestication (Turcotte, Turley, & Johnson, 2014; Whitehead, Turcotte, & Poveda, 2017), suggesting that the phenomenon of plants being "disarmed by domestication" may not be as widespread as first thought.

Cereals are grass species grown for their edible seed, and were among the first plants to be domesticated, around 12,000 years ago (Zohary & Hopf, 2000). There are c. 30 cereal species (Bouchenak-Khelladi et al., 2008), which are grown world-wide at a huge scale, with an estimated production of over 2,500 million tonnes for 2014/2015 (FAO, 2015). Three of these species (maize, rice and wheat) alone represent the world's most important staple crops, together providing 60% of human food energy intake (FAO, 1995). Through the domestication process, cereals developed a shared suite of traits that are strikingly different to their wild relatives (Harlan, Wet, & Price, 1973). Traits associated with harvesting (non-shattering of mature seeds), seed production (increased seed size and yield), regeneration (loss of seed dormancy), development (reduced branching and synchronous maturation) and growth (increased seedling size and vigour) were selected for, consciously or unconsciously, through domestication.

The impacts of these major phenotypic changes on cereal resistance to herbivory have been explored in several studies (Bellota et al., 2013; Maag et al., 2015; Rosenthal & Dirzo, 1997; Turcotte et al., 2014). Cereal anti-herbivore defences consist of chemical defences, including the presence of distasteful phenolic compounds in leaves and stems, and physical defences, such as tough leaves (Vicari & Bazely, 1993). Domestication appears to have had mixed effects on these defences. For example, a reduction in leaf toughness through maize domestication was associated with lower resistance to a specialist herbivore (Bellota et al., 2013). However, in a comparison of 29 crops, including seven cereals, Turcotte et al. (2014) found that most domestication events did not cause differences in leaf phenolic concentrations or resistance to two common herbivores. However, none of these studies investigated the effect of domestication on the principal defence of grasses—silicon.

A widespread characteristic of grasses is tissue silicification, in which silicon is taken up in unusually large amounts from the soil and deposited in the leaves, in the form of spines and as sharp granules called phytoliths (Hartley, Fitt, McLarnon, & Wade, 2015; Hodson, White, Mead, & Broadley, 2005). Plant silicon has multiple functional

roles in plants (Cooke, DeGabriel, & Hartley, 2016), including resistance to abiotic stresses, such as drought and salinity, and to biotic stresses such as pests and diseases (Cooke & Leishman, 2011; Ma & Yamaji, 2008). In addition, silicon may have important structural roles in plants, as a compression-resistant stiffening material (Schoelynck et al., 2010; Strömberg, Di Stilio, & Song, 2016). One key function of silicon is as the main physical anti-herbivore defence in grasses (Cooke & Leishman, 2011; Hartley & DeGabriel, 2016). By reducing digestibility and the absorption of essential nutrients, leaf silicon can detrimentally affect vertebrate and invertebrate herbivore performance (Massey & Hartley, 2006, 2009; Reynolds, Keeping, & Meyer, 2009; Wieczorek, Zub, Szafranska, Ksiazek, & Konarzewski, 2015). Phytoliths also make leaves highly abrasive, which wears down the mouthparts of herbivores (Massey & Hartley, 2009). In addition to being a constitutive defence, silicon-based defences can also be induced, with leaf silicon levels being raised substantially in response to herbivore damage (Hartley et al., 2015; Massey, Ennos, & Hartley, 2007a; Wieczorek et al., 2015). This reactive increase in silicon levels implies active control over silicon uptake and/or allocation within the leaf, and hence a potential fitness cost of this defence. Although, the relative contributions of active and passive processes to silicon uptake remain uncertain and are likely to vary between species (Faisal, Callis, Slot, & Kitajima, 2012; Hartley, 2015; Kumar, Milstein, Bami, Elbaum, & Elbaum, 2016; Quigley & Anderson, 2014), the existence of an active proton pump required for efflux of silicon into the xylem in many crop species suggests an energetic cost to silicon uptake (Ma & Yamaji, 2015; Ma et al., 2007). However, costs associated with silicon-based defences have not been conclusively demonstrated, and whether these defences trade-off with growth or other physiological process is currently unknown.

This study aimed to determine how anti-herbivore defences were modified through cereal domestication and agronomic selection. Through comparisons of cereal landraces with their wild progenitors and with modern cultivars, we established the effects of domestication and agronomic selection on silicon- and phenolic-based constitutive defences. To determine whether any changes in leaf silicon or phenolic concentrations arose from resource reallocation, plant growth rate was measured as a performance trait that is expected to trade-off against defence. We predicted that wild progenitors would be the best defended and slowest growing, modern cultivars would be

the fastest growing and poorest defended, and landraces would have intermediate trait values.

2 | MATERIALS AND METHODS

2.1 | Species selection and plant growth conditions

We chose 21 cereal species, representing eight independent domestication events, from multiple centres of domestication, and five periods of modern agronomic selection. Of these, eight species are landraces, and eight their wild progenitors (Table 1). Modern cultivars were chosen for five major crops (excluding einkorn, *Triticum monococcum*, and the two millets, *Pennisetum glaucum* and *Setaria italica*). Seeds for all species were obtained from a variety of germplasm holdings (see Table S1).

Seeds germinated in Petri dishes within a growth chamber set to 25°C and a 16-hr photoperiod. After 17 seeds of a species had germinated, the seedlings were transplanted into 1-litre pots filled with M3 compost (Levington Horticulture Ltd., Ipswich, UK), supplemented with fertilizer (Scotts M3-4 Osmocote Exact Standard fertilizer, Maryville, OH, USA), so that nutrient supply was not limiting to growth. Plants were grown in a randomized block design within a controlled environment chamber (Conviron BDW 40; Winnipeg, Canada). Conditions were: 16/8 hr day/night cycle, 25/16°C (day/night temperature), relative humidity of 80%, and photosynthetic photon flux density (PPFD) of 500 $\mu\text{mol m}^{-2} \text{s}^{-1}$. Plants were top watered to saturation three times a week with deionised water and the blocks were re-randomized twice weekly.

2.2 | Growth rate analysis

To examine the effect of domestication on plant growth, we determined the relative growth rate for each species when plants were at a common mass (standardized growth rate, or SGR, Rees et al., 2010). Two randomly selected plants of each species were destructively harvested at six time points over a 30-day period. At each harvest, plants were removed from pots, washed clean of growth medium, and dried at 50°C for a week, before weighing the total biomass (both above- and below-ground) using a four-point balance (PA413/1, Ohaus, NJ, USA).

