



Conservation Agriculture Enhances Soil Fauna Richness and Abundance in Low Input Systems: Examples From Kenya

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Conservation agriculture (CA) (zero tillage + organic inputs as surface residue) is believed to improve soil nutrient status, soil structure, control soil erosion, and also enhance soil fauna diversity. Despite the widespread interest in CA, empirical evidence of the benefits of CA on soil fauna diversity is limited, especially in low-input systems of sub-Saharan Africa (SSA). Consequently, the magnitude and effect by CA on soil fauna remains unquantified. The aim of this study was to evaluate the effect of CA and associated management practices on soil fauna richness and abundance. We hypothesized that CA and mixed cropping would positively influence soil fauna richness and abundance. We compared CA with conventional till (CT; with or without residues) in sole maize and maize-bean cropping systems. Soil macrofauna and mesofauna were sampled across the treatments in medium-term (6 years) trials in Embu, Central Kenya, and Kakamega (6 years) and a long-term trial in Nyabeda (15 years) using soil monoliths and core samplers, respectively. In agreement with our hypothesis, higher macrofauna taxonomic richness and mesofauna was recorded in CA than in CT without residues. This study demonstrated that: (1) medium to long-term addition of organic residues enhances soil fauna richness and abundance, (2) CA increases soil fauna taxonomic richness and abundance compared with CT, and (3) CA under maize-bean intercropping, rotation and sole maize cropping systems promote soil fauna richness and abundance compared with sole legume (common beans). We conclude that adoption of CA is important in enhancing richness of soil fauna. Given the numerous challenges faced by smallholder farmers of SSA in the adoption of CA, who in most cases rarely practice all the three CA principles simultaneously, we propose a further study that will determine the effects and interactions between each of the CA components on soil fauna richness and abundance.

Keywords: tillage, organic resources, soil fauna, richness, abundance

INTRODUCTION

Conservation agriculture (CA), which encompasses minimum soil disturbance, soil cover, and crop diversification as its three main principles [with a fourth principle on fertilizer application being recently suggested by Vanlauwe et al. (2014)], has been successfully promoted in different parts of the world such as the temperate zone and parts of Latin America (Lal, 2007; Wall, 2007; Derpsch et al., 2010). Despite its low adoption in sub-Saharan Africa (SSA), the rapidly changing demographics in the region including a transformation of its economies to middle class, and the ongoing large initiatives to promote CA present opportunities of increasing its adoption. As such, evidence is needed around the various sustainability indicators of CA including soil biological indices (Ayuke et al., 2011a,b; Paul et al., 2015). Indeed, CA has been shown to stimulate soil fauna that play important role in soil aggregation, soil C sequestration, soil nutrient and water use efficiencies, and influence crop yields (Nhamo, 2007; Castellanos-Navarrete et al., 2012; Paul et al., 2015).

In many parts of the world, especially areas of SSA, the challenge of feeding an ever increasing population is persistent, due to prevalent land degradation and low soil productivity. Farmers' efforts to restore and maintain soil fertility are constrained by various challenges, the major ones being accessibility, affordability and knowledge on input management (Karanja et al., 2006; Vanlauwe et al., 2015). Historically, increasing food production in Africa has largely been attributed to land intensification and there has been efforts to intensify by increasing productivity per unit area. Sustainable intensification (SI) attempts to promote increased crop productivity by small-holder farmers in SSA (Garnett et al., 2013). Although SI has addressed the yield component, other aspects of sustainability such as soil biological functions have not been sufficiently addressed. SI advocates for management practices that reconcile environmental conservation and sustainability, and food security (Brussaard et al., 2010).

Despite the widespread interest in CA, empirical evidence of the benefits of CA in SSA is limited (Paul et al., 2013). For instance, the magnitude and direction of effect by CA on soil fauna, however, remains unquantified, especially in low-input systems of Sub-Saharan Africa. Response of soil fauna to soil tillage, available crop residues, and cropping practices in the region is largely unclear yet such knowledge is imperative for environmental conservation, sustainability and improved ecosystem services. Therefore, the aim of this study was to evaluate the effect of CA and associated management practices on soil fauna richness and abundance. Specifically, the study was to assess how conservation agriculture and its principal elements that encompass zero tillage, application of organic or inorganic inputs and cropping system, affect soil fauna richness and abundance. It is hypothesized that CA, organic or inorganic inputs and crop rotation and or mixed cropping would positively influence soil fauna richness and abundance.

MATERIALS AND METHODS

Study Sites

The studies were conducted at three sites (Plate 1). The first included a medium-term (6 years) trial, based at the Kenya Agricultural and Livestock Research Institute (KALRO) in Embu County, about 130 km north of Nairobi, Kenya. The trial is located in the sub-humid Central highlands of Kenva on the southern slopes of Mt Kenya at: latitude 0° 32' S, longitude 37° 37' E and an altitude of 1,480 m (Table 1). The average temperature is 19.5°C. The area receives bimodal rains with a mean total of 1,450 mm in two distinct seasons: long rains (March to August) and short rains (October to January). Agroecological zone is Upper mid land (UM3) (Jaetzold et al., 2007). The Soils are mainly Humic Nitisols (FAO, 1989) derived from basic volcanic rocks (FAO-UNESCO, 1997). They are deep, wellweathered with a clay texture (% sand, clay, silt: 3, 22, and 75) with moderate to high inherent fertility. The second is a medium-term trial (6 years) based at the Kenya Agricultural and Livestock Research Institute (KALRO) Kakamega (also in sub-humid western Kenya, Kakamega County), while the third is Nyabeda, a sub-humid site in western Kenya, within Siaya County where a long-term trial (15 years) is located. The Kakamega and Nyabeda sites are characterized by two rainy seasons: a long rainy season between March and August and short rainy season between September and January (Jaetzold et al., 2007). In all the sites, maize is the main staple crop and is normally grown either as a monocrop or in association with legumes, mainly common beans and groundnuts. Soybean, a cash crop, is also grown by farmers in the western Kenya region. All sites have predominantly smallholder settlements, with land sizes ranging from 0.3 to 3 ha per household. The study sites represent different soil types with varied physical and chemical characteristics. The study was conducted in December 2015 for the sites in eastern Kenya and June-July 2016 for the two trials in western Kenya.

