

Animal board invited review: OneARK: Strengthening the links between animal production science and animal ecology

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OneARK: Strengthening the links between animal production science and animal ecology

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Abstract

Wild and farmed animals are key elements of natural and managed ecosystems that deliver functions such as pollination, pest control and nutrient cyclingwithin the broader roles they play in contributing to biodiversity and to every category of ecosystemservices. They are subjected to global changeswith a profound impact on the natural range and viability of animal species, the emergence and spatial distribution of pathogens, land use, ecosystem services and farming sustainability. We urgently need to improve our understanding of how animal populations can respond adaptively and therefore sustainably to these new selective pressures. In this context, we explored the common points between animal production science and animal ecology to identify promising avenues of synergy between communities through the transfer of concepts and/or methodologies, focusing on seven concepts that link both disciplines. Animal adaptability, animal diversity (bothwithin and between species), selection, animalmanagement, animalmonitoring, agroecology and viability risks were identified as key concepts that should serve the cross-fertilization of both fields to improve ecosystem resilience and farming sustainability. The need for breaking down interdisciplinary barriers is illustrated by two representative examples: i) the circulation and reassortment of pathogens betweenwild and domestic animals and ii) the role of animals in nutrient cycles, i.e. recycling nitrogen, phosphorus and carbon through, for example, contribution to soil fertility and carbon sequestration. Our synthesis identifies the need for knowledge integration techniques supported by programmes and policy tools that reverse the fragmentation of animal research toward a unification into a single Animal Research Kinship, OneARK, which sets new objectives for future science policy. At the interface of animal ecology and animal production science, our article promotes an effective application of the agroecology concept to animals and the use of functional diversity to increase resilience in both wild and farmed systems. It also promotes the use of novel monitoring technologies to quantify animalwelfare and factors affecting fitness. These measures are needed to evaluate viability risk, predict and potentially increase animal adaptability and improve the management of wild and farmed systems, thereby responding to an increasing demand of society for the development of a sustainable management of systems

Keywords

Adaptation; Agroecosystem; Livestock sustainability; Resilience; Zoonotic disease

Implication

We are in a world where the boundaries between natural and managed ecosystems are increasingly unclear with the climate, and other consequences of human activity impacting both wild and farmed animals. In this context, a strong scientific basis is needed to improve the management of all types of animal ecosystem but the disciplines involved have traditionally been fragmented. This synthesis shows that there is much common ground between animal production science and animal ecology, and much to be gained from initiatives to integrate them to provide the urgently needed scientific basis for sustainable management of animals across diverse ecosystems.

Introduction

Our planet is undergoing major global environmental changes mainly caused by a rapid increase in human population and the concomitant agriculture industrialization (specialization, concentration and intensification). These changes have a profound impact on biodiversity, on land use due to modified resource availability, as well as on emergence and spatial distribution of pathogens (Keesing et al., 2010). A primary concern is the extremely rapid rate of these changes, which apply strong and often novel selective pressures on animals, at rates rarely encountered over evolutionary time scales. These challenges are placing new demands on physiological and adaptive capacities (particularly phenotypic plasticity which allows for the compensation of rapid environmental changes when genetic adaptation is too slow), on the interactions among species and ultimately on species persistence and biodiversity. The consequences are major in terms of conservation of biodiversity but will also have impacts on every category of ecosystem services: support (e.g. soil formation), production (e.g. milk, eggs and meat), regulation (e.g. pest control) and cultural, or on their combination (e.g. biodiversity-related ecotourism (Fuller et al., 2007)). Thus, we have a responsibility to find new ways to better understand and preserve the functional diversity of ecosystems. These have been, and will continue to be, a major support of human endeavours.

Animals represent an enormous part of biodiversity, contributing 1.12 million species from a total of 1.43 million catalogued species throughout eukaryotic kingdoms (Mora et al., 2011). Only a very limited number of species are farmed, but they contribute a significant amount of biomass with, for example, mammalian livestock accounting for 60% of global mammalian biomass (Bar-On et al., 2018). Wild and farmed animals are landscape shapers and ecosystem engineers that control the availability of resources by causing changes in biotic or abiotic materials. However, animals are also important vectors, intermediate hosts and reservoirs for microorganisms causing major infectious diseases (Woolhouse et al., 2005). Additionally, wild and farmed animals have always been a major source of proteins for human consumption.

It is increasingly recognized that there is a continuum between animals in managed ecosystems and animals in natural environments. No production system whatever its level of biosecurity is completely isolated from the surrounding environment. Likewise, today, no ecosystem is completely isolated from human influence, and increasingly ecosystems are subject to some degree of human management or have limits imposed on them by human activity. Therefore, it is highly relevant to consider what the cross-fertilization between the two communities of animal production science and animal ecology can bring.

A number of basic concepts appear at first sight to be fundamentally different between animal production science and ecology. However, when these concepts are given due consideration, it transpires that they are actually more similar and not really in opposition. The aim of this paper is to explore the common points between animal production science and animal ecology. Better recognizing the similarities between the two communities will identify promising avenues of synergy by concept and/or methodology transfers between communities. We first discuss seven topics that are common to both communities but viewed from differing perspectives, in order to show their potential for synergy and then highlight these points using two examples. This prospective thinking for a community unification into a single Animal Research Kinship, i.e. OneARK, sets new objectives for future science policy.

Artificial selection vs natural selection

Selection denotes the fact that, among individuals born at a given generation, those that will survive to mate and procreate a new generation can be considered as "chosen" according to some of their characteristics. These characteristics typically impact on their survival, mating probability and their number of descendants. For domestic species, artificial selection depends on decisions taken by humans (breeding managers). For wild species, natural selection emerges from interactions with conspecifics, other species and the abiotic and stochastic environment.

Natural selection can act simultaneously on multiple traits, so that trade-offs are an important part of understanding adaptation and response to selection: natural selection maximizes average fitness of the population, not trait values (Stearns, 1977). Another fundamental aspect is that natural selection varies spatially and temporally depending on the environment (Siepielski et al., 2013 and 2017) so that traits may be positively selected in one environment and counter-selected in another. Investigating selection is thus complex notably because we not only need to assess the actual target of selection but also make sure that the covariances between trait and fitness are not only due to environmental covariance (Morrissey et al., 2010).

It is generally admitted that artificial selection started in the early stages of domestication, the first selected traits being favourable to the domestication process itself, e.g. docility. During the last three centuries, and especially during the last six decades, this artificial selection has become more organized and intense, targeting and maximizing specific traits (e.g. dairy production and growth rate). Another consequence of domestication was to decrease the natural selection pressure because humans increasingly controlled the environment of animals. This is typified by the strong intensification of animal production.

After domestication, selection in different places and with different goals first led to a huge increase in diversity between populations (Darwin, 1859). However, the recent changes in livestock breeding led to the opposite, with (i) a decrease in the number of breeds for a given species (Sherf, 2000) and (ii) a reduction of within-population genetic variability in intensively selected populations (Danchin-Burge et al., 2012), which means a lower adaptive potential in the long run. In the short run, this selection of highly specialized and rather homogeneous "elite" breeding animals led to (i) the unwanted evolution of some functional traits due to unfavourable genetic correlations (e.g. milk yield and female fertility) (Oltenacu and Broom, 2010) and (ii) reduced robustness and flexibility i.e., lower resilience to environmental variability, particularly to new stress and disease challenges. The multivariate nature of selection acknowledged by animal ecologists (Lande and Arnold, 1983) has promoted the development of artificial selection programmes which include the use of selection on multiple traits (Puillet et al., 2016). Indeed, current livestock selection programmes are increasingly seeking to optimize animal fitness in the production environment by putting more emphasis on functional traits and including robustness and adaptability traits alongside production (Berghof et al., 2019). Taking into account such trade-offs is particularly important in the context of global changes where resource availability and variability will be strongly affected.

Collaborative efforts are increasingly needed because the rapid and strong changes of environmental conditions generate strong selective pressures, so much so that humans are now considered as the greatest evolutionary force (Palumbi, 2001; Sarrazin and Lecomte, 2016). Understanding how populations respond to these new selective pressures (e.g. evolution, plasticity or change in distribution area), which means understanding the interrelationships between rates of environmental change and the selection pressure this exerts on animal populations, is a key issue in applied evolution and conservation (e.g. Siepielski et al., 2017). It is also a key issue for artificial selection since global changes are altering the environmental conditions under which artificial selection is operating. For example, because genotypes can perform differently under different environmental conditions (gene by environment interactions, G*E), there is a strong risk that individuals with high breeding values for production traits in protected environments will tend to be negatively impacted by adverse environments, leading to poorer breeding values for those animals that are most environmentally sensitive. Conversely, animals with poorer breeding values for production traits may be the individuals best equipped to deal with environmental perturbations, so that the selection criteria ought to be multivariate and in multiple environments. Animal ecology will benefit from the rapid advances in quantifying the genetic bases of phenotypic/ performance robustness of animals to environmental

3

variability (quantitative genetics, epigenetic regulation), a field that is likely to advance much more rapidly in animal production science because of easier access to controlled genetic materials, advanced control of environmental backgrounds, rapid expansion of multivariate massive phenotyping (including omics) and the ability to account for social interactions between conspecifics (Wade et al., 2010). A major challenge is to understand how global environmental changes are going to affect selective pressures acting on both wild and domesticated populations. Determining the theoretical bases of how natural and artificial selections actually modulate adaptive (and therefore, sustainable) responses of these populations to these new selective pressures is a corner-stone objective. This will pave the way of resolving how we may improve (i) our management of agro- and wild ecosystems by increasing biodiversity and/ or within populations' genotypic/phenotypic diversity, (ii) thereby improving resilience capacity of individuals, populations and systems and (iii) reducing viability risks of our farmed and wild environments.

