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Reproductive damage and compensation of wild earthworm *Metaphire californica* from contaminated fields with long-term heavy metal exposure

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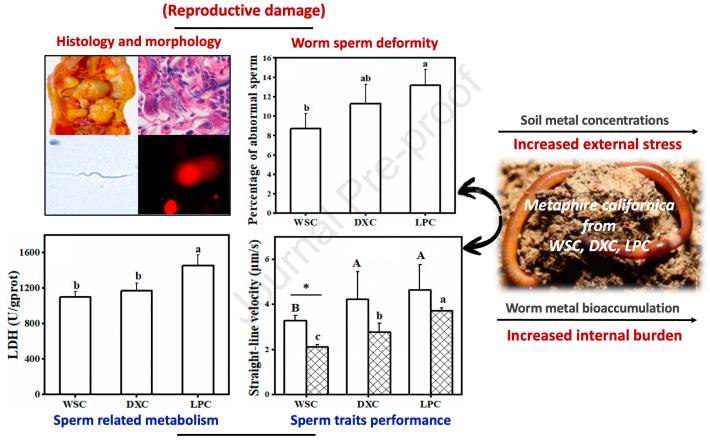
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## Author contribution statement

Caide Huang: Conceptualization, Writing-original draft. Zhiqiang Shen: Methodology.
Liang Li: Methodology. Shizhong Yue: Writing - review & editing. Li Jia: Writing - review
& editing. Kun Wang: Writing - review & editing. Wenhao Zhou: Methodology. Yuhui Qiao:
Conceptualization, Funding acquisition, Supervision.

...on, Supervision.

## **Graphical abstract:**



(Reproductive compensation)

1	Reproductive damage and compensation of wild earthworm Metaphire
2	californica from contaminated fields with long-term heavy metal exposure
3	Caide Huang <sup>a, b</sup> , Zhiqiang Shen <sup>a</sup> , Liang Li <sup>a</sup> , Shizhong Yue <sup>a, c</sup> , Li Jia <sup>a, d</sup> , Kun Wang <sup>a, e</sup> ,
4	Wenhao Zhou <sup>a</sup> , Yuhui Qiao <sup>a, f,</sup> *
5	<sup>a</sup> College of Resources and Environmental Science, China Agricultural University, Beijing 100193, China
6	<sup>b</sup> Centre for Ecology and Hydrology, Maclean Building, Benson Lane, Crowmarsh Gifford, Wallingford,
7	Oxon, OX10 8BB, UK
8	<sup>c</sup> Shandong Provincial Key Laboratory of Biophysics, Institute of Biophysics, Dezhou University,
9	Dezhou 253023, China
10	<sup>d</sup> Centre National de la Recherche Scientifique, Institut des Sciences de la Terre D'Orleans, Université
11	d'Orleans-Brgm, UMR 7327, France
12	<sup>e</sup> State Key Laboratory of North China Crop and Regulation, College of Resources and Environmental
13	Science, Hebei Agricultural University, Baoding 071001, China
14	<sup>f</sup> Beijing Key Laboratory of Biodiversity and Organic Farming, China Agricultural University, Beijing
15	100193, China
16 18	
19	*Corresponding author: Yuhui Qiao, E-mail: qiaoyh@cau.edu.cn, Tel.: +86-10-62731166, Fax: +86-
20	10-62731165, Address: Yuan-Ming-Yuan West Rd. 2#, Beijing 100193, China
21	

# 22 Abstract

23	Reproduction is a significant biological process for organisms responding to
24	environmental stresses, however, little is known about the reproductive strategies of
25	invertebrates under long-term exposure to contaminations. In this study, earthworm
26	Metaphire californica (Kinberg, 1867) from contaminated fields with an increased
27	metal gradient were collected to investigate their reproductive responses. The results
28	showed heavy metals (Cd, Cu, Zn, and Pb) induced histological damage to earthworms'
29	seminal vesicles, including tissue disorders and cavities, and decreases in mature
30	spermatozoa. Sperm morphology analysis indicated deformity rates were up to13.2%
31	(e.g. head swollen or missing) for worms from the most contaminated site, which
32	coincided with DNA damages. Furthermore, the computer-assisted sperm analysis
33	(CASA) system was employed for the evaluation of sperm kinetic traits. Results
34	suggested earthworms exposed to higher contamination showed a lower sperm viability
35	rate but faster sperm velocity after re-exposure with Cd solution (like the curvilinear
36	velocity and straight-line velocity paraments) compared with those from relatively
37	clean sites. The activities of lactate dehydrogenase and sorbitol dehydrogenase showed
38	the highest 32.5% and 12.5% up-regulation respectively with the increased metal
39	gradient. In conclusion, this study elucidated the earthworm reproductive toxicity,
40	underlying reproductive compensation, metal stress-induced damages, and adaptive
41	responses caused by heavy metal exposure, while also providing the possibility of
42	sperm trait analysis (CASA) for related earthworm toxicological studies.

43 *Capsule:* Earthworms under long-term heavy metal stresses coordinate more energy

44 metabolism for sperm kinetic traits to compensate for reproductive damage.