Experimental Design

Embu Trial

The medium-term (6 years) field trial was established in 2010. Treatments tested for soil fauna include tillage (conventional till: CT, zero till: CA), organic residues (maize or bean residues: CR) and cropping systems (sole maize: SM, sole bean: SB and maize-bean intercrop: MBi) (Table 2). During the cropping seasons, plots are planted with maize (Zea mays Hybrid 512) and beans (Phaseolus vulgaris, variety Rose Coco) as per treatments. Mineral N fertilizer is applied targeting 60 kg N ha⁻¹ for sole maize (SM) and 20 kg N ha⁻¹ for sole beans (SB), applied as calcium ammonium nitrate (CAN), one-third at planting, and two-third at stage V6 of maize. Maize-bean intercrop (MBi) received 80 kg N ha⁻¹. Irrespective of tillage methods, SM, SB, and MBi received 60, 51, and 111 kg P_2O_5 ha⁻¹ from [N23P23], [N18P46], and combination of [N23P23] + [N18P46] fertilizer materials, respectively (Micheni et al., 2016). For the treatments receiving organic inputs, all residues from the preceding crop were incorporated into the soil, thus providing about 39.2 t C



ha⁻¹ and 67.4 t of SOM, respectively. Residues were removed from those treatment not receiving residues. The treatments were replicated three times in a randomized complete block design with plots of 10 m long by 8 m wide. Conventional till was done by hand-hoeing to a maximum depth of 15 cm, whereas no tillage was conducted in the zero till plots. Sole maize (SM): Maize spacing was 75 cm between rows and 50 cm within. Three seeds were sown per hill and thinned 1 week after crop emergence to two plants per hill, to give a plant population of 53,333 plants ha^{-1} . Sole bean (SB): Sole bean were spaced at 50 cm between rows and 15 cm within rows while maintaining one plant per hill giving plant population of 133,330 plants ha^{-1} . Maize bean intercrop (MBi): Maize spacing was maintained like in maize sole crop but bean spacing was slightly adjusted to 50 cm between rows and 20 cm within rows and two plants per hill. This gave plant population of 133,330 plants ha⁻¹, which was the same as in SB configuration. This was done to minimize confounding effects due to plant population differences (Micheni et al., 2016). Under CT, two weed control events were conducted manually using hand tools (machete and hoes) within a given season. The first weeding was done 1-2 weeks after crop emergence and the second was conducted in approximately one and half months after the first weeding. In the CA treatments, weeds were controlled using pre and post-emergence herbicides. Roundup (Glyphosate), a post-emergence herbicide, was applied at the rate of 3.0 liters (L) ha^{-1} to kill weeds at the beginning of the seasons. Dual Gold (960 g L^{-1} Metolachlor), a pre-emergence herbicide, was applied at the rate of 2.0 L ha⁻¹ on relatively moist soil surface after planting but before emergence of crops and weeds. One month after the crop emergence, selective Basagran post-emergence herbicide was applied at the rate of 2.0 Lha⁻¹

to manage actively growing grass and broad-leafed weeds in maize-bean intercrop.

The herbicide is effective mainly through contact action and therefore care was taken to have all weeds thoroughly covered with the herbicide sprays while avoiding maize and bean leaves.

Kakamega

The experimental set up, design and management of this trial was the same as that of Embu medium-term. The trial tested the effect of different treatments on arable crop production and these include: tillage (conventional till, zero till), organic (maize residues) and cropping systems (continuous maize, maize-bean rotation and intercrop; Table 2). However, fertilizer was applied at a rate of 50 kg N ha⁻¹ and 25 kg P ha⁻¹. The organic inputs are applied onto the soil surface every season at 2 t ha^{-1} . The residues, however, were removed from those treatment not receiving residues. The treatments were replicated three times in a randomized complete block design with plots of 10 m long by 8 m wide. Treatments selected for soil fauna studies included: maize-bean intercrop (MBi) under conventional (CT) and zero (CA) till, but all with crop residues (CR) applied. A no input treatment that typically represented farmer practice (or control) was also sampled (Table 2).

Nyabeda Trial

The field experiment was established in March 2003, and has been managed by the International Center for Tropical Agriculture (CIAT). The details of the trial set-up and management are documented in Kihara et al. (2012). The treatments used to study soil fauna diversity include a farmer practice under sole maize, CT with residues added, CA under maize-soybean

TABLE 1	Location,	climatic,	and soil	characteristics	of the	study sites
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Embu	Kakamega	Nyabeda
MT-2010	MT-2010	LT-2003
Humid	Sub-humid	Sub-humid
Upper midland 3	Upper midland 1	lower midland 2
00° 33.18′ S	0° 16.96' N	0° 07′ N
037° 53.27′ E	34 46.07' E	34° 24′ E
1,420	1,534	1,420
1,250	1,978	1,800 ^a
20	21	23.2
16–21	11	14
21–28	26	31
Humic Nitisols ^b	Eutric Nitisol ^c	Ferralsol ^c
3:22:75	13:34:53	15:21:64
3.88	5.40	5.08
0.27	0.70	0.10
16.13	3.40	2.99
2.15	0.93	4.69*
0.45	0.05	1.68*
3.70	4.10	1.35
0.37	0.30	0.15
	Embu MT-2010 Humid Upper midland 3 00° 33.18' S 037° 53.27' E 1,420 1,250 20 16–21 21–28 Humic Nitisols ^b 3:22:75 3.88 0.27 16.13 2.15 0.45 3.70 0.37	Embu Kakamega MT-2010 MT-2010 Humid Sub-humid Upper midland 3 Upper midland 1 0° 33.18' S 0° 16.96' N 037° 53.27' E 34 46.07' E 1,420 1,534 1,250 1,978 20 21 16–21 11 21–28 26 Humic Nitisols ^b Eutric Nitisol ^c 3:22:75 13:34:53 3.88 5.40 0.27 0.70 16.13 3.40 2.15 0.93 0.45 0.05 3.70 4.10

*Value obtained in meq 100 g^{-1} of soil.

^a2002–2008 period.

^bSee Jaetzold et al. (2007) for details.

^cSee Jaetzold et al. (2006) for details.

MT, medium-term; LT, long-term.

rotation and intercropping. There were 3 replicates included and individual plots measured 7 \times 4.5 m. The crop rotation since trial establishment consisted of soybean (Glycine max L.) during short rains and maize (Zea mays L.) during long rains. All plots were fertilized with 60 kg ha⁻¹ N (urea), 60 kg ha⁻¹ P (Triple Super Phosphate), and 60 kg ha⁻¹ K (Muriate of Potash) per season (Table 2). To control stem borer, 5 kg ha^{-1} of granulated Bulldock (beta-cyfluthrin) was applied in the funnel of the maize plants during the 5th week after planting in all treatments. Under CT, the seedbed was prepared by hand-hoeing to 15 cm soil depth. Weeding was performed three times per season using hand hoe. Under CA, a 3 cm deep seedbed was prepared with the hand hoe. Weeding was performed three times per season by hand pulling until the long rainy season of 2009. Thereafter, herbicides (glyphosate and 2, 4-dichlorophenoxyacetic acid) have been applied to all CA treatments before planting and subsequent weeding done by hand pulling. Maize residues were collected after crop harvest, dried, chopped and stored during the dry season. At the time of soybean planting, the residues were reapplied at a rate of 2 h ha⁻¹ for those treatments receiving crop residues. Since soybeans drop leaves prior to grain maturity, soybean residues (leaves and stems) always remained in the field after harvesting, irrespective of treatment. These soybean residues were then either incorporated in CT or remained at the soil surface in CA. In our fauna sampling regime, only a few of the trial treatments were selected, and a no input treatment was also sampled as in Kakamega (Table 2).