Viability risks for farmed systems vs natural ecosystems

Global changes pose a viability risk for both natural and farmed systems, although the "currencies" by which viability is judged have traditionally differed; it is largely about economics for farmed systems and about biodiversity and population persistence for natural ecosystems. The framework of ecosystem services links both types of systems by considering them as essential for sustainable development, but viability of natural populations for their own sake also needs to be integrated (Martin et al., 2016). The most commonly used currency to assess viability in wild populations is the probability of extinction of a population over an arbitrarily chosen time period (e.g. 100 years in the International Union for Conservation of Nature red list) or the median time to extinction. Several components of global change will affect viability of both natural and farmed systems.

The impacts of climate change emerge through both long-term changes in average conditions within local environments and an increase in the frequency of extreme events (Ummenhofer and Meehl, 2017). The former has received more attention so far. The effects of climate change can be mediated through many indirect effects such as the disruption of interaction between species because of changes of phenology or morphology (van Gils et al., 2016). A typical example is the earlier breeding of insectivorous birds so that the peak of offspring energetic needs coincides with the peak of food abundance (caterpillars, Visser et al., 1998): if the timing is mismatched, then breeding success is low. These effects are more likely to be encountered in wild than farmed system where longterm changes in average environmental conditions will more frequently be experienced in terms of direct effects that alter resource availability. In farmed systems, the impact on animals will be less direct but in the longer term will impact farm management systems, e.g. impacting the stocking densities of animals that are sustainable in extensive systems, and incurring greater costs for intensive systems (e.g. cooling systems). In managed populations, extreme events such as drought or flooding require the farmer to make costly, unplanned interventions (buying food, transporting animals) where possible. These clearly have economic consequences especially if possible interventions are limited and loss of animals occurs (e.g. rangeland grazing). In wild populations, extreme events reduce individual fitness both through lower survival (e.g. die-offs, McKechnie and Wolf, 2010) and reduced breeding success (Jenouvrier et al., 2015). Extreme events may generate very strong selection pressures leading to marked evolutionary shifts in wild populations (Grant et al., 2017). However, the impact of extreme events is particularly complex to anticipate, as they engage non-linear shifts in multi-species interactions.

Introduced exotic species, which may be pathogens, pathogen carriers, predators or directly competing species, represent another major viability risk to both farmed and wild populations (Bellard et al., 2016; Paini et al., 2016; see section on circulation of zoonotic pathogens). They are likely to be more prevalent and successful in highly anthropized habitats such as peri-urban and agricultural lands, and species of tropical origin benefit from the warming climate in temperate and boreal regions (Hufbauer et al., 2012; Bellard et al., 2013).

Land use is another class of viability risks. There are direct economic impacts of human movement in terms of (i) the value of land or other shared resources such as water in zones where agricultural land is in competition with urban development, and (ii) in terms of rural depopulation (difficulties in recruiting labour, human isolation, costly supply chains) affecting ecological function of agro-landscapes (Sabatier et al., 2014). Extinction risks are further

increased for wild populations due to competition with urban and agricultural land (e.g. palm oil, cocoa) and non-sustainable harvesting (Maxwell et al., 2016). To fully understand viability risks, all these factors and their interactions need to be taken into account.

There are also viability risks due to rigidity of human behaviour. For wild animals, one example is how human habits of farming landscape may evolve in response to recolonization by wild animal species like large carnivores, a question for which some straightforward solutions may exist (Kuijper et al., 2019). In farming, an example of rigidity of human behaviour is the continued use of inappropriate animal genetics through a failure to recognize the traits needed for sustainability in new conditions. Indeed, the loss of genetic diversity of domesticated breeds due to rigid selection of a very few breeds is a major issue being addressed by the FAO (Food and Agriculture Organization (FAO), 2015). Rigidity in farm management, such as failing to adapt fodder cropping practices to changing seasonal patterns, can also increase the viability risks for the animals that depend on this fodder. Rigidity of behaviour can apply not just to humans but also to animal species when one considers differences between generalist/specialist or plastic/non-plastic species (Clavel et al., 2011). For example, one issue is the existence of ecological traps where species respond to cues that were supposed to signal a high-quality environment but that got uncorrelated from this environment, such as asphalt roads that may reflect light in the same manner as water bodies attracting some insects to breed (Schlaepfer et al., 2002). Ultimately, population viability will depend on the ability of organisms to respond adaptively to complex environmental changes inducing novel selective pressures.

Both farmed and wild populations share some of the same viability risks and ultimately must respond by adaptation (microevolution and/ or plasticity). The degree of management of the animal populations within a given ecosystem will mainly affect the extent to which risks can be buffered by human intervention, e.g. deploying reproductive technologies developed in animal production science to aid in rewilding and to overcome habitat fragmentation. Biodiversity and economics are connected across the spectrum from farmed to natural ecosystems. Tools developed at the frontier between ecology and economics, such as coviability analyses (Mouysset et al., 2013), which aim at finding compromises where viability of both farmed and natural systems can co-exist by coupling economic and biodiversity models, will be important for the future.

Agro-ecosystems and farmed animal management vs ecosystems and wild animal management

In contrast to wild animals in natural ecosystems that are fully in interaction with the environment, the magnitude of interactions of farmed animals with the environment covers a spectrum, ranging from agro-ecosystems to landless livestock production. This gradient is driven by the form of the feeding system, ranging from land sharing to land sparing, and the level of interaction the livestock population has vis-a-vis agricultural and natural system components (crops, forest, water, wildlife, etc.). Livestock agro-ecosystems are defined by a high dependence of livestock on local resources, like land and water (pastoralism being its apogee). At the opposite end of the scale, landless livestock systems maximize their direct independence from environmental constraints by means of feed trade, thus establishing production systems with almost no direct relation (excluding by the market) between the places and times where livestock are reared, where their feed is produced, and where their products are consumed.

Gradients in degree of human intervention are also a common element of wild animal and natural ecosystem management. Indeed, not a single natural ecosystem is human proof, at least since climate change started. More direct wild animal ecosystem management profiles can range from biodiversity reserves through natural parks, run as wildlife sanctuaries, to wildlife areas managed by local communities, which recognize combined wildlife, livestock and rangeland services as essential for human groups, a vision emphasized in Southern Africa (Chomba and Nyirenda, 2014; Jones et al., 2015).

In the latter case, there is a strong interaction between agricultural activity and ecosystem management. More generally, the frontier between the "wild" and the "farmed" animals is progressively being eroded, changing to situations where more coexistence and interactions are inevitable if we wish to reconcile preserving biodiversity and better resource sustainability. Achieving this in the design of these reexpanding agro-ecosystems imposes a tightening of the collaboration between animal production scientists and animal ecologists to reconcile opposing

interests. Some examples of this are studies on heathlands or the policy of "Natura 2000" to preserve biodiversity in Europe, often in human-made ecosystems. The governance mode of Natura 2000 brings together land users and civil society in decision-making. It also includes both animal scientists and animal ecologists on its scientific committees, valuing their role in providing evidence through qualitative and quantitative evaluation of benefits, i.e. finding the balance between provisioning services to local farming systems, and markets, and conservation services to the society (McCauley, 2008; Morán-Ordóñez et al., 2013). Furthermore, and in line with societal considerations, there is a visible shift in livestock and wildlife policy dialogue, moving beyond the simple support of resource sufficiency and food provision to now provide incentives for conservation and rehabilitation of functional integrity, and payment for environment services in production areas, and at a global Earth scale (Frost and Bond, 2008; Kammili et al., 2011). Both animal ecology and animal production scientists are then forced to converge when it becomes time to inform politics and the society about solutions to reach the sustainable development objectives (e.g. McCauley, 2008).

The key role of animal adaptability toconnect evolutionary and animal production sciences

Adaptation processes are multifaceted, taking place at different biological levels with different temporal modalities (Gould and Lloyd, 1999). Physiologists, who deal with laboratory and farmed strains, have focused on within lifetime reversible processes that allow individuals to adjust to their environment, with less focus on their heritability. These biological processes depend on the variability of the environment, and adaptation can be described by the following continuum:

(i) phenotypic flexibility of individuals leading to temporary/reversible changes, (ii) developmental plasticity leading to more permanent changes of phenotypes through physiological and/or epigenetic mechanisms and (iii) intergenerational modification of allele frequencies through natural selection (Chevin and Beckerman, 2011). Integrating these different adaptive mechanisms has to be developed together at the interface with animal production science. Studying performance and behavioural changes induced by modifications in the farming environment would provide a great opportunity for evolutionary biologists to investigate the key mechanisms allowing individuals to maintain their performances over different abiotic conditions, complementing and providing a bridge between approaches in the laboratory and in the wild.