*Keywords:* Earthworms; seminal vesicles; morphology; sperm traits; reproductive
metabolism

47

## 48 **1. Introduction**

49 In recent decades, existing and emerging environmental issues have been 50 enhanced by rapid urbanization and industrialization, from the pathways of atmospheric 51 deposition, mining activities, chemical waste emission, pesticide and fertilizer abuses 52 (Navarro et al., 2016; Gupta et al., 2021). Heavy metals are one of the most consistent 53 and ubiquitous pollutants posing significant ecological risks and therefore requiring careful consideration (Wu et al., 2022). Once entering the environment, non-54 55 biodegradable metals can be bioaccumulated, transferred to trophic levels, cause 56 biological toxicity, and eventually interfere with normal biogenesis and public health 57 (Huang et al., 2020; Yaashikaa et al., 2022). Consequently, using terrestrial and aquatic 58 organisms (like earthworms and zebrafishes) to carry out metal risk assessments is 59 always the foreword research in environmental science and toxicology (Chen et al., 60 2021; Sivakumar, 2015).

Earthworms are regarded as "soil ecological engineers" because of their critical
dwelling and burrowing roles in soil improvements (Morgan and Morgan, 1999).

3

63	Furthermore, earthworms are typical sentinel invertebrates and have been widely
64	recommended as model test species in OECD No. 207 and ISO guidelines for risk
65	assessments of various soil pollutants (ISO, 2012; Morgan et al., 2007; OECD, 1984).
66	At present, various individual indexes and biomarkers have been applied to assess the
67	fitness and response of earthworms under hazardous exposure. These include $LC_{50}$ (50%)
68	lethal concentration) and EC50 (50% effective concentration) values for survival rate
69	and individual growth, antioxidant enzyme activities and gene expression for biological
70	processes (Spurgeon et al., 2020; Spurgeon and Hopkin, 1995). Unlike other organisms,
71	earthworms are hermaphrodites, simultaneously keeping male and female reproductive
72	organs and systems, which provides an excellent model to evaluate reproductive
73	toxicity (Domínguez and Velando, 2013; Kwak and An, 2021a). However, up to now
74	related indicators for reproductive studies on earthworms and other invertebrates are
75	very limited (Yesudhason et al., 2013). For instance, Yang et al. (2022) reported that the
76	earthworm cocoon numbers decrease by 16.7% after exposure to 100 mg/kg carbon
77	nanotubes, whilst other studies also used conventional recordings like juveniles, and
78	hatching rates for reproduction assessments (Huang et al., 2021a).

The reproductive system is essential for organisms under unfavorable conditions to ensure their specie and population continuation (Taslima et al., 2022). Related indicators like gene expression, reproductive organ changes, sperm morphology and mobility have been proven relatively more sensitive for assessing the reproductive toxicity of some vertebrates exposed to different environmental stressors (Kime et al.,

84	1996; Kime et al., 2001; Kwak and An, 2021b; Yang et al., 2019). Studies using these
85	novel biomarkers within earthworm toxicological studies are still rare, although the
86	development of microscopic and computer-assisted sperm analysis (CASA) systems
87	provides more novel determinations. CASA has been the dominant method for sperm
88	quality estimation and provides possibilities for rapid in vivo monitoring and
89	comparison of sperm kinetic traits of different animals like rats and birds (Yang et al.,
90	2019; Waberski et al., 2021). Despite this, as most animal spermatozoa will be rapidly
91	inactivated in vitro, appropriate dilution and capacitation liquid must be maintained to
92	ensure the activities and movement of spermatozoa before CASA detection (Travis et
93	al., 2004), presenting one of the main challenges for the detection and assessment of
94	earthworm sperm quality.

Furthermore, most previous reproductive toxicity studies were validated via 95 laboratory microcosm exposures, while few studies were devoted to considering those 96 organisms under the sustained stresses of *in-situ* field contaminations (Gao et al., 2015; 97 98 Kwak and An, 2021b; Yang et al., 2019). Normally, reproductive regulation is an 99 alternative response for certain organisms under stress from long-term exposure 100 through increased reproductive efforts with energy trade-offs to allow reproductive success (Gowaty et al., 2007; Orton and Tyler, 2015; Yang et al., 2020a). The stress-101 102 induced resource redistribution and reproductive compensation that is driven by 103 environmental tolerance, phenotypic plasticity, and even parental inheritance have not been clearly elucidated (Huang et al., 2022). Accordingly, it is of importance to 104

107 earthworm *Metaphire californica* exposed to heavy metal contaminations over several
108 decades, possessed more considerable population dominance, bioaccumulation, and
109 better detoxification capacity, especially for Cd resistance and detoxification in
110 comparison to other wild species, providing an advisable research candidate (Huang et
111 al., 2021b; Huang et al., 2020; Wang et al., 2020; Wang et al., 2018a).

112 Overall, it could be hypothesized that wild earthworms under persisting 113 contamination can respond to heavy metal stress by reproductive compensation. The 114 present study was carried out to systematically investigate the reproductive damage and 115 compensation in wild earthworm *M.californica* from different long-term heavy metal 116 exposure histories (Cd, Cu, Zn, and Pb). The detailed purposes of this study were: (1) 117 to employ the CASA system to evaluate the sperm traits and performance of earthworm 118 M.californica; (2) to determine the reproductive toxicity of heavy metals on 119 earthworms via histology, morphology and DNA damage analyses; (3) to explore the 120 reproductive compensation by assessing the sperm traits and metabolic levels. These 121 findings can shed light on the dynamic reproductive strategies of those organisms living 122 under persistent stresses.