TABLE 1 Cereal wild progenitor and landrace comparisons

Landrace	Wild progenitor	Centre of domestication	Reference
<i>Hordeum vulgare</i> (Barley)	<i>Hordeum spontaneum</i>	Southwest Asia	Zohary and Hopf (2000)
<i>Oryza sativa</i> subsp. <i>indica</i> (Asian rice)	<i>Oryza rufipogon</i>	South China	Cheng et al. (2003)
<i>Pennisetum glaucum</i> (Pearl millet)	<i>Pennisetum violaceum</i>	Africa (sub-Saharan)	De Wet (1995)
<i>Setaria italica</i> (Foxmail millet)	<i>Setaria viridis</i>	North China	Doust and Kellogg (2002)
<i>Sorghum bicolor</i> (Sorghum)	<i>Sorghum arundinaceum</i>	Africa (sub-Saharan)	Aldrich and Doebley (1992)
<i>Triticum monococcum</i> (Einkorn)	<i>Triticum boeoticum</i>	Fertile Crescent	Feldman, Lupton, and Miller (1995)
<i>Triticum dicoccon</i> (Emmer wheat)	<i>Triticum dicoccoides</i>	Fertile Crescent	Feldman et al. (1995)
<i>Zea mays</i> (Maize)	<i>Zea mays</i> subsp. <i>parviglumis</i>	Meso-America	Doebley (2004)

Growth curves were fitted to log-transformed total dry mass data over time for each species in R (R Core Development Team, 2013) using nonlinear mixed effects models (NLME package; Pinheiro, Bates, Debroy, & Sarkar, 2016). The four-parameter logistic growth function was used with a self-starting routine. Species-specific, independent random effects were fitted for maximum plant mass (m_{\max}), minimum plant mass (m_{\min}) and the time point when plant mass was midway between these (x_{mid}). These species-specific growth curves were then used to estimate SGR (see Rees et al. 2010 for further details).

2.3 | Plant defence measurements

We examined whether domestication and modern agronomic selection had consistently altered leaf phenolic and silicon concentrations in cereals, and how the latter trait relates to another physical defence, leaf tensile strength. Seven plants of each species were harvested 30 days after transplanting (five plants in addition to the two plants from the final harvest of the growth rate analysis). Leaf tensile strength was tested on freshly harvested leaf material following the methods of Cornelissen et al. (2003). The force required to tear two leaf sections of known width was determined for each plant, and an average value calculated. All leaf material was then oven dried at 60°C for 48 hr, before being ground into a fine powder.

The Folin-Ciocalteu method was used to determine total foliar phenolic concentration (Kerslake, Woodin, & Hartley, 1998). A sample (c. 10 mg) from each individual was extracted in methanol at 80°C for 30 min, before being centrifuged. 100 μ l of the supernatant was added to 0.25 ml Folin-Ciocalteu reagent and 1 ml saturated sodium carbonate. After 1 hr, the absorbance of each sample was measured on a spectrophotometer at a wavelength of 760 nm. Absorbance values were converted to tannic acid equivalents using a tannic acid standard curve (0–2 mg tannic acid), and results were expressed in % per dry leaf mass.

For measurements of leaf silicon concentration, two oven-dried, ground samples (>0.1 g) of leaf material from each individual were pressed into 13 mm-diameter pellets. Following the methods of Reidinger, Ramsey, and Hartley (2012), foliar silicon concentration (% dry mass) was determined using a Niton XL3t XRF analyzer (Thermo Fisher Scientific, Inc., MA, USA), for a measurement time of 30 s. The machine was calibrated using a standard curve of methyl cellulose, validated with certified plant reference material of known silicon concentration (Garbuzov, Reidinger, & Hartley, 2011).

2.4 | Statistical analysis

The effects of domestication and agronomic selection on SGR, leaf silicon concentration, phenolic concentration and tensile strength were determined by fitting linear mixed-effects models to the trait data ("LME4" package; Bates, Maechler, Bolker, & Walker, 2014). The fixed effect was "domestication status", and "comparison" was added as a random effect to account for the phylogenetic non-independence of species within each comparison. "Species" was an additional random effect in

the silicon, phenolic and tensile strength models to account for multiple measurements per species. The effects that domestication status had on SGR, leaf silicon concentration, phenolic concentration and tensile strength were determined through model comparison (complete model vs. grand mean model) using a parametric bootstrapping method ("PBKRT-EST" package; Halekoh & Højsgaard, 2014) with 10,000 simulated generations. As there was a significant effect of domestication status on leaf silicon concentration, the same model comparison approach was used to determine where in the domestication/agronomic selection process this significant change occurred.

To determine whether silicon- and phenolic-based defences represent a metabolic cost to plants, two approaches were taken. Firstly, the relationships between growth and defence traits were investigated to see if they exhibited a trade-off. Linear mixed effects models were fitted to the species-level SGR and defence (leaf silicon and phenolic concentration) data, with "comparison" as the random effect. Model comparison was carried out as above. Secondly, the effect of varying leaf silicon and phenolic concentrations on relative growth rate (RGR) was modelled by allowing m_{\max} to be a linear function of silicon/phenolic concentration. Using this, the relationship between RGR and plant size was predicted for each species, based on the species-specific average silicon and phenolic concentrations.

To establish the relationships between the measured anti-herbivore defences across species, linear mixed effects models were fitted to the species-level leaf silicon concentration data, and both the leaf tensile strength and leaf phenolic concentration data in turn. Comparison was included as a random effect. To determine the relationships between silicon concentration and tensile strength/phenolic concentration within each species, linear models were fitted to these data for each species.

