

Soil biota in vineyards are more influenced by plants and soil quality than by tillage intensity or the surrounding landscape (vol 7, 2017)

Jacob Buchholz, Pascal Querner, Daniel Paredes, Thomas Bauer, Peter Strauss, Muriel Guernion, Jennifer Scimia, Daniel Cluzeau, Francoise Burel, Sophie Kratschmer, et al.

To cite this version:

Jacob Buchholz, Pascal Querner, Daniel Paredes, Thomas Bauer, Peter Strauss, et al.. Soil biota in vineyards are more influenced by plants and soil quality than by tillage intensity or the surrounding landscape (vol 7, 2017). Scientific Reports, 2018, 7 (1), pp.17445. 10.1038/s41598-018-24797-y. hal-01833143

HAL Id: hal-01833143 <https://univ-rennes.hal.science/hal-01833143v1>

Submitted on 12 Jul 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

[Distributed under a Creative Commons Attribution 4.0 International License](http://creativecommons.org/licenses/by/4.0/)

SCIENTIFIC REPERTS

[Correction: Author Correction](https://doi.org/10.1038/s41598-018-24797-y)

Received: 30 June 2017 Accepted: 29 November 2017 Published online: 12 December 2017

Soil biota in vineyards are more infuenced by plants and soil quality than by tillage intensity or the surrounding landscape

Jacob Buchholz1, Pascal Querner¹, Daniel Paredes², Thomas Bauer3, Peter Strauss3, Muriel Guernion4, Jennifer Scimia4, Daniel Cluzeau4, Françoise Burel5, Sophie Kratschmer6, Silvia Winter⁶, Martin Potthof7 & Johann G. Zaller ¹

Tillage is known for its adverse efects on soil biota, at least in arable agroecosystems. However, in vineyards efects might difer as tillage is often performed during dry periods or only in every other inter-row allowing species to re-colonise disturbed areas. We examined the response of earthworms (lumbricids), springtails (collembola) and litter decomposition to periodically mechanically disturbed (PMD) and permanently green covered (PGC) vineyard inter-rows and assessed whether site efects are altered by the surrounding landscape. In commercial vineyards in Austria we sampled earthworms by handsorting, springtails by soil coring and pitfall trapping and installed litter decomposition bags. Earthworm species diversity increased with plant biomass under PMD but not under PGC; earthworm density was unafected by tillage but increased with plant biomass mainly at high soil quality (soil fertility index). Springtail species diversity was unafected by tillage; springtail densities (mainly larger species) were reduced under PGC. Litter decomposition was little afected by investigated parameters. Landscape heterogeneity afected the functional diversity of surface springtails, but did not infuence soil-dwelling springtails, earthworms or litter decomposition. We conclude that efects on soil biota of periodical tillage in vineyards need not necessarily be detrimental and will be modifed by plant biomass and soil quality.

Viticulture is an ancient and globally widespread agricultural production system. Smallholder wine-growers contribute to multifunctional landscapes that, in addition to grape and wine production, provide a variety of ecosystem services¹. Vineyards are often intensively managed involving frequent tilling and/or pesticide treatments². Inter-row soil management practices are performed for weed control, water conservation or to prevent erosion. These soil tillage regimes include permanent bare soil as a result of intensive tillage, moderate tillage of every second inter-row only (also called alternating tillage) or no tillage, leaving permanently vegetated inter-rows³. Soil tillage has been shown to negatively afect soil biodiversity which is directly linked to associated ecosystem functions such as nutrient cycling or soil formation⁴. The compostion and biomass of vegetation may also modify the food source for soil organisms thereby altering their community composition^{5,6}. However, only a few studies have investigated effects of vineyard soil management practices on earthworms⁷⁻¹¹, springtails^{12,13} or soil microbial communities^{14,15}. These studies mainly compared very contrasting management measures such as intensive

1 Institute of Zoology, University of Natural Resources and Life Sciences Vienna (BOKU), Gregor Mendel Straße 33, 1180, Vienna, Austria. ² Grupo de Protección Vegetal, Departamento de Protección Ambiental, Estación Experimental de Zaidín, CSIC, Profesor Albareda n◦ 1, 18008, Granada, Spain. 3Institute for Land and Water Management Research, Austrian Federal Agency for Water Management, Pollnbergstraße 1, A-3252, Petzenkirchen, Austria. 4 Université de Rennes I, OSUR, UMR CNRS 6553 'EcoBio', Station Biologique de Paimpont, 35380, Paimpont, France. 5 Université de Rennes I, OSUR, UMR CNRS 6553 'EcoBio', Avenue du Général Leclerc Campus de Beaulieu, F-35042, Rennes Cedex, France. ⁶Institute of Integrative Nature Conservation Research, University of Natural Resources and Life Sciences Vienna, Gregor Mendel Straße 33, 1180, Vienna, Austria. ⁷Centre of Biodiversity and Sustainable Land Use (CBL), University of Göttingen, Grisebachstraße 6, 37077, Göttingen, Germany. Correspondence and requests for materials should be addressed to J.G.Z. (email: johann.zaller@boku.ac.at)

tillage, or herbicide applications vs. permanent vegetation cover. To the best of our knowledge nothing is known on the efects on soil biota in vineyards with alternating tillage vs. no tillage.

In vineyards, as in many other agroecosystems, the role of earthworms (Oligochaeta: Lumbricidae) in promoting soil fertility, aggregate formation and soil organic matter protection is important¹⁶. Generally, earthworms are sensitive to soil tillage¹⁷, and have been used as bioindicators for sustainable soil use and soil quality^{18–20}. Earthworms can be divided into three ecological groups according to their habitat and functional role in the agroecosystem²¹. Epigeic species live in the top soil and litter layer and usually do not build permanent burrows. Anecic species drill vertical burrows reaching from the soil surface down to subsoil layers. Endogeic species burrow mainly horizontally in the upper mineral soil layers. In vineyards, earthworm abundance has been shown to reach 241 worms m⁻², but this can vary considerably⁸. Other important indicators of soil quality and sustainable land use are springtails (Hexapoda: Collembola²²⁻²⁷). By feeding on fungi, bacteria and dead organic matter, as well as living plant tissue they mobilize locked-up nutrients^{28,29}, contribute to supporting ecosystem services, and are sensitive to land use changes^{30,31}. Springtails are among the most abundant microarthropods in soil^{32,33}, with densities of up to 60.000 ind. m^{−2} in grasslands³⁴. Springtails can be grouped into the three life forms: eu- and hemiedaphic species live mainly in the soils while epedaphic species are active on the soil surface^{32,35}. Earthworm and springtail community changes are considered to be powerful indicators for ecosystem service provision in vineyard soils. Another important process integrating the activity of soil biota is litter decomposition 36 . For example, mesofaunal contribution to litter decomposition is directly involved in the degradation of material and indirectly by feeding on microorganisms. Microorganisms in soil have also been shown to be sensitive to tillage practices³⁷. Yet, also other practices such as pesticide applications can modify litter decomposition rates^{38,39}.

There is a growing body of studies showing that both plant and aboveground animal biodiversity in vineyards are influenced by the diversity and structure of the surrounding landscape^{40–45}. For example, plant communities in vineyards have been shown to be positively affected by nearby semi-natural habitats⁴³, specialized ambush spiders in vineyards increased with landscape heterogeneity based on land use categories⁴⁰, sedentary butterfly species in vineyards were related to remnant patches of native vegetation in the surrounding landscape⁴¹. We also know from studies in arable crops that both earthworms and springtails were afected by the surrounding landscape structure and land-use intensity within a radius of several hundred meters⁴⁶⁻⁵². However, to the best of our knowledge it has never been examined whether soil biota inhabiting vineyards are also infuenced by the surrounding landscape structure.

The aim of this study was to examine the response of earthworm and springtail communities and litter decomposition to periodically mechanically-disturbed (PMD) or permanently green covered (PGC) inter-row management in vineyards and whether this response is infuenced by the surrounding landscape structure. We expected that (i) earthworms will be more vulnerable to soil disturbance than springtails due to their body size, (ii) both earthworms and springtails will be positively related to plant diversity measures and (iii) landscape complexity will alter potential effects of springtails or earthworms. These interactions were tested in 16 viticultural landscapes in Austria.

