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Research Article

Effect of Mercury Contamination on the Diversity of Soil Arthropods in Poboya Gold Mining

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Abstract

Background and Objective: Gold mining activities which use mercury have negative impacts on the diversity of soil arthropods. The present study aimed to analyze the diversity of soil arthropods in Poboya gold mining. **Methodology:** This study was an explorative descriptive analytical study. The research location was divided into 6 sampling points from the center of the gold mining area. One kilogram of soil was collected from every sampling point to analyze mercury, C-organic and nitrogen fiber contents of vegetation types. The sampling of soil arthropods used pitfall trap and core sampler. The analysis of difference of diversity, evenness, dominance and richness of arthropod species in every location used one-way ANOVA with level of significance $p < 0.05$. **Results:** There were two classes of soil arthropods found, i.e., insecta and arachnida classes. Class insecta was the most dominant with 21,835 individuals from the total of 23,111 individuals. There was a negative correlation between the diversity of soil arthropods and mercury content of the soil. The higher the mercury contents in the soil, the lower the diversity of soil arthropods. Canonical correspondence analysis (CCA) showed that mercury content and vegetation types affected the diversity of arthropods in the mining areas (Km0, Km1, Km2) and nitrogen and C organic contents affected the diversity of arthropods in Km3, Km4 and Km5. **Conclusion:** Mercury contamination in the soil reduced the diversity of insect and arachnida classes arthropods in the gold mining area.

Key words: Arthropod diversity, class insect, class arachnida, mercury contamination, gold mining area

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Data Availability: All relevant data are within the paper and its supporting information files.

INTRODUCTION

Land use change, especially agricultural lands, leads to reduced biodiversity¹. Increased gold mining activities which use mercury in Indonesia, especially in Central Sulawesi, have caused agricultural land damage and reduced physical and chemical qualities of soil²⁻⁴.

Mercury contamination in soil can change the physical, chemical and biological conditions of soil^{5,6}. It also will affect soil biotas, including soil arthropods because mercury is also known as an active ingredient of synthetic pesticide which can kill insects, thus directly changing composition and community of soil arthropods⁷⁻⁹.

There have been limited studies on the relationship between mercury content in the soil and diversity of soil arthropods⁷. Therefore, it is important to examine the relationship between mercury contamination of the soil and diversity of arthropods because changes of composition, species richness and diversity of soil arthropods will hamper the biological process of the soil as soil arthropods, including insects, collembola, mites, are the first decomposers destroying organic materials into small fragments to be decomposed further by bacteria or fungi.

This study aimed to analyze the diversity of soil arthropods in several levels of mercury contamination around Poboya gold mining area.

MATERIALS AND METHODS

Research type: The research was an explorative descriptive analytical study. The main focus of the study was the correlation between the level of mercury contamination in the soil and diversity of arthropods around Poboya gold mining. This study consisted of field research and laboratory research. The research location was determined purposefully (purposive sampling) by determining sampling points based on the data of previous research indicating that mercury contamination levels in the soil will gradually decrease the farther it is from Poboya goldmine¹⁰.

The research location comprised 6 sampling points:

Km0 = The first location was an area 0 km from the center of the gold mining area (active tromol location; contamination level: highly contaminated)

Km1 = The second location was 1 km (contaminated) from the center of the mining area

Km2 = The third location was 2 km (slightly contaminated) from the center of the mining area

Km3 = The fourth location was 3 km from the center of the mining area

Km4 = The fifth location was 4 km from the center of the mining area

Km5 = The sixth location was 5 km (safe) from the center of the mining area

The sampling points in the field were determined using GPS (global positioning system).