Faunal Sampling Techniques

Two techniques were employed to sample fauna for richness and abundance.

Macrofauna

Using a monolith of size $25 \text{ cm} \times 25 \text{ cm} \times 30 \text{ cm}$, samples were taken 8 weeks (in December 2015 for the sites in eastern Kenya and June-July 2016 two experimental trials in western Kenya) after planting crops in the season (Swift and Bignell, 2001; Bignell et al., 2008): At each observation, one sample was taken randomly from each plot. The monolith was situated over a randomly selected spot and dug with a spade and hoe to a 30 cm soil depth (Plate 2). The soil from the monolith was removed by hand depth-wise (0-15 and 15-30 cm) into plastic buckets. The soil sample from each depth was placed in different plastic trays (20 cm by 30 cm) and gently sorted out to locate the animals. The animals were separated into major taxonomic groups, recorded and then collected in plastic bottles. The soil fauna collected were preserved in 75% alcohol for subsequent identification at the Soil microbiology laboratory of CIAT, ICIPE Duduville Campus, Nairobi, Kenya. Earthworms were killed in 75% alcohol and fixed in 4% formaldehyde. In the laboratory, counting was done. Species richness, and number of different categories of animals were expressed per meter square.

Mesofauna

At the same time of sampling for macrofauna soil samples were collected for mesofauna observations using a metallic core of 10 cm diameter up to 30 cm depth (and at same 0–15 or 15–30 cm depths as macrofauna). One sample was taken in each plot at each sampling. The samples were taken to the CIAT laboratory where mesofauna groups were extracted using the behavioral or dynamic method with Berlese-Tullgren as the basic apparatus (**Plate 3**) (Southwood, 1995). This apparatus was originally designed by an Italian entomologist, Berlese, A. and later modified by a Swede, Tullgren, A. who used a light bulb as the source of heat. The apparatus has since been modified by many workers (Southwood, 1995). For this study the apparatus was designed and constructed locally.

Basically, the collected soil was poured into the perforated soil sample containers of the "Berlese-Tullgren" funnel apparatus over a funnel. Heat was applied to the soil using a 75 watts bulb placed above the sample container. Heat supplied was regulated by upward adjustments of the sample containers. Thus, the animals were exposed to a controlled gradient of high to low temperature and light, and low to high humidity from top to bottom, so that the animals are driven gradually downwards and out into the collection jars filled with 75% alcohol. The set up was left for 24 h. After this period, the animals collected in the jars were sorted out and counted under a light microscope. The preserved animals were also stored and later taken to the laboratory for taxonomic analyses.

Soil Sampling and Nutrient Analysis

Immediately after handpicking the soil macrofauna, soils from each monolith was mixed thoroughly to make

TABLE 2 | Treatment selected and descriptions.

Treatment Tillage		Cropping	Organic input	Inorganic input
Embu medium-term trial				
1. CTMBi-CR	Conventional	Maize-beans intercrop	None	80 kg N, 111 P ₂ O ₅
2. CASB+CR	Zero	Sole beans	Beans residues	20 kg N, 51 kg P ₂ O ₅
3. CASM+CR	Zero	Sole maize	Maize residues	60 kg N, 60 kg P ₂ O ₅
4. CAMBi+CR	Zero	Maize-beans intercrop	Maize and beans residues	80 kg N, 111 P ₂ O ₅
Kakamega medium-term tr	ial			
1. FP (Farmer practice)	Conventional	Sole maize	None	None
2. CTMBi+CR	Conventional	Maize-bean intercrop	2 t/ha maize residues	50N, 25P
3. CAMBi+CR	Zero	Maize-bean intercrop	2 t/ha maize residues	50N, 25P
Nyabeda long-term trial				
1. FP (Farmer practice)	Conventional	Sole maize	None	None
2. CTMSr+CR	Conventional	Maize-soybean rotation	2 t/ha maize residues	60 kg N/ha-Urea
3. CAMSr+CR	Zero	Maize-soybean rotation	2 t/ha maize residues	60 kg N/ha-Urea
4. CAMSi+CR	Zero	Maize-soybean intercrop	2 t/ha maize residues	60 kg P/ha-TSP

CT, Conventional till; CA, Conservation agriculture (Zero till); FP, Farmer practice; SM, Sole maize; SB, Sole beans; MBi, Maize bean intercrop; MSi, Maize-soybean intercrop; MSr, Maize-soybean rotation CR, Crop residue; N, Nitrogen; +/- denotes with or without crop residues and with or without nitrogen.



PLATE 2 | Soil monolith excavation and macrofauna sampling.

a composite sample of about 500 g for analysis. Soil parameters measured included: soil pH total organic C and N, available P, and exchangeable bases (Na, K, Ca, and Mg). Soil pH was determined using a pH meter with soil-water ratio of 1:2.5 (Anderson and Ingram, 1993). Total organic C and N were determined using a CN-analyser, while P and the bases were extracted by the Mehlich-3 procedure (Mehlich, 1984) and measured through inductively coupled plasma atomic emission spectroscopy (Isaac and Johnson, 1998).

Statistical Analyses

The data obtained on soil fauna richness and abundance and soil chemical properties were subjected to analysis of variance (ANOVA) with Genstat 17.1 (2015). Levene's test was used to test for homogeneity of variances (Field, 2005). In case of non-homogeneity of variances, data were square root $(x + 0.5)^{1/2}$ transformed before further analysis. Fauna data were

analyzed separately for each depth (0-15 and 15-30 cm). Linear Mixed Model was fitted by Restricted Maximum Likelihood (RELM) procedure using the Genstat package. This procedure allows for inclusion of both fixed- and random-effects terms in the model such that profiled deviance of RELM criterion is optimized for the parameter estimates (Kuznetsova et al., 2014; Bates et al., 2015). Treatments were included in the model as fixed factors, whereas block was defined as a random factor. The statistical significance was determined at $p \le 0.05$ and levels of significance among the different treatments were evaluated using Fischer's least significance difference (LSD). Correlation analysis (Pearson correlations), was conducted to establish the significance of the relationships between soil fauna and soil factors. Because fauna and soil variables had different units of measurement, they were standardized first so that each variable received equal weight in the analysis and also to make the coefficient (r) values comparable (Cao et al., 1999; Jongman et al., 2005).