Results

Earthworms. In our vineyards we found a total of 564 individuals of earthworms, comprising 13 species (5 anecic and 8 endogeic spp.; a full species list is provided in Supplementary Table S1). The mean earthworm density was 146 ± 132 ind. m⁻² (mean \pm sd) across all samples; the maximum of 550 ind. m⁻² and 286 g m⁻² was reached in a PMD inter-row (Supplementary Table S1). Overall, 83% of earthworms were endogeics (most abundant spp. was *Aporrectodea caliginosa*), 17% belonged to anecics (most abundant spp was *Lumbricus terrestris*); no epigeics were found.

Earthworm species richness, density and biomass were best explained by soil quality (expressed by the soil feld index) and plant biomass (richness: $R_m^2 = 0.27$; density: $R_m^2 = 0.30$; biomass: $R_m^2 = 0.26$; Table 1). Both soil quality and plant biomass increased the number of earthworm species and earthworm density (Fig. 1a,b). Similar patterns were observed for earthworm biomass, however data are not shown. Lowest species richness of earthworms was predicted for vineyards with low soil quality (feld index 23), and highest earthworm species richness for vineyards with a high soil quality (field index 67; Fig. 1a). The effect of plant biomass on earthworms was small in sites with low soil quality, but increased with increasing soil quality (Fig. 1a). Similar patterns were found for earthworm density (Fig. 1b), with the lowest density in vineyards with low soil quality, and highest density in vineyards with high soil quality. Again, the efect of plant biomass on earthworm density was higher in sites with high soil quality.

Response of anecic earthworm density was best explained by soil quality, plant biomass and the interaction between those two variables (R_{m}^{2} = 0.33; Table 1). On sites with low soil quality, fewer anecics were found when plant biomass was low (Fig. 1c). Anecic density started to increase above 180 gm^{−2} plant biomass and reached a maximum with plant biomass of 350 gm⁻². On sites with high soil quality, highest anecic densities (57 ind. m⁻²) were found with low inter-row plant biomass, and decreased with increasing plant biomass.

Endogeic species richness was best explained by soil management, plant biomass, interaction between management and plant biomass and soil quality, $(R^2_{adj}=0.31;$ Table 1). Under PGC management the effect of plant biomass on the number of endogeic species was negative (Fig. 1d). However, under PMD efect of plant biomass was positive and higher with a threefold increase in the species numbers along the plant biomass gradient. Responses of endogeic species richness and density to plant biomass and soil quality was similar to overall earthworm richness and density ($R_{m}^{2}=0.22$).

Neither anecic nor endogeic earthworms were afected by the surrounding landscape.

Springtails. We collected 12784 ind. of springtails comprising 60 different species. Mean density of the soil dwelling springtails was 37315 ± 35322 ind. m⁻² (most abundant spp. *Parisotoma notabilis*) and mean activity density of surface dwelling springtails was 70±52 ind. pitfall sample[−]¹ (most abundant spp. *Isotoma viridis*; a full species list is provided in Supplementary Table S2).

Table 1. Comparison of alternative models (using AICc) for earthworms (richness...spp. m^{-2} , density...ind. m^{-2} , biomass...g m[−]²) and litter decomposition (S value…stabilisation index, k…decomposition rate) in vineyards. The best models (lowest AICc and \triangle AICc \leq 2) are indicated in boldface type. R² refers to the best models. For clarity, only the most parsimonious (i.e., lowest AICc) of all possible models are presented. Random efect indicates whether the individual vineyard was included or not.

Soil-dwelling springtail species richness was affected by plant biomass and soil quality ($R^2_{adj} = 0.32$; Table 2; Fig. 2a). The effect of plant biomass on springtail species richness was low with a slight decrease of springtail species numbers with increasing plant biomass. Soil quality efect on springtail species numbers was very small, however, a positive efect was seen along an increasing gradient of soil quality.

Soil-dwelling springtail density was best explained by soil management and plant biomass ($R^2_{\text{adj}} = NA$; Table 2; Fig. 2b). Springtail density was noticeable higher in vineyards with PMD, compared to PGC. Under PGC springtail density decreased with increasing plant biomass, while under PMD springtail density increased with increasing plant biomass. The dominant morphological trait value in the springtail community (CWM) was best explained by plant biomass and plant biomass interacting with management and soil quality (R_m^2 = 0.15; Table 2). In vineyards with PGC the CWM was higher compared to vineyards under PMD and increased more than double when plant biomass increased (Fig. 2c). Under PMD, CWM increased almost twofold with increasing plant biomass. There was a strong interaction between plant biomass and the soil quality. On sites with low soil quality, plant biomass noticeably reduced springtail CWM, while in moderate soil qualities no relationship between CWM and plant biomass was seen. By contrast, on sites with high soil quality, CWM increased with increasing plant biomass.

Functional diversity of soil-dwelling springtails was afected by plant biomass, soil quality and their interaction (R_m^2 = 0.12; Table 2; Fig. 2d). Similarly to the response of springtail CWM to plant biomass and soil quality, plant biomass adversely efected springtail functional diversity when soil quality was low. However, at sites with high soil quality, functional diversity of springtails increased notably along a gradient of increasing plant biomass. When soil quality was moderate, plant biomass showed a slighty negative efect on functional diversity

Surface-dwelling springtails species richness was best explained by plant biomass and soil management $(R^2_{adi.}=0.43;$ Table 2; Fig. 3a). Under PGC, plant biomass had a small effect on springtail species richness. However, increasing plant biomass had a negative efect on springtail species richness. Soil management efect was negligible when plant biomass was low. Nevertheless, the efect of plant biomass was higher under PMD and increased the richness of surface springtails noticeably, with high plant biomass.

Surface-dwelling springtail activity was affected by plant biomass and management (R_{m}^{2} = NA; Table 2; Fig. 3b). Similar to the results for surface-dwelling springtail species richness, the efect of plant biomass difered noticeably between the management regimes. While density was higher under PGC than under PMD, when plant biomass was low, increasing plant biomass decreased springtail activity under PGC considerably, whereas the response was positive under PMD.

Surface springtail CWM was affected by plant biomass, soil quality, the their interactions (R^2 _m = 0.39; Table 2, Fig. 3c). The CWM was similar across soil quality levels when there was little plant biomass in the vineyard interrow. While CWM increased with plant biomass in sites with moderate soil quality, the efect size was almost twofold on sites with low soil quality. By contrast, on sites with high soil quality increasing plant biomass reduced the CWM.

Functional diversity of surface-dwelling springtails was also infuenced by soil quality and mean patch size in the surrounding landscape (MPS; R_m^2 = 0.30, Table 2; Fig. 3d). Functional diversity increased with increasing

MPS, however, the efect was marginal. Also, the efect of soil quality on functional diversity surface-dwelling springtails was small.

Litter decomposition. Litter decomposition rate (k) was 0.017 ± 0.009 for PMD and 0.016 ± 0.007 for PGC, stabilisation factor (S) was 0.442 ± 0.067 for PMD and 0.454 ± 0.072 for PGC. Decomposition rate was influenced by plant biomass, soil management and soil quality (R_m^2 = 0.06; Table 1). Stabilization factor was influenced by plant biomass, management, soil quality and their interactions (R_m^2 = 0.06; Table 1). However, R^2 obtained with these models were very low.

Discussion

Overall, our results showed that site factors such as plant biomass and soil quality were more decisive for earthworms and springtails than inter-row soil management. The surrounding landscape structure appeared to affect only the functional diversity of surface dwelling springtails but had no impact on earthworms or soil-dwelling springtails. Litter decomposition remained little afected by the investigated factors and was not related to earthworm or springtail parameters.