Arthropod sampling:

- Soil arthropods were sampled in each research plot in two ways, i.e. pitfall trap and core sampler
- Sampling and extraction techniques referred to modified Standard Methods for Assessment of Soil Biodiversity and Land Use Practice¹¹
- Pitfall trap was 7×10 cm plastic cup containing 70% alcohol filled 1/3 of it and detergent as necessary. It was plated in the ground until the surface was on the same level as the ground. Pitfall trap was set for 1×24 h. Iron sheet was installed above the pitfall to protect it from rain
- In every transect, there were 10 pitfall traps. After 24 h, the collected arthropods were taken and put in micro tubes containing 70% alcohol, to be taken to laboratory for identification
- Core sampler technique was used to collect arthropods in the ground. Core sampler was plastic tube with 5 cm diameter and 7 cm height. In every transect, there were 10 core samplers
- Soil sample was collected by pressing the tube onto the ground surface until the tube was level with the ground
- The collected soil sample was put in a cloth bag to maintain soil moisture and then taken to laboratory to be processed further using modified barlese tullgren
- All collected specimens were stored in micro tubes containing 70% alcohol

Supporting data observation: The observed supporting data included ground mercury level, nitrogen level, organic material and vegetation type and in each sampling location.

Arthropod identification: All arthropods collected from sampling were identified to species or morphospecies level for any unidentifiable sample. Identification was performed using arthropod identification book¹². This study used two laboratories namely Laboratory of Plant Pests and Diseases of the Faculty of Agriculture and Laboratory of Agrotechnology Faculty of Agriculture, Tadulako University. This study was conducted for 8 months in 2017 from January-August, 2017.

Data analysis: Data analysis of species richness, abundance, dominance and evenness as well as diversity of soil arthropods used Shannon Wiener Index and Simpson Index. The relationships between various soil environmental factors such as soil mercury level, nitrogen level, organic material and vegetation, in affecting the diversity of arthropods were analyzed by Canonical Correspondence Analysis (CCA), while similarity of arthropod species in every location was analyzed by cluster analysis based on Morista similarity index. The entire analysis used PAST software.

The analysis of differences of diversity, evenness, dominance and species richness of arthropods in every location used one-way-ANOVA dan DUNCAN (DNMRT) with level of significance $p < 0.05$.

RESULTS

The soil arthropods found in the present study consisted of two classes, i.e., Insecta and Arachnida. From the two classes, insecta was the most dominant with 21,835 individuals from the total of 23,111 individuals. There were 62 species from 28 families and 10 orders. *Entomobrya sp1.* is a species that was most commonly found at the mine site, which was 4,260 individuals and followed by *Solenopsis geminata* with 2,524 individuals (Table 1).

Table 1 also showed a trend that the further from the center of the mine, the higher the numbers of individuals, species and families. However, the number of the orders found tended to be similar in all sampling locations.

Further analysis showed that species richness, total individual, dominance, evenness and diversity of insects also decreased along with distance from the center of the mine. In the mine location (Km0), species richness and total individual were lower and significantly different from other locations while evenness was lower and dominance was higher. The diversity index was also lower in Km0 and different from other locations when using Shannon Wiener index, as well as Simpson index (Table 2).

Soil analysis showed that mercury level in the mining mine location was higher than that in locations further from the center of the mine (Table 3). There was a negative correlation between the diversity of soil arthropods and soil mercury content. The higher the mercury level in the soil, the lower the diversity of soil arthropods (Fig. 1). Besides mercury, there might had been other environmental factors which affected the diversity of soil arthropods. This was evident in multidimensional scaling analysis by Canonical Correspondence Analysis (CCA) which showed that mercury level and vegetation type affected arthropod diversity in mine locations (Km0, Km1 and Km2). Meanwhile, nitrogen and C organic contents affected arthropod diversity 3.4 and 5 km away from the mining location is the right term (Fig. 2).

Table 1: Class, order, family, species and total individual of every arthropod species by distance from gold processing location