RESULTS

Soil Fauna Composition

Overall, a total of 58 macrofauna species classified into 14 major groups were sampled across the three trials in eastern and western Kenya. However, it should be noted that, the numbers given are estimations because it is difficult (or even impossible) to be sure that these additional categories consist of just one species (already listed or not) or more species. The medium trial of Embu had relatively higher species richness compared to the mediumterm trial of Kakamega and long-term trial of Nyabeda, and the highest number of species (37) was recorded in the Embu medium-term trial followed by Nyabeda (32) > Kakamega (27)(Data is available as Supplementary Tables). The macrofauna groups were dominated by Oligochaeta (earthworms), Isoptera (termites), Hymenoptera (ants), and Coleoptera (beetles), with these groups constituting over 80% of mean totals across the different trials (Table 3). In Embu medium-term trial, Hymenoptera were the most abundant of these faunal groups, constituting about 39% of the mean total followed by Isoptera (28%), Coleoptera (12%), and Oligochaeta (10%) (Table 3). In Kakamega trial, Oligochaeta (64%) was the most dominant of all the macrofauna groups followed by Isoptera (7%) >Hymenoptera (6%) > Coleoptera (5%), whereas in Nyabeda, Isoptera was dominant (55%) followed by Oligochaeta (21%) > Hymenoptera > Coleoptera (6%) in that order. In all the trials, however, the other macrofauna groups were observed in very low numbers, with each group constituting \leq 5% (**Table 3**). Generally, macrofauna were more abundant in the eastern Kenya than in western Kenya sites, with the former recording a mean total between 875 and 1,386 numbers per m^{-2} compared to that between 565 and 784 numbers per m^{-2} in the latter (Table 3).

For mesofauna groups, 18 species classified into seven major groups were observed across the four trials, and the western Kenya sites (Nyabeda and Kakamega) had relatively higher mesofauna richness compared to the eastern Kenya (Embu) sites (Data is available as a **Supplementary Tables**). Mesofauna was equally relatively more abundant in the western Kenya sites compared to the eastern Kenya site (**Table 3**). In the eastern Kenya (Embu) trial, mesofauna groups were dominated by Acarina, which constituted > 50% followed by Collembolla (16–17%) and Enchytraeids (10–15%), with each of the other groups constituting < 6% (**Table 3**). Western Kenya trials were also dominated by Acarina (40–59%) and Collembolla (35–46%) and the other groups each constituted \leq 5% (**Table 3**).

Soil Fauna Taxonomic Richness

The combinations of tillage practice, organic residues and cropping system had significant effect on macrofauna taxonomic richness (p < 0.05) largely at the top 0–15 cm soil depth than at the lower 15-30 cm soil depth in most of the study sites (Figure 1). At 0-15 cm soil depth of the Embu medium-term trial, mean macrofauna taxonomic richness was significantly higher in both conservation agriculture under sole maize (CASM+CR) cropping and conservation agriculture under maize-bean intercrop (CAMBi+CR) treatments than in either conventional till minus crop residues under maize-bean intercrop (CTMBi-CR) or conservation agriculture under sole beans (CASB+CR) (Figure 1A). At 15-30 cm, mean macrofauna taxonomic richness did not differ among the treatments (p =0.503) (Figure 1B). At the Kakamega, no significant differences were noted for macrofauna mean richness among the treatments at both 0-15 and 15-30 cm soil depths (Figures 1C,D). At Nyabeda, macrofauna richness was significantly lower in conventional (typical farmer's practice without residues) till without inputs than in the other treatments (Figure 1E). At 15-30 cm, mean macrofauna taxonomic richness did not differ among the treatments (p = 0.370) (Figure 1F).

In addition, no significant differences were noted for mesofauna mean richness among the treatments at both 0-15 cm and 15-30 cm soil depths in all study sites (**Figure 2**). As

TABLE 3 | Mean total population and percent composition of macrofauna and mesofauna within each group across medium-term (MT) and long-term (LT) trials in Embu, Kakamega, and Nyabeda, Kenya.

	Embi	u-MT	Kakame	ega-MT	Nyabeda-LT		
Macrofauna group	Mean total	% of total	Mean total	% of total	Mean total	% of total	
Oligochaeta	136	9.8	499	63.6	120	21.2	
Isoptera	381	27.5	58	7.4	312	55.2	
Hymenoptera	545	39.3	46	5.9	56	9.9	
Coleoptera	171	12.3	39	5.0	36	6.4	
Diptera	15	1.1	4	0.5	4	0.7	
Lepidoptera	33	2.4	16	2.0	4	0.7	
Diplopoda	24	1.7	38	4.8	1	0.2	
Chilopoda	34	2.4	39	5.0	7	1.2	
Orthoptera	21	1.5	4	0.5	3	0.5	
Araneae	11	0.8	20	2.5	17	3.1	
Odonata	3	0.2	-	-	4	0.7	
Hemiptera	4	0.3	22	2.8	1	0.2	
Blattoidea	5	0.4	-	-	-	_	
Isopoda	3	0.2	-	-	-	_	
Total	1,386	100	784	100	565	100	
Mesofauna group							
Acarina	592	57.9	1,196	39.5	2,900	58.9	
Arachnida	3	0.2	-	-	-	-	
Collembolla	172	16.9	1,392	46.0	1,741	35.3	
Diplura	50	4.9	155	5.1	32	0.6	
Enchytraeidae	150	14.7	14	0.5	11	0.2	
Protura	55	5.4	-	-	-	-	
Symphyla	-	_	268	8.9	243	4.9	
Total	1,022	100	3,025	100	4,927	100	

expected, soil fauna richness reduced with depth where these were nearly \leq 50% that of top soil for each of the treatments.

Soil Fauna Abundance

Across all sites, and in both (0–15 and 15–30 cm) soil depths, no significant effect of tillage, cropping and organic inputs on soil macrofauna abundance were observed (**Figure 3**). Equally no significant treatment effects on soil mesofauna abundance were observed in Embu medium-term (**Figures 4A,B**) and Nyabeda long-term trials for both soil depths (**Figures 4E,F**). Although a similar observation was made on mesofauna abundance at the 15–30 cm soil depth of Kakamega trial (**Figure 4D**), at 0–15 cm soil depth, maize-bean intercrop system under conservation agriculture (CAMBi+CR) had significantly higher mesofauna abundance than the convention till with similar management (CTMBi+CR) practices or conventional till (farmers practice) without any inputs (**Figure 4C**).

Soil Fauna Group Abundance Across Treatments

Faunal abundance were analyzed and assessment made depthwise across treatments for the various groups.

Macrofauna

In the long-term trial, tillage, residue application and cropping system affected only a few of the macrofauna groups such as Oligochaeta, Chilopoda, and Araneae, but only at the top soil depth. This is unlike medium-term trials where significant effects were noted in either depths (**Table 4**).