We observed a higher species richness of endogeic earthworms in periodically mechanically disturbed (PMD), compared to permanent green covered (PGC) inter-rows, while anecic species number remained unafected. Tis is in contrast to other studies showing sensitivity of earthworm species richness to mechanical vineyard inter-row management $8-10$. However, the unresponsiveness of anecic species numbers is in line with reports showing a higher tolerance to soil tillage, compared to endogeic species because of a better ability to escape from disturbance through their vertical burrows8. We interpret our findings as follows. First, earthworm populations could have recovered quickly after the last tillage event in the PMD vineyards⁹ as in our study, the last tillage events in the PMD plots had occurred 1–5 years before sampling. Second, we ofen found an interaction between plant parameters and earthworm species numbers indicating that plants established afer tillage compensated for the detrimental mechanical efects. Indeed, plant diversity has been shown to correlate positively with earthworm

Table 2. Comparison of alternative models (using AICc) for springtails in vineyards (richness...spp. m⁻², density...ind. m[−]² , CWM...community weighted morphological trait value, FD...functional diversity Rao's quadratic entropy). The best models (lowest AICc and $\Delta AICc \leq 2$) are indicated in boldface type. R² refer to the best models. For clarity, only the most parsimonious (i.e., lowest AICc) of all possible models are presented. Random efect indicates whether the individual vineyard was included or not. CWM… community weighted morphological trait value, FD…functional diversity.

biomass and density^{5,53,54}. Third, tilling in our vineyards is performed at an average depth of 17 cm mainly during dry soil conditions when earthworms might be active in deeper soil horizons and therefore not affected. The lack of epigeic earthworm species in our vineyards refects the sensitivity of this group to frequent disturbances (e.g., pruning activities, tractor traffic during pesticide applications) taking place in vineyards during the whole season^{8,11,55}. Therefore, we conclude, that potential negative effects of soil tillage on earthworms appeared to be levelled out in the next season provided that inter-rows are then covered with vegetation.

Across treatments, earthworm species richness, density and biomass, as well as anecic species richness was positively affected by soil quality underlining their role as bioindicators for soil quality⁵⁶. It was interesting to see that in sites with high soil quality the density of anecic earthworms decreased with increasing plant biomass as this ecological group usually feeds on plant litter at the soil surface⁵⁷. As earthworms differentially respond to plant species this fnding suggests that plant nutritional quality at high soil quality with more grasses and less legumes, was not as attractive for earthworms as under lower soil quality^{5,7,54,51}

The finding that densities of soil dwelling springtails were lower in the soils of PGC compared to PMD vineyards concurs with other studies conducted in vineyards¹³. Because in the current study both management regimes had plant cover during sampling, we assume that these efects are mainly due to soil compaction rather than habitat disturbance⁵⁹ or the lacking vegetation cover under $PMD¹³$. This interpretation is supported by the results of CWM in the soil samples, which were lower in PMD vineyards. The CWM value indicates which life-form dominates a springtail community. Small springtails with reduced pigmentation and furca, absence of ocelli and small antennae, indicated by a small morphological trait value, are more adapted to life in soil where it is difficult to disperse. In contrast larger and pigmented species, with eyes, longer antennae and fully developed furca are better adapted to favourable conditions for dispersal on the soil surface^{60,61}. Hence, our results suggest that PGC management provided a less favourable habitat for larger euedaphic species. In support it was reported that springtail life-form types difer in their response to tillage depending on soil texture, i.e., soil dwelling species were harmed by tillage in loamy soils, while surface dwelling species remained relatively unaffected⁵⁹. Our fndings for a 54%-increased activity density and a 3.5%-increased species richness of surface dwelling springtails under PMD compared to PGC seemed to disagree with those studies. A likely explanation for these fndings can be found in the interaction between tillage and plant biomass. The activity of surface dwelling springtails increased under PMD and decreased under PGC when plant biomass increased. There are two possible explanations for this outcome. First, it could be a consequence of the manifold functions of plant biomass which could serve as a source of nutrients, creating a favourable microclimate and function as a refuge against predation for edaphic springtails13,62. Contrary, plant cover also hinders rapid movement of springtails increasing the likelihood of capture63. Predators like spiders and carabids also beneft from vegetation cover in vineyard inter-rows64,65. It is

Figure 2. Soil springtail species numbers (**a**), density (**b**), community weighted morphological trait value (CWM, (**c**) and functional diversity (**d**) in response to plant biomass, soil quality and inter-row soil management (PGC…permanent green cover, PMD…periodical mechanical disturbance. O…PGC, Δ…PMD, $N=4$.

possible that there was a higher presence of predators in PGC, compared to PMD, where every second row represented a disturbed habitat. However, we did not evaluate the direct relationship between springtails and predators in the current study. Second, in our case the relation between springtail activity, tillage and plant biomass might refect the microclimate created by the litter layer in PGC, which altered the aboveground springtail community composition and density66,67. Additionally, plant biomass promoted larger surface-dwelling species in sites with low and moderate soil quality, while in vineyards of high soil quality, plant biomass production promoted smaller soil-dwelling species. In this context, soil-dwelling species afect the soil organic matter decomposition, whereas surface-dwelling species also feed on aboveground plant material and afect the dynamics of the frst stages of litter decomposition68. Interestingly, CWM and functional diversity of soil dwelling community showed the opposite response to soil quality and plant biomass. In high-quality soils, plant biomass increased springtail functional diversity, probably due to enhanced food resources available for springtails⁶⁶. Species richness of soil dwelling springtails was only marginally afected by site conditions and plant biomass, a fnding which is in line with the results from other vineyards¹³ and arable fields²³.

The only soil biota parameter influenced by the surrounding landscape was the functional diversity of surface dwelling springtails. Springtail functional diversity increased with increasing mean patch size (MPS), where a high MPS indicates low fragmentation and low heterogeneity of the landscape. This is in contrast with the general assumption that heterogeneous landscapes positively affect springtail density and species richness^{52,69}. Reasons for these contrasting patterns could be that diferent landscape matrices and land use histories were studied: agro-forest mosaics^{31,49} versus viticultural landscapes in the current study. Another reason could be that the very small-structured viticultural landscapes of the study regions precluded the build-up of functionally diverse springtail communities or the 2-m high grapevine trellis negatively affected springtail dispersal. The absence of landscape efects on earthworms suggests little connection between non-crop habitats and vineyards. However, to

draw a more general conclusion on the infuence of viticultural landscapes on earthworms more detailed analyses including earthworm tracking would be necessary.

Litter decomposition was unafected by soil management and was only weakly predicted by the other site and landscape parameter we assessed in this study. Because the mesh size of the decomposition bags excluded larger springtails and all earthworms the contribution of smaller springtails to litter decomposition appeared to be marginal. Since vineyards are always treated with several pesticides a lack of response of decomposition could also mean that pesticide efects on non-target soil biota overrode potential tillage or landscape efects on litter decomposition^{38,70,71}. Clearly, it would need more specific experimental approaches to disentangle underlying interrelationships.

In conclusion, our study suggests that response patterns of soil biota to soil management in vineyards difer from those known from arable agroecosystems, where negative efects of intensifcation on soil biota are common72. Earthworms and springtails in vineyard inter-rows difered in their sensitivity to soil disturbance and were modifed by plant biomass and soil quality. While the size and diversity of earthworm populations were little afected by soil tillage, mainly larger-sized surface and soil dwelling springtails were reduced in vineyards with permanent green cover. Our results suggest that efects of inter-row tillage on the diversity and functionality of soil biota could be compensated by establishing a plant cover during the rest of the year. We conclude that soil biota in vineyards situated in heterogeneous landscapes are mainly infuenced by site and management factors.

Methods

Site description and study design. The study sites are located in neighbouring wine regions Carnuntum (48° 4′N, 16° 47′E, province of Lower Austria) and Leithaberg (47° 54′N, 16° 41′E, province of Burgenland), about 40km south east of Vienna, Austria. In these regions, wine cultivation has a long tradition dating back to the Roman Empire in the $1st$ century B.C. Typically, vineyards in these regions are small $(0.4-1)$ ha) belonging to numerous wine growers.

Vineyards are interspersed within a more or less heterogeneous landscape adjoined by arable crops and several noncrop landscape elements such as fallows, forests or hedges. The climate in the regions is continental with annual average temperature of 11.5°C and annual rainfall of 530mm (between 1990 and 2004; weather data for the city of Bruck/Leitha which is located in the centre of the study region). In the study region the most frequent soil types are Chernozems and Cambisols. Vineyards are cropped with red or white grape varieties cultivated in a vertical trellising system with within-row grapevine distance of 0.75–1.30m and inter-row distance varying between 2.15–3.00m.