3 | RESULTS

Domestication status had a significant effect on leaf silicon concentration (Likelihood ratio test, LRT = 7.82, $df = 2$, $p = .02$), but not on leaf phenolic concentration (LRT = 0.90, $df = 2$, $p = .64$) or leaf tensile strength (LRT = 4.75, $df = 2$, $p = .09$). Cereal domestication caused an average 10% reduction in leaf silicon concentration (LRT = 4.78, $df = 1$, $p = .03$), but agronomic selection did not reduce leaf silicon significantly (LRT = 0.88, $df = 1$, $p = .34$; Figure 1). Species average leaf silicon concentration ranged between 0.5 and 1.2% of leaf dry mass (see Table S2), and was significantly positively correlated with leaf tensile strength between species (LRT = 16.49, $df = 1$, $p < .001$; Figure 2 and Table S2). Within species, the relationship between leaf silicon concentration and tensile strength was not significant for 19/21 species ($p > .05$ in all cases). In the two cases where the relationship was significant, it was positive in one species (*Sorghum bicolor*, slope = 0.692, $p = .007$) and negative in the other (*Triticum dicoccon*, slope = -0.253, $p = .01$). Species average leaf phenolic concentration ranged from 0.2% and 2.1% of leaf dry mass (Table S2), and was significantly negatively related to leaf silicon concentration across species (LRT = 4.49, $df = 1$, $p = .034$; Figure 2), but was only

significantly negatively correlated within one species (*Hordeum spontaneum*, slope = -0.693 , $p = .006$).

Domestication status had no effect on SGR (LRT = 2.80; $df = 2$; $p = .24$), with average values differing little between wild progenitors ($0.29 \text{ g g}^{-1} \text{ day}^{-1}$), landraces ($0.32 \text{ g g}^{-1} \text{ day}^{-1}$) and modern cultivars ($0.26 \text{ g g}^{-1} \text{ day}^{-1}$). Domestication status also had a marginally significant influence on the minimum mass (m_{\min}), consistent with the larger seed size of domesticated crops than their wild progenitors, but there was no effect on the other parameters used to fit the growth curves ($p > .05$; Figure S1).

We found no direct evidence for a trade-off between SGR and leaf silicon concentration (LRT = 0.64, $df = 1$, $p = .42$), or between SGR and leaf phenolic concentration (LRT = 2.74, $df = 1$, $p = .12$), when SGR was compared among species at a common size of 0.2 g. However, the modelled relationship between leaf silicon concentration and growth parameters throughout plant development provided evidence for a cost of silicon-based defences that emerged at larger plant sizes: increasing foliar silicon concentration was linked to a decrease in both the final plant size and the maximum relative growth rate ($p = .035$ for the fitted model, Figure 3). The species with the highest silicon concentration (*Triticum boeoticum*) was predicted to have 18% lower maximum relative growth rate (0.235 vs. $0.288 \text{ g g}^{-1} \text{ day}^{-1}$) and 15% lower final plant size (taken at RGR = 0; 7.47 g vs. 8.81 g), when compared to the species with the lowest silicon concentration (*Sorghum bicolor*).

In contrast, the modelled relationship between leaf phenolic concentration and growth parameters showed that increasing phenolic concentrations were associated with a greater plant size at maturity, and faster maximum relative growth rates ($p = .001$ for the fitted model, Figure 3). *Zea mays* subsp. *parviglumis*, the species with the highest average foliar phenolic concentration, was predicted to have a 25% greater final size (9.47 g vs. 7.15 g) and a 30% higher maximum relative growth rate (0.315 vs. $0.222 \text{ g g}^{-1} \text{ day}^{-1}$) than *Triticum aestivum* cv. "Hereward", the species with the lowest phenolic concentration.

4 | DISCUSSION

This study is the first to determine the impacts of domestication and modern agronomic selection on silicon-based anti-herbivore defences in cereals, which include some of our most economically important staple crops. While a handful of studies have explored the influence of domestication on other cereal defences (Bellota et al., 2013; Chen et al., 2015; Rosenthal & Dirzo, 1997; Turcotte et al., 2014), none have investigated silicon, the key anti-herbivore defence in grasses (Massey, Ennos, & Hartley, 2007b; Reynolds et al., 2009; Vicari & Bazely, 1993). The finding of a small but significant reduction in cereal silicon-based defence levels here suggests that this defence has been "disarmed" by domestication, but to only a limited extent (Massey & Hartley, 2000; Meyer et al., 2012), whilst modern agronomic selection had no detectable effect on cereal silicon-based defences.

A high level of accumulated silicon acts as an effective plant anti-herbivore defence. Multiple detrimental impacts on herbivore performance are well-documented (Hunt, Dean, Webster, Johnson, & Ennos, 2008; Ma & Takahashi, 2002; Massey & Hartley, 2009). In addition, we found that species with high leaf silicon concentrations also tended to have high tensile strength. Like silicon, this leaf trait is an important physical defence against herbivores, reducing their performance by increasing the metabolic costs of leaf digestion (Clissold, Sanson, Read, & Simpson, 2009; Roces & Lighton, 1995). Interestingly, within species there is little evidence of significant correlations between leaf silicon concentration and tensile strength, and neither was there evidence of changes in leaf tensile strength through domestication and agronomic selection as found elsewhere (Maag et al., 2015). The 10% reduction in silicon concentration through domestication we detected could imply that cereal landraces may be more susceptible to herbivory than their wild progenitors. However, whether such changes in silicon are sufficient to influence herbivore behaviour is unknown; studies demonstrating impacts on herbivore preference and performance have involved much larger changes in silicon concentration (e.g. Massey, Ennos, & Hartley, 2006; Massey et al., 2007a).

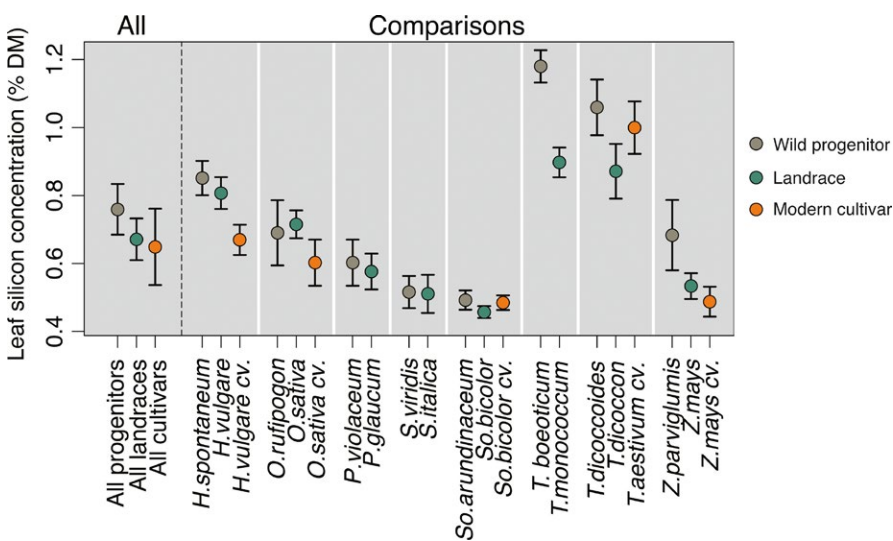


FIGURE 1 The effects of domestication and agronomic selection on cereal silicon-based defence. Points represent mean values of leaf silicon concentration for each species, and these are grouped within comparisons. Silicon concentration was significantly reduced through domestication ($p = .02$) but not modern agronomic selection ($p = .34$). Error bars = $\pm SE$. DM, dry mass; cv, cultivar