At 0–15 cm soil depth, in the Embu medium-term trial, Oligochaete, and Araneae were significantly more abundant in both conservation agriculture, maize-bean intercropping system with residues (CAMB+CR) and conservation agriculture, sole maize with residues (CASM+CR) treatments than zero till, sole beans with residues (CASB+CR) and conventional till, maize bean system without residues treatments (CTMB-CR) (**Table 4**). Chilopoda, however, were significantly more abundant in CASM+CR than in the other treatments. The other macrofauna groups did not differ among the treatments, and at 15–30 cm, all the macrofauna groups did not differ among the treatments as well.

In Kakamega, management practices affected only Oligochaeta and Chilopoda groups but only at 0–15 and 15–30 cm soil depths, respectively. At 0–15 cm depth, maize-bean intercrop system under conservation agriculture with residue applied (CAMBi+CR) had significantly higher Oligochaete abundance than the convention till with similar management (CTMBi+CR) practices or conventional till (farmers practice: FP) without any inputs. At 0–15 cm, the predacious Chilopod group, was on the other hand, significantly higher in famers practice (FP-conventional till without any inputs) than in the other treatments.



FIGURE 1 [Soil macrotauna diversity (richness) across trials of Embu, Kakamega, and Nyabeda. M1, Medium-term; L1, Long-term; C1, Conventional till; CA, Conservation agriculture (Zero till); SM, Sole maize; SB, Sole beans; MBi, Maize bean intercrop; MSi, Maize-soybean intercrop; MSr, Maize-soybean rotation; CR, Crop residue. (A) Embu MT (0-15 cm, $p = 0.027^{+}$); (B) Embu MT (15-30 cm, p = 0.503); (C) Kakamega MT (0-15 cm, p = 0.329); (D) Kakamega MT (15-30 cm, p = 0.414); (E) Nyabeda LT (0-15 cm, $p = 0.029^{+}$); (F) Nyabeda LT (15-30 cm, p = 0.379). Bars with different lower case letters are statistically significantly different at p < 0.05.

At 0–15 cm soil depth of Nyabeda, cropping system and residue addition seem to be the influencing factors for Hymenoptera abundance as this group was significantly higher under maize-soybean rotation system for both conventional (CTMSr+CR) and conservation agriculture systems with crop residue applied (CAMSr+CR) than in the conventional till under continuous maize without inputs (FP) and conservation agriculture under maize soybean intercrop but with crop residues (CAMSr+CR). No significant treatment effects were observed for the other macrofauna groups at this top soil depth and for all the macrofauna groups at 15–30 cm soil depth.

Mesofauna

In the medium-term trial of Embu, and in both soil depths, all the mesofauna groups did not differ among the treatments indicating lack of significant influence of tillage, organic residue application

and crop management (**Table 5**). However, significant effect of tillage on mesofauna abundance were noted at top 0-15 cm soil depths for both Kakamega and Nyabeda trials. At Kakamega, Collembolan group was significantly higher in the conservation agriculture practices than in the conventional till practices, and the same pattern was observed for Symphyla group in the Nyabeda trial. As expected, mesofauna groups were less abundant in the lower soil depths compared to the upper depths.

Management Practices and Effect on Soil Chemical Properties

Soil management practices had significant effects on soil chemical properties (**Table 6**). In Embu site, soil TOC was significantly lower in conservation tillage under sole maize (CASM+CR) than in all the other treatments. Total organic N, on the other hand, was significantly higher in conventional tillage under



Conservation agriculture (Zero till); SM, Sole maize; SB, Sole beans; MBi, Maize bean intercrop; MSi, Maize-soybean intercrop; MSr, Maize-soybean rotation; CR, Crop residue. (A) Embu MT (0-15 cm, p = 0.864; (B) Embu MT (15-30 cm, p = 0.515); (C) Kakamega MT (0-15 cm, p = 0.058); (D) Kakamega MT (15-30 cm, p = 0.525); (E) Nyabeda LT (0-15 cm, p = 0.405); (F) Nyabeda LT (15-30 cm, p = 0.125).

maize-bean intercrop without crop residues (CTMBi-CR) than in the conservation tillage practices under sole maize and maize-bean intercrop (**Table 6**). It however, did not differ from conservation tillage under sole beans. In Kakamega site, soil pH was significantly higher in conventional tillage under maize-bean intercrop with crop residues than in conservation tillage under intercrop (CAMBi+CR) and farmer practice (FP), and a trend similar to that of soil pH was observed for Ca and Mg (**Table 6**). Tillage influenced TOC and P contents, and were significantly higher in conservation tillage under intercrop (CAMBi+CR) than in the conventional tillage under intercrop (CA MBi+CR) and farmer practice (FP). Although TON was significantly higher in CAMBi+CR than FP, it did not differ from CTMBi+CR (Table 6). At Nyabeda, treatment effect was observed only for soil pH, Ca, and Mg. Soil pH was higher in CTMSr+CR than either CAMSi+CR or FP although it did not differ from CAMSr+CR. Calcium content was highest in CAMSr+CR > CTMSr+CR > FP, but lowest in CAMSi +CR. Magnesium content on the other hand was highest in CTMSr+CR > CAMSr+CR > FP, but again lowest in CAMSi+CR (Table 6).

Correlation Between Soil Fauna and Soil Chemical Properties

Over 60% of the macrofauna, and over 70% of mesofauna showed significant correlation with soil parameters (**Table 7**). Oligochaeta, Diplopoda, and Hemiptera significantly positively



Nyabeda LT (0-15 cm, p = 0.285); (F) Nyabeda LT (15-30 cm, p = 0.843).

correlated with soil organic matter (TOC, TON) and P, whereas Hymenoptera, Coleoptera and Blattodea positively correlated with nearly all the exchangeable bases (Na, K, Ca, and Mg). Diptera, Lepidoptera, and Odonata on the other hand significantly correlated with all exchangeable bases measured (except K and Mg). Whereas, Orthoptera positively correlated with Mg, and Isopoda positively with K, Chilopoda negatively correlated with soil pH (**Table 7**). Mesofauna groups on the other hand showed significant correlations with exchangeable bases (Na, K, Ca, and Mg) only. Whereas, Enchytraeid and Protura positively correlated with all the exchangeable bases and Arachnida with Mg, Acarina, Collembolla, and Symphila negatively correlated with all the bases.

DISCUSSION

The total number of macrofauna taxa of between 25 and 37 recorded in our study sites conforms with the range of 35–38 taxa

recorded by Ayuke (2000) and Ayuke et al. (2009) across arable fields within western and eastern Kenya, respectively. However, these taxa were much lower than 75 taxa recorded by Karanja et al. (2009) across arable sites of a different ecozone, the Coastal region of Kenya. Mesofauna total taxa were, however, three times higher than that recorded under biomass transfer agroforestry technology at Maseno, Western Kenya (Ayuke, 2000).