The complex phenotypes underlying adaptability are forcing scientists to develop an integrated approach looking at multiple characters. The recent expansion of genomics, and other – omic data, offers new avenues to understand the mechanisms that shape adaptability (Valcu and Kempenaers, 2014). Studying organisms as a whole, taking into account functional links between traits, is now made possible by combining – omic data with the characterization of physiological and performance traits (Prunet et al., 2012). This should uncover cell or physiological processes important for adaptability in both wild and farmed animals. However, such approaches often produce complex data on cell and physiological pathways that are concomitantly affected. Building an integrated phenotyping (Headon, 2013) that sorts the mechanisms underlying adaptability in order of importance now needs to combine biological knowledge of the processes involved, bioinformatics and statistical knowledge.

Important questions remain regarding the role of transgenerational adaptation pathways in fitting, in the long term, populations to their environment. Such phenotypic modulation has a predictive power and may help the offspring to be better adapted to future environmental conditions. Intergenerational plasticity encompasses various mechanisms, including epigenetic changes. These mechanisms are likely to sustain rapid adaptation and promote survival of the next generation (Rey et al., 2016). Their understanding is also a key element for animal production science: it opens an innovative way to optimize productivity, via the modulation of farming conditions during reproduction and offspring growth.

This is not an exhaustive list of the research of interest that remains to be conducted on animal adaptability. However, it emphasizes that promoting the understanding of the link between adaptation and fitness (survival or health state) and of the inheritance of related processes will enhance our ability to predict adaptability of animal populations, living in the wild or under farming conditions.

The importance of animal diversity for system resilience

Ecological resilience focuses on the adaptive capacity of an ecosystem and is defined as the amount of disturbance this system can absorb while remaining within the same stability range and retaining the same function(s), achieved through reinforcing within-system structures, processes and reciprocal feedbacks (Holling, 1996; Kaarlejärvi et al., 2015; Gladstone-Gallagher et al., 2019).

Resilience strongly depends on the initial composition of the local ecological assemblage and the degree of disturbance (Sasaki et al., 2015). In highly disturbed areas, differences in the recovery trajectory of assemblages have been related to differences in the composition and the dispersal capacities of the surrounding species pool of colonists and the level of connectivity among populations, species and ecosystems (Allison, 2004). These factors influence both probability of species persistence by increasing the genetic diversity of local populations (Bach and Dahllöf, 2012) and capacity for recovery by providing sources of propagating organisms (de Juan et al., 2013).

Biodiversity, a key factor for improving the long-term resilience of ecosystems (Awiti, 2011; Mori et al., 2013; Oliver et al., 2015a), is frequently associated with high functional redundancy (i.e. presence of several species able to perform similar functions) (Sasaki et al., 2015; Kaiser-Bunbury et al., 2017) and high species complementarity (Lindegren et al., 2016). Both taxonomic and functional diversities, but not species richness, adequately capture the aspects of biodiversity most relevant to ecosystem stability and functionality (Mori et al., 2013). Taxonomic diversity enhances resilience because most of the rare species within an assemblage are considered as functionally similar to the dominant ones and able to compensate their potential loss under changing environmental conditions, thus maintaining ecosystem functions. However, the maintenance of a particular assemblage is not a necessary requirement for the resilience of ecosystem functions (Oliver et al., 2015b). Functions could be resistant to change or recovered following disturbance with taxonomically different assemblages of species, while exhibiting rather similar sets of traits (Gladstone-Gallagher et al., 2019) or maintaining interactions with sufficient resemblance to the previous system so as to allow it to be recognizably similar (Bregman et al., 2017). Functional diversity improves resilience because a more diverse set of traits increases the variety of potential responses to disturbance (Messier et al., 2019). This then increases the likelihood that species can compensate function(s) lost during disturbance events (Moretti et al., 2006; Kühsel and Blüthgen, 2015). However, resilience is also likely to be scale-dependent (Schippers et al., 2015; GladstoneGallagher et al., 2019), i.e. a combination of traits providing resilience to smallscale disturbance can be ineffective against disturbance acting at largest scale. As a result, the link between biodiversity and resilience is sometimes weak (Bellwood et al., 2003). If the trait structure of highly diverse animal assemblages remains rather stable after moderate stress, further intensification of human pressure can substantially reduce the variety of traits and results in significant alteration of functional diversity (Bregman et al., 2017). This raises the question of how to manage resilience and ecosystem services (i.e. the varied benefits that humans freely gain from the natural environment and from properly functioning managed ecosystems, including provisioning, regulating, cultural and habitat and ecosystem functioning services) in socio-ecological systems?

Conceptual frameworks, tools and indicators (Sasaki et al., 2015; Oliver et al., 2015a) have been defined for quantifying the resilience of coastal fisheries, estuaries or agricultural landscapes (de Juan et al., 2013; Mijatović et al., 2013) based on structural and functional attributes, e.g. ecosystem elasticity or sensitivity and adaptive capacity (López et al., 2013). Trends in the frequency of animal species that provide key ecosystem functions in Great Britain have highlighted that they are not equally impaired by global change, and conservation actions should focus on the functional groups for which there is clear evidence of resilience erosion (Oliver et al., 2015b). Moreover, community field experiments have clearly shown that vegetation restoration can improve pollination, suggesting that the degradation of ecosystem functions is at least partially reversible (Kaiser-Bunbury et al., 2017) and that severe disturbance-driven reduction in ecosystem function does not preclude rapid ecosystem recovery, at least when the ecosystem has not been pushed beyond a tipping point.

Several pattern- or process-oriented strategies have been suggested (Pauly et al., 2002; Fischer et al., 2006) to enhance biodiversity and ecosystem resilience for an improved management of marine and terrestrial production systems including: (i) promoting structurally complex patches of resources throughout the system, and species of

particular concern for functional diversity, but (ii) controlling over-abundant and alien species and minimizing threatening ecosystem processes. Implementing those strategies will result in more heterogeneous production areas, with structurally more complex mosaics of habitats. The resulting production areas are likely to sustain higher levels of animal diversity and will be more resilient to external disturbances.

The concept of animal diversity can be applied in various ways within livestock farming systems. The first aspect of animal diversity is the diversity of species, with for instance a mixed farm exploiting sheep and cattle, or horses and cattle, or an aquaculture farm exploiting different fish species. The benefit of species diversity in the farm is generally based on the ability of various species to exploit different resources. Sheep, cattle and horses in grazing systems are using different patches of grass, with different plants favoured by the different selection strategies. The same type of complementarity is used in recirculated aquaculture systems with fish that feed in different levels of the water column. Complementarity of species can also go beyond complementarity of resources used, with farming systems based on the complete trophic chain such as integrated multi-trophic aquaculture systems (IMTA). The benefit of species diversity in a farm can also rely on the diversity of products that are commercialized. For instance, small ruminants can be used as cash flow, while larger ruminants have a role of savings.

A second aspect of animal diversity is the diversity of individuals of the same species. Animals may be diverse in terms of their adaptive profiles, with for instance a type of cow that copes with heat stress and another type that copes with feed shortage. Having these two types of individuals in a herd can enlarge the range of perturbations that the livestock system can absorb and thereby increase the resilience of system. Animals can also be diverse in terms of their lifetime trajectories, with for instance females that have different types of reproductive rhythms (e.g. extended lactation in dairy production, accelerated lambing in sheep production). This diversity of trajectories within the herd can be useful to cope with environmental challenges (portfolio effect) or to have different types of products answering to different market needs (e.g. heavy/light lambs). Since the 1970's, an increasing emphasis has been put on the preservation of domestic animal diversity. Surveys and field research have been conducted at regional, national or world levels, to identify and characterize all livestock breeds or populations (e.g., Food and Agriculture Organization (FAO), 2015). The preservation of rare or endangered breeds is mainly managed in situ (on farm), and this management involves a huge diversity of actors (Lauvie et al., 2011): farmers, breeding organizations, value chains actors, territorial bodies, natural parks, research institutions, NGOs, governments, etc. Gene banks have been developed in many countries and are used in a complementary way with in situ devices.

The concept of agro-ecology as a sustainable and responsible way forwards

Agro-ecology, a concept originally defined as "the application of ecological theory to the design and management of sustainable agricultural systems" Altieri, 1987, has recently become a hot topic with the aim to optimize economic, ecological and social dimensions to achieve sustainable food production. Understanding the mechanisms underlying the resilience of agro-ecosystems is critical for conserving biodiversity and ecosystem functions in the face of disturbances (Moretti et al., 2006) and for securing the production of essential ecosystem services. Surprisingly, the majority of research on agro-ecology has been done in plant production. This concept now calls scientists from animal ecology and animal production domains to readily interact by developing more interdisciplinarity.

Thus, five key ecological processes were proposed to be adapted to the animal context (Dumont et al., 2013): 1) adopting management practices, including breeding, to improve animal resilience and health; 2) decreasing the external inputs needed for production, particularly use of resources that are directly useable by humans; 3) decreasing pollution by optimizing the metabolic functioning of farming systems, including consideration of animal manure as a resource; 4) enhancing diversity within animal production systems to strengthen farm resilience and 5) preserving biological diversity in agroecosystems.