123 **2.** Materials and methods

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106

124 2.1 Description of sampling sites

125	All three sampling sites Wanshou (WSC, 112° 40' 05"; 26° 44' 49"), Daxing (DXC,
126	112° 47' 18"; 26° 54' 21"), and Lupu (LPC, 112° 79' 50"; 27° 00' 07") (Fig. 1) were
127	located in the "hometown of nonferrous metals", Hengyang City, Hunan Province, the
128	subtropical areas of southern China. These sites showed different distances away from
129	mining or tailing areas with over 30 years of pollution history, which were all confirmed
130	as long-term observing and sampling points by our field survey studies began in 2018
131	(Huang et al., 2020; Wang et al., 2018b; Wang et al., 2018c).
132	The five-spot-sampling method (50 cm $\times$ 50 cm $\times$ 25 cm) was applied to sample
133	earthworms and soils (0-25 cm) from each site. The heavy metal contaminations totally
134	followed an increasing gradient from WSC to DXC and LPC sites, especially for soil
135	Cd, which was 1.50, 5.08, and 12.6 mg/kg, respectively. According to our previous
136	studies (Huang et al., 2020; Huang et al., 2021b), only heavy metals of Cd, Cu, Zn, and
137	Pb exceed the Chinese Environmental Quality Standard for Soils (GB 15618-2018),
138	which were mainly considered in our study. Detailed site information, soil properties,
139	and metal concentration are provided in Table S1, and related descriptions of these three
140	sites also can be found in our previous studies.

141 2.2 Test earthworms

Adult *Metaphire californica* with similar weight (500-550 mg) and fully developed clitellum were collected from each site. Earthworms were kept in *in-situ* soils and fed with their original surface litter (leaves and humus), and all subsequent analysis was done within two weeks to reduce the interferences. Part of the worms was

depurated on the Petri dish with moist filter paper for 24 h, washed with deionized water,

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147 frozen in liquid nitrogen, and then dried in a freeze dryer (TF-FD-18S, China) to 148 analyze metal bioaccumulation. Rest living earthworms were used for further dissection 149 and reproductive analysis. Meanwhile, M.californica was ever confirmed in our former 150 studies for species classification with morphological analysis and the sequencing of 151 cytochrome oxidase subunit first (COI) genes (Huang et al., 2021b; Xu and Xiao, 2010). 2.3 The analysis of soil properties and heavy metals 152 153 Soil pH, soil organic carbon (SOC), total nitrogen (TN), cation exchange capacity (CEC), and heavy metals (Cd, Cu, Zn, and Pb) were analyzed to assess the overall soil 154 conditions of different sampling sites, following the protocols in our previous study and 155 meeting the certificated requirements of detection limits and recovery rates (Huang et 156 al., 2020). 157 158 Freeze-dried worms were ground into powder and digested in a microwave with 159 a mixture of HNO<sub>3</sub>-H<sub>2</sub>O<sub>2</sub>, and metal concentration were analyzed through inductively 160 coupled plasma-mass spectrometry (ICP-MS, 7700ce, USA). Shrimp powder (GBW 161 10050) was used as standard reference material and all of the recovery rates were over 162 95%. Besides, the bioaccumulation factor (BAF) value was calculated by dividing worm metal concentration into soil metal contents, which represents the ability of 163 164 earthworms to bioaccumulate different heavy metals (Huang et al., 2022).

165 2.4 Histology and morphology examination

166	Depurated M.californica was fixed with pins on a smooth foam and two pairs of
167	spermathecal pores were located in the 7th to 9th lateroventral intersegment (Nguyen
168	et al., 2020; Xu and Xiao, 2010). Dissected and gently tore the septa along the
169	abdominal midline of worms from anterior to clitellum segments, separated out the
170	seminal vesicles, then rinsed with ultrapure water to remove adhering mucus, finally
171	fixed in 10% formaldehyde solution for 48h. Fixed vesicles were dehydrated with a
172	graded series of ethanol with different concentration and cleared with a xylene agent to
173	completely displace ethanol. Afterward, all vesicles were embedded with paraffin,
174	sliced into films, and fully stained with haematoxylineosin (HE). All prepared slides
175	were finally observed using an optical microscope (ZEISS-AX10, Zeiss, GmbH,
176	Germany) and image-recorded under different magnifications to determine histological
177	changes (Adeel et al., 2019; Huang et al., 2021a; Kwak and An, 2021a).
178	Separated seminal vesicles of at least 10 earthworms from the same treatment were
179	put into a 1.5 ml centrifuge tube to prepare earthworm seminal suspension by exerting
180	pressure and forcing out the sperm from vesicles with fine-tipped forceps. 10 $\mu L$
181	earthworm semen and 10 $\mu$ L phosphate-buffered saline (pH=6.8) was placed in the
182	center of slides, then dispersed and smeared by overlapping another slide. All slides
183	were immersed in 10% formaldehyde solution for 15 min fixing and then stained for
184	25 min using 10% Giemsa staining stock (Solarbio, China), and finally a slow flow was
185	used to rinse off the excess staining solutions prior to air-drying. Prepared slides were
186	observed using an optical microscope (ZEISS-AX10, Zeiss, GmbH, Germany) under

9

400× magnification (Gao et al., 2015; Yesudhason et al., 2013; Zheng and Li, 2009).
All earthworms from each site were tested as above and photo-recorded. The sperm
deformity rate was calculated by dividing the number of abnormal sperm into total
numbers, and over 100 spermatozoa were randomly selected to examine the abnormal
rates of each treatment.