Within these regions we selected 16 commercial vineyards along an altitudinal range of 120–260 m a.s.l. In 8 vineyards the inter-rows had permanent green cover (PGC) and in 8 vineyards every second inter-row was periodically mechanically disturbed (PMD). Both management regimes were practiced for at least 5 years. Management data was collected with a questionnaire among the participating wine growers (Supplementary Table S3). Inter-row vegetation originates from a cover crop seed mixture or from the spontaneous vegetation. Vegetation was mulched 1 to 5 times per year. Application of organic (shredded pruning material, compost, green manure) or inorganic nitrogen-phosphorous-potassium-fertilizers, was done according to good wine-growers practice. Among the wine-growers, three were certifed organic farmers; the remaining vineyards were managed conventionally. As inter-row soil management does not difer between conventional or organic vineyards, we included both farming practices.

At each vineyard, we selected one inter-row sampling area (approximately 2×40 m) in the middle of the vineyard; this area had at least 5m distance to the vineyard edge. In PMD sites, samples were taken in the inter-rows that had been cultivated in the previous year.

Sampling and measurements. *Earthworm sampling*. From April 13–21, 2015 we sampled earthworm communities by hand-sorting of 6 soil blocks ($20 \times 20 \times 25$ cm, $L \times W \times H$) along a transect in the middle of the vineyard inter-row; distance between excavated soil blocks was 5–7m. Earthworms were conserved in 4% formaldehyde and identifed at the Station Biologique of the University of Rennes, Paimpont, France.

In the laboratory, earthworms were identifed to the species level, counted, weighed, and categorized into the ecological groups epigeics, anecics and endogeics according to Bouché^{73,74}.

Springtail sampling. Soil dwelling springtail communities were sampled by taking 4 soil cores ($5.5 \times 5.5 \times 10$ cm, $L \times W \times H$) between April 13-21, 2015 in about 50 cm distance of the installed pitfall traps at each vineyard site. Soil samples were immediately sealed in plastic bags and transported in a cooling box to the laboratory for extraction using a Berlese-Tullgren extractor for 4 days. In the extractor, the animals were collected in plastic jars flled with 10% benzoic acid solution. Afer extraction, samples were washed with tap water in a 0.063mm sieve in order to clear from adherent soil particles and then stored in 80% ethanol alcohol. Springtails were sorted and determined to species level, except of *Mesaphorura* sp, which were determined to genus level only. Identifcation of springtail was done using several determination keys⁷⁵⁻⁸³.

Surface dwelling springtail species were collected using six pitfall traps (diameter 17 mm, length 60 mm) installed on April 21, 2015 at each vineyard site⁸⁴. Traps were inserted with about 6 m distance to each other along a transect in the middle of inter-rows and flled with ethylene glycol and a drop of odourless detergent. Afer an exposure of 13 days in the feld, the traps were removed, sorted for springtails and determined to species level and counted. The number of trapped springtails during the sampling period is referred to as activity density.

Additionally to the taxonomic identification, springtails were assigned morphological trait values^{61,85}. The trait value attributes a measurement or a score to each of the following traits: number of ocelli, body size, body pigmentation level and pattern, presence of modifed hairs or scales, furca development and antennae length (Supplementary Table S4). Traits were extracted from literature used for identifcation. For body size we chose the maximum value found in the most recent literature. When there was more than one specifcation for an attribute, we used the dominant expression of the respective species in our samples as reference (e.g. *Proisotoma notabilis* can have 2+2 to 5+5 ommatidia (compound eyes)⁸⁰, however we counted $4+4$ ommatidia on the individuals in our samples).

We used the number of species and densities as references to taxonomic diversity and calculated the community weighted morphological trait value (CWM) 85 , as an index of the dominant trait value in the springtail community. Further we calculated the springtail functional diversity (FD) by computing the Rao's quadratic entropy using the "FD" package⁸⁶ in the software R^{87} .

Litter decomposition. Litter decomposition was assessed using the teabag index (TBI)⁸⁸. Therefore, we buried 10 pairs of green tea and rooibos teabags (EAN: 87 22700 05552 5 and EAN: 87 22700 18843 8, for green and rooibos tea, respectively; Lipton Tea, Washington St, USA) between April 13–21, 2015. Teabags were inserted at a soil depth of 8 cm along a transect in the middle of the inter-row; distance between teabags was about 5–7m. The mesh size of the tea bags of 0.25 mm allows micro-, and mesofauna to enter, but excludes macrofauna⁸⁹. After approx. 90 days the teabags were recovered, dried, and weighed. About 15% of the teabags were destroyed by mice or farming machinery during exposition in the field. The TBI consists of the decomposition rate (k) and the stabilisation factor (S); k refers to the frst phase of decomposition; S accounts for the labile fraction of the material which is not decomposed, but stabilized. We used the average S value for calculating k.

Vegetation and soil quality parameters. Vegetation cover of each vascular plant species was measured once in April and June 2015 on four 1×1 m plots located in the centre of each vineyard inter-row. Cover percentage of each vascular plant species was estimated following the scale of Londo⁹⁰. For statistical analysis, we calculated the plant cover percentages of legumes, grasses and herbs and the Shannon diversity index⁹¹. In May, aboveground plant biomass production was assessed by cutting all vegetation on four 1×0.5 m plots, drying at 60 °C for 48hours and weighing the dry matter (Table 3).

Table 3. Overview of site, vegetation and landscape variables used for the statistical analyses. Means \pm SD. n.a. not applicable.

As a proxy of soil quality in the vineyards we used the Austrian soil feld index in our analysis (Table 3). Tis soil feld index is assigned to all agriculturally used parcels of land and is used to calculate agricultural land taxes in Austria. It represents the natural yield capacity of a feld in relation to the highest yielding capacity of the country; values from 0 to 100 points, where 100 points stands for the highest yield capacity. The soil index integrates soil type, humus content, soil depth, soil texture, bulk density, soil structure, lime content, gleying and soil congregation as well as climatic and hydrologial properties and the slope of the site 92 .

Landscape parameters. Field mapping was done in July 2015. As a reference we used the official Austrian land utilization map from 2012 (Integriertes Verwaltungs- und Kontrollsystem, INVEKOS). Landscape elements within a 750 m-radius were classifed into habitat types according to CORINE Land Cover and EUNIS Habitat Classification^{93,94}. Habitat types were then clustered into two levels of classification. The first level classification is composed by semi-natural elements (SNE: hedges, tree rows, grass stripes, natural grassland, pasture, fallow, heathland, wetland and woodlots), viticulture, other agricultural land (annual crops), open land, wood, water entities, and artificial/constructed entities (urban area, buildings and roads). The more detailed landscape classifcation was used for calculating the Shannon diversity Index which represents landscape composition, the mean patch size (MPS), which respresents landscape confguration and the distances to closest SNE and crop. Mapping and analysis was done using the programs ArcGis 10.2.1⁹⁵, QGis 2.8.1, FRAGSTATS 4.2⁹⁶ and CHLOE2012⁹⁷.

Statistical analysis. For earthworm analyses data from all 96 soil excavations were used; for collembolan analyses we used the data from 64 soil cores (i.e. 4 soil cores for each vineyard) and 89 pitfall traps (usually 6 traps per vineyard, except 7 traps that were flled with soil during exposure in the feld).

Because of Poisson error distributions of the response variables, we ftted generalized linear mixed models (GLMM) for our nested experimental design to analyse the efects of management and soil quality on earthworms, collembolan and decomposition and how these efects are altered by site and landscape variables. Vineyard was used as a random factor in the models. Further, we used model selection as an alternative to traditional hypothesis testing^{64,98}. Prior to fitting GLMMs we excluded several explanatory variables following a selection procedure based on scatterplots between responsive and explanatory variables: first we selected the site and landscape variables correlating highest $(r > 0.18)$ with the response variables. Provided there was linear correlation $(r>0.3)$ with vineyard management or soil quality, we removed these variables from the analysis (e.g., farming system organic or conventional). Remaining variables were then tested on co-linearity with subsequent exclusion of one of the correlating variables from the analysis. As a result, plot parameters (plant richness, plant cover, grass, herb cover) and landscape parameters (SNE, viticulture, other agriculture, open land, wood, water entities, artifcial/constructed entities, distances to next crop and Shannon landscape diversity) were excluded from the GLMM analysis. Aferwards, we built two sets of models (in total 46 models), one with management and one with soil quality as fxed factors with the selected plant (plant biomass, legume cover) and landscape variables (MPS, distance to next SNE) as covariables and vineyard as random efect variable. To be able to test for effects of different soil quality levels we considered plots with a field index < 23 as low, plots with a field index > 67 as high; the remaining plots were considered with medium soil quality.