Results of this study have demonstrated the benefits of conservation agriculture in enhancing soil fauna richness and abundance. Higher macrofauna taxonomic richness and abundance of mesofauna in CA treatments than in CT without residue application are related to an improved microclimate and access to food in the CA system. Disturbances caused by tillage operations and residue removal (CT) are known to negatively affect sensitive fauna (Ayuke et al., 2011a,b), except for the predating and foraging groups like Hymenoptera (ants), Chilopoda (centipedes) and some species of Isoptera (termites) e.g., *Microtermes* sp. Earthworm species, among them, the



Dichogaster sp. are also sensitive to disturbance (Ayuke et al., 2011a,b), and so were lacking under conventional till as opposed to conservation agriculture system.

Microclimate modification, food resource availability, and land Management practices (e.g., tillage, organic resource use, crop rotation, and application of agrochemicals such as pesticides, herbicides, and inorganic fertilizers) are known to either positively or negatively influence the diversity and abundance of soil fauna communities (Dangerfield, 1993; Beare et al., 1997; Nhamo, 2007; Gianessi, 2010; Isenring, 2010). Tillage influences soil fauna in several ways: the mechanical and physical disturbance due to tillage can cause habitat destruction for some of the macrofauna groups. When vegetation are removed, the resultant change in habitat structure, the reduced range and abundance of food resources and the extreme climate at the soil surface, most likely combine to create harsh environment that may be intolerable to most soil fauna groups among them earthworms. Consequently, the population of these soil fauna groups could be suppressed, possibly explaining why some species of earthworms, springtails (Collembolla) and Symphyla were absent from the conventional till systems, hence the lower diversity and abundances observed. However, predacious fauna groups such as ants (Hymenoptera) were favored by tillage as this exposed their likely prey among them termites, on to the soil surface hence abundant food.

Conservation agriculture favored those fauna groups e.g., earthworms (Oligochaeta) and predacious spiders (Araneae) that are sensitive to disturbance caused by tillage. Many known Araneae (spider) species live on the soil surface or in soil crevices, invade the natural pore system of soils or are in some way closely associated with the soil systems. Higher Araneae recorded under CA practices could be attributed to less disturbance TABLE 4 | Mean abundance of soil macrofauna groups across treatments in medium-term and long term trials of Embu, Kakamega, and Nyabeda, Kenya.

Group	0–15 cm					15–30 cm					
	Treatment				P-value	Treatment				P-value	
Group	CTMBi-CR	CASB+CR	CASM+CR	CAMBi+CR	P-value	CTMBi-CR	CAMBi+CR	CASB+CR	CASM+CR	P-value	
Embu medium-term trial											
Oligochaeta	80b	Ob	192a	197a	0.029*	11	48	0	16	0.225	
Isoptera	272	43	283	187	0.810	165	37	107	432	0.420	
Hymenoptera	16	501	901	251	0.283	21	91	144	256	0.388	
Coleoptera	139	59	139	160	0.215	85	21	43	37	0.064	
Diptera	27	43	0	5	0.091	11	0	0	0	0.070	
Lepidoptera	59	32	0	37	0.412	5	0	0	0	0.455	
Diplopoda	11	0	27	37	0.070	0	16	0	5	0.572	
Chilopoda	Ob	Ob	64a	5b	0.031*	21	5	5	21	0.349	
Orthoptera	0	48	21	5	0.566	0	11	0	0	0.455	
Araneae	Ob	Ob	16a	21a	0.035*	5	5	5	5	0.455	
Odonata	5	0	0	5	0.654	0	0	0	0	_	
Hemiptera	11	5	0	0	0.189	0	0	0	0	_	
Blattoidea	0	5	0	16	0.216	0	0	0	0	_	
Isopoda	5	0	5	0	0.455	0	0	0	0	_	
Group	FP	CTMBi+CR	CAMBi+CR		P-value	FP	CTMBi+CR	CAMBi+CR		P-value	
Kakamega medium-term	trial										
Oligochaeta	112b	213b	869a		0.016*	64	91	149		0.493	
Isoptera	27	5	5		0.444	69	16	53		0.476	
Hymenoptera	16	48	32		0.444	16	16	11		0.790	
Coleoptera	27	16	32		0.815	5	21	16		0.444	
Diptera	0	5	0		0.444	0	0	5		0.444	
Lepidoptera	0	0	43		0.218	0	0	5		0.444	
Diplopoda	0	5	101		0.444	0	0	5		0.444	
Chilopoda	27	16	32		0.744	37a	Ob	5b		0.048*	
Orthoptera	0	0	0		-	0	11	0		0.111	
Araneae	11	16	0		0.373	0	16	16		0.617	
Hemiptera	0	11	48		0.444	0	0	5		0.444	
Group	FP	CTMSr+CR	CAMSi+CR	CAMSr+CR	P-value	FP	CTMSr+CR	CAMSi+CR	CAMSr+CR	P-value	
Nyabeda long-term trial											
Oligochaeta	91	27	17	112	0.736	32	64	21	75	0.628	
Isoptera	5b	Ob	224a	5b	0.005**	139	11	43	64	0.667	
Hymenoptera	5	85	37	69	0.161	16	43	16	0	0.175	
Coleoptera	5	80	11	21	0.495	0	11	5	5	0.572	
Diptera	5	405	0	261	0.542	5	0	0	0	0.455	
Lepidoptera	0	11	5	0	0.455	5	0	0	0	0.455	
Diplopoda	0	0	0	0	-	5	0	0	0	0.455	
Chilopoda	0	11	0	0	0.591	0	0	0	0	-	
Orthoptera	0	5	0	0	0.455	0	0	0	0	-	
Araneae	0	43	21	5	0.392	0	0	5	5	0.654	
Hemiptera	0	5	0	0	0.455	0	5	5	0	0.654	
Blattoidea	0	0	0	5	0.455	0	0	0	0	-	

Means followed by the same lowercase letters across rows are not significantly different at p < 0.05. Significant p-values are indicated in bold. For some of the groups with zeros at lower depths, ANOVA did not calculate the p-value. For treatment abbreviations (see **Table 2**). *Significant; **Highly significant; **Very highly significant.

that might have affected the populations of their likely prey. It has been shown that, reduced- or no-tillage agriculture is beneficial for soil conservation in that, soil-surface accumulation

of crop residues as a result of minimal soil disturbance protects soil from water and wind erosion (Stinner and House, 1990). Unlike in conventional tillage, litter and soil organic matter TABLE 5 | Mean abundance of soil mesofauna groups across treatments in long- and short-term trials of Embu, Nyabeda, and Kakamega.