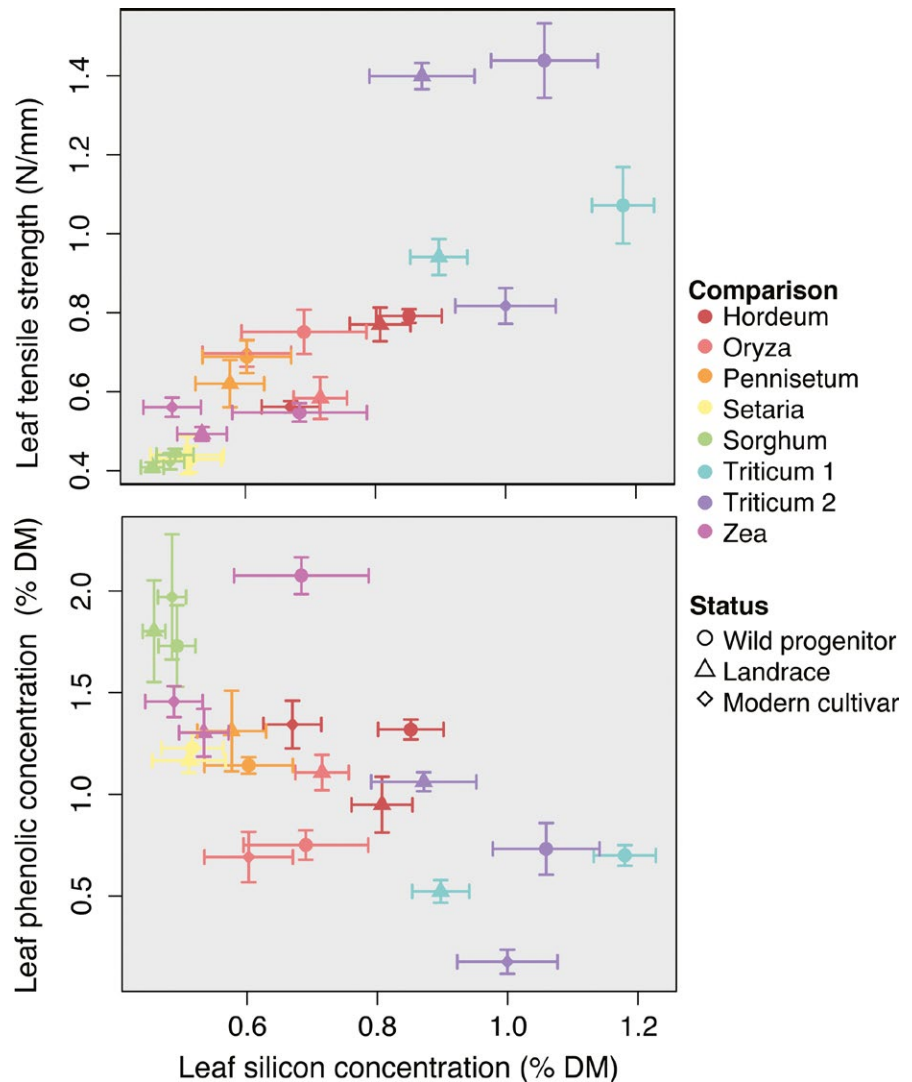


FIGURE 2 The relationships of leaf silicon concentration with leaf tensile strength (top) and leaf phenolic concentration (bottom). Values represent $M \pm SE$ bars. Across species, leaf silicon concentration correlated positively with leaf tensile strength ($p < .001$) but negatively with leaf phenolic concentration ($p = .034$). Triticum 1 is the comparison between Einkorn wheat (*Triticum monococcum*) and its progenitor (*Triticum boeoticum*); Triticum 2 is the comparison between Emmer wheat (*Triticum dicoccon*), its progenitor (*Triticum dicoccoides*) and modern cultivar (*Triticum aestivum*). DM, dry mass

The negative relationship between silicon- and phenolic-based defences is consistent with several studies that have found that silicon accumulation is associated with reduced concentrations of phenolic compounds (Cooke & Leishman, 2011; Frew, Powell, Sallam, Allsopp, & Johnson, 2016). A proposed explanation for this trade-off is that plants partly substitute carbon-based defences for silicon-based defences (Cooke & Leishman, 2011; Schaller, Brackhage, & Dudel, 2012). This may result in an increased availability of carbon for both defence and growth, which would be consistent with the predicted positive association here between phenolic concentration, plant growth rate and final size. We found no reduction in phenolic concentration through cereal domestication, which mirrors the findings of other studies investigating cereal defences (Maag et al., 2015; Turcotte et al., 2014), possibly because allocation to phenolics is known to be particularly variable in response to a range of environmental and other parameters (Jones & Hartley, 1999).

Agronomic selection has not significantly reduced constitutive chemical or physical defence levels in cereals, a result consistent with previous findings for maize (Rosenthal & Dirzo, 1997) and cranberry (Rodríguez-Saona et al., 2011). An insufficient change in selection

pressure between cereal landraces and modern cultivars may explain these non-significant results. Many modern cultivars are grown outside their natural range, so may experience reduced selection pressure for anti-herbivore defences, due to potential “escape” from herbivore attack (Keane & Crawley, 2002; Müller-Schärer, Schaffner, & Steinger, 2004; Schaffner et al., 2011). Alternatively, breeders may have selected for structural traits linked to high silicon concentrations through agronomic selection. Silicon-rich stiff straw and strong leaves may be associated with desirable properties such as resistance to lodging (Shimoyama, 1958) and enhanced photosynthesis (Okuda & Takahashi, 1962). Silicon offers multiple, often unrelated, benefits to plants simultaneously, so distinguishing between the adaptive significance of increased structural support and resistance against herbivores is challenging. Indeed, the result of a recent attempt to do this in wild grasses was inconclusive and suggested that both herbivore pressure and structural support could be important evolutionary drivers of silicon accumulation (Strömberg et al., 2016).