Response variables were square-rooted or log transformed to account for non-normal distribution when necessary. The analyses were performed using the "lme4" package⁹⁹ and the "glmmADMB" package¹⁰⁰ in the software R^{87} .

For model selection we used the Akaike Information Criterion corrected for small sample sizes (AIC_c), a measure of the quality of a statistical model. The most parsimonious model was considered the best model to describe the significant effect on the response variable. Models where AIC_c was less than 2 units above the AIC_c min were considered identical good fts as the best model. In such a case the most complex model (number of variables, interaction, random factor) was plotted and used for further interpretation. In plots containing two continuous variables, soil biota response was plotted against one variable at the 5%, 50% and 95% levels of the second variable. Model assumptions were verified using diagnostic plots and the Package "DHARMa"¹⁰¹

We calculated the R^2 for the best models to account for the variability explained by mixed models with Gaussian or Poisson error distributions¹⁰². Though this approach cannot be applied to models with a negative binomial error distribution, it allows for two components of R^2 to be calculated: (i) a marginal $R^2 (R^2_m)$ that only includes the variance explained by fixed effects; and (ii) a conditional $R^2 (R^2)$ that represents the variance explained by both fixed and random effects. In case of two or more best fitting models, $R_{\rm m}^2$ and $R_{\rm c}^2$ were calculated for the more complex model. For models without random effects we calculated the adjusted \mathbb{R}^2 .