Even if agro-ecosystem resilience has been considered as a key driver of sustainable agriculture under increasing environmental uncertainty, only a very few studies have explicitly tested the resilience of productivity to disturbance. Taking agroecology forward as a shared discipline needs a number of challenges to be overcome; these relate to scientific problems (Dumont et al., 2013; Carlisle, 2014) and cultural issues. From an ecologist perspective,

agroecosystems are often seen as being a special case study that offers the opportunity to test ecological principles in conditions that are less complex and more clearly controlled than purely natural ecosystems. From the perspective of an animal production scientist, agroecology is often perceived as a constraint problem, i.e. how to achieve economic performance without breaking some environmental "rules". An important objective to better understand the interactions between environmental and biological processes that control community resistance and resilience will be to move beyond these viewpoints and exploit the synergies that the biodiversity within agroecosystems can bring (Tabacchi et al., 2009; TixierBoichard et al., 2015). One example of a useful synergy is to view climatic events as manageable phenomena resulting from processes whose effects could be much more mitigated through the use of integrated ecosystem management and flexible diversification than through adaptation to severe stress (Carlisle, 2014).

Thus, the notion of eco-efficiency may be a powerful tool (Keating et al., 2010). This implies enlarging traditional production-related efficiency definitions to include environmental (land, water, energy), ecological (biodiversity, resilience, conservation) and economic (labour, capital) dimensions. This eco-efficiency approach creates significant challenges for the integration of these multiple dimensions, but there are promising avenues of research tackling this issue (Soteriades et al., 2016).

The commonality in the use of advanced technologies to monitor animals

In the context of agro-ecology, understanding the variability with which individuals respond to their environment is a key entry point for understanding most of the issues raised above. Similarly, study of this variability will also help to assess animal welfare at individual level, an issue which is now a necessary response to the societal demand to improve animal welfare. Animal ecology and production science are both interested in explaining the variability with which individuals respond to their environment and have a lot to win from merging methodological approaches for quantifying this variability.

Recent technological advances allow ecologists studying freeranging animals access to multiple parameters encompassing foraging patterns, social interactions, physiological parameters but also to monitor environmental variables or entire ecological communities (e.g, Rutz and Hays, 2009). These bio-logging technologies, recording from a distance several variables many times per second over periods up to years, now allow the quantification of energetic and behavioural variability between individuals (e.g. accelerometry, Gleiss et al., 2011).

Bio-logging is extensively used, as well, in animal production science and now recognized as field in its own right, in precision livestock farming (Wathes et al., 2008). It permits the monitoring of animals for signs of health problems, allowing timely intervention by the farm manager. The broad nature of the bio-logging data is increasingly useful, particularly with respect to phenotyping complex traits such as resilience and efficiency. Being able to achieve a sustainable balance between resilience and efficiency is a key goal of selection programmes for agroecology. For instance, the efficiency with which farmed animals transfer energy towards body mass production could be evaluated from biologging measurements based on the time-budget devoted to feeding, locomotion, sleeping or social interactions at a daily scale. Such proxy measurements allow the phenotyping of efficiency (and other complex traits) in large populations, and thereby open up for incorporation of such traits in genomic selection (e.g. www.gentore/eu). From a husbandry perspective, finding fine-tuned modifications of farming environment to positively influence this productivity is also conceivable, e.g. detection of circadian optimal conditions in food access or ambient temperature. Those methodologies may change our view of how farmed animals are able to adapt their energy balance in response to changes in farming environments, as they did for wild animals or humans (Villars et al., 2012).

This offers the potential to integrate multiple markers over long time scales to quantify factors affecting overall fitness. One promising step will be to combine diverse biomarkers to evaluate how environmental variations impact fitness and productivity over ages (a fundamental factor for selection in the wild) or over life stages (a key parameter to improve animal productivity). The use of non-invasive methodologies (using hairs, feathers, etc.) including biosensors raises the issue of integrating all this information in a valuable way. Consider, for example, animal

resilience, the capacity to cope with short-term environmental fluctuations. There is no direct measure that encompasses all the facets of resilience; in other words, it is a latent variable that can only be deduced by combining multiple (proxy) measures of its different aspects (see Højsgaard and Friggens, 2010 for a health-related example). This issue of accessing latent variables from multiple proxies is the focus of much research using signal processing methods and will be extremely useful for quantifying the ultimate consequences of within and between individual differences in ecology (e.g. habitat use) and physiology (i.e. energy demands over different time scales).

An important challenge for ecology and animal production science is to safeguard animal welfare and thus health status across the wide range of husbandry and production environments, and also among individuals of different sizes and/or ages. This can range from the surveillance of animals scattered across very extensive rangelands to the monitoring of stress within groups in indoor environments. Currently, most protocols for welfare assessment rely on human observation (i.e. limited duration and potentially subjective). In this context, biologging technologies developed to be implemented in large or small animals have considerable potential to provide continuous monitoring of welfare status, allowing early and rapid identification of changes in behavioural and physiological components (Sadoul et al., 2014; Borchers et al., 2016; Ripperger et al., 2016). We suggest that combining these different types of parameters offers a more complete way to quantify animal welfare, which better integrates animal coping ability to changing environments both in wild and farmed conditions.

Two topical examples of breaking down the interdisciplinary barriers

Elaboration of the above points, and the commonalities that emerge, reinforces the call to more explicitly link these two disciplines for a better understanding of animals as systems, and animals within ecosystems. The importance of making such links, and the benefits arising, is illustrated by considering the following examples:

Circulation andreassortment of potential zoonotic pathogens between wild and domestic populations

Historically, animal domestication has indirectly mediated the transfer of infectious agents between wildlife and humans (Morand et al., 2014). If cases of domestic emergence are not refuted (Pearce-Duvet, 2006), almost threequarters of emerging infectious diseases significant in terms of public health originate in wild animals (Woolhouse et al., 2005). The ongoing covid-19 pandemic is a typical example of a coronavirus (SARS-CoV-2) circulating in wildlife whose evolution has led to human infection (Zhou et al., 2020). To date, information is lacking on where, when, and why it circulates to fully understand the evolution and transmission of SARS-CoV-2 to humans. Transmission of SARSCoV-2 in domestic cats was recently evidenced (Halfmann, 2020), and targets of the virus have been found not only in cats but also in domestic pigs, and to a lower extent goats and hamster (Chen et al., 2020), which indicates that the virus also has the potential to circulate in domestic animals. Earlier in the recent past, the outbreak of highly pathogenic avian influenza (HPAI) H5N8 clade 2.3.4.4 in both wild and domestic birds in Europe is a major example of the "round trips" of viruses between wild and domestic populations. The ancestor of the H5N8 virus was first identified in January 2014 in domestic poultry in South Korea, then adapted to wild migrating aquatic birds and rapidly spread in 2014-2015 (Lycett et al., 2016). This virus affected poultry worldwide from fall 2016 to spring 2017. It caused a few domestic cases in northern Europe, mainly in gallinaceous populations and more rarely in domestic or wild ducks and geese population, which are commonly more resistant to HPAI. A H5N8-related virus appeared in June 2016 in Touva Republic (southern Siberia) causing high mortality in waterfowl (OIE, 2016).

Crossing the species barrier favours transmission and circulation of pathogens and constitutes a major advantage for multi-host pathogens (generalists). Host switches rely on genetic changes including nucleotide substitutions, acquisition of mobile genetic elements or important genome rearrangements through recombinations and reassortments. Influenza viruses are a remarkable example of genetic material exchange between viruses issued from domestic and wild animals. H5N8 is itself a long lasting descendant of the HPAI H5N1 virus, first detected in China in 1996 and responsible for epizootics in domestic birds and some human cases since 2003 (Lycett et al., 2016). The complete sequence of the H5N8 Siberian strain isolated from wild birds in June 2016 revealed many reassortments with other poultry viruses. This virus infected northern European wild and domestic species, whereas

other reassortants infected birds in southern Europe birds in fall 2016 to spring 2017 (Anses, 2017). The emergence of novel pathogenic strains within a region concentrating high densities of a receptive population (fat liver ducks) made possible (i) the dissemination of the virus within domestic and wild bird populations (abundant opportunities for cross-species transmission) and (ii) its reassortment with other low pathogenic strains of influenza virus circulating in the domestic and wild bird populations, thereby creating high levels of genetic diversity that can in turn broaden host-spectra. This example of massive spreading of a wildlife virus within a domestic population is emblematic of the risk induced by the industrialization of production methods, which increased the rearing production of ducks with a number of birds per flock frequently higher than 10 000 and with a higher density of ducks in the free-range pens. These increases in number and density of susceptible birds (without recourse to special sanitary protection measures) are certainly risk factors for a higher spreading of avian influenza.

Production of genetic variants is a mechanism predicted to favour the emergence of zoonotic strains and is difficult to prevent but could be minimized by avoiding passages of the virus from bird to bird or between animal species. Fortunately, most of the time this has not led to pandemic viruses as avian influenza strains do not transfer easily from human to human due to the absence of important receptors in human bronchial tubes. Pigs are an exception to that as they are receptive to influenza viruses specific for pigs, humans and birds (Kaplan et al., 2017). As a consequence, when pigs are co-infected with viruses from different animal origins, they become gene reservoirs with the potential to facilitate reassortments and the emergence of pandemic viruses. Therefore, traditional farming systems mixing free range poultry and pigs in the same backyard close to human populations presents a risk for the emergence of new reassortants of influenza virus able to spread within human populations as pandemic viruses.