		0–15 cm					15–30 cm			
		Treatment					Treatment			
Group	CTMBi-CR	CASB+CR	CASM+CR	CAMBi+CR	P-value	CTMBi-CR	CASB+CR	CASM+CR	CAMBi+CR	P-value
Embu medium	-term trial									
Acarina	1,100	233	533	233	0.415	33	33	133	67	0.142
Arachnida	0	167	0	167	0.654	0	0	0	0	-
Collembola	67	100	167	133	0.931	67	0	133	33	0.514
Diplura	67	33	33	0	0.714	0	0	33	33	0.654
Enchytraeidae	0	67	100	167	0.136	133	67	33	33	0.285
Protura	33	33	0	67	0.572	100	0	33	0	0.285
Group	FP	CTMBi+CR	CAMBi+CR		P-value	FP	CTMBi+CR	CAMBi+CR		P-value
Kakamega me	dium-term tria	al								
Acarina	464	506	1,477		0.122	84	295	759		0.322
Collembola	127b	42b	3,249a		0.012*	42	422	295		0.391
Diplura	0	84	42		0.444	84	211	42		0.432
Enchytraeidae	42	0	0		0.444	0	0	0		-
Symphyla	0	211	169		0.132	127	295	0		0.273
Group	FP	CTMSr+CR	CAMSi+CR	CAMSr+CR	P-value	FP	CTMSr+CR	CAMSi+CR	CAMSr+CR	P-value
Nyabeda long-	-term trial									
Acarina	1,350	2,110	2,363	2,363	0.538	464	1,772	676	506	0.199
Collembola	338	2,110	2,012	295	0.224	380	1,224	338	253	0.441
Diplura	0	0	0	0	-	84	42	0	0	0.613
Enchytraeidae	42	0	0	0	0.455	0	0	0	0	-
Symphyla	84b	0b	295a	295a	0.025*	42	42	211	0	0.150

Means followed by the same lowercase letters across rows are not significantly different at p < 0.05. Significant p values are indicated in bold. For treatment abbreviations (see **Table 2**). *Significant; **Highly significant; **Very highly significant.

(and also nutrients) tend to concentrate near the soil surface of conservation-tillage systems, and these condition are likely to favor soil fauna within the CA. Although crop or organic residues get incorporated into the soil, albeit slowly, through invertebrate activity (Ayuke et al., 2009; Kihara et al., 2015), the litter layer is a very important factor in ameliorating soil temperature and water content of soil, thus providing a more stable environment for soil- and litter-dwelling invertebrates. This may explain the expected higher macrofauna richness and mesofauna abundance observed in top-soils than sub-soils.

The important role of organic amendments in influencing presence and abundance of specific faunal groups have been discussed (Ayuke, 2000) and their role explains why, *P. annulatus* earthworm species that thrive best under high organic matter environments were only recorded under sole maize and maizebean CA systems where crop residues were retained, and detritus groups such as Collembolla and Symplyla were few under conventional till without residues. Equally no *P. annulatus* worms were recorded under CA with sole bean system where highly transient bean residues had been applied. Indeed, we observed during fauna sampling (eight weeks after cropping), that none of the trash remained at the soil surface, probably due to fast disappearance and rapid decomposition of bean trash. Soil fauna were deprived of food resources as a result hence were low or absent in such systems. Oligochaeta, Araneae,

and Chilopoda were favored with long-term addition of organic residues while Isoptera were favored in the medium-term. In our study sites, earthworms were dominated by the epigeics (those living on top soil layers but forage primarily on plant residuese.g., Dichogaster sp. and P. annulatus) and the endogeics (those living and feeding in the soil but foraging on soil organic matter). The main source of organic matter as food for earthworms include litter from aboveground plant parts, but dead roots and rhizodeposition can also be important food sources (Curry, 2004). Because earthworms are often food limited, their population can increase following organic amendments (Lowe and Butt, 2002; Ayuke et al., 2011b). Higher numbers of Oligochaeta (earthworms) in treatments with organic residues could be attributed to abundant food resources due to crop residues. Importance of soil organic matter as food and energy sources is reinforced by the positive correlations observed between TOC and TON as well as P with some of the macrofauna groups such as Oligochaeta, Diplopoda, and Hemiptera. Exchangeable bases (Na, K, Ca, and Mg) are equally important for some of the fauna groups where they are used in the formation of body parts such as the exoskeletons.

In natural ecosystems, spiders constitute the main invertebrate predatory group hence play an important ecological function in pest control. Araneae being mostly polyphagous predators, can significantly affect the population dynamics of many

TABLE 6 Soil chemical characteristics (0-30 cm soil depth) across	treatments of Embu, Nyabeda, and Kakamega.
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Site	Treatment	Soil pH	тос	TON	Р	Na	к	Ca	Mg
Embu-MT	CT MBi -CR	4.75	2.43a	0.23a	15.87	0.27	0.58	2.23	4.30
	CA SB +CR	4.83	2.45a	0.22ab	14.20	0.24	0.67	2.27	5.25
	CA SM +CR	4.56	2.20b	0.20b	16.67	0.16	0.41	1.77	5.49
	CA MBi +CR	4.54	2.47a	0.18b	16.67	0.17	0.50	1.83	5.46
	P-value	0.428	0.040*	0.016*	0.946	0.491	0.377	0.515	0.362
Kakamega-MT	FP	4.63b	3.26b	0.26b	36.10b	0.02	0.07	0.68b	0.08b
	CT MBi +CR	5.22a	3.57b	0.29ab	4.74b	0.02	0.16	1.34a	0.18a
	CA MBi +CR	4.59b	4.07a	0.32a	93.30a	0.02	0.11	0.59b	0.08b
	P-value	0.023*	0.012*	0.020*	0.011*	0.791	0.162	0.018*	0.014*
Nyabeda-LT	FP	4.95b	2.39	0.18	36.57	0.03	0.09	0.73c	0.10c
	CT MSr +CR	5.39a	2.36	0.18	30.10	0.03	0.05	0.85b	0.20a
	CA MSr +CR	5.18ab	2.40	0.19	27.07	0.02	0.11	0.93a	0.14b
	CA MSi +CR	4.67b	2.35	0.18	30.69	0.03	0.07	0.57d	0.08d
	P-value	0.011*	0.954	0.856	0.181	0.735	0.114	0.004**	<0.001***

For treatment abbreviations (see **Table 2**). TOC, total organic carbon; TON, total organic nitrogen; P, phosphorus; N, sodium; K, potassium; Ca, calcium; Mg, magnesium. Significant p-values are in bold. *Significant; **Highly significant; **Very highly significant.