References

- 1. Winkler, K. J., Viers, J. H. & Nicholas, K. A. Assessing ecosystem services and multifunctionality for vineyard systems. *Front. Environ. Sci.* **5**, [https://doi.org/10.3389/fenvs.2017.00015](http://dx.doi.org/https://doi.org/10.3389/fenvs.2017.00015) (2017).
- 2. Jackson, R. *Wine Science: Principles and Applications*. 4th ed edn, 978 pp (Academic Press, 2014).
- 3. Bauer, K., Fox, R. & Ziegler, B. *Moderne Bodenpfege im Weinbau*. (Oesterreichischer Agrarverlag and Eugen Ulmer Verlag, 2004).
- 4. Brussaard, L., de Ruiter, P. C. & Brown, G. G. Soil biodiversity for agricultural sustainability. *Agric. Ecosyst. Environ.* **121**, 233–244 (2007).
- 5. Zaller, J. G. & Arnone, J. A. Earthworm responses to plant species' loss and elevated CO₂ in calcareous grassland. *Plant Soil* 208, 1–8 (1999).
- 6. Milcu, A., Partsch, S., Langel, R. & Scheu, S. The response of decomposers (earthworms, springtails and microorganisms) to variations in species and functional group diversity of plants. *Oikos* **112**, 513–524 (2006).
- 7. Paoletti, M. G. *et al*. Earthworms as useful bioindicators of agroecosystem sustainability in orchards and vineyards with diferent inputs. *Appl. Soil Ecol.* **10**, 137–150, [https://doi.org/10.1016/S0929-1393\(98\)00036-5](http://dx.doi.org/10.1016/S0929-1393(98)00036-5) (1998).
- 8. Schreck, E., Gontier, L., Dumat, C. & Geret, F. Ecological and physiological efects of soil management practices on earthworm communities in French vineyards. *Europ. J. Soil Biol.* **52**, 8–15, [https://doi.org/10.1016/j.ejsobi.2012.05.002](http://dx.doi.org/10.1016/j.ejsobi.2012.05.002) (2012).
- 9. Virto, I. *et al*. Soil quality evaluation following the implementation of permanent cover crops in semi-arid vineyards. Organic matter, physical and biological soil properties. *Span. J. Agricult. Res.* **10**, 1121–1132, [https://doi.org/10.5424/sjar/2012104-613-11](http://dx.doi.org/10.5424/sjar/2012104-613-11) (2012).
- 10. Vrsic, S. Soil erosion and earthworm population responses to soil management systems in steep-slope vineyards. *Plant Soil Environ.* **57**, 258–263 (2011).
- 11. Faber, F., Wachter, E. & Zaller, J. G. Earthworms are little afected by reduced soil tillage methods in vineyards. *Plant Soil Environ.* **63**, 257–263 (2017).
- 12. Sturm, M., Sturm, M. & Eisenbeis, G. Recovery of the biological activity in a vineyard soil afer landscape redesign: A three-year study using the bait-lamina method. *Vitis* **41**, 43–45 (2002).
- 13. Renaud, A., Poinsot-Balaguer, N., Cortet, J. & Le Petit, J. Infuence of four soil maintenance practices on Collembola communities in a Mediterranean vineyard. *Pedobiologia* **48**, 623–630 (2004).
- 14. Zehetner, F. *et al*. Soil organic carbon and microbial communities respond to vineyard management. *Soil Use Managem.* **31**, 528–533 (2015).
- 15. Castañeda, L. E. & Barbosa, O. Metagenomic analysis exploring taxonomic and functional diversity of soil microbial communities in Chilean vineyards and surrounding native forests. *PeerJ* **5**, e3098, [https://doi.org/10.7717/peerj.3098](http://dx.doi.org/10.7717/peerj.3098) (2017).
- 16. Lavelle, P. Faunal activities and soil processes: Adaptive strategies that determine ecosystem function. *Adv. Ecol. Res.* **27**, 93–132 (1997).
- 17. Briones, M. J. I. & Schmidt, O. Conventional tillage decreases the abundance and biomass of earthworms and alters their community structure in a global meta-analysis. *Global Change Biol*. **23**, 4396–4419, [https://doi.org/10.1111/gcb.13744](http://dx.doi.org/https://doi.org/10.1111/gcb.13744) (2017).
- 18. Coll, P., Le Cadre, E., Blanchart, E., Hinsinger, P. & Villenave, C. Organic viticulture and soil quality: A long-term study in Southern France. *Appl. Soil Ecol.* **50**, 37–44, [https://doi.org/10.1016/j.apsoil.2011.07.013](http://dx.doi.org/10.1016/j.apsoil.2011.07.013) (2011).
- 19. Fründ, H.-C., Graefe, U. & Tischer, S. In *Biology of Earthworms* Vol. 24 (ed Ayten Karaca) 19–39 (Springer Verlag, 2011).
- 20. Peres, G., Piron, D., Bellido, A., Goater, C. & Cluzeau, D. Earthworms used as indicators of agricultural managements. *Fresenius Environ Bull* **17**, 1181–1189 (2008).
- 21. Bouché, M. B. In *Soil O*rganis*ms* as C*omponents of Ecosystems* Vol. Ecological Bulletins 25, Stockholm (eds U Lohm & T Persson) 122–133 (1977).
- 22. Alvarez, T., Frampton, G. K. & Goulson, D. Epigeic Collembola in winter wheat under organic, integrated and conventional farm management regimes. *Agric Ecosyst Environ* **83**, 95–110, [https://doi.org/10.1016/S0167-8809\(00\)00195-X](http://dx.doi.org/10.1016/S0167-8809(00)00195-X) (2001).
- 23. Brennan, A., Fortune, T. & Bolger, T. Collembola abundances and assemblage structures in conventionally tilled and conservation tillage arable systems. *Pedobiologia* **50**, 135–145, [https://doi.org/10.1016/j.pedobi.2005.09.004](http://dx.doi.org/10.1016/j.pedobi.2005.09.004) (2006).
- 24. Filser, J., Fromm, H., Nagel, R. F. & Winter, K. Efects of previous intensive agricultural management on microorganisms and the biodiversity of soil fauna. *Plant Soil* **170**, 123–129, [https://doi.org/10.1007/BF02183060](http://dx.doi.org/10.1007/BF02183060) (1995).
- 25. Gardi, C., Tomaselli, M., Parisi, V., Petraglia, A. & Santini, C. Soil quality indicators and biodiversity in northern Italian permanent grasslands. *Europ. J. Soil Biol.* **38**, 103–110, [https://doi.org/10.1016/S1164-5563\(01\)01111-6](http://dx.doi.org/10.1016/S1164-5563(01)01111-6) (2002).
- Heisler, C. & Kaiser, E. A. Influence of agricultural traffic and crop management on collembola and microbial biomass in arable soil. *Biol. Fertil. Soils* **19**, 159–165, [https://doi.org/10.1007/BF00336153](http://dx.doi.org/10.1007/BF00336153) (1995).
- 27. Rebek, E. J., Hogg, D. B. & Young, D. K. Efect of four cropping systems on the abundance and diversity of epedaphic springtails (Hexapoda: Parainsecta: Collembola) in Southern Wisconsin. *Environm Entomol* **31**, 37–46, [https://doi.org/10.1603/0046-](http://dx.doi.org/https://doi.org/10.1603/0046-225X-31.1.37) [225X-31.1.37](http://dx.doi.org/https://doi.org/10.1603/0046-225X-31.1.37) (2002).
- 28. Rusek, J. Biodiversity of Collembola and their functional role in the ecosystem. *Biodiv. Conserv.* **7**, 1207–1219, [https://doi.](http://dx.doi.org/https://doi.org/10.1023/A:1008887817883) [org/10.1023/A:1008887817883](http://dx.doi.org/https://doi.org/10.1023/A:1008887817883) (1998).
- 29. Filser, J. The role of Collembola in carbon and nitrogen cycling in soil. *Pedobiologia* 46, 234-245, [https://doi.org/10.1078/0031-](http://dx.doi.org/10.1078/0031-4056-00130) [4056-00130](http://dx.doi.org/10.1078/0031-4056-00130) (2002).
- 30. Chauvat, M. & Wolters, V. & Dauber, J. Response of collembolan communities to land-use change and grassland succession. *Ecography* **30**, 183–192, [https://doi.org/10.1111/j.2007.0906-7590.04888.x](http://dx.doi.org/10.1111/j.2007.0906-7590.04888.x) (2007).
- 31. Martins da Silva, P., Berg, M. P., Serrano, A. R. M., Dubs, F. & Sousa, J. P. Environmental factors at diferent spatial scales governing soil fauna community patterns in fragmented forests. *Landscape Ecol* **27**, 1337–1349, [https://doi.org/10.1007/s10980-012-9788-2](http://dx.doi.org/https://doi.org/10.1007/s10980-012-9788-2) (2012). 32. Hopkin, S. P. *Biology of the springtails (Insecta: Collembola)*. (Oxford University Press, 1997).
- 33. Bardgett, R. D. & der Putten, V. W. H. Belowground biodiversity and ecosystem functioning. *Nature 515* **505**, 505–511, [https://doi.](http://dx.doi.org/10.1038/nature13855) [org/10.1038/nature13855](http://dx.doi.org/10.1038/nature13855) (2014).
- 34. Gange, A. C. & Bower, E. In *Multitrophic interactions in terrestrial systems* (eds A. C. Gange & V. K. Brown) pp. 115–132 (Blackwell, 1997). 35. Gisin, H. *Ökologie und Lebensgemeinschafen der Collembolen im schweizerischen Excursionsgebiet Basels*. Bd. 50 edn, (Revue Suisse
- de Zoologie, 1943). 36. Hättenschwiler, S., Tiunov, A. & Scheu, S. Biodiversity and litter deomposition in terrestrial ecosystems. *Ann. Rev. Ecol. Evol.* **36**, 191–218, [https://doi.org/10.1146/annurev.ecolsys.36.112904.151932](http://dx.doi.org/https://doi.org/10.1146/annurev.ecolsys.36.112904.151932) (2005).
- Wardle, D. A. Impacts of disturbance on detritus food webs in agro-ecosystems of contrasting tillage and weed management practices. *Adv. Ecol. Res.* **26**, 105–185, [https://doi.org/10.1016/S0065-2504\(08\)60065-3](http://dx.doi.org/https://doi.org/10.1016/S0065-2504(08)60065-3) (1995).
- 38. Zaller, J. G. *et al*. Pesticide seed dressings can afect the activity of various soil organisms and reduce decomposition of plant material. *BMC Ecol*. **16**, [https://doi.org/10.1186/s12898-016-0092-x](http://dx.doi.org/10.1186/s12898-016-0092-x) (2016).
- 39. Gaupp-Berghausen, M., Hofer, M., Rewald, B. & Zaller, J. G. Glyphosate-based herbicides reduce the activity and reproduction of earthworms and lead to increased soil nutrient concentrations. *Sci. Rep.* **5**, 12886, doi:12810.11038/srep12886 (2015).
- 40. Isaia, M., Bona, F. & Badino, G. Infuence of landscape diversity and agricultural practices on spider assemblage in Italian vineyards of Langa Astigiana (Northwest Italy). *Environm Entomol.* **35**, 297–307 (2006).
- 41. Gillespie, M. & Wratten, S. D. Te importance of viticultural landscape features and ecosystem service enhancement for native butterfies in New Zealand vineyards. *J Insect Conserv* **16**, 13–23, [https://doi.org/10.1007/s10841-011-9390-y](http://dx.doi.org/10.1007/s10841-011-9390-y) (2012).
- 42. Kelly, R. M., Kitzes, J., Wilson, H. & Merenlender, A. Diversity promotes bat activity in a vineyard landscape. *Agric Ecosyst Environ* **223**, 175–181, [https://doi.org/10.1016/j.agee.2016.03.010](http://dx.doi.org/10.1016/j.agee.2016.03.010) (2016).
- 43. Nascimbene, J., Zottini, M., Ivan, D., Casagrande, V. & Marini, L. Do vineyards in contrasting landscapes contribute to conserve plant species of dry calcareous grasslands? *Sci. Tot. Environ.* **545–546**, 244–249, [https://doi.org/10.1016/j.scitotenv.2015.12.051](http://dx.doi.org/https://doi.org/10.1016/j.scitotenv.2015.12.051) (2016).
- 44. Saatkamp, A. *et al*. Decrease of Plant Alpha and Beta-Diversity with Management Intensity in Vineyards and the Infuence of Landscape Context. *Ecologia Mediterranea* **40**, 17–27 (2014).
- Tanadini, M., Schmidt, B. R., Meier, P., Pellet, J. & Perrin, N. Maintenance of biodiversity in vineyard-dominated landscapes: a case study on larval salamanders. *Animal Conserv.* **15**, 136–141, [https://doi.org/10.1111/j.1469-1795.2011.00492.x](http://dx.doi.org/https://doi.org/10.1111/j.1469-1795.2011.00492.x) (2012).
- 46. Flohre, A., Rudnick, M., Traser, G., Tscharntke, T. & Eggers, T. Does soil biota beneft from organic farming in complex vs. simple landscapes? *Agric. Ecosyst. Environ.* **141**, 210–214 (2011).
- 47. Lindberg, N. & Bengtsson, J. Population responses of oribatid mites and collembolans afer drought. *Appl. Soil Ecol.* **28**, 163–174, [https://doi.org/10.1016/j.apsoil.2004.07.003](http://dx.doi.org/10.1016/j.apsoil.2004.07.003) (2005).
- 48. Martins da Silva, P. *et al*. Soil fauna through the landscape window: factors shaping surface-and soil-dwelling communities across spatial scales in cork-oak mosaics. *Landscape Ecol.* **30**, 1511–1526, [https://doi.org/10.1007/s10980-015-0206-4](http://dx.doi.org/https://doi.org/10.1007/s10980-015-0206-4) (2015).
- Ponge, J. F., Dubs, F., Gillet, S., Sousa, J. P. & Lavelle, P. Decreased biodiversity in soil springtail communities: the importance of dispersal and landuse history in heterogeneous landscapes. *Soil Biol. Biochem.* **38**, 1158–1161, [https://doi.org/10.1016/j.](http://dx.doi.org/10.1016/j.soilbio.2005.09.004) [soilbio.2005.09.004](http://dx.doi.org/10.1016/j.soilbio.2005.09.004) (2006).