We found no consistent change in size-standardized growth rate through cereal domestication or agronomic selection. This adds to a growing body of studies that have failed to find growth rate alterations

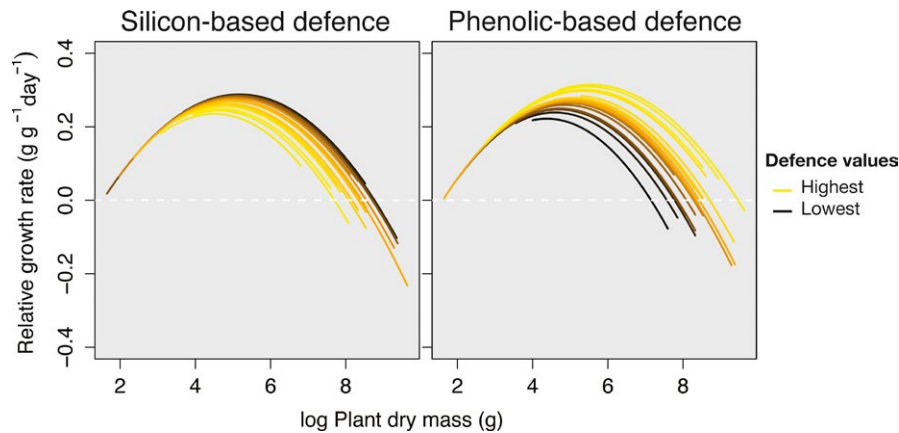


FIGURE 3 Silicon-based defence is linked to slower growth whereas phenolic-based defence is associated with faster growth. The modelled effect of varying levels of leaf silicon concentration (left) and phenolic concentration (right) on plant relative growth rate. Final plant size (m_{\max}) is modelled as a linear function of silicon/phenolic concentration. Each line represents the predicted relationship between relative growth rate and plant size for one species based on the species-specific average silicon and phenolic concentrations. The line colour reflects defence concentration values. Plant dry mass values were multiplied by 1,000 before log-transformation to ensure positive end values

through domestication (Cook & Evans, 1983; Preece et al., 2017; Welter, 2000). For example, Evans (1993) found no consistent trends in growth rate between wild and domesticated wheats, whilst Preece et al. (2017) attributed the greater yield in domesticated wheat and barley landraces to larger initial and final sizes, and changes in allocation, compared with their wild relatives. Some differences in traits relating to seed size and dormancy were consistent with the domestication syndrome (i.e. an increase in initial plant mass and shorter seedling emergence times); however, these traits are not directly related to growth rate. In our SGR analysis, we therefore find no direct evidence that the small reduction in silicon-based defence through domestication was caused by a reallocation of resources towards faster growth despite our modelled prediction that higher silicon concentrations are associated with lower growth rates and smaller final plant sizes. Through our modelling approach, we found that the adverse effect of allocation to silicon defences on growth rate increases with plant size, implying that the costs of silicon defences are relatively greater for larger plants. This could be due to the greater costs of uptake, mobilization and deposition of silicon in larger plants, or because plants with higher potential maximal growth rate and final size suffer most from the costs associated with silicon uptake. The predictions from our model may contrast with the results from the SGR analysis because we conducted this species comparison with plants of only 0.2 g, possibly too small to show any effects of silicon on growth rate. However, as the plants increase in size, the effect is predicted to become more pronounced.

The deposition of silicon in plant tissues is an energetically expensive process involving active efflux transporters (Ma et al., 2007; Ma & Yamaji, 2015). Our overall analysis is consistent with this: the species with the highest silicon concentration was modelled to be 15% smaller than the species with the lowest silicon concentration, and was predicted to grow more slowly, suggesting silicon uptake is costly. Silicon uptake in the roots occurs via a combination of passive uptake in the transpiration stream through aquaporin-type transporters and energy-demanding processes via proton pumps (Cooke & Leishman, 2011; Hartley, 2015; Kumar et al., 2016; Ma & Yamaji, 2006). The balance

between active and passive processes, which may depend upon transpiration rates, plant silicon demands and soil silicon availability (Faisal et al., 2012), could determine the impact of silicon uptake on plant growth rate. Despite such potential costs and associated impacts on growth, increased silicon concentration may also benefit plant growth under different abiotic and biotic stresses (Cooke & Leishman, 2011; Gong, Zhu, Chen, Wang, & Zhang, 2005; Guntzer, Keller, & Meunier, 2012; Ma, 2004), such that the benefits from silicon uptake must sometimes outweigh the negative impacts of actively transporting additional silicon.

Our finding that silicon, the principal defence in grasses, has been unchanged or reduced by only a small degree through domestication and modern agronomic selection, suggests that cereal crops have not been “disarmed” in this part of their armoury (Massei & Hartley, 2000). Silicon also has an important role in alleviating other biotic and abiotic plant stresses (Ma & Yamaji, 2006; Zhu & Gong, 2014); in grasses for example, silicon increases resistance to fungal and bacterial pathogens (Rodrigues, Polanco, Silveira Duarte, Resende, & Ribeiro Do Vale, 2015; Seebold, Kucharek, Datnoff, Correa-Victoria, & Marchetti, 2001), and to drought (Hattori et al., 2005). The small decrease in silicon concentration associated with domestication is unlikely to have a major effect on the ability of cereal plants to withstand a range of abiotic and biotic stresses. Modern agricultural systems are required to produce more food, cope with the impacts of climate change and reduce their use of synthetic chemicals with negative impacts on the environment and human health, so the need for sustainable means of pest control in these systems is particularly acute. Our findings suggest silicon is one such means which could be further exploited, particularly as we demonstrate modern crop varieties have retained their ability to deploy it.

ACKNOWLEDGEMENTS

We thank Georg Frenck, Catherine Preece, Rebecca Atkinson, Millie Mockford, Chris Bennett, Irene Johnson and Stefan Reidinger for their assistance in the lab. Seeds were kindly provided by a number of

germplasm resources. Funding was provided by Natural Environment Research Council grants awarded to CO (NE/H022716/1) and SH (NE/F003137/1).

AUTHORS' CONTRIBUTIONS

K.S., C.O. and S.H. conceived the ideas and designed methodology. K.S. collected the data. K.S., R.W., C.O., S.H. and M.R. analysed the data. K.S. and S.H. led the writing of the manuscript. All authors contributed critically to the drafts and gave final approval for publication.