Together with emblematic examples of emerging and re-emerging vector-borne diseases in which wild and domestic animals play a key role as vectors, intermediate hosts and/or reservoirs (Boissier et al., 2016), influenza highlights the increasing globalization of health risks and the importance of the human—animal-ecosystem interface in the evolution and emergence of pathogens. It illustrates how a better knowledge of causes and consequences of certain human activities, lifestyles and behaviours in ecosystems is crucial for understanding disease dynamics and driving public policies. Therefore, health security must be understood on a global scale integrating human health, animal health, plant health, ecosystems health and biodiversity, as defined by the One Health concept (Gibbs, 2014; Cunningham et al., 2017; Destoumieux-Garzón et al., 2018). This ambition requires breaking down the interdisciplinary barriers that separate human and veterinary medicine from ecological, evolutionary and environmental science. It calls upon the development of integrative approaches linking the study of proximal factors underlying pathogen emergence and host physiological and adaptive responses to stress to their consequences on ecosystem functioning and evolution (Destoumieux-Garzón et al., 2018).

In that sense, several points discussed in this article may be considered to tackle epizootic diseases and zoonotic diseases. This starts with a required knowledge on the ecology of pathogens of interest (environmental niches, hosts, reservoirs and vectors), which may be complex for multi-host pathogens. While reliable and efficient tools for pathogen monitoring are usually rapidly available, complex pathogen transmission routes are often poorly characterized. New technologies for the monitoring animal contact data, including social networks, give now access to this knowledge. Network modelling should help understanding transmission dynamics in wild animal and livestock populations, which is needed to predict and reduce pathogen transmission (Craft, 2015). Adapting livestock management according to ecological principles is also an important avenue to improve animal health. By reducing contacts, low density farming has been shown to limit pathogen transmission (Tendencia et al., 2011). Introducing genetic diversity in livestock should also be considered as a sustainable way to reduce disease spread. Indeed, genetically homogenous populations (monocultures) are more vulnerable to infection than genetically diverse populations, which have the potential to buffer populations against epidemics in nature (King and Lively, 2012; Ekroth et al., 2019). Finally, new avenues remain to be explored to increase the adaptability of farmed animals. If selective breeding (artificial selection) remains largely used in animal farming, recent studies have shown that new prophylaxes that increase animal adaptability can be envisioned to confer resistant phenotypes to otherwise susceptible animals without affecting the genetic diversity of the livestock. Indeed, several invertebrates (e.g.

oysters, shrimp, honey bees) can be protected from pathogen infections by immune priming, which confers the potential to control infections and limit pathogen transmission, even in species that cannot be vaccinated (Lafont et al., 2017, 2020). A high interest is currently paid to immune priming, which has proven to be trans-generational in a series of cultured invertebrate species (Tetreau et al., 2019). However, the epidemiological consequences of transgenerational immune priming and its impact on the evolution of parasite/pathogen virulence are still debated (Tidbury et al., 2012) and remain to be studied.

The role of animals in the nutrient cycles in terrestrial and aquatic agroecosystems

Pushed by a dynamic political agenda on climate change, the roles of animals on biogeochemical cycles, the livestock sector contribution to global anthropogenic greenhouse gas (GHG) emissions (14.5% of CO₂, CH₄ and N₂O emission) and mitigation options were highlighted (Gerber et al., 2013). This incited animal production research to collaborate with environment science. Initial studies were restricted to closed farm systems, and animals were seen as "a system" emitting nutrients and gases in the atmosphere. Moreover, some effort was given to modelling nutrient emissions associated with waste management (Génermont and Cellier, 1997), proposing some treatment options (Martinez et al., 2009) and practices (Thien Thu et al., 2012).

However, this first era of research focused on partial and segmented analysis of systems, neglecting more complex sets of interactions and flows between ecosystem compartments (not only exchanges with the atmosphere). Research somehow neglected the role of wild and farmed animals in contributing to nutrient and carbon recycling to other compartments of the ecosystem like soil or crops, i.e. considering "animals in their systems", and yet there are clear examples. In Australia, changing dung resources thanks to import of bovine animals has altered the provision of ecosystem services by local population of dung beetles, highlighting again the fact that ecological processes have to be studied in an holistic manner (Nichols et al., 2008). This case study provides evidence of the importance of considering interactions between wild and farmed animals and the need for collaboration, in this case between beetle ecologists and animal scientists.

More recently, there has been a marked increase of holistic and interdisciplinary research addressing biomass, nutrient and carbon recycling in soil—crop—animal systems at various scales, and their ecological, agronomic, environmental and economic impacts (Vayssières et al., 2009). Accordingly, animal science has adopted more holistic models, developing multi-dimensional impact assessment with metrics and methods derived from other disciplines including ecology, biogeochemistry, sociology and economics. Meanwhile, animal ecology and animal science have increasingly stressed the importance of considering the role of humans in their research, i.e. addressing sustainability and functioning of social ecological systems, a concept derived from new institutional economics (Ostrom, 2009).

In the terrestrial production context, research is now addressing animal effects on nutrient and carbon cycles in diverse agroecosystems Altieri, 1987. There are studies of the influence of specific management factors (e.g. ruminantgrazingintensity) on nutrient recycling pathways, soil compaction and carbon stocks (de Faccio Carvalho et al., 2010). In systems research on carbon balance, the use of pasture as the main source of feed was shown to be a non-negligible carbon sink under both semi-arid (e.g. Sahel) and humid environments (e.g. Amazonia). Some authors have addressed the importance of developing an ecosystem approach to better assess the real contribution of livestock (Assouma et al., 2017; Stahl et al., 2017). Enteric methane from ruminants, emission from manure deposition, emission by termites and savannah fire have been accounted for as well as carbon sink function of soils and perennial ligneous vegetation in an annual cycle. The carbon balance was ultimately found to be slightly negative, i.e. emissions due to livestock activities are compensated by carbon sequestration in soil and trees at landscape level. Thus, when environmental impact assessments integrate all the compartments of the agroecosystem (biomass, soil, plants and animals in relation to the atmosphere), and both emission and sequestration, the results contrast with partial analysis that classed African pastoral ecosystems as high GHG contributors. Finally, recent work showed that the use of various metrics would slightly change the evaluated impact of ruminant's methane emission on global warming (Allen et al., 2018). These results, largely to do with a better understanding of

GHG physics, come from another community, and they also stress the need to include other disciplines, i.e. climate and atmospheric science for evaluating environmental impact of animals GHG emissions on global warming.

In the aquatic production context, waste accounts for up to 75% of the nutrient discharge for Nitrogen and Phosphorus in conventional salmon and shrimp aquaculture. Therefore, biological and chemical filters have been developed to partially remove dissolved nutrients from waste. These various pathways of nutrient bioremediation have been increasingly embedded in diverse IMTA, which are mostly adapted for land-based intensive aquaculture (fish, shrimp in ponds) (Troell et al., 2003). In such systems, the addition of extractive organisms like seaweeds (macroalgae, culture of microalgae) (Milhazes-Cunha and Otero, 2017) or bivalves (shellfish) as biofilters to recycle wastewater and reduce discharge and particulate and dissolved nutrient concentration was found promising (from 35 to 100% nitrogen removal). In open culture systems (fish cages), the setting up of IMTA is more complex and results are less clear. Accordingly, research is still on-going.

Such research needs continuity on the long term and design of new models (Lamprianidou et al., 2015). In particular, study of factors influencing reduction efficiency (seaweed species, capacity to uptake beyond physiological requirements, characteristics of production system and the environment, etc.) requires an interdisciplinary research approach (Troell et al., 2003). Similarly, increasing biomass recycling in terrestrial systems, or increasing carbon sequestration by soils and crops, is a long run and complex effort that argues for more global scientific collaboration.

Conclusion

This review highlights seven basic concepts that require crossfertilization between animal ecology and animal production science in order to respond to important societal challenges such as ecosystem resilience and farming sustainability. At the interface of animal ecology and animal production science, our article promotes an effective application of the agroecology concept to animals and the use of functional diversity to increase resilience in both wild and farmed systems. It also promotes the use of novel monitoring technologies to quantify animal welfare and factors affecting fitness. These measures are needed to evaluate viability risk, predict and potentially increase animal adaptability and improve the management of wild and farmed systems, thereby responding to an increasing demand of Society for the development of a sustainable management of systems.

This ambition requires interdisciplinary research: we need a new era of translational research before application of results. Animal ecology has particular strengths in the study of interactions between species, biodiversity, adaptive evolution in natural populations and ecosystem resilience, but in-situ experiments considering broader system impacts are relatively rare. Animal production science has disciplinary strengths in selective breeding, production chains, economics and management. It also has a heritage of methods for combining these at farm or regional systems levels. Therefore, the two disciplines have many complementary skills, but a stronger synergy is lacking due to old habits, i.e. perceived differences in viewpoints on the goal of each discipline, different knowledge and scientific vocabulary (e.g. in quantitative genetics), and different policy masters. Nevertheless, there are substantial advantages to be gained for animal-related research and for society's interaction with animals, from an enhanced cross-fertilization between disciplines.