- 50. Querner, P. *et al*. Landscape and site efects on Collembola diversity and abundance in winter oilseed rape felds in eastern Austria. *Agric. Ecosyst. Environ.* **164**, 145–154, [https://doi.org/10.1016/j.agee.2012.09.016](http://dx.doi.org/https://doi.org/10.1016/j.agee.2012.09.016) (2013).
- 51. Sousa, J. P. *et al*. Changes in Collembola richness and diversity along a gradient of land-use intensity: A pan European study. *Pedobiologia* **50**, 147–156, [https://doi.org/10.1016/j.pedobi.2005.10.005](http://dx.doi.org/10.1016/j.pedobi.2005.10.005) (2006).
- 52. Vanbergen, A. J. *et al*. Scale-specifc correlations between habitat heterogeneity and soil fauna diversity along a landscape structure gradient. *Oecologia* **153**, 713–725, [https://doi.org/10.1007/s00442-007-0766-3](http://dx.doi.org/10.1007/s00442-007-0766-3) (2007).
- 53. Arnone, J. A., Zaller, J. G., Hofer, G., Schmid, B. & Körner, C. Loss of plant biodiversity eliminates stimulatory efect of elevated CO2 on earthworm activity in grasslands. *Oecologia* **171**, 613–622, [https://doi.org/10.1007/s00442-012-2585-4](http://dx.doi.org/https://doi.org/10.1007/s00442-012-2585-4) (2013).
- 54. Eisenhauer, N. *et al*. Plant community impacts on the structure of earthworm communities depend on season and change with time. *Soil Biol. Biochem.* **41**, 2430–2443, [https://doi.org/10.1016/j.soilbio.2009.09.001](http://dx.doi.org/10.1016/j.soilbio.2009.09.001) (2009).
- 55. Paoletti, M. G. *et al*. Earthworms as useful bioindicators of agroecosystem sustainability in orchards and vineyards with diferent inputs. *Appl. Soil Ecol.* **10**, 137–150 (1998).
- 56. Paoletti, M. G. Te role of earthworms for assessment of sustainability and as bioindicators. *Agric. Ecosyst. Environ.* **74**, 137–155, [https://doi.org/10.1016/S0167-8809\(99\)00034-1](http://dx.doi.org/https://doi.org/10.1016/S0167-8809(99)00034-1) (1999).
- 57. Curry, J. P. & Schmidt, O. Te feeding ecology of earthworms A review. *Pedobiologia* **50**, 463–477, [https://doi.org/10.1016/j.](http://dx.doi.org/10.1016/j.pedobi.2006.09.001) [pedobi.2006.09.001](http://dx.doi.org/10.1016/j.pedobi.2006.09.001) (2007).
- 58. Arnone, J. A., Zaller, J. G., Hofer, G., Schmid, B. & Körner, C. Loss of plant biodiversity eliminates stimulatory efect of elevated CO2 on earthworm casting activity in grasslands. *Oecologia* **171**, 613–622 (2013).
- 59. van Capelle, C., Schrader, S. & Brunotte, J. Tillage-induced changes in the functional diversity of soil biota A review with a focus on German data. *Europ. J. Soil Biol.* **50**, 165–181, [https://doi.org/10.1016/j.ejsobi.2012.02.005](http://dx.doi.org/10.1016/j.ejsobi.2012.02.005) (2012).
- 60. Makkonen, M. *et al*. Traits explain the responses of a sub-arctic Collembola community to climate manipulation. *Soil Biol. Biochem.* **43**, 377–384, [https://doi.org/10.1016/j.soilbio.2010.11.004](http://dx.doi.org/10.1016/j.soilbio.2010.11.004) (2011).
- 61. Martins da Silva, P. *et al*. Traits of collembolan life-form indicate land use types and soil properties across an European transect. *Appl. Soil Ecol.* **97**, 69–77, [https://doi.org/10.1016/j.apsoil.2015.07.018](http://dx.doi.org/10.1016/j.apsoil.2015.07.018) (2015).
- 62. Favretto, M. R. *et al*. Invertebrates and nutrients in a Mediterranean vineyard mulched with subterranean clover (*Trifolium subterraneum* L.). *Biol. Fertil. Soils* **14**, 151–158, [https://doi.org/10.1007/BF00346055](http://dx.doi.org/10.1007/BF00346055) (1992).
- 63. Bauer, T. & Christian, E. Habitat dependent diferences in the fight behaviour of Collembola. *Pedobiologia* **30**, 233–239 (1987).
- 64. Paredes, D., Cayuela, L. & Campos, M. Synergistic efects of ground cover and adjacent vegetation on natural enemies of olive insect pests. *Agric. Ecosyst. Environ.* **173**, 72–80, [https://doi.org/10.1016/j.agee.2013.04.016](http://dx.doi.org/https://doi.org/10.1016/j.agee.2013.04.016) (2013).
- 65. Tomson, L. J. & Hofmann, A. A. Efects of ground cover (straw and compost) on the abundance of natural enemies and soil macro invertebrates in vineyards. *Agric. Forest Entomol.* **9**, 173–179, [https://doi.org/10.1111/j.1461-9563.2007.00322.x](http://dx.doi.org/https://doi.org/10.1111/j.1461-9563.2007.00322.x) (2007).
- 66. Eisenhauer, N., Sabais, A. C. W. & Scheu, S. Collembola species composition and diversity efects on ecosystem functioning vary with plant functional group identity. *Soil Biol. Biochem.* **43**, 1697–1704, [https://doi.org/10.1016/j.soilbio.2011.04.015](http://dx.doi.org/10.1016/j.soilbio.2011.04.015) (2011).
- 67. Salmon, S. & Ponge, J. F. Species traits and habitats in springtail communities: A regional scale study. *Pedobiologia* **55**, 295–301, [https://doi.org/10.1016/j.pedobi.2012.05.003](http://dx.doi.org/10.1016/j.pedobi.2012.05.003) (2012).
- 68. Potapov, A. A., Semenina, E. E., Korotkevich, A. Y., Kuznetsova, N. A. & Tiunov, A. V. Connecting taxonomy and ecology: Trophic niches of collembolans as related to taxonomic identity and life forms. *Soil Biol. Biochem.* **101**, 20–31, [https://doi.org/10.1016/j.](http://dx.doi.org/10.1016/j.soilbio.2016.07.002) [soilbio.2016.07.002](http://dx.doi.org/10.1016/j.soilbio.2016.07.002) (2016).
- 69. Chust, G., Pretus, J. L., Ducrot, D., Bedòs, A. & Deharveng, L. Response of soil fauna to landscape heterogeneity: determining optimal scales for biodiversity modeling. *Conserv. Biol.* **17**, 1712–1723, [https://doi.org/10.1111/j.1523-1739.2003.00564.x](http://dx.doi.org/https://doi.org/10.1111/j.1523-1739.2003.00564.x) (2003).
- 70. van Hoesel, W. *et al*. Single and combined efects of pesticide seed dressings and herbicides on earthworms, soil microorganisms, and litter decomposition. *Front. Plant Sci.* **8**, 215, [https://doi.org/10.3389/fpls.2017.00215](http://dx.doi.org/https://doi.org/10.3389/fpls.2017.00215) (2017).
- 71. Pelosi, C., Barot, S., Capowiez, Y., Hedde, M. & Vandenbulcke, F. Pesticides and earthworms. A review. *Agron. Sustain. Developm.* **34**, 199–228 (2014).
- 72. Tsiafouli, M. A. *et al*. Intensive agriculture reduces soil biodiversity across Europe. *Global Change Biol.* **21**, 973–985, [https://doi.](http://dx.doi.org/10.1111/gcb.12752) [org/10.1111/gcb.12752](http://dx.doi.org/10.1111/gcb.12752) (2015).
- 73. Bouché, M. B. Lombriciens de France. Ecologie et systématique. *INRA Publications 72-2. Institut National de la Recherche Agronomique, Paris, France* **72**, 671pp (1972).
- 74. Bouché, M. B. Strategies lombriciennes. *Ecological Bulletins*, 122–132 (1977).
- 75. Bretfeld, G. *Synopses on Palaearctic Collembola, vol. 2 Symphypleona*. (Staatliches Museum für Naturkunde, 1999).
- 76. Dunger, W. & Schlitt, B. *Synopses on Palaearctic Collembola, Volume 6, Part 1: Onychiuroidea: Tullbergiidae*. (State Museum of the Natural History Museum of Gorlitz, 2011).
- 77. Fjellberg, A. *Te Collembola of Fennoscandia and Denmark, Part II: Entomobryomorpha and Symphypleona*. (Brill, 2007).
- 78. Gisin, H. *Collembolenfauna Europas*. (Museum d'Histoire Naturelle, 1960).
- 79. Hopkin, S. P. *A Key to the Collembola (Springtails) of Britain and Ireland*. (Field Studies Council, 2007).
- 80. Potapow, M. S*ynopses on Palaearctic Collembola, Volume 3: Isotomidae*. (State Museum of the Natural History Museum of Gorlitz, 2001).
- 81. Stach, J. *Te Apterygotan Fauna of Poland in Relation to the World-Fauna of this Group of Insects. Tribe: Orchesellini*. (Polska Akademia Nauk, 1960).
- 82. Stach, J. *Te Apterygotan Fauna of Poland in Relation to the World-Fauna of this Group of Insects. Tribe: Entomobryini*. (Polska Akademia Nauk, 1963).
- 83. Tibaud, J. M., Schulz, H.-J. & da Gama Assalino, M. M. *Synopses on Palaearctic Collembola, vol. 4. Hypogastruridae*. (Staatliches Museum für Naturkunde, 2004).
- 84. Querner, P. & Bruckner, A. Combining pitfall traps and soil samples to collect Collembola for site scale biodiversity assessments. *Appl. Soil Ecol.* **45**, 293–297, [https://doi.org/10.1016/j.apsoil.2010.05.005](http://dx.doi.org/10.1016/j.apsoil.2010.05.005) (2010).
- 85. Vandewalle, M. *et al*. Functional traits as indicators of biodiversity response to land use changes across ecosystems and organisms. *Biodiv. Conserv.* **19**, 2921–2947, [https://doi.org/10.1007/s10531-010-9798-9](http://dx.doi.org/https://doi.org/10.1007/s10531-010-9798-9) (2010).
- 86. Laliberté, A. E., Legendre, P., Shipley, B. & Laliberté, M. E. FD: measuring functional diversity from multiple traits, and other tools for functional ecology. *R package 1.0–12*, 0–12, <http://cran.r-project.org/web/packages/FD/FD.pdf>(2014).
- 87. R Development Core Team. R: A Language and Environment for Statistical Computing Version 3.3.1. *R Foundation for Statistical Computing* **1**, 409–409 (2012).
- 88. Keuskamp, J. A., Dingemans, B. J. J., Lehtinen, T., Sarneel, J. M. & Hefing, M. M. Tea Bag Index: A novel approach to collect uniform decomposition data across ecosystems. *Methods Ecol. Evol.* **4**, 1070–1075, [https://doi.org/10.1111/2041-210X.12097](http://dx.doi.org/10.1111/2041-210X.12097) (2013).
- 89. Setälä, H., Marshall, V. G. & Trofymow, J. A. Infuence of body size of soil fauna on litter decomposition and 15N uptake by poplar in a pot trial. *Soil Biol. Biochem.* **28**, 1661–1675, [https://doi.org/10.1016/S0038-0717\(96\)00252-0](http://dx.doi.org/10.1016/S0038-0717(96)00252-0) (1996).
- 90. Londo, G. Te decimal scale for releves of permanent quadrats. *Vegetatio* **33**, 61–64, [https://doi.org/10.1007/BF00055300](http://dx.doi.org/10.1007/BF00055300) (1976).
- 91. Shannon, C. E. & Weaver, W. Te Mathematical Teory of Communication. *Te mathematical theory of communication* **27**, 117–117, [https://doi.org/10.2307/3611062](http://dx.doi.org/10.2307/3611062) (1949).
- 92. ÖBG. Bodenaufnahmesysteme in Österreich. *Mitt. Österr. Bodenkundl. Ges.* **62**, 219 pp (2001).
- 93. European Commission. CORINE land cover contents. *Methodology*, 1–94 (1994). 94. Davies, C. E., Moss, D. & Hill, M. O. EUNIS Habitat Classifcation Revised 2004. *European Topic Centre on Nature Protection and*
- *Biodiversity, Paris*, 310–310 (2004). 95. ESRI. *ArcGIS Desktop: Release 10.2*. (Environmental Systems Research Institute, 2013).
- 96. McGarigal, K., Cushman, S. A. & Ene, E. FRAGSTATSv4: Spatial Pattern Analysis Program for Categorical and Continuous Maps. Computer sofware program produced by the authors at the University of Massachusetts, Amherst. Available at the following web site:<http://www.umass.edu/landeco/research/fragstats/fragstats.html>(2012).
- 97. Boussard, H. & Baudry, J. *Documentation utilisateur pour le logiciel Chloe2012* (2014).
- 98. Johnson, J. B. & Omland, K. S. Model selection in ecology and evolution. *TREE* **19**, 101–108, [https://doi.org/10.1016/j.](http://dx.doi.org/10.1016/j.tree.2003.10.013) [tree.2003.10.013](http://dx.doi.org/10.1016/j.tree.2003.10.013) (2004).
- 99. Bates, D., Maechler, M., Bolker, B. & Walker, S. Packagelme4. *J Statist Sofw* **67**, 1–91, <http://lme4.r-forge.r-project.org> (2015).
- 100. Skaug, H., Fournier, D., Nielsen, A., Magnusson, A. & Bolker, B. Generalized Linear Mixed Models using 'AD Model Builder'. *R package ver. 0.8.4* (2012).
- 101. Hartig, F. Residual Diagnostics for Hierarchical (Multi-Level / Mixed) Regression Models Version 0.1.2 (2016).
- 102. Nakagawa, S. & Schielzeth, H. A general and simple method for obtaining R2 from generalized linear mixed-efects models. *Methods Ecol. Evol.* **2**, 133–142, [https://doi.org/10.1111/j.2041-210x.2012.00261.x](http://dx.doi.org/10.1111/j.2041-210x.2012.00261.x) (2013).