192 2.5 Assessment of earthworm sperm trait performance

In this study, glucose (Glc) was proven as the best sperm capacitation liquid for *M.californica* compared with more common-used PBS and HBSS (Hank's Balanced Salt Solution) (Travis et al., 2004; Williams and Ford, 2001; Yang et al., 2020a), which successfully capacitated earthworm sperms and make them move in straight or curved lines (Fig. S1, A). Earthworm sperms were diluted by 20 times with 2.5% Glc solutions (30°C, adjusted pH to 7) to avoid sperm overlap and ensure the appropriate sperm amount in each viewing field for trait analysis.

Specifically, 10 µL seminal suspension was diluted into 190 µL Glc and blown 200 201 with a pipette until all sperm was completely diffused into the surrounding medium. 202 After 10-min dissociation and thermostat, 10 µL suspension was dropped on the slide 203 center (pre-warmed under 30°C) and then applied with coverslips for later analysis 204 (Kime et al., 1996; Kime et al., 2001; Niu et al., 2022; Yang et al., 2019). All earthworm 205 sperm traits were assessed under a computer-assisted sperm analysis (CASA, LINE-HY, China) system, provided by the College of Animal Science and Technology, China 206 207 Agricultural University, which has been successfully applied for sperm analysis on

different animals. A total of 300 earthworm sperms from at least 5 nonconsecutive
viewing fields of each treatment were analyzed to comprehensively evaluate the sperm
traits.

The reproductive performances of earthworms from WSC, DXC, and LPC sites were determined by comparing sperm density, sperm viability, and also several movement parameters: (a) velocity parameters, including curvilinear velocity (VCL), straight-line velocity (VSL), and average path velocity (VAP); (b) trajectory parameters, including linearity (LIN) and wobble (WOB) percentages (Kime et al., 2001; Yang et al., 2019).

In order to directly assess the impacts of heavy metals on earthworm sperm traits, as well as to further confirm the elevated Cd resistance of earthworms reported in our previous studies (Huang et al., 2020; Huang et al., 2021b), another lab re-exposure was carried out to conduct the comparative analysis and evaluate the sperm traits continuity by using 0.1 mg/L Cd-spiked Glc, which could make sperms visibly affected but not wholly inactivated (Fig. S1).

223 2.6 DNA damage and metabolism evaluation

DNA damage of earthworm sperms was determined by the single-cell gel electrophoresis assay (also entitled comet assay), which was described in detail in our and other studies (Huang et al., 2021a; Li et al., 2020; Wang et al., 2018d). Briefly, dissected seminal vesicles were suspended with cold PBS buffer, then centrifuged to obtain the sperm cell suspensions for slide preparation by employing "three-steps"

229	agarose and electrophoresis preparing methods. Processed slides were dyed with
230	propidium iodide buffer, finally observed and photographed under a fluorescence
231	microscope. Randomly selected DNA cells were analyzed by using Comet Assay
232	Software Project (CASP) software with the values of tail DNA proportion (Tail DNA %)
233	and olive tail moment (OTM). Damaged DNA will show typical comet tails with the
234	Tail DNA% $> 5$ and OTM $> 2$ values (Fig. S2).
235	Sperm-related metabolism was evaluated by determining the activities of lactate
236	dehydrogenase (LDH) and sorbitol dehydrogenase (SDH), which were both significant
237	biomarker enzymes involved in energy acquisition, sperm production, and germ cell
220	development (Veneration 2020h, 7hz et al. 2010). I DIL et d'ODU

Sperm-related metabolism was evaluated by determining the activities of lactate dehydrogenase (LDH) and sorbitol dehydrogenase (SDH), which were both significant biomarker enzymes involved in energy acquisition, sperm production, and germ cell development (Yang et al., 2020b; Zhu et al., 2019). LDH and SDH were determined at 450 and 600 nm via a microplate reader (XS2, BioTeK, USA), respectively, by using seminal vesicle cells and commercial enzyme kits (Nanjing Jiancheng Bioengineering Institute, Nanjing, China), following the manufacturer's specifications.

## 242 2.7 Statistical analysis

All data analyses were performed with IBM SPSS 24.0 by using Tukey's honest significant difference (HSD) test. Data were expressed as means  $\pm$  standard deviations with *P* < 0.05 representing significant differences. Pearson's correlations analyses were employed to measure the association between worm metal bioaccumulation and each reproductive parameter (Fig. S3). Tables and figures were completed in Excel 2016 and Origin 2020.