DATA ACCESSIBILITY

Trait data: Species average values of leaf silicon concentration, phenolic concentration and tensile strength are uploaded as online supporting information (Table S2). Individual level data for defence and growth rate are deposited in the Dryad Digital Repository: <https://doi.org/10.5061/dryad.6nd96> (Simpson, Wade, Rees, Osborne, & Hartley, 2017).

REFERENCES

- Aldrich, P. R., & Doebley, J. (1992). Restriction fragment variation in the nuclear and chloroplast genomes of cultivated and wild *Sorghum bicolor*. *Theoretical & Applied Genetics*, *85*, 293–302.
- Bates, D., Maechler, M., Bolker, B., & Walker, S. (2014). *lme4: Linear mixed-effects models using Eigen and S4*. R package version 1.0–6.0. <http://lme4.r-forge.r-project.org>
- Bellota, E., Medina, R. F., & Bernal, J. S. (2013). Physical leaf defences – Altered by *Zea* life-history evolution, domestication, and breeding – Mediate oviposition preference of a specialist leafhopper. *Entomologia Experimentalis et Applicata*, *2*, 185–195.
- Bouchenak-Khelladi, Y., Salamin, N., Savolainen, V., Forest, F., Van Der Bank, M., Chase, M. W., & Hodgkinson, T. R. (2008). Large multi-gene phylogenetic trees of the grasses (Poaceae): Progress towards complete tribal and generic level sampling. *Molecular Phylogenetics & Evolution*, *47*, 488–505.
- Chen, Y. H., Gols, R., & Benrey, B. (2015). Crop domestication and its impact on naturally selected trophic interactions. *Annual Review of Entomology*, *60*, 35–58.
- Cheng, C. Y., Motohashi, R., Tsuchimoto, S., Fukuta, Y., Ohtsubo, H., & Ohtsubo, E. (2003). Polyphyletic origin of cultivated rice: Based on the interspersed pattern of SINEs. *Molecular Biology & Evolution*, *20*, 67–75.
- Clissold, F. J., Sanson, G. D., Read, J., & Simpson, S. J. (2009). Gross vs. net income: How plant toughness affects performance of an insect herbivore. *Ecology*, *90*, 3393–3405.
- Coley, P. D., Bryant, J. P., & Chapin, F. S. (1985). Resource availability and plant antiherbivore defence. *Science*, *230*, 895–899.
- Cook, M. G., & Evans, L. T. (1983). Some physiological-aspects of the domestication and improvement of rice (*Oryza* spp). *Field Crop Research*, *6*, 219–238.
- Cooke, J., DeGabriel, J. L., & Hartley, S. E. (Eds) (2016). Special feature: The functional role of silicon in plant biology. *Functional Ecology*, *30*, 1267–1474.
- Cooke, J., & Leishman, M. R. (2011). Is plant ecology more siliceous than we realise? *Trends in Plant Science*, *16*, 61–68.
- Cornelissen, J. H. C., Lavorel, S., Garnier, E., Diaz, S., Buchmann, N., Gurvich, D. E., ... Poorter, H. (2003). A handbook of protocols for standardised and easy measurement of plant functional traits worldwide. *Australian Journal of Botany*, *51*, 335–380.
- De Wet, J. M. J. (1995). *Pearl millet*. New York, NY: John Wiley & Sons.
- Doebley, J. (2004). The genetics of maize evolution. *Annual Review of Genetics*, *38*, 37–59.
- Doust, A. N., & Kellogg, E. A. (2002). Inflorescence diversification in the panicoid “bristle grass” clade (Paniceae, Poaceae): Evidence from molecular phylogenies and developmental morphology. *American Journal of Botany*, *89*, 1203–1222.
- Evans, L. T. (1993). *Crop evolution, adaptation, and yield*. Cambridge, UK: Cambridge University Press.
- Faisal, S., Callis, K. L., Slot, M., & Kitajima, K. (2012). Transpiration-dependent passive silica accumulation in cucumber (*Cucumis sativus*) under varying soil silicon availability. *Botany-Botanique*, *90*, 1058–1064.
- Feldman, M., Lupton, F. G. H., & Miller, T. E. (1995). Wheats. In J. Smartt, & N. W. Simmonds (Eds.), *Evolution of crop plants* (pp. 184–192). Harlow, UK: Longman Scientific and Technical.
- Food and Agriculture Organisation of the United Nations. (1995). *Dimensions of need: An atlas of food and agriculture*. Rome, Italy: FAO.
- Food and Agriculture Organisation of the United Nations. (2015). *FAO cereal supply and demand brief*. Rome, Italy: FAO.
- Frew, A., Powell, J. R., Sallam, N., Allsopp, P. G., & Johnson, S. N. (2016). Trade-offs between silicon and phenolic defenses may explain enhanced performance of root herbivores on phenolic-rich plants. *Journal of Chemical Ecology*, *42*, 768–771.
- Garbuzov, M., Reidinger, S., & Hartley, S. E. (2011). Interactive effects of plant-available soil silicon and herbivory on competition between two grass species. *Annals of Botany*, *108*, 1355–1363.
- Gepts, P. (2004). Crop domestication as a long-term selection experiment. J. Janick (Ed.), *Plant breeding reviews* (Vol. 24, pp. 1–44). New York, NY: John Wiley & Sons.
- Gong, H., Zhu, X., Chen, K., Wang, S., & Zhang, C. (2005). Silicon alleviates oxidative damage of wheat plants in pots under drought. *Plant Science*, *169*, 313–321.
- Guntzer, F., Keller, C., & Meunier, J. D. (2012). Benefits of plant silicon for crops: A review. *Agronomy for Sustainable Development*, *32*, 201–213.
- Halekoh, U., & Hojsgaard, S. (2014). A Kenward-Roger approximation and parametric bootstrap methods for tests in linear mixed models – The R package pbkrtest. *Journal of Statistical Software*, *59*, 1–30.
- Hammer, K. (1984). The domestication syndrome. *Kulturpflanze*, *32*, 11–34.
- Harlan, J. R. (1971). Agricultural origins – Centers and noncenters. *Science*, *174*, 468–474.
- Harlan, J. R., Wet, J., & Price, E. G. (1973). Comparative evolution of cereals. *Evolution*, *27*, 311–325.
- Hartley, S. E. (2015). Round and round in cycles? Silicon-based plant defences and vole population dynamics. *Functional Ecology*, *29*, 151–153.
- Hartley, S. E., & DeGabriel, J. L. (2016). Herbivore-induced silicon defences in grasses. *Functional Ecology*, *30*, 1311–1322.
- Hartley, S. E., Fitt, R. N., McLarnon, E. L., & Wade, R. N. (2015). Defending the leaf surface: Intra- and inter-specific differences in silicon deposition in grasses in response to damage and silicon supply. *Frontiers in Plant Science*, *6*, 35.
- Hattori, T., Inanaga, S., Araki, H., An, P., Morita, S., Luxova, M., & Lux, A. (2005). Application of silicon enhanced drought tolerance in *Sorghum bicolor*. *Physiologia Plantarum*, *123*, 459–466.
- Hermes, D. A., & Mattson, W. J. (1992). The dilemma of plants – To grow or defend. *Quarterly Review of Biology*, *67*, 283–335.
- Hodson, M. J., White, P. J., Mead, A., & Broadley, M. R. (2005). Phylogenetic variation in the silicon composition of plants. *Annals of Botany*, *96*, 1027–1046.
- Hunt, J. W., Dean, A. P., Webster, R. E., Johnson, G. N., & Ennos, A. R. (2008). A novel mechanism by which silica defends grasses against herbivory. *Annals of Botany*, *102*, 653–656.
- Jones, C. G., & Hartley, S. E. (1999). A protein competition model of phenolic allocation. *Oikos*, *86*, 27–44.