Order	Family	Number species	Sampling points							Total
			Km0	Km1	Km2	Km3	Km4	Km5		
Class insecta										
1 Hymenoptera	1 Formicinae	1 <i>Camponotus sp1.</i>	0	13	0	0	0	55	68	
		2 <i>Camponotus sp2</i>	0	3	2	4	3	0	12	
		3 <i>Oeophylla smaragdina</i>	0	0	89	32	15	37	173	
		4 <i>Paratrechina longicornis</i>	382	302	362	322	286	683	2337	
	2 Ponerinae	5 <i>Diacamma rugosum</i>	0	8	36	20	9	90	163	
		6 <i>Odontomachus sp.</i>	0	17	29	4	0	0	50	
	3 Dolichoderinae	7 <i>Dolichoderus thoracicus</i>	0	16	0	12	8	12	48	
		8 <i>Iridomyrmex sp1</i>	0	22	0	0	0	88	110	
		9 <i>Iridomyrmex sp2</i>	0	11	25	5	11	54	106	
		10 <i>Tapinoma melanocephalum</i>	11	0	12	5	410	186	624	
		11 <i>Tapinoma sp.</i>	0	45	11	88	473	0	617	
		12 <i>Technomyrmex sp.</i>	0	0	21	0	13	24	58	
		13 <i>Monomorium floricola</i>	30	86	143	50	29	190	528	
	4 Myrmicinae	14 <i>Monomorium sp1.</i>	0	25	8	0	0	0	33	
		15 <i>Monomorium sp2.</i>	0	0	37	0	0	153	190	
		16 <i>Oligomyrmex sp.</i>	0	0	22	0	0	5	27	
		17 <i>Pheidole sp1</i>	0	15	25	0	43	33	116	
		18 <i>Pheidole sp2</i>	0	0	0	0	9	53	62	
		19 <i>Solenopsis geminata</i>	294	396	253	194	335	1052	2524	
		20 <i>Tetramorium bicarinatum</i>	0	116	223	382	398	276	1395	
		21 <i>Tetramorium sp.</i>	0	0	15	0	0	61	76	
	5 Aenictinae	22 <i>Aenictus sp.</i>	0	0	0	135	105	153	393	
22 <i>Aenictus sp.</i>		0	0	0	135	105	153	393		

Table 1: Continue

Order	Family	Number species	Sampling points							
			Km0	Km1	Km2	Km3	Km4	Km5	Total	
2 Coleoptera	6 Cerapachyinae	23 <i>Cerapachys</i> sp.	0	0	0	110	111	128	349	
	7 Pompilidae	24 <i>Priocnemis</i> sp.	0	53	173	0	0	43	269	
	8 Encyrtidae	25 Encyrtidae sp1.	0	18	11	52	77	31	189	
	9 Reduviidae	26 Reduviidae sp1.	15	27	65	0	32	0	139	
	10 Carabidae	27 <i>Amara</i> sp.	20	19	0	36	0	0	75	
		28 <i>Carabidae</i> sp1	0	18	27	12	2	0	59	
		29 <i>Carabidae</i> sp2.	5	10	9	31	0	11	66	
		11 Scarabaeidae	30 <i>Clivina</i> sp.	23	14	41	0	21	0	99
	3 Diptera	12 Sciaridae (agas)	31 <i>Sciara</i> sp.	0	15	47	28	21	39	150
		13 Cecidomyiidae	32 <i>Lestremia</i> sp.	0	22	85	0	0	104	211
	14 Muscidae	33 <i>Musa domestica</i>	31	42	49	0	21	0	143	
4 Hemiptera	15 Cicadidae	34 <i>Tibicen</i> sp. (nymph)	0	19	25	71	22	0	137	
	16 Aphididae	35 <i>Aphis fabae</i>	18	28	56	41	17	32	192	
5 Dermaptera	17 Cacinophorida	36 <i>Chelisoches</i> sp.	1	21	0	0	0	77	99	
6 Orthoptera	18 Gryllidae	37 <i>Allonemobius</i> sp.	0	52	158	10	0	49	269	
7 Isoptera	19 Rhinotermitidae	38 <i>Captotermes</i> sp1	14	5	23	128	0	145	315	
		39 <i>Captotermes</i> sp2.	0	15	16	19	32	57	139	
8 Collembola	20 Entomobryidae	40 <i>Entomobrya</i> sp1.	529	421	589	857	878	986	4260	
		41 <i>Entomobrya</i> sp2.	0	21	40	89	129	247	526	
		42 <i>Entomobrya</i> sp3.	12	0	120	147	48	187	514	
		43 <i>Entomobrya</i> sp4.	0	73	69	49	109	54	354	
		44 <i>Lepidocyrtus</i> sp.	12	27	12	20	0	123	194	
		45 <i>Coenalestes</i> sp12.27	22	12	11	137	102	85	369	
		46 <i>Pogognatelus</i> sp.	0	10	21	8	23	0	62	
		47 <i>Homidia</i> sp.	0	23	65	22	0	12	122	
	21 Isotomiidae	48 <i>Isotoma viridis</i>	24	99	145	87	82	122	559	
		49 <i>Isotoma</i> sp.	8	29	80	50	64	68	299	
		50 <i>Folsomia candida</i>	11	58	65	25	34	69	262	
	22 Neanuridae	51 <i>Anurida granaria</i>	0	34	124	98	103	173	555	
	23 Hypogastruidae	52 <i>Podura acuatica</i>	0	48	92	71	0	71	282	
		53 <i>Hypogastura</i> sp.	38	77	95	26	45	49	330	
		54 <i>Ceratophysella</i>	0	0	29	11	22	59	121	
		55 Unidentified	20	25	99	94	12	0	250	
9 Diplura	24 Japygidae	56 <i>Metajapyx</i> sp.	0	21	43	0	79	23	166	
Class arachnida										
10 Araneae	25 Oxyptidae	57 <i>Pardosa</i> sp.	0	0	0	0	22	105	127	
	26 Salticidae	58 <i>Cormophasis</i> sp.	8	0	0	31	0	89	128	
	27 Loxoscelidae	59 <i>Loxosceles</i> sp.	34	12	105	74	21	45	291	
	28 Acari	60 <i>Diaptorebates notatus</i>	20	42	75	112	110	122	481	
		61 <i>Tetranychus</i> sp.	0	12	22	8	22	0	64	
		62 Unidentified	17	0	11	65	68	24	185	
Number of individual (Class Insecta)			1520	2431	3797	3582	4233	6249	21835	
Number of individual (Class Arachnida)			79	66	213	290	243	385	1276	
Total individual			1599	2497	4010	3872	4476	6634	23111	
Number of orders			8	10	8	8	8	10		
Number of families			14	21	19	14	18	20		
Number of species			25	48	52	45	44	49		