Acknowledgements

We are grateful to Johann Grassl and to the winegrowers in the Leithaberg and Carnuntum region for allowing us to run this study in their vineyards. Tanks to Edith Gruber, Friederike Hegselmann, Barbara Messerer, Thomas Müllner, Yoko Muraoka and Norbert Schuller for assistance during field sampling. Comments by Andrea Bryndum Buchholz, Rasmus Jonathan Buchholz, Georg Iwanov and three anonymous reviewers helped to improve the text. Tis work is part of the project VineDivers [\(www.vinedivers.eu](http://www.vinedivers.eu)) funded through the 2013–2014 BiodivERsA/FACCE JPI joint call with the national funders Bundesministerium für Bildung und Forschung (BMBF, Germany), Agence Nationale de la Recherche (ANR, France), Executive Agency for Higher Education, Research, Development and Innovation Funding (UEFISCDI, Romania), Ministerio des economía y competitividad (MINECO, Spain) and FWF-Der Wissenschaftsfonds (Austria).

Author Contributions

S.W., P.S., J.G.Z., D.C., M.G., M.P. and F.B. conceived and planned the experiment; J.B. and P.Q. identified Collembola samples; M.G., J.S. and D.C. identifed earthworm samples; S.K., M.G. and F.B. did the landscape mapping and analysis; T.B. and P.S. did the soil analyses; J.B. and D.P. statistically analysed the data; J.B., P.Q., D.P., T.B., P.S., M.G., S.K., S.W., M.P. and J.G.Z. wrote and reviewed the manuscript.

Additional Information

Supplementary information accompanies this paper at [https://doi.org/10.1038/s41598-017-17601-w.](https://doi.org/10.1038/s41598-017-17601-w)

Competing Interests: The authors declare that they have no competing interests.

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional afliations.

CO Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit [http://creativecommons.org/licenses/by/4.0/.](http://creativecommons.org/licenses/by/4.0/)

 $© The Author(s) 2017$