## 249 **3. Results**

## 250 3.1 Heavy metal bioaccumulation in M.californica

251	The internal metal concentration of <i>M.californica</i> and bioaccumulation factor
252	values (BAF) are shown in Table 1. The metal bioaccumulation in earthworms showed
253	consistency with their exposure history. M.californica from the LPC site showed the
254	highest Zn and Pb bioaccumulation compared with those from WSC and DXC sites,
255	which were 239 and 38.4 mg/kg, respectively. Furthermore, LPC M.californica also
256	suffered the highest body burden for soil Cd, which was 3.64 and 1.43 times higher
257	than that of WSC and DXC worms, respectively. M.californica also bioaccumulated
258	other heavy metals (Cu and Pb), but differences were found to be non-significant
259	between WSC and LPC worms ( $P > 0.05$ ; Table 1). For bioaccumulation factor values,
260	almost only BAF <sub>cd</sub> values were larger than 1.0, indicating the significant ability of
261	earthworms to bioaccumulate soil Cd. Specifically, the BAF <sub>cd</sub> values were 11.2, 8.43,
262	and 4.83 for WSC, DXC, and LPC worms, respectively, showing a significant decrease
263	with the increase of heavy metal concentration ( $P < 0.05$ ; Table 1).

264 *3.2 Histological and morphological changes* 

The histological changes in earthworm seminal vesicles are shown in Fig. 2. Pairs of the spermathecal were located in the 7/8/9 lateroventral segments within the clitellum area and the intact one showed a spherical structure and cream color (Fig. 2A, B). Seminal vesicles aid in sperm nourishment and spermatogenesis, in which the initial

development of anucleate spermatocytes occurs and finally separate as sperm bundles and mature spermatozoa (Fig. 2C - Fig. 2H). The comparison of histological images revealed that the increased metal concentration from WSC to LPC sites coincided with the increases in tissue cavities and disordered germ cells and a decrease in bundles of mature sperms in seminal vesicles.

274 The sperm morphology of *M.californica* and the abnormal sperm rates are presented in Fig. 3. Unlike human sperm, normal earthworm sperm was more like a 275 276 smooth straight or curved line with a coarser nucleus head and a flagellum that is several 277 times longer. Sperm deformities occur mainly in the head, including swollen heads, helix heads, bent heads, and missing heads (Fig. 3A). The sperm deformity rates of 278 WSC, DXC, and LPC worms showed a significant increase from 8.73% to 13.2%, with 279 280 the increased gradient of heavy metal contaminations, especially for soil Cd concentration (Fig. 3B). 281

282 3.3 Earthworm sperm motility traits

The sperm trait parameters under Cd and no Cd treatments are shown in Fig. 4. The density and viability of unprocessed sperms showed a slight decrease but nonsignificant difference from WSC to LPC worms (P > 0.05; Fig. 3C and Fig. 4A). Besides, without the Cd re-exposure, no significant difference was found for the sperm LIN, WOB, VCL, and VAP values among three worms (P > 0.05). Only the VSL values were significantly increased, which were 3.29, 4.23, and 4.64 µm/s for WSC, DXC, and LPC worms, respectively (P < 0.05; Fig. 4). After re-exposure with Cd-spiked Glc, most

290 sperm trait values were significantly inhibited and decreased to varying degrees (P <291 0.05; Fig. 4). In contrast, the LIN, WOB, and VSL values showed significant 292 differences among WSC, DXC, and LPC worms, presenting significant increases with 293 the increased metal gradient from WSC to LPC and a maximum of 102% increase for

- 294 LIN performance (P < 0.05; Fig. 4).
- 295 3.4 Sperm DNA damage and enzyme activities

296 The DNA damage and metabolic enzyme activities of M.californica sperms are shown in Fig. 5. The Tail DNA% and OTM values were 3.46 and 1.24, 3.53 and 2.02 297 for WSC and DXC worms, respectively. This showed no DNA damage because it didn't 298 299 exceed any threshold value limits. Compared with those results, LPC worm sperm showed slight DNA damage with higher Tail DNA % and OTM values of 5.83 and 3.85, 300 respectively (Fig. 5A, B; Fig. S2). The LDH activities of LPC worms (1453 U/gprot) 301 were significantly higher than that of WSC and DXC worms (1097 and 1170 U/gprot, 302 respectively) (P < 0.05; Fig. 5C). The SDH activities were 11.2, 11.9, and 12.6 303 304 U/mgprot for WSC, DXC, and LPC worms respectively, and a non-significant 305 difference was found among them (P > 0.05; Fig. 5D).