Table 2: Species richness, total individual, dominance, evenness and diversity arthropod in all sampling point from the center of poboya gold mining area

Parameters	Sampling point (km)					
	0	1	2	3	4	5
Species richness	25 ^a	48 ^b	52 ^c	45 ^c	44 ^{bc}	49 ^d
Total individual	1,599 ^a	2,497 ^a	4,010 ^{ab}	3,872 ^b	4,476 ^b	6,634 ^c
Dominance index	0.27 ^a	0.14 ^b	0.082 ^c	0.11 ^{bc}	0.11 ^{bc}	0.08 ^c
Evenness Index	0.60 ^a	0.66 ^{ab}	0.75 ^b	0.60 ^a	0.63 ^a	0.60
Shannon-wiener diversity index	1.67 ^a	2.37 ^b	2.77 ^{cd}	2.64 ^{bc}	2.55 ^{bc}	2.92 ^d
Simpson diversity index	0.73 ^a	0.86 ^b	0.91 ^c	0.88 ^{bc}	0.88 ^{bc}	0.91 ^c

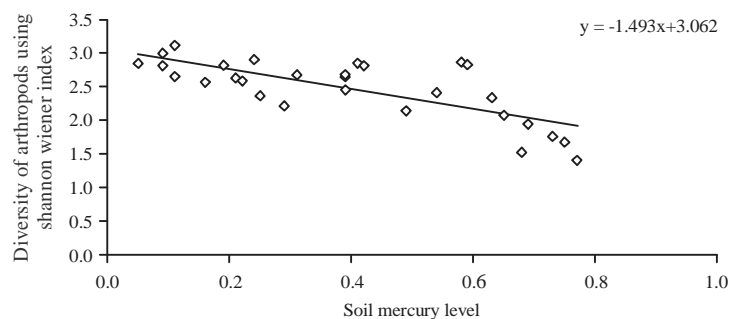


Fig. 1: Linear regression showing the relation between soil mercury level and diversity of arthropods using Shannon Wiener Index
 → Nilai regression coefficient (R²)