Modelling approaches have the power to integrate disciplinary visions and knowledge and to translate them into actionable research. However, so far, research has not reached the level of operationality required to fully "pilot" animal systems and agroecosystems. Further, implementation often involves socio-economic factors and innovation processes, which hampers the adoption of any proposed changes. Integration of knowledge holders from the society in the process of research is also needed to tackle anticipated challenges at the interface between science, policy and society. This needs the development of knowledge integration techniques and enhanced collective expertise backed by participatory modelling and science. Such a process begins by breaking down the disciplinary boundaries and promoting cross-fertilization between the animal ecology and animal production science disciplines. This should be accompanied by scientific vision, programmes and policy tools that reverse the fragmentation of animal research

across other themes and instead create critical mass for animal science. The analogy to the emergence of One Health seems highly relevant; it is time for One Animal Research Kinship, OneARK!

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References

Allen, M.R., Shine, K.P., Fuglestvedt, J.S., Millar, R.J., Cain, M., Frame, D.J., Macey, A.H., 2018. A solution to the misrepresentations of CO2-equivalent emissions of shortlived climate pollutants under ambitious mitigation. NPJ Climate and Atmospheric Science 1, 16.

Allison, G., 2004. The influence of species diversity and stress intensity on community resistance and resilience. Ecological Monographs 74, 117–134.

Altieri, 1987. Agroecology: the scientific basis of alternative agriculture. Westview Press, Boulder, USA.

Anses, 2017. Point sur le virus émergent d'influenza aviaire H5N8. https://www.anses.fr/ fr/content/point-sur-le-virus-%C3%A9mergent-d%E2%80%99influenza-aviaire-h5n8.

Assouma, M.H., Serça, D., Guérin, F., Blanfort, V., Lecomte, P., Touré, I., ... Vayssières, J., 2017. Livestock induces strong spatial heterogeneity of soil CO2, N2O and CH4 emissions within a semi-arid sylvo-pastoral landscape in West Africa. Journal of Arid Land 9, 210–221.

Awiti, A.O., 2011. Biological diversity and resilience: lessons from the recovery of cichlid species in Lake Victoria. Ecology and Society 16, 9.

Bach, L., Dahllöf, I., 2012. Local contamination in relation to population genetic diversity and resilience of an arctic marine amphipod. Aquatic Toxicology 114, 58–66.

Bar-On, Y.M., Philipps, R., Milo, R., 2018. The biomass distribution on Earth. Proceedings of the National Academy of Sciences of the USA 115, 6506–6515.

Bellard, C., Thuiller, W., Leroy, B., Genovesi, P., Bakkenes, M., Courchamp, F., 2013. Will climate change promote future invasions? Global Change Biology 19, 3740–3748.

Bellard, C., Cassey, P., Blackburn, T.M., 2016. Alien species as a driver of recent extinctions. Biology Letters 12, 20150623.

Bellwood, D.R., Hoey, A.S., Choat, J.H., 2003. Limited functional redundancy in high diversity systems: resilience and ecosystem function on coral reefs. Ecology Letters 6, 281–285.

Berghof, T.V.L., Poppe, M., Mulder, H.A., 2019. Opportunities to improve resilience in animal breeding programs. Frontiers in Genetics 9, 692.

Boissier, J., Grech-Angelini, S., Webster, B.L., Allienne, J.F., Huyse, T., Mas-Coma, S., ... Mitta, G., 2016. Outbreak of urogenital schistosomiasis in Corsica (France): an epidemiological case study. The Lancet Infectious Diseases 16, 971–979.

Borchers, M.R., Chang, Y.M., Tsai, I.C., Wadsworth, B.A., Bewley, J.M., 2016. A validation of technologies monitoring dairy cow feeding, ruminating, and lying behaviors. Journal of Dairy Science 99, 7458–7466.

Bregman, T.P., Lees, A.C., MacGregor, H.E.A., Darski, B., de Moura, N.G., Aleixo, A., ... Tobias,

J.A., 2017. Using avian functional traits to assess the impact of land-cover change on ecosystem processes linked to resilience in tropical forests. Proceedings of the Royal Society B 283, 20161289.

Carlisle, L., 2014. Diversity, flexibility, and the resilience effect: lessons from a social ecological case study of diversified farming in the northern Great Plains, USA. Ecology and Society 19, 45.

Chen, D., Sun, J., Zhu, J., Ding, X., Lan, T., Zhu, L., Xiang, R., et al., 2020. Single-cell screening of SARS-CoV-2 target cells in pets, livestock, poultry and wildlife. bioRxiv 2020.06.13.149690.

Chevin, L.M., Beckerman, A.P., 2011. From adaptation to molecular evolution. Heredity 108, 457–459.

Chomba, C., Nyirenda, V., 2014. Game ranching: a sustainable land use option and economic incentive for biodiversity conservation in Zambia. Open Journal of Ecology 4, 571–581.

Clavel, J., Julliard, R., Devictor, V., 2011. Worldwide decline of specialist species: toward a global functional homogenization? Frontiers in Ecology and the Environment 9, 222–228.

Craft, M.E., 2015. Infectious disease transmission and contact networks in wildlife and livestock. Philosophical Transactions of the Royal Society, B: Biological Sciences 370, 1669.

Cunningham, A.A., Daszak, P., Wood, J.L.N., 2017. One health, emerging infectious diseases and wildlife: two decades of progress? Philosophical Transactions of the Royal Society, B: Biological Sciences 372, 20160167.

Danchin-Burge, C., Leroy, G., Brochard, M., Moureaux, S., Verrier, E., 2012. Evolution of the genetic variability of eight French dairy cattle breeds assessed by pedigree analysis. Journal of Animal Breeding and Genetics 129, 206–217.

Darwin, C., 1859. On the origin of species by means of natural selection, or the preservation of favoured races in the struggle for life. John Murray, London, UK.

de Faccio Carvalho, P.C., Anghinoni, I., De Moraes, A., De Souza, E.D., Sulc, R.M., Lang, C.R., ... Bayer, C., 2010. Managing grazing animals to achieve nutrient cycling and soil improvement in no-till integrated systems. Nutrient Cycling in Agroecosystems 88, 259–273. de Juan, S., Thrush, S.F., Hewitt, J.E., 2013. Counting on β -diversity to safeguard the resilience of estuaries. PLoS ONE 8, e65575.

Destoumieux-Garzón, D., Mavingui, P., Boetsch, G., Boissier, J., Darriet, F., Duboz, P., ... Voituron, Y., 2018. The One Health concept: 10 years old and a long road ahead. Frontiers in Veterinary Science. 5, 14.

Dumont, B., Fortun-Lamothe, L., Jouven, M., Thomas, M., Tichit, M., 2013. Prospects from agroecology and industrial ecology for animal production in the 21st century. Animal 7, 1028–1043.

Ekroth, A.K.E., Rafaluk-Mohr, C., King, K.C., 2019. Host genetic diversity limits parasite success beyond agricultural systems: a meta-analysis. Proceedings of the Royal Society B: Biological Sciences 286, 20191811.

Fischer, J., Lindenmayer, D.B., Manning, A.D., 2006. Biodiversity, ecosystem function, and resilience: ten guiding principles for commodity production landscapes. Frontiers in Ecology and the Environment 4, 80–86.

Food and Agriculture Organization (FAO), 2015. Second state of the world's animal genetic resources for food and agriculture. Rome, Italy, FAO.

Frost, P.G.H., Bond, I., 2008. The CAMPFIRE programme in Zimbabwe: payments for wildlife services. Ecological Economics 65, 776–787. https://doi.org/10.1016/j. ecolecon.2007.09.018.

Fuller, R.A., Irvine, K.N., Devine-Wright, P., Warren, P.H., Gaston, K.J., 2007. Psychological benefits of greenspace increase with biodiversity. Biology Letters 3, 390–394.

Génermont, S., Cellier, P., 1997. A mechanistic model for estimating ammonia volatilization from slurry applied to bare soil. Agricultural and Forest Meteorology 88, 145–167.

Gerber, P.J., Steinfeld, H., Henderson, B., Mottet, A., Opio, C., Dijkman, J., ... Tempio, G., 2013. Tackling climate change through livestock – A global assessment of emissions and mitigation opportunities. Food and Agriculture Organization of the United Nations (FAO), Rome.

Gibbs, E.P.J., 2014. The evolution of One Health: a decade of progress and challenges for the future. Veterinary Record 174, 85–91.

Gladstone-Gallagher, R.V., Pilditch, C.A., Stephenson, F., Thrush, S.F., 2019. Linking traits across ecological scales determines functional resilience. Trends in Ecology & Evolution 34, 1080–1091.

Gleiss, A.C., Wilson, R.P., Shepard, E.L., 2011. Making overall dynamic body acceleration work: on the theory of acceleration as a proxy for energy expenditure. Methods in Ecology and Evolution 2, 23–33.