TABLE 7 Pearson correlation matrix of soil fauna and soil parameters.										
Fauna	Soil pH	тос	TON	Р	Na	к	Ca	Mg		
Macrofauna										
Oligochaeta	-0.20	0.69***	0.62***	0.66***	-0.23	-0.15	-0.28	-0.23		
Isoptera	-0.06	-0.24	-0.30	-0.07	0.11	-0.01	0.04	0.16		
Hymenoptera	-0.23	-0.22	-0.26	-0.19	0.58***	0.39*	0.50**	0.59***		
Coleoptera	-0.17	-0.25	-0.05	-0.10	0.60***	0.56***	0.57***	0.59***		
Diptera	-0.17	-0.19	0.10	-0.12	0.41*	0.26	0.42*	0.31		
Lepidoptera	-0.14	0.08	0.33	0.24	0.47***	0.30	0.38*	0.32		
Diplopoda	-0.18	0.39*	0.38*	0.41*	0.04	0.14	0.03	0.06		
Chilopoda	-0.37*	0.32	0.27	0.06	0.02	0.25	0.11	0.17		
Orthoptera	-0.21	-0.22	-0.13	-0.13	0.16	0.11	0.20	0.44*		
Areneae	-0.09	0.09	-0.07	0.26	-0.18	-0.05	-0.23	-0.16		
Odonata	-0.04	-0.11	0.00	-0.08	0.38*	0.32	0.38*	0.23		
Hemiptera	-0.13	0.44*	0.39*	0.45**	-0.13	-0.04	-0.18	-0.14		
Blattoidea	0.03	-0.19	-0.11	-0.22	0.44*	0.39*	0.34	0.40*		
Isopoda	-0.21	-0.11	-0.05	-0.11	0.22	0.64***	0.28	0.34		
Mesofauna										
Acarina	0.25	-0.14	-0.23	0.12	-0.43*	-0.46**	-0.48**	-0.49**		
Arachnida	-0.07	-0.19	-0.05	-0.12	0.25	0.16	0.23	0.42*		
Collembolla	0.10	0.21	0.08	0.34	-0.40*	-0.37*	-0.41*	-0.42*		
Diplura	-0.06	0.24	0.29	-0.01	-0.10	-0.15	-0.07	-0.14		
Enchytraeidae	-0.20	-0.32	-0.14	-0.31	0.73***	0.48**	0.71***	0.72***		
Protura	-0.13	0.23	0.04	0.19	0.68***	0.61***	0.65***	0.55**		
Symphyla	0.06	0.18	0.09	0.19	-0.49**	-0.47**	-0.50**	-0.53**		

For abbreviations (see **Table 6**). Significant *p*-values are in bold. *Significant; **Highly significant; ***Very highly significant.

phytophagous and saprophagous invertebrates (Ekschmitt et al., 1997; Ziesche and Roth, 2008). In this study, CA most likely provided a conducive and prey-rich environment for Araneae. Chilopods (Centipedes) are among the oldest extant terrestrial arthropods and are an ecologically important group of soil and leaf litter predators (Undheim and King, 2011). In our study, higher Chilopod numbers was observed where maize

stover residues had been applied in medium and long-term. It is possible that maize stover residues, apart from providing a moist conducive environment, might have favored availability of prey for Chilopods. Isoptera (termites) on the other hand, were favored by medium-term addition of organic residues and were therefore more abundant under CA than under CT without residues. Most of the isopteran species sampled in our study sites were all group II trophic members that are fungal growers, and typical feeders of wood, litter and grass. These species build subterranean nests and are sensitive to tillage disturbances (Ayuke et al., 2011b). Being foragers of wood, litter and grass, they were likely favored by addition of crop residues, and to avoid disturbance, they moved to deeper soil depths hence the significant differences observed at 15–30 cm soil depth.

Applications of plant residues have been shown to increase the population of earthworms (Ayuke et al., 2003; Fonte et al., 2009). Although other studies have demonstrated that short-term additions of organic residues increase macrofauna populations, but have little effect on their diversity (Mando, 1998; Ayuke et al., 2003; Ou'edraogo et al., 2006), our study has demonstrated that conservation agriculture and medium to long-term application of organic residues can enhance the richness and abundance of soil fauna, which can in turn promote their activity, hence important soil functions like organic matter retention, stable aggregation and water infiltration (Castellanos-Navarrete et al., 2012; Paul et al., 2015). We speculate that higher soil fauna taxonomic richness in the medium and long-term trials could be due to longterm build-up of soil organic matter as food for the soil fauna groups. Studies conducted by Ayuke et al. (2009, 2011a) showed that earthworm taxonomic richness was higher in high carbon soils than in low carbon soils and this was attributed to longterm application of the various organic amendments across the long-term trials that resulted in corresponding build-up of soil organic matter. The relatively lower macrofauna abundance in the long-term trial of Nyabeda could be attributed to removal of crop residues during soybean planting, which possibly reversed the gains of CA that could otherwise promoted soil organic matter build-up thus food source for the soil fauna. Application of agrochemicals (e.g., pesticides and herbicides) could have also affected the soil organisms as these have been shown to directly affect the organisms through toxicity or indirectly by altering habitat structure and food chain (Gianessi, 2010; Isenring, 2010).

The soil fauna invertebrate community responses to the environmental (soil parameters) changes induced by land management practices and associated ecosystem disturbance reinforce the benefits of conservation agriculture and associated practices (e.g., the use of organic inputs). Soil invertebrates are important determinants of soil chemical and physical characteristics. As such, their potential beneficial role in biodegradation and humification of organic residues, SOM incorporation and soil aggregation in agricultural soils is well-established (Bossuyt et al., 2005; Pulleman et al., 2005a,b; Coq et al., 2007; Fonte et al., 2009; Ayuke et al., 2011b; Brussaard, 2012; Kihara et al., 2015). In view of this, we reiterate that knowledge such as of this study, that demonstrate impact of management practices among them tillage, organic resource use, crop rotation and mixed cropping, and application of inorganic fertilizers on soil fauna is important in enhancing ecosystem functioning and environmental sustainability.

CONCLUSIONS

This study assessed how conservation agriculture, and application of organic or inorganic inputs and cropping system, affected soil fauna richness and abundance. Results showed that: (1) Conservation agriculture enhances soil macrofauna taxonomic richness and mesofauna abundance than conventional agriculture, and that with addition of residues, both diversity and abundance are enhanced, both under CA and CT, (2) Rotation and mixed cropping (intercropping) such as maize legume systems, and sole maize systems coupled with organic residue addition are best bet practices that promote soil fauna diversity and abundance, and (3) Long-term addition of organic residues also enhances soil fauna diversity and abundance more than medium-term addition of organic residues.

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Given the numerous challenges faced by smallholder farmers of SSA in the adoption of CA, who in most cases rarely practice all the three CA principles simultaneously, we propose a further study that will determine how each of the CA components (tillage, organic inputs and cropping systems and their interactions affect soil fauna diversity and abundance.

DATA AVAILABILITY

The datasets generated for this study can be found in **Supplementary Tables** uploaded in the site.

AUTHOR CONTRIBUTIONS

FA carried out the field activities, data analysis, and write up of the manuscript. JK financed the research activities through SIMLESA II project. He is the manager of the Nyabeda trial and also participated in the write up of the manuscript. GA managed the Kakamega medium-term trial and also contributed in the data generation and write up of the manuscript. AM managed the Embu medium-term trials also contributed in the data generation and write up of the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fenvs. 2019.00097/full#supplementary-material

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

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