- Keane, R. M., & Crawley, M. J. (2002). Exotic plant invasions and the enemy release hypothesis. *Trends in Ecology & Evolution*, *17*, 164–170.
- Kerslake, J. E., Woodin, S. J., & Hartley, S. E. (1998). Effects of carbon dioxide and nitrogen enrichment on a plant–insect interaction: The quality of *Calluna vulgaris* as a host for *Operophtera brumata*. *New Phytologist*, *140*, 43–53.
- Kneitel, J. M., & Chase, J. M. (2004). Trade-offs in community ecology: Linking spatial scales and species coexistence. *Ecology Letters*, *7*, 69–80.
- Kumar, S., Milstein, Y., Bami, Y., Elbaum, M., & Elbaum, R. (2016). Mechanism of silica deposition in sorghum silica cells. *New Phytologist*, *213*, 791–798.
- Lind, E., Borer, E., Seabloom, E., Adler, P., Bakker, J. D., & Blumenthal, D. M. (2013). Life history constraints in grassland plant species: A growth–defence tradeoff is the norm. *Ecology Letters*, *16*, 513–521.
- Lindig-Cisneros, R., Dirzo, R., & Espinosa-García, F. J. (2002). Effects of domestication and agronomic selection on phytoalexin antifungal defence in Phaseolus beans. *Ecology Research*, *17*, 315–321.
- Ma, J. F. (2004). Role of silicon in enhancing the resistance of plants to biotic and abiotic stresses. *Soil Science & Plant Nutrition*, *50*, 11–18.
- Ma, J. F., & Takahashi, E. (2002). *Soil, fertilizer and plant silicon research in Japan*. Amsterdam, the Netherlands: Elsevier.
- Ma, J. F., & Yamaji, N. (2006). Silicon uptake and accumulation in higher plants. *Trends in Plant Science*, *11*, 392–397.
- Ma, J. F., & Yamaji, N. (2008). Functions and transport of silicon in plants. *Cellular & Molecular Life Sciences*, *65*, 3049–3057.
- Ma, J. F., & Yamaji, N. (2015). A cooperative system of silicon transport in plants. *Trends in Plant Science*, *20*, 435–442.
- Ma, J. F., Yamaji, N., Mitani, N., Tamai, K., Konishi, S., Fujiwara, T., ... Yano, M. (2007). An efflux transporter of silicon in rice. *Nature*, *448*, 209–212.
- Maag, D., Erb, M., Bernal, J. S., Wolfender, J. L., Turlings, T. C., & Glauser, G. (2015). Maize domestication and anti-herbivore defences: Leaf-specific dynamics during early ontogeny of maize and its wild ancestors. *PLoS ONE*, *10*, e0135722.
- Massei, G., & Hartley, S. E. (2000). Disarmed by domestication? Induced responses to browsing in wild and cultivated olive. *Oecologia*, *122*, 225–231.
- Massey, F. P., Ennos, A. R., & Hartley, S. E. (2006). Silica in grasses as a defence against insect herbivores: Contrasting effects on folivores and a phloem feeder. *Journal of Animal Ecology*, *75*, 595–603.
- Massey, F. P., Ennos, A. R., & Hartley, S. E. (2007a). Herbivore specific induction of silica-based plant defences. *Oecologia*, *152*, 677–683.
- Massey, F. P., Ennos, A. R., & Hartley, S. E. (2007b). Grasses and the resource availability hypothesis: The importance of silica-based defences. *Journal of Ecology*, *95*, 414–424.
- Massey, F. P., & Hartley, S. E. (2006). Experimental demonstration of the antiherbivore effects of silica in grasses: Impacts on foliage digestibility and vole growth rates. *Proceedings of the Royal Society of London B*, *273*, 2299–2304.
- Massey, F. P., & Hartley, S. E. (2009). Physical defences wear you down: Progressive and irreversible impacts of silica on insect herbivores. *Journal of Animal Ecology*, *78*, 281–291.
- Mayrose, M., Kane, N. C., Mayrose, I., Dlugosch, K. M., & Rieseberg, L. H. (2011). Increased growth in sunflower correlates with reduced defences and altered gene expression in response to biotic and abiotic stress. *Molecular Ecology*, *20*, 4683–4694.
- Meyer, R. S., Duval, A. E., & Jensen, H. R. (2012). Patterns and processes in crop domestication: An historical review and quantitative analysis of 203 global food crops. *New Phytologist*, *196*, 29–48.
- Milla, R., Osborne, C. P., Turcotte, M. M., & Violle, C. (2015). Plant domestication through an ecological lens. *Trends in Ecology & Evolution*, *30*, 463–469.
- Mondolot, L., Marlas, A., Barbeau, D., Gargadennec, A., Pujol, B., & Mckey, D. (2008). Domestication and defence: Foliar tannins and C/N ratios in cassava and a close wild relative. *Acta Oecologica*, *34*, 147–154.
- Müller-Schärer, H., Schaffner, U., & Steinger, T. (2004). Evolution in invasive plants: Implications for biological control. *Trends in Ecology & Evolution*, *19*, 417–422.
- Okuda, A., & Takahashi, E. (1962). Studies on the physiological role of silicon in crop plant: VIII. Some examination on the specific behavior of low land rice in silicon uptake. *Journal on the Science of Soil Science Manure Japan*, *33*, 217–221.
- Pinheiro, J., Bates, D., Debroy, S., Sarkar, D., & R Development Team. (2016). *nlme: Linear and nonlinear mixed effects models*. R package version 3.1-124. <https://CRAN.R-project.org/package=nlme>
- Preece, C., Livarda, A., Christin, P.-A., Wallace, M., Martin, G., Charles, M., ... Osborne, C. P. (2017). How did the domestication of Fertile Crescent grain crops increase their yields? *Functional Ecology*, *31*, 387–397.
- Quigley, K. M., & Anderson, T. M. (2014). Leaf silica concentration in Serengeti grasses increases with watering but not clipping: Insights from a common garden study and literature review. *Frontiers in Plant Science*, *5*, 568.
- R Core Development Team. (2013). *R: A language and environment for statistical computing*. Vienna, Austria: R Foundation for Statistical Computing.
- Rees, M., Osborne, C. P., Woodward, F. I., Hulme, S. P., Turnbull, L. A., & Taylor, S. H. (2010). Partitioning the components of relative growth rate: How important is plant size variation? *American Naturalist*, *176*, 152–161.
- Reidinger, S., Ramsey, M. H., & Hartley, S. E. (2012). Rapid and accurate analyses of silicon and phosphorus in plants using a portable X-ray fluorescence spectrometer. *New Phytologist*, *195*, 699–706.
- Reynolds, O. L., Keeping, M. G., & Meyer, J. H. (2009). Silicon-augmented resistance of plants to herbivorous insects: A review. *Annals of Applied Biology*, *155*, 171–186.
- Roces, F., & Lighton, J. R. B. (1995). Larger bites of leaf-cutting ants. *Nature*, *373*, 392–393.
- Rodrigues, F. A., Polanco, L. R., Silveira Duarte, H. S., Resende, R. S., & Ribeiro Do Vale, F. X. (2015). Photosynthetic gas exchange in common bean submitted to foliar sprays of potassium silicate, sodium molybdate and fungicide and infected with *Colletotrichum lindemuthianum*. *Journal of Phytopathology*, *163*, 554–559.
- Rodriguez-Saona, C., Vorsa, N., Singh, A. P., Johnson-Cicalese, J., Szendrei, Z., Mescher, M. C., & Frost, C. J. (2011). Tracing the history of plant traits under domestication in cranberries: Potential consequences on anti-herbivore defences. *Journal of Experimental Botany*, *62*, 2633–2644.
- Rosenthal, J. P., & Dirzo, R. (1997). Effects of life history, domestication and agronomic selection on plant defence against insects: Evidence from maize and wild relatives. *Evolutionary Ecology*, *11*, 337–355.
- Schaffner, U., Ridenour, W. M., Wolf, V. C., Bassett, T., Müller, C., Müller-Schärer, H., ... Callaway, R. M. (2011). Plant invasions, generalist herbivores, and novel defence weapons. *Ecology*, *92*, 829–835.
- Schaller, J., Brackhage, C., & Dudel, E. G. (2012). Silicon availability changes structural carbon ratio and phenol content of grasses. *Environmental Experimental Botany*, *77*, 283–287.
- Schoelynck, J., Bal, K., Backx, H., Okruszko, T., Meire, P., & Struyf, E. (2010). Silica uptake in aquatic and wetland macrophytes: A strategic choice between silica, lignin and cellulose? *New Phytologist*, *186*, 385–391.
- Seebold, K. W., Kucharek, T. A., Datnoff, L. E., Correa-Victoria, F. J., & Marchetti, M. A. (2001). The influence of silicon on components of resistance to blast in susceptible, partially resistant, and resistant cultivars of rice. *Phytopathology*, *91*, 63–69.
- Shimoyama, S. (1958). Effect of silicon on lodging and wind damage in rice. Report for the Research Funds Granted by Ministry of Agriculture, Japan.
- Siemens, D. H., Garner, S. H., Mitchell-Olds, T., & Callaway, R. M. (2002). Cost of defence in the context of plant competition: *Brassica rapa* may grow and defend. *Ecology*, *83*, 505–517.
- Simpson, K. J., Wade, R. N., Rees, M., Osborne, C. P., & Hartley, S. E. (2017). Data from: Still armed after domestication? Impacts of domestication