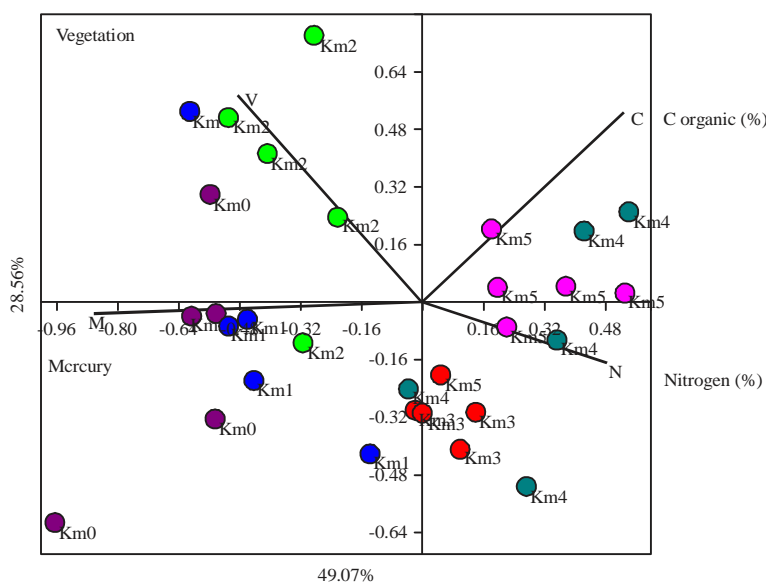


Fig. 2: Canonical correspondence analysis (CCA) diagram showing the effects of various environmental factors on the diversity and abundance of soil arthropods in different distances from the center of the mine

Table 3: Analysis results of mercury, nitrogen, c organic contents of soil and vegetation diversity in every arthropod sampling point

Sampling points	Environmental parameters			
	Mercury (mg L ⁻¹)	Nitrogen (%)	C organic (%)	Vegetation diversity*
Km0	0.73	0.08	0.91	4.2
Km1	0.58	0.16	1.44	8.0
Km2	0.42	0.21	2.62	12.8
Km3	0.26	0.21	2.01	5.4
Km4	0.21	0.18	2.04	4.6
Km5	0.11	0.21	2.32	5.6

Figure 3 confirmed that mercury contamination in the soil strongly affected the diversity of soil arthropods. Arthropod diversity 5 km (Km5) away from the center of the mine or location with low mercury level clustered separately, while locations 1-4 km (Km1, Km2, Km3, Km4) away from the center of the mine had similarities with arthropods in the mine location (Km0).

DISCUSSION

Gold processing activities which use mercury in the amalgamation process in Poboya goldmine is the main source of pollutant of the soil around the mine. It has high mercury contamination which exceeds the safety threshold. Soil analysis in the sampling locations showed that the soil