### 306 4. Discussion

307 With the increasing emergence of environmental pollutants and developments in 308 chemical risk assessment, more test endpoints are used to conduct toxicological studies (Madia et al., 2021; Robinson et al., 2008). Various indicators represent different 309

310	biological processes of the test model organisms, while among which, reproduction is
311	one of the most important evaluations because it's decisive in population continuation
312	for all organisms (Mirmonsef et al., 2017). In the present study, several different
313	endpoints including individual, histological, and biological ones were comprehensively
314	used to determine the effects of <i>in situ</i> metals-contaminated soils on the reproductive
315	system in wild earthworm <i>M.californica</i> . The following discussion detailly organized
316	and talked about the reproductive toxicity and the relationship with exposure histories,
317	then the influence of reproductive compensation on earthworm adaptation and
318	evolution, and finally prospects and disadvantages of the CASA method.
319	In this work, test worms were collected from an increasing gradient of metals-
320	contaminated soils with decades of pollution histories (Huang et al., 2020). Indubitably,
321	earthworms under such persistent stresses suffered different adverse effects including
322	reproductive damages. Analyses here showed the ultrastructural changes in earthworms'
323	seminal vesicles and significant internal spermatocyte decrease based on histological
324	observation (Fig. 2). These results were in agreement with the study of Kwak and An.
325	(2021a) which reported that the endocrine-disrupting chemicals induced significant
326	damage to seminal vesicle cells and also inhibited normal internal spermatogenesis after
327	a 7-day exposure.
220	A coordine to marriesse studies, soil motels his communities diverse where wis the

According to previous studies, soil metals bioaccumulated by earthworms via the epidermal and intestinal pathways can suppress cell proliferation, alter osmotic pressure and affect the synthesis and secretion of reproductive hormones (Orton and Tyler, 2015;

Warner et al., 2020). Additionally, the statistical data on sperm deformity and DNA 331 damage provided more direct evidence for reproductive toxicity (Fig. 3; Fig. 5A, B; Fig. 332 333 S2). With the increase of heavy metal contamination, larger abnormal sperm rates and 334 DNA damage levels were revealed here. These findings were also formerly proved by 335 several studies with various chemicals and different earthworm species (Gao et al., 336 2015; Huang et al., 2021a; Yesudhason et al., 2013; Zheng and Li, 2009). Arguably, it may be easier to use some above reproductive parameters, such as 337 cocoon and offspring recordings to assess the reproductive toxicity impacts, which have 338 been confirmed and widely applied in toxicological fields (Sivakumar, 2015; Zheng 339 and Li, 2009). Nevertheless, it is almost impossible to generalize these in field studies 340 because of the complexity of *in-situ* fields and the difficulty of breeding wild organisms 341 342 in the laboratory. It is also challenging sometimes to extrapolate laboratory results of model organisms to fields and universal species, which may under- or overestimate the 343 344 acute toxicity (Jaskulak et al., 2021; Spurgeon et al., 2020). Therefore, it is necessary 345 to develop individual test metrics rather than those across generations. This was also 346 one of our main motivations to employ the CASA system to evaluate sperm traits, while 347 the current study successfully demonstrated the feasibility of this approach in earthworm reproductive studies. 348

The mobility and velocity of earthworm sperms were affected to different degrees and showed especially significant inhibition in our "re-exposure" tests, while adverse effects of the lab Cd compound on the sperm traits also supported the other field

352	findings of sperm deformities and DNA damage. However, the results concurrently
353	confirmed that earthworms from DXC and LPC sites showed good sperm traits
354	continuity, with better movement trends and faster swimming traits compared to those
355	from relatively clean WSC sites under the same re-exposure conditions (Fig. 4; Fig.S1).
356	Existing studies have pointed out that sperm quality and kinetic performance directly
357	determine whether they can overcome reproductive barriers and successfully fertilize
358	within the entire reproductive process (Kime et al., 2001; Yang et al., 2020b). In
359	addition, available biochemical energy resources also determine sperm traits. Some
360	involved enzymes, such as LDH and SDH, have been reported as essential metabolism
361	enzymes that participate in glycolysis and oxidative phosphorylation processes (van der
362	Horst, 2020; Zhu et al., 2019). The results of the current study also showed that the
363	LDH and SDH activities of LPC worms were slightly higher than that of WSC and
364	DXC ones.

Earthworms under long-term stress tend to coordinate more resources for sperm 365 kinetics to compensate for morphological damage, while these changes were probably 366 highly-depend on their exposure histories. The correlation analysis between the 367 reproductive changes and the initial exposure history implied some positive 368 relationships between sperm parameters (including trait values and enzyme activities) 369 and metal concentration in earthworms (Fig.S3). Increased body burdens of heavy 370 metals may promote the occurrence of phenotypic plasticity in earthworms, which 371 regulates their physiological distribution in coping with environmental stresses. In 372

373 addition, these "adaptive changes" may also be explained by the reproductive compensation hypothesis proposed by Gowaty et al. (2007), which suggests that parents 374 375 attempt to increase reproductive effort for producing competitive offspring in order to 376 make up for lowered offspring viability. Similarly, Yang et al. (2020a) found that tree 377 sparrows under persistent exposure evolved to grow longer flagella to obtain greater 378 mitochondrial load volumes and achieve better reproductive efforts. However, such compensation is also the result of long-term exposure, while acute toxicity experiments 379 380 in the laboratory may induce higher adaptive costs and result in challenges needed to 381 make related adaptive responses and changes (Huang et al., 2022). The reproductive 382 compensation of *M.californica* also supports our previous results and further confirmed its metal tolerance based on population surveys and bioaccumulation analyses (Huang 383 384 et al., 2021a; Huang et al., 2020; Wang et al., 2020). Certainly, such compensation may also attribute to the ability to evolutionary genetic adaptation after decades of exposure. 385 386 The study of population genetics and omics mechanisms is further explored by ourselves now in pursuit of more proof. 387