Gould, S.J., Lloyd, E.A., 1999. Individuality and adaptation across levels of selection: how shall we name and generalize the unit of Darwinism? Proceedings of the National Academy of Sciences 96, 11904–11909.

Grant, P.R., Grant, B.R., Huey, R.B., Johnson, M.T.J., Knoll, A.H., Schmitt, J., Grant, P.R., 2017. Evolution caused by extreme events. Philosophical Transactions of the Royal Society, B: Biological Sciences 372, 20160146.

Halfmann, P.J., et al., 2020. Transmission of SARS-CoV-2 in domestic cats. The New England Journal of Medicine 383, 592–594 NEJMc2013400.

Headon, D., 2013. Systems biology and lifestock production. Animal 7, 1959–1963.

Højsgaard, S., Friggens, N.C., 2010. Quantifying degree of mastitis from common trends in a panel of indicators for mastitis in dairy cows. Journal of Dairy Science 93, 582–592.

Holling, C.S., 1996. Engineering resilience versus ecological resilience. In: Schulze, P. (Ed.), Engineering within ecological constraints. National Academy Press, Washington, DC, USA, pp. 31–44.

Hufbauer, R.A., et al., 2012. Anthropogenically induced adaptation to invade (AIAI): contemporary adaptation to human-altered habitats within the native range can promote invasions. Evolutionary Applications 5, 89–101.

Jenouvrier, S., Péron, C., Weimerskirch, H., 2015. Extreme climate events and individual heterogeneity shape life-history traits and population dynamics. Ecological Monographs 85, 605–624.

Jones, B.T.B., Diggle, R.W., Thouless, C., 2015. From exploitation to ownership: wildlifebased tourism and communal area conservancies in Namibia. In: Van Der Duim, R., Lamers, M., Van Wijk, J. (Eds.), Institutional arrangements for conservation, development and tourism in Eastern and Southern Africa. Springer Netherlands, Dordrecht, The Netherlands, pp. 17–37.

Kaarlejärvi, E., Hoset, K.S., Olofsson, J., 2015. Mammalian herbivores confer resilience of Arctic shrub-dominated ecosystems to changing climate. Global Change Biology 21, 3379–3388.

Kaiser-Bunbury, C.N., Mougal, J., Whittington, A.E., Valentin, T., Gabriel, R., Olesen, J.M., Blüthgen, N., 2017. Ecosystem restoration strengthens pollination network resilience and function. Nature 542, 223–229.

Kammili, T., Hubert, B., Tourrand, J.F., 2011. A paradigm shift in livestock management: from resource sufficiency to functional integrity. Editions Cardère, Avignon, France.

Kaplan, B.S., Torchetti, M.K., Lager, K.M., Webby, R.J., Vincent, A.L., 2017. Absence of clinical disease and contact transmission of North American clade 2.3.4.4 H5NX HPAI in experimentally infected pigs. Influenza and Other Respiratory Viruses 11, 464–470.

Keating, B.A., Carberry, P.S., Bindraban, P.S., Asseng, S., Meinke, H., Dixon, J., 2010. Ecoefficient agriculture: concepts, challenges, and opportunities. Crop Science 50, S109–S119.

Keesing, F., Belden, L.K., Daszak, P., Dobson, A., Harvell, C.D., Holt, R.D., ... Ostfeld, R.S., 2010. Impacts of biodiversity on the emergence and transmission of infectious diseases. Nature 468, 647–652.

King, K.C., Lively, C.M., 2012. Does genetic diversity limit disease spread in natural host populations. Heredity. 109, 199–203.

Wathes, C.M., Kristensen, H.H., Aerts, J.M., Berckmans, D., 2008. Is precision livestock farming an engineer's daydream or nightmare, an animal's friend or foe, and a farmer's panacea or pitfall? Computers and Electronics in Agriculture 64, 2–10.

Kühsel, S., Blüthgen, N., 2015. High diversity stabilizes the thermal resilience of pollinator communities in intensively managed grasslands. Nature Communications 6, 7989.

Kuijper, D.P.J., Churski, M., Trouwborst, A., Heurich, M., Smit, C., Kerley, G.I.H., Cromsigt, J. P.G.M., 2019. Keep the wolf from the door: How to conserve wolves in Europe's human-dominated landscapes? Biological Conservation 235, 102–111.

Lafont, M., Petton, B., Vergnes, A., Pauletto, M., Segarra, A., Gourbal, B., Montagnani, C., 2017. Long-lasting antiviral innate immune priming in the Lophotrochozoan Pacific Oyster Crassostrea gigas. Scientific Reports 7, 13143.

Lafont, M., Vergnes, A., Vidal-Dupiol, J., de Lorgeril, J., Gueguen, Y., Haffner, P., et al., 2020. A sustained immune response supports long-term antiviral immune priming in the Pacific Oyster, Crassostrea gigas. mBio 11, e02777-19.

Lamprianidou, F., Telfer, T., Ross, L.G., 2015. A model for optimization of the productivity and bioremediation efficiency of marine integrated multitrophic aquaculture. Estuarine, Coastal and Shelf Science 164, 253–264.

Lande, R., Arnold, S.J., 1983. The measurement of selection on correlated characters. Evolution 37, 1210-1226.

Lauvie, A., Audiot, A., Couix, N., Casabianca, F., Brives, H., Verrier, E., 2011. Diversity of rare breed management programs: Between conservation and development. Livestock Science 140, 161–170.

Lindegren, M., Checkley Jr., D.M., Ohman, M.D., Koslow, J.A., Goericke, R., 2016. Resilience and stability of a pelagic marine ecosystem. Proceedings of the Royal Society B: Biological Sciences 283, 20151931.

López, D.R., Brizuela, M.A., Willems, P., Aguiar, M.R., Siffredi, G., Bran, D., 2013. Linking ecosystem resistance, resilience, and stability in steppes of North Patagonia. Ecological Indicators 24, 1–11.

Lycett, S.J., Bodewes, R., Pohlmann, A, Banks, J., Bányai, C., Boni, M.J., ... Kuiken, T., 2016. Role for migratory wild birds in the global spread of avian influenza H5N8. The Global Consortium for H5N8 and Related Influenza Viruses. Science 354, 213–217.

Martin, J.-L., et al., 2016. The need to respect nature and its limits challenges society and conservation science. Proceedings National Academy Science USA 113, 6105–6112.

Martinez, J., Dabert, P., Barrington, S., Burton, C., 2009. Livestock waste treatment systems for environmental quality, food safety, and sustainability. Bioresource Technology 100, 5527–5536.

Maxwell, S.L., Fuller, R.A., Brooks, T.M., Watson, J.E.M., 2016. The ravages of guns, nets and bulldozers. Nature 536, 143–145.

McCauley, D., 2008. Sustainable development and the 'governance challenge': the French experience with Natura 2000. European Environment 18, 152–167.

McKechnie, A.E., Wolf, B.O., 2010. Climate change increases the likelihood of catastrophic avian mortality events during extreme heat waves. Biology Letters 6, 253–256.

Messier, C., Bauhus, J., Doyon, F., Maure, F., Sousa-Silva, R., Nolet, P., Mina, M., Aquilué, N., Fortin, M.J., Puettmann, K., 2019. The functional complex network approach to foster forest resilience to global changes. Forest Ecosystems 6, 21.

Mijatović, D., Van Oudenhoven, F., Eyzaguirre, P., Hodgkin, T., 2013. The role of agricultural biodiversity in strengthening resilience to climate change: towards an analytical framework. International Journal of Agricultural Sustainability 11, 95–107.

Milhazes-Cunha, H., Otero, A., 2017. Valorisation of aquaculture effluents with microalgae: the integrated multi-trophic aquaculture concept. Algal Research 24, 416–424.

Mora, C., Tittensor, D.P., Adl, S., Simpson, A.G.B., Worm, B., 2011. How many species are there on earth and in the ocean? PLoS Biology 9, 1–8.

Morand, S., McIntyre, K.M., Baylis, M., 2014. Domesticated animals and human infectious diseases of zoonotic origins: domestication time matters. Infection, Genetics and Evolution 24, 76–81.

Morán-Ordóñez, A., Bugter, R., et al., 2013. Temporal changes in socio-ecological systems and their impact on ecosystem services at different governance scales: a case study of heathlands. Ecosystems 16, 765–782.

Moretti, M., Duelli, P., Obrist, M.K., 2006. Biodiversity and resilience of arthropod communities after fire disturbance in temperate forests. Oecologia 149, 312–327.

Mori, A.S., Furukawa, T., Sasaki, T., 2013. Response diversity determines the resilience of ecosystems to environmental change. Biological Reviews 88, 349–364.

Morrissey, M.B., Kruuk, L.E.B., Wilson, A.J., 2010. The danger of applying the breeder's equation in observational studies of natural populations. Journal of Evolutionary Biology 23, 2277–2288.

Mouysset, L., Doyen, L., Jiguet, F., 2013. From population viability analysis to coviability. Conservation Biology 28, 187–201.

Nichols, E., Spector, S., et al., 2008. Ecological functions and ecosystem services provided by Scarabaeinae dung beetles. Biological Conservation 141, 1461–1474.