- and agronomic selection on silicon defences in cereals *Dryad Digital Repository*, <https://doi.org/10.5061/dryad.6nd96>
- Strömberg, C. A. E., Di Stilio, V. S., & Song, Z. (2016). Functions of phytoliths in vascular plants: An evolutionary perspective. *Functional Ecology*, *30*, 1286–1297.
- Turcotte, M. M., Turley, N. E., & Johnson, M. T. J. (2014). The impact of domestication on resistance to two generalist herbivores across 29 independent domestication events. *New Phytologist*, *204*, 671–681.
- Vicari, M., & Bazely, D. R. (1993). Do grasses fight back – The case for anti-herbivore defences. *Trends in Ecology & Evolution*, *8*, 137–141.
- Welter, S. C. (2000). Contrasting plant responses to herbivory in wild and domesticated habitats. In R. K. D. Peterson, & L. G. Higley (Eds.), *Biotic stress and yield loss*. Boca Raton, FL: CRC Press.
- Whitehead, S., Turcotte, M. M., & Poveda, K. (2017). Domestication impacts on plant-herbivore interactions: A meta-analysis. *Philosophical Transactions of the Royal Society B*, *327*, 20160034.
- Wieczorek, M., Zub, K., Szafranska, P. A., Ksiazek, A., & Konarzewski, M. (2015). Plant-herbivore interactions: Silicon concentration in tussock sedges and population dynamics of root voles. *Functional Ecology*, *29*, 187–194.
- Zhu, Y., & Gong, H. (2014). Beneficial effects of silicon on salt and drought tolerance in plants. *Agronomy for Sustainable Development*, *34*, 455–472.
- Zohary, D., & Hopf, M. (2000). *Domestication of plants in the old world*. Oxford, UK: Oxford University Press.

SUPPORTING INFORMATION

Additional Supporting Information may be found online in the supporting information tab for this article.

How to cite this article: Simpson KJ, Wade RN, Rees M, Osborne CP, Hartley SE. Still armed after domestication? Impacts of domestication and agronomic selection on silicon defences in cereals. *Funct Ecol*. 2017;*31*:2108–2117. <https://doi.org/10.1111/1365-2435.12935>