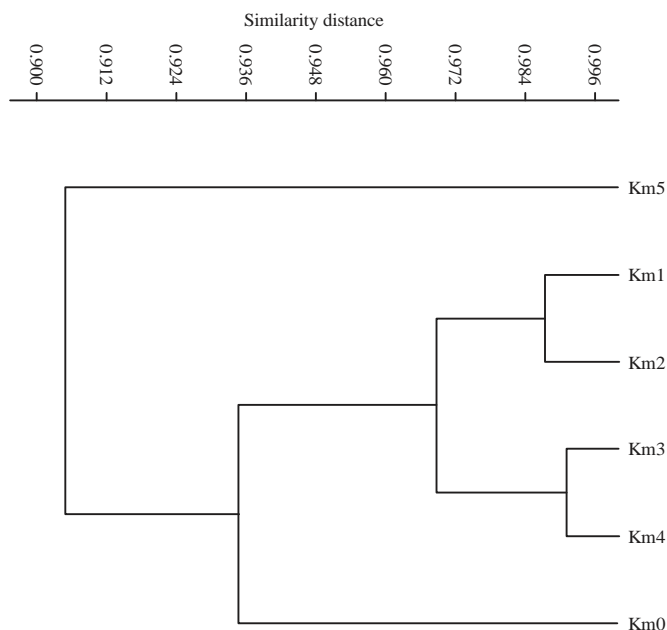


Fig. 3: Cluster analyses of arthropods by distance from the center of the mine

Cluster I: Km0 = The first location was an area 0 km from the center of the gold mining area (active tromol location; contamination level: highly contaminated), Cluster II: Km1 = The second location was 1 km (contaminated) from the center of the mining area, Km2 = The third location was 2 km (slightly contaminated) from the center of the mining area, Km3 = The fourth location was 3 km from the center of the mining area, Km4 = The fifth location was 4 km from the center of the mining area, Cluster III: Km5 = The fifth location was 5 km (safe) from the center of the mining area

mercury levels were 0.05-0.77 mg kg⁻¹, exceeding 0.01 mg kg⁻¹ of soil. The same thing is reported by Sari *et al.*¹⁰ and Mirdat *et al.*⁴. Leiva and Morales¹³ and Luo *et al.*¹⁴ also reported that gold mining activities which use amalgamation technology are sources of mercury contamination of the soil.

Mercury contamination in the ground can last a long time because it can be bound with other elements to create new compounds which remain toxic or even more toxic and dangerous to life, e.g., organic mercury methylmercury¹⁵⁻¹⁷. The formation of methylmercury was caused by bacterial activities, but it was reported that termites can also methylate mercury into methylmercury¹⁸.

Several species of soil arthropods can live or tolerate soil with high mercury level, especially ground-surface arthropods¹⁹. However, from 62 species found, only several species from the family formicidae (ant) and family entomobryidae (Collembola) were found in abundance in the mining location. *Paretrechina longicornis*, *monomorium floricola* and *Solenopsis geminata* ants were collected in large numbers. The presence of the three ant species in high soil mercury level showed that they have high adaptability compared to other ant species. A study by Hasriyanty *et al.*²⁰ showed that the three ant species can be found in habitats contaminated by mercury.

Another arthropod species found in abundance was collembola. There were 16 species from the order of

collembola which were found and 10 of them were found in the mining location. However, only one species from the family entomobryidae was found in abundance. The high number of collembola species found might be because the species have high adaptability and tolerance to mercury contamination or because there were many species with different roles in the ecosystem. Besides the family entomobryidae, some species from the family isotomiidae and hypogastridae were also found in the mine location.

The effects of mercury on arthropods, particularly insects, are often used to study bioaccumulation process on terrestrial insects. Some of them are toxicology effects on *Locusta migratoria* (Coleoptera: Acrididae)²¹, *Folsomia candida* and *Proisotoma minuta* (Collembola: Isotomidae)⁷; as well as *Blatella germanica* (Blattaria: Blattellidae)²². It has been reported that in sicada, bioaccumulation of mercury in male insects is higher than that in female insects^{23,24}.

This study has shown that gold mining activities reduce the diversity of soil arthropods. Thus, it can be proposed that rehabilitation is needed to reduce mercury pollution in the mining area. One alternative is the bioremediation by planting multiplying plants that can accumulate mercury to the soil. Isrun *et al.*³ showed that the tithonia diversifolia compost can be used for the recovery of agricultural land and plants contaminated by Hg²⁺. This study has a limitation that is the wide range of the observation areas.

CONCLUSION

Based on the findings, it can be concluded that mercury contamination in the soil reduced the diversity of soil arthropods. There was a correlation between the diversity of arthropods and soil mercury content. The closer to the center of the mine (higher soil mercury level), the lower the diversity of arthropods. From 62 species found, only 25 species were collected from the gold mining area.

SIGNIFICANCE STATEMENT

This study indicates that mercury usage in gold mining activities reduced classes of insecta and arachnida arthropods. This study will help the researchers uncover critical areas of soil arthropod types which can live or are tolerant to soil with high level of mercury contamination, especially ground-surface arthropods. Thus, the high mercury level of soil showed that ant species of *Paratrechina longicornis*, *Monomorium floricola* and *Solenopsis geminata* have a high adaptability level.

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