OIE, 2016. Highly pathogenic avian influenza, Russia. http://www.oie.int/wahis_2/public/wahid.php/Reviewreport/Review?page_refer=MapFullEventReport&reportid= 20335.

Oliver, T.H., Heard, M.S., Isaac, N.J.B., Roy, D.B., Procter, D., Eigenbrod, F., ... Bullock, J.M., 2015a. Biodiversity and resilience of ecosystem functions. Trends in Ecology & Evolution 30, 673–684.

Oliver, T.H., Isaac, N.J.B., August, T.A., Woodcock, B.A., Roy, D.B., Bullock, J.M., 2015b. Declining resilience of ecosystem functions under biodiversity loss. Nature Communications 6, 10122.

Oltenacu, P.A., Broom, D.M., 2010. The impact of genetic selection for increased milk yield on the welfare of dairy cows. Animal Welfare 19, 39–49.

Ostrom, E., 2009. A general framework for analyzing sustainability of social-ecological systems. Science 325, 419–422.

Paini, D.R., Sheppard, A.W., Cook, D.C., De Barro, P.J., Worner, S.P., Thomas, M.B., 2016. Global threat to agriculture from invasive species. Proceedings of the National Academy of Sciences of the USA 113, 7575–7579.

Palumbi, S.R., 2001. Humans as the world's greatest evolutionary force. Science 293, 1786–1790.

Pauly, D., Christensen, V., Guenette, S., Pitcher, T.J., Sumaila, U.R., Walters, C.J., ... Zeller, D., 2002. Towards sustainability in world fisheries. Nature 418, 689–695.

Pearce-Duvet, J.M., 2006. The origin of human pathogens: evaluating the role of agriculture and domestic animals in the evolution of human disease. Biological Review Cambridge Philosophical Society 81, 369–382.

Prunet, P., Overli, O., Douxfils, J., Bernardini, G., Kestemont, P., Baron, D., 2012. Fish welfare and genomics. Fish Physiology and Biochemistry 38, 43–60.

Puillet, L., Réale, D., Friggens, N.C., 2016. Disentangling the relative roles of resource acquisition and allocation on animal feed efficiency: insights from a dairy cow model. Genetics Selection Evolution 48, 1–16.

Rey, O., Danchin, E., Mirouze, M., Loot, C., Blanchet, S., 2016. Adaptation to global change: a transposable element—epigenetics perspective. Trends in Ecology & Evolution 31, 514–526.

Ripperger, S., Josic, D., Hierold, M., Koelpin, A., Weigel, R., Hartmann, M., ... Mayer, F., 2016. Automated proximity sensing in small vertebrates: design of miniaturized sensor nodes and first field tests in bats. Ecology and Evolution 6, 2179–2189.

Rutz, C., Hays, G.C., 2009. New frontiers in biologging science. Biological Letters 5, 289–292.

Sabatier, R., Doyen, L., Tichit, M., 2014. Heterogeneity and the trade-off between ecological and productive functions of agro-landscapes: A model of cattle-bird interactions in a grassland agroecosystem. Agricultural Systems 126, 38–49.

Sadoul, B., Evouna Mengues, P., Friggens, N.C., Prunet, P., Colson, V., 2014. A new method for measuring group behaviours of fish shoals from recorded videos taken in near aquaculture conditions. Aquaculture 430, 179–187.

Sarrazin, F., Lecomte, J., 2016. Evolution in the Anthropocene. Science 351, 922–923.

Sasaki, T., Furukawa, T., Iwasaki, Y., Seto, M., Mori, A.S., 2015. Perspectives for ecosystem management based on ecosystem resilience and ecological thresholds against multiple and stochastic disturbances. Ecological Indicators 57, 395–408.

Schippers, P., van der Heide, C.M., Koelewijn, H.P., Schouten, M.A.H., Smulders, R.M.J.M., Cobben, M.M.P., ... Verboom, J., 2015. Landscape diversity enhances the resilience of populations, ecosystems and local economy in rural areas. Landscape Ecology 30, 193–202.

Schlaepfer, M.A., Runge, M.C., Sherman, P.W., 2002. Ecological and evolutionary traps. Trends in Ecology & Evolution 17, 474–480.

Sherf, B., 2000. World watch list for domestic animal diversity. 3rd edition. FAO, Rome, Italy.

Siepielski, A.M., Gotanda, K.M., Morrissey, M.B., Diamond, S.E., DiBattista, J.D., Carlson, S.M., 2013. The spatial patterns of directional phenotypic selection. Ecology Letters 16, 1382–1392.

Siepielski, A.M., Morrissey, M.B., Buoro, M., Carlson, S.M., Caruso, C.M., Clegg, S.M., Coulson, T., Dibattista, J., Gotanda, K.M., Francis, C.D., Hereford, J., Kingsolver, J.G., Sletvold, N., Svensson, E.I., Wade, M.J., Maccoll, A.D.C., 2017. Precipitation drives global variation in natural selection. Science 355, 959–962.

Soteriades, A.D., Stott, A.W., Moreau, S., Charroin, T., Blanchard, M., Liu, J., Faverdin, P., 2016. The relationship of dairy farm eco-efficiency with intensification and selfsufficiency. Evidence from the French dairy sector using life cycle analysis, data envelopment analysis and partial least squares structural equation modelling. PLoS ONE 11, e0166445.

Stahl, C., Fontaine, S., Klumpp, K., Picon-Cochard, C., Grise, M.M., Dezécache, C., ... Blanfort, V., 2017. Continuous soil carbon storage of old permanent pastures in Amazonia. Global Change Biology 23, 3382–3392.

Stearns, S.C., 1977. The evolution of life history traits: a critique of the theory and a review of the data. Annual Review of Ecology, Evolution, and Systematics 8, 145–171.

Tabacchi, E., Steiger, J., Corenblit, D., Monaghan, M.T., Planty-Tabacchi, A.-M., 2009. Implications of biological and physical diversity for resilience and resistance patterns within highly dynamic river systems. Aquatic Sciences 71, 279–289.

Tendencia, E.A., BosmabJohan, R.H., Verreth, A.J., 2011. White spot syndrome virus (WSSV) risk factors associated with shrimp farming practices in polyculture and monoculture farms in the Philippines. Aquaculture 311, 87–93.

Tetreau, G., Dhinaut, J., Gourbal, B., Moret, Y., 2019. Trans-generational Immune Priming in Invertebrates: current knowledge and future prospects. Frontiers in Immunology 10, 1938.

Thien Thu, C.T., Cuong, P.H., Hang, L.T., Chao, N.V., Anh, L.X., Trach, N.X., Sommer, S.G., 2012. Manure management practices on biogas and non-biogas pig farms in developing countries – using livestock farms in Vietnam as an example. Journal of Cleaner Production 27, 64–71.

Tidbury, H.J., Best, A., Boots, M., 2012. The epidemiological consequences of immune priming. Proceedings of the Royal Society B: Biological Sciences 279, 4505–4512.

Tixier-Boichard, M., Verrier, E., Rognon, X., Zerjaln, T., 2015. Farm animal genetic and genomic resources from an agroecological perspective. Frontiers in Genetics 6, 153.

Troell, M., Halling, C., Neori, A., Chopin, T., Buschmann, A.H., Kautsky, N., Yarish, C., 2003. Integrated mariculture: asking the right questions. Aquaculture 226, 69–90.

Ummenhofer, C.C., Meehl, G.A., 2017. Extreme weather and climate events with ecological relevance: a review. Philosophical Transactions of the Royal Society, B: Biological Sciences 372, 20160135.

Valcu, C.M., Kempenaers, B., 2014. Proteomics in behavioral ecology. Behavioral Ecology 26, 1–15.

van Gils, J.A., Lisovski, S., Lok, T., Meissner, W., Ozarowska, A., de Fouw, J., ... Klaassen, M., 2016. Body shrinkage due to Arctic warming reduces red knot fitness in tropical wintering range. Science 352, 819–821.

Vayssières, J., Guerrin, F., Paillat, J.-M., Lecomte, P., 2009. GAMEDE: a global activity model for evaluating the sustainability of dairy enterprises. Part I – Whole-farm dynamic model. Agricultural Systems 101, 128–138.

Villars, C., Bergouignan, A., Dugas, J., Antoun, E., Schoeller, D.A., Roth, H., ... Simon, C., 2012. Validity of combining heart rate and uniaxial acceleration to measure free-living physical activity energy expenditure in young men. Journal of Applied Physiology 113, 1763–1771.

Visser, M.E., Noordwijk, A.J.V., Tinbergen, J.M., Lessells, C.M., 1998. Warmer springs lead to mistimed reproduction in great tits (Parus major). Proceedings of the Royal Society B: Biological Sciences 265, 1867–1870.

Wade, M.J., Bijma, P., Ellen, E.D., Muir, W., 2010. Group selection and social evolution in domesticated animals. Evolutionary Applications 3, 453–465.

Woolhouse, M.E.J., Haydon, D.T., Antia, R., 2005. Emerging pathogens: the epidemiology and evolution of species jumps. Trends in Ecology & Evolution 20, 238–244.

Zhou, P., Yang, X.L., Wang, X.G., et al., 2020. A pneumonia outbreak associated with a new coronavirus of probable bat origin. Nature 579, 270–273.