Previous studies also frequently used other different biomarkers, like oxidative stress, membrane integrity, and 3-[4,5-dimethylthiazol-2-yl]-2,5 diphenyl tetrazolium bromide (MTT) measurements to indirectly assess the sperm quality and reproductive toxicity induced by chemicals (Cuccaro et al., 2021; De Marchi et al., 2020). Related studies also establish some molecular indicators, like the earthworm annetocin gene that shows significant down-regulation after Cd exposure, and in turn influencing the

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cocoon production processes (Mo et al., 2012). Nevertheless, the CASA system was
gradually applied to gain more direct data for sperm evaluation on various animals,
such as zebrafish, sparrows, and domestic animals, highlighting such possibilities for
earthworm sperm analysis (Hong et al., 2022; van der Horst, 2020; Yang et al., 2020a;

398 Yang et al., 2019).

399 To the best of our knowledge, the present study is the first to successfully employ the CASA system for earthworm reproduction analysis, providing direct evidence for 400 the evaluation of reproductive strategies in earthworms. However, it is important to note 401 402 that some limitations still remain. We employed the most appropriate system conditions (30°C, pH=7, and 10-min dissociation) integrated from several previous references for 403 the CASA tests (Travis et al., 2004; Yang et al., 2020b; Zhu et al., 2019) because of the 404 405 difficulty to raise wild earthworms in a laboratory as well as minimizing the impacts of environmental changes. Future studies could further explore the best system for model 406 407 earthworm sperm tests and different sperm capability liquids for different earthworm 408 species considering the potential species-specific limitations. To date, the 409 histopathological examination can only be achieved by image-overlay and step-410 zooming comparison as in previous studies. Hence, more likewise artificial-intelligence 411 technology and tools should be developed for more accurate analysis. Moreover, further 412 research should aim to study the reproductive impacts of environmental stress on invertebrates, from the phenotype, physiology, genes and whole perspective in the 413 414 context of biodiversity loss.

## 415 **Conclusion**

416	In summary, various endpoints were determined to explore the reproductive
417	strategies of wild earthworms <i>M.californica</i> under the long-term stress of heavy metals.
418	Continued exposure indeed induced different reproductive damages to earthworms,
419	however, earthworms also showed elevated reproductive compensation effects. These
420	trade-offs of earthworm reproduction are likely the processes of adaptive regulation and
421	phenotypic plasticity. The current study developed a novel bioassay, which employed
422	CASA to determine the kinetic traits of earthworm sperms, and further confirmed the
423	metal tolerance of <i>M.californica</i> from reproductive perspectives together with our
424	previous results, providing the information and data for related toxicological studies,
425	especially for reproductive evaluations. In addition, the reproductive study also aimed
426	to comprehensively understand the stress responses and promote biodiversity
427	conservation in view of increasing genetic erosion, and even biodiversity extinction of
428	existing organisms.

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## **Conflict of interest**

There are no conflicts to declare. ound 

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## Tables:

Sites		Internal conce	entrations		Bioaccumulation factor				
	Cd	Cu	Zn	Pb	Cd	Cu	Zn	Pb	
WSC	16.8±1.12c	26.0±2.82b	166±6.48b	34.0±6.13a	11.2±0.75a	0.57±0.06b	1.13±0.04a	0.32±0.06a	
DXC	42.8±8.08b	43.0±6.73a	156±16.7b	19.9±3.90b	8.43±1.59b	0.52±0.08b	0.88±0.09b	0.28±0.06a	
LPC	61.1±8.56a	31.7±1.82b	239±10.8a	38.4±10.3a	4.83±0.68c	0.85±0.05a	0.83±0.04b	0.40±0.11a	

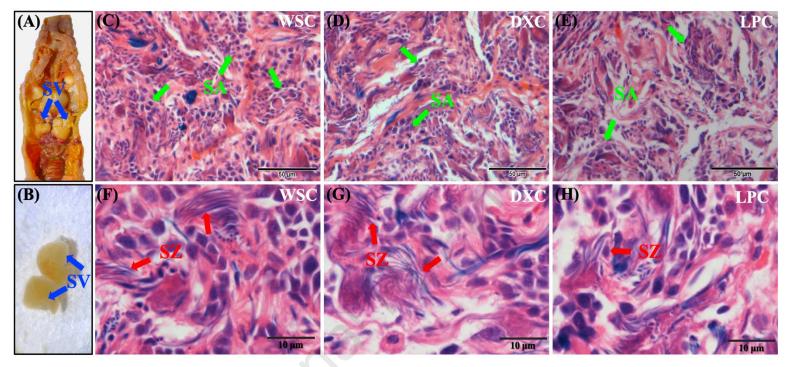
**Table 1.** Internal metal concentrations in earthworms (mg/kg) and bioaccumulation factors (means  $\pm$  SD, n=3)

Note: Different letters (a-c) indicate significant differences (P < 0.05) among WSC, DXC, and LPC sampling plots.

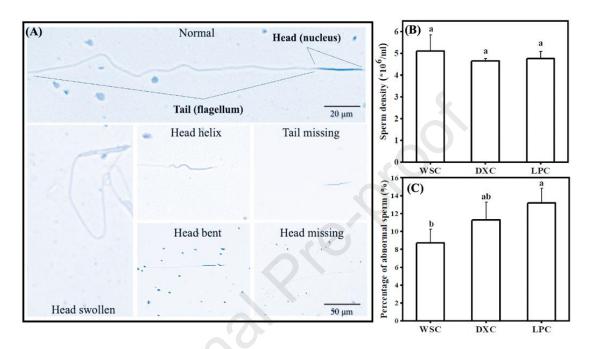




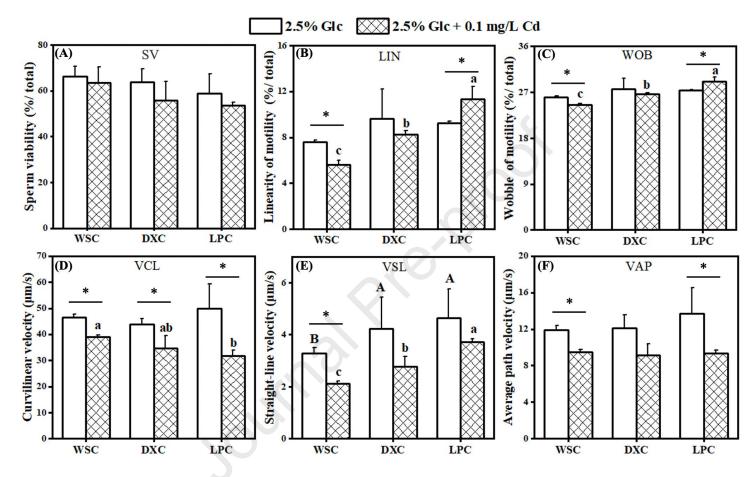
**Fig. 1.** Site location of the earthworm sampling in Hengyang, Hunan province, China. The green to red sampling site labels represents the increased gradient of soil metal contaminations (WSC: Wangshou; DXC: Daxing; LPC: Lupu). The five-spot sampling method was applied for each sampling site (n=5).



**Fig. 2.** Ultrastructural histology changes in the seminal vesicles of *M.californica*. (A-B) abdominal dissection near clitellum for worms, SV (blue arrows): seminal vesicles; (C-E) spermatocyte morphology in seminal vesicles, SA (green arrows): spermatocyte; (F-H) mature sperms within the spermatogenesis processes in seminal vesicles, SZ (red arrows): spermatozoon. The hematoxylin-eosin staining showed increases in disordered germ cells and tissue cavities and a decrease in mature sperm in the SV of worms from WSC to LPC sites.



**Fig. 3.** Sperm morphology, density, and the abnormal sperm rates of *M.californica*. (A) the morphology of normal and abnormal earthworm sperms, the abnormal performance including the head swollen, head helix, head bent, head or tail missing; (B) sperm density (determined using a 20-fold dilution); (C) percentage of abnormal sperms (n=100). Bars are the mean  $\pm$  standard deviation (SD), and different letters (a-b) indicate significant differences among the treatments (*P* < 0.05).



**Fig. 4.** Parameters of sperm performance traits under the treatments with pure sperm capacitation liquid (Glucose, Glc) or Cd-spiked ones reported by the computer-assisted sperm analysis (CASA) system (n = 300). Bars are the mean  $\pm$  standard deviation (SD), and different letters (A-C) indicate significant differences among treatments with pure capacitation liquid while (a-b) indicate significant differences among treatments with Cd-spiked liquid, \* represents significant differences between treatments with or without Cd spiking (P < 0.05), and unmarked ones represent no significance.

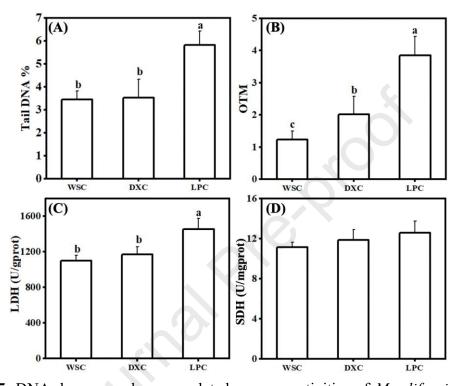


Fig. 5. DNA damage and sperm-related enzyme activities of *M.californica* seminal vesicles. (A) and (B) are Tail DNA% and OTM values, which represent the levels of sperm DNA damage (n = 50); (C) and (D) are the activities of enzyme LDH and SDH, which are both related to the metabolism of sperms (n = 4). Bars are the mean  $\pm$  standard deviation (SD), different letters (a-b) indicate significant differences among the treatments (*P* < 0.05), and unmarked one represents no significance.

## Highlights

- (1) Earthworms with different exposure history were collected for reproduction tests.
- (2) The CASA system was employed for assessing the earthworm sperm kinetic traits.
- (3) Histological changes, sperm deformities, and DNA damage were observed.
- (4) Sperm velocity and metabolism were increased for reproductive compensation.

## **Declaration of interests**

 $\boxtimes$  The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

□The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: