

Co-applied AMF and biogas digestate biochar in Pb-polluted soil: Efficient strategy for minimizing Pb mobility in soil-plant system and improving dietary value of oats

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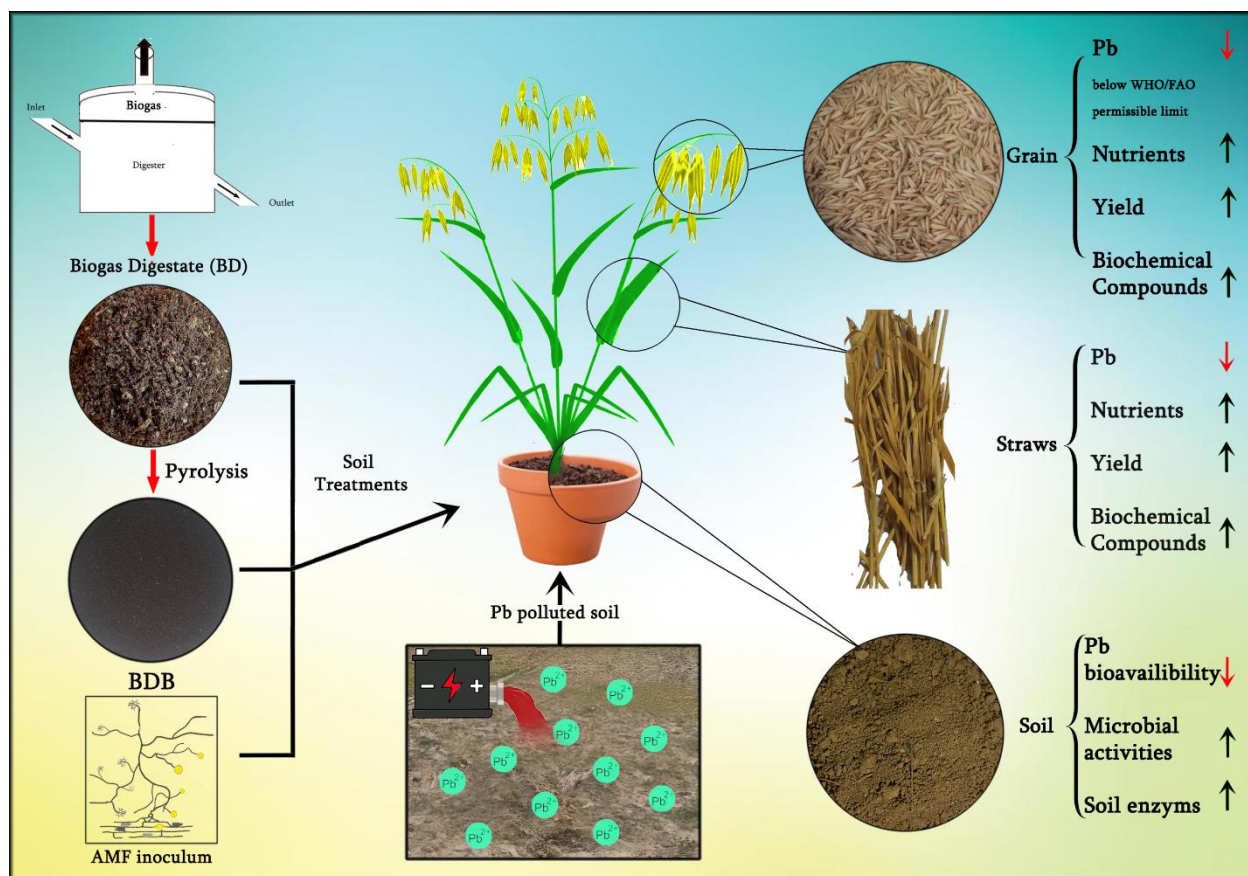
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Abstract

Biochar (BCR) coupled with suitable microorganisms can reduce lead (Pb) mobility in soil-plant system. Biogas plant digestate (BD), an abundant, inexpensive, and year-round available waste, can yield low-cost and nutrient-rich biochar (BDB). This pot experiment involves sole BD, BDB, and arbuscular mycorrhizal fungi (AMF) soil treatments and combining AMF with BD and BDB for Pb immobilization in Pb-polluted soil. Treatment effects on Pb uptake in oat plants, plant growth, and yield were assessed. Moreover, enzymatic activities, AMF root colonization, easily extractable glomalin (EEG), microbial biomass carbon (MBC), and microbial numbers in soil were also measured. BDB+AMF showed the highest soil Pb immobilization index (62.1%), leading to maximum Pb reductions in oats grain (80%), straw (66%), roots (45%), and its soil available fraction (62%), with grain Pb concentrations ($3.27 \text{ mg kg}^{-1} \text{ DW}$) below the WHO/FAO critical limit, than control. This treatment also maximally enriched grain and straw with nutrients (N, P, K, Ca, Mg, Fe, Zn, and Mn).

than control. Moreover, topmost soil enzymatic activities (>68%), MBC (73%), EEG (165%), AMF root colonization (110%), and microbial numbers (>74%) were achieved with BDB+AMF treatment. Conclusively, BDB+AMF is a cost-effective soil amendment that can remediate Pb-polluted soil and produce safer and nutrient-rich food.

Keywords: Lead, Biogas plant digestate, Immobilization, Nutrients, Colonization

55 1. Introduction

56 Due to the poor economic status of several countries, the recycling and repairing of defunct Pb–
57 acid batteries is widespread (Poudel *et al.*, 2023). With high effluent treatment costs and ignoring
58 environmental regulations, small-scale enterprises that recycle and repair these batteries discharge
59 untreated effluents into the nearby water bodies and land, resulting in Pb pollution of soil and water
60 (Boorboori *et al.*, 2022). Resultantly, a higher accumulation of Pb in agricultural fields has been
61 reported, leading to Pb entry into the food chain (Adeyemi *et al.*, 2021; Kumar *et al.*, 2022). Before
62 releasing Pb-containing effluents into water bodies, Internet of Things (IoT) sensors and an HG-RNN
63 model can be used to ensure safe water quality. Herein, the IoT sensor can detect pollutants. At the
64 same time, the Hierarchical Gated Recurrent Neural Network (HG-RNN) model can optimize the
65 treatment, thereby protecting aquatic systems from Pb pollution (Maruthai *et al.*, 2025). Interestingly,
66 as an alternative to growing food crops in Pb-polluted soils, another viable option for obtaining
67 pollution-free food is a hydroponic system with nutrient film technology, powered by the IoT and
68 automation, which can enhance plant productivity and sustainability compared to traditional
69 agricultural practices (Venkatraman and Surendran, 2023). However, for Pb-polluted soils, cost-
70 effective and eco-friendly remediation approaches are the most effective ways to prevent Pb entry into
71 the food chain (Afzal *et al.*, 2024; Boorboori *et al.*, 2022). In parallel, the increasing emphasis on
72 sustainable resource management and eco-innovative practices across sectors has elevated the role of
73 financial strategies and coordinated development models in addressing environmental degradation (Ma
74 and Appolloni, 2025; Ma *et al.*, 2024; Ma *et al.*, 2025a). In agriculture, cost-effective bio-based
75 amendments such as BDB offer practical solutions for remediating Pb-contaminated soils. These
76 environmentally friendly approaches reflect a growing alignment between ecological conservation and
77 regional economic development goals, as highlighted by recent studies on coordinated ecological-
78 economic models (Wang *et al.*, 2025).

Previously, phytoremediation and chemical fixation have been evaluated for the remediation of soils contaminated with heavy metals (HMs). However, phytoremediation has several drawbacks, including being a slow process that depends on the life cycle of the selected phytoremediation crop, low HMs removal efficiency due to the limited extent and depth of root growth, being labour-intensive, and requiring continuous monitoring (Shen *et al.*, 2022). Unfortunately, high concentrations of HMs in soil can also lead to poor plant establishment for phytoremediation purposes, hindering the achievement of the full potential of this approach (Mankè *et al.*, 2024). To accelerate the phytoremediation process, HMs bioavailability is increased by several chemical agents, which also cause secondary pollution of groundwater and surface water through leaching and runoff (Tauqeer *et al.*, 2024). After the harvest of phytoremediation crops, the HMs-contaminated biomass also presents severe disposal issues (Shen *et al.*, 2022). Whereas, the in situ chemical fixation remediation technique involves the addition of chemical substances like lime, limestone powder, various apatites, and phosphate fertilizer to the soil, which reduces the solubility, mobility, and phytoavailability of HMs (Xu *et al.*, 2024). Unfortunately, the use of single amendments partially immobilize HMs, posing a risk of HMs dissolution due to changes in environmental conditions (Xu *et al.*, 2021). Furthermore, these chemical binders pose a risk to the survival and functionalities of soil microorganisms, compromising soil quality and incurring higher costs (Xu *et al.*, 2024).

Biochar (BCR) is a carbon (C) and nutrient-rich product obtained after the pyrolysis of various bioresidues. It has been successfully used to minimize Pb uptake in different crops after Pb immobilization in the soil via raising soil pH, precipitation, ion exchange, and adsorption reactions (Siles *et al.*, 2022; Pandey *et al.*, 2022). Interestingly, BCR also boosts plant growth and yield by improving soil fertility, enzymology, and water holding capacity (WHC) (Afzal *et al.*, 2024; Ayaz *et al.*, 2022). Despite these advantages, the steady demand for vast amounts of BCR for its commercial success remains a question due to the inconsistent availability of cheap feedstock around the year (Hu

103 *et al.*, 2021). Secondly, to fulfil the increased demand for BCR for ecological restoration and plant
104 growth enhancement, using poor-quality feedstock and reduced pyrolysis time yields low-quality BCR.
105 Lastly, BCR cost greater than producing charcoal (100%), like algae BCR (1500%), exotic crops BCR
106 (800%), and nano-biochars (1100%), are not commercially effective for achieving various
107 environmental advantages (Maroušek *et al.*, 2023).

108 Biogas production through anaerobic digestion of bio-residues is cheap and has covered the
109 energy shortage in several nations (Dutta *et al.*, 2021; Mickan *et al.*, 2022). The biogas plant digestate
110 (BD) is a waste which is produced in huge quantities, is rich in nutrients, and can be used as organic
111 fertilizer. However, it contains various pathogenic bacteria (*Escherichia coli* and *Salmonella* sp.),
112 negatively affecting soil and plant attributes and limiting its safe disposal (Baştabak and Koçar, 2020;
113 Tsai *et al.*, 2018). Converting BD into digestate-derived biochar (BDB) can overcome problems, such
114 as 1) cheap feedstock availability all year around, 2) nutrient richness of BDB, and 3) disposal issues
115 (Mickan *et al.*, 2022; Dutta *et al.*, 2021). The BDB is superior to BD due to its nutrient richness, larger
116 surface area, surface charges, and abundant functional groups, which depict its suitability for crop
117 nutrition and the remediation of HMs-polluted soils (Ayaz *et al.*, 2022; Tsai *et al.*, 2018; Wang *et al.*,
118 2022a).

119 A probiotic microorganism, arbuscular mycorrhizal fungi (AMF), coexist with plant roots
120 through symbiosis (Boorboori *et al.*, 2022). Nowadays, different AMF inocula containing multiple
121 AMF species are cheap and available in powder, granular, tablet, and liquid forms (Yang *et al.*, 2023)
122 for their usage in forestry, horticulture, agriculture, and soil rehabilitation (Bisht and Garg, 2022;
123 Ramírez-Zamora *et al.*, 2022). AMF improve plant growth and yield through enhancing water use
124 efficiency and mineral acquisition, and reduce Pb uptake by the plants by producing glomalin,
125 phytochelatins (PCs), extracellular polymeric substances (EPS) and trapping Pb into fungal parts (Li *et*
126 *al.*, 2023; Farhad *et al.*, 2024; Gu *et al.*, 2025).

127 The interaction between BCR and AMF is quite appealing for the sustainability of soil
128 environment and favoring plant growth by improving soil quality, promoting beneficial
129 microorganisms, enhancing water retention in the soil, minimizing HM mobility in the soil-plant
130 system and improving plant growth through nutrient provision (Jia *et al.*, 2024; Vejvodová *et al.*,
131 2020). BCR+AMF reduces the mobility of HMs in soil and their plant uptake through several
132 processes, thereby lowering the dissolution risk of HMs in the soil and leaching (Li *et al.*, 2023; Farhad
133 *et al.*, 2024; Gu *et al.*, 2025; Zhao *et al.*, 2024). Herewith, AMF also play an essential role in
134 immobilizing HMs in their vesicles, spores, mycelium and hyphae (Gujre *et al.*, 2021; Zhao *et al.*,
135 2024). In the BCR+AMF association, BCR supports AMF perpetuation in the soil via improving spore
136 germination and hyphal branching and significantly protects the AMF from different grazers (Gujre *et*
137 *al.*, 2021; Jia *et al.*, 2024). Moreover, this combination also enhances the rhizosphere
138 microenvironment, creating a favorable niche for soil indigenous microorganisms and promoting their
139 activities (Gujre *et al.*, 2021). In this combined treatment, AMF actively interacts with plant growth-
140 promoting rhizobacteria (PGPR), thereby enhancing soil nitrogen fixation and increasing plant growth
141 and yield (Khaliq *et al.*, 2022). In a Pb polluted soil, AMF+BCR was regarded as best compared to
142 AMF and BCR treatments alone for reducing Pb bioavailability in soil by 30.94% and maize shoot Pb
143 concentration by 32.94%, compared to unamended soil (Jia *et al.*, 2024). In another pot experiment, the
144 combined treatment of dry olive residue BCR and AMF reduced Pb concentrations in roots (47%),
145 shoots (16%), and grain (58%) of *Triticum aestivum* grown on Pb-polluted soil (Vejvodová *et al.*,
146 2020). Besides, amending Cd polluted soil with BCR and AMF decreased water-extractable Cd by 24%
147 while reduced Cd accumulation in shoots by 26% and its translocation in rice grain (Zhao *et al.*, 2024).

148 Oats, being high in protein, lipids, minerals, fat, carbohydrate, thiamine, riboflavin, pyridoxine,
149 and fiber contents, are significant in cereal crops due to their economic, medicinal, and dietary
150 importance for livestock and humans (Chawla *et al.*, 2022; Shahidi *et al.*, 2021). Interestingly, oat

151 straw is softer than other cereals, nutrient-rich, and is widely used as an animal ration (Chawla *et al.*,
152 2022). However, the presence of Pb in soil reduces growth, yield, mineral nutrition, and biochemical
153 compounds, as well as high Pb concentrations in various edible plant portions of cereals, which results
154 in malnutrition and Pb toxicity in animals and humans (Khan *et al.*, 2020; Poudel *et al.*, 2023; Wang *et*
155 *al.*, 2022b).

156 The literature lacks information regarding the use of BDB with AMF inoculum for the
157 restoration of Pb-polluted soils, reducing Pb uptake in oat plants, improving the mineral nutrition of oat
158 grain and straw, and enhancing soil health. Our hypothesis is that co-applying AMF with either BD or
159 BDB can have additive effects on reducing Pb concentrations in the edible portions of oat plants,
160 thereby enhancing their dietary value and improving soil health. Thus, we conducted a pot study with
161 the objectives 1) to evaluate the capacity of BD, BDB, and AMF as solitary soil treatment and integrate
162 AMF with BD and BDB for Pb fixation in soil and its restricted accumulation in edible plant tissues, 2)
163 effects of soil additives on grain and straw dietary value and 3) additives effects on soil health.

164 **2. Materials and methods**

165 **2.1. Soil acquisition and its analysis**

166 The soil was gathered from cropland getting irrigation with canal water, having raw effluents
167 originating from Pb–acid battery refurbishing enterprises. The upper layer (0–20 cm) of soil was collected
168 and carefully delivered to the laboratory. Later, the soil samples were blended to achieve a homogenized
169 sample, sieved (2-mm) to remove foreign bodies, and air-dried. Further, the physicochemical traits of the
170 soil were assessed, as presented in Table 1, according to the standard procedures (Zubair *et al.*, 2021).

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177 **Table 1:** Soil properties after its analysis.

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Characteristics	Units	Values
Texture	—	Clay loam
Sand	g kg^{-1}	390
Silt	g kg^{-1}	290
Clay	g kg^{-1}	320
Soil pH	—	6.60
Electrical conductivity (EC)	dSm^{-1}	1.99
Cation exchange capacity (CEC)	$\text{cmol}_c \text{ kg}^{-1}$	12.9
Organic matter	g kg^{-1}	6.80
CaCO_3	g kg^{-1}	37.0
Available P	mg kg^{-1}	8.50
Exchangeable K	mg kg^{-1}	103
N	g kg^{-1}	0.17
Ca	g kg^{-1}	1.24
Fe	mg kg^{-1}	109
Zn	mg kg^{-1}	13.1
Mg	mg kg^{-1}	1490
Mn	mg kg^{-1}	7.4
Cu	mg kg^{-1}	45
Bioavailable Pb	mg kg^{-1}	2.90
Total Pb	mg kg^{-1}	639

179 2.2. AMF consortium

180 A commercial AMF consortium, "Endo Roots Soluble", was purchased from
181 <https://www.bioglobal.com.tr/>, containing nine species of AMF (*Glomus intraradices*, *Glomus*
182 *aggregatum*, *Glomus mosseae*, *Glomus clarum*, *Glomus monosporum*, *Glomus deserticola*, *Glomus*
183 *brasilianum*, *Glomus etunicatum*, and *Gigaspora margarita*), having concentration of 1×10^4 CFU g⁻¹ of
184 inoculum.

185 2.3. Digestate and its biochar

186 The BD was obtained from a local methane biogas facility in Faisalabad, Pakistan. This facility was
187 using buffalo manure near a livestock farm having 1237 heads, producing about 800 kWh day⁻¹. After
188 getting the BD, it was carefully transferred to the laboratory. To prevent odour, the wet BD was oven-dried
189 for multiple days at 60 °C to acquire the proper state for soil amendment. Further, half of the BD was
190 converted into BDB. For this purpose, the BD was pyrolyzed in a local charcoal retort at 600 °C to produce
191 BDB. Later, BDB was cooled at room temperature and ground (2 mm). To examine the surface
192 morphology of the digestate-derived biochar (BDB), scanning electron microscopy (SEM) was performed
193 using a ZEISS GeminiSEM 300 (Germany). The specific surface area and porosity characteristics of the
194 sample were evaluated using a Micromeritics TriStar II Plus 2.02 instrument based on the Brunauer-
195 Emmett-Teller (BET) and Barrett-Joyner-Halenda (BJH) approaches, respectively, under standardized
196 operating conditions. The surface functional groups of BDB were analyzed using an FTIR spectrometer
197 (Nicolet, Thermo Scientific). Samples were scanned in the range of 4000–400 cm⁻¹ at a resolution of 4 cm⁻¹
198 using the KBr pellet method. The BD exhibited properties such as pH = 8.6, EC = 1.22 dS m⁻¹, C =
199 42.7%, H = 5.91%, O = 37.2%, N = 2.91%, P = 1.02%, K = 2.77%, Ca = 2.14%, Mg = 1.14%, S = 0.37%,
200 Fe = 784.6 mg kg⁻¹, Zn 217.1 mg kg⁻¹, Mn = 194.2 mg kg⁻¹, and Cu = 43.4 mg kg⁻¹. Likewise, BDB
201 exhibited properties as pH = 9.8, BET surface area = 52.37 m² g⁻¹, EC = 1.49 dS m⁻¹, CEC = 32.1 cmolc
202 kg⁻¹, C = 51.4%, H = 4.86%, O = 23.4%, average pore diameter 19.25 nm, cumulative pore volume 0.045

203 $\text{cm}^3 \text{g}^{-1}$, N = 2.14%, P = 1.13%, K = 3.53%, Ca = 2.27%, Mg = 1.33%, S = 0.49%, Fe = 1013.1 mg kg^{-1} ,
204 Zn 232.2 mg kg^{-1} , Mn = 213.4 mg kg^{-1} , and Cu = 47.3 mg kg^{-1} .

205 **2.4. Pot experiment and economic analysis of soil remediation**

206 Six soil treatments were prepared in separate plastic vats, for three pots per treatment, each
207 containing 15 kg of soil. These treatments were signified as control, BD, BDB, AMF, BD+AMF, and
208 BDB+AMF. Soils where BD or BDB were to be mixed received 5% (w/w of soil) of these soil
209 amendments. For the treatments receiving AMF inoculum, each corresponding vat received 0.57 g of
210 inoculum per kg of soil. This dose of AMF is based on the manufacturer's recommendation and has been
211 used in our previous study, where it was found to be effective for Pb immobilization in the soil (Farhad *et*
212 *al.*, 2024). In contrast, the control treatment did not receive any additives. The soil of each plastic vat was
213 uniformly mixed, manually watered (WHC 65%), and incubated (dark place, 6 weeks, room temperature).
214 After incubation, three replicates of each treatment were prepared by filling three plastic planters (length =
215 30 cm, diameter = 24.5 cm) with 15 kg of soil. After filling, all 18 planters were placed in the greenhouse
216 (light \approx 10–12 h, temperature \approx 25 °C, and humidity \approx 50%) of Government College University
217 Faisalabad, Pakistan, in a completely randomized order. The plastic planters were irrigated until suitable
218 moisture was gained for planting oat seedlings. Three healthy seedlings of oats (3 cm long, grown in
219 perlite) were planted in each planter. A recommended dose of a balanced plant fertilizer [Osmocote classic
220 (14–14–14)] was applied in each planter. The distilled water was used to irrigate plants while considering
221 the atmospheric conditions. Plants grew for 143 days until the whole plants and ears became solid yellow,
222 indicating plant maturity.

223 The production cost of the BDB (energy consumption, labour, and equipment amortization) in
224 Pakistan ranged from PKR 20–30 kg^{-1} , which is approximately USD (\$) 0.07–0.10 kg^{-1} . The
225 application rate of BDB was 5%, representing 50 g of BDB kg^{-1} soil. At the upper bound of production
226 cost (\$0.10 kg^{-1}), this corresponds to a BDB input cost of \$0.005 kg^{-1} of soil. Likewise, the AMF

inoculum was applied at a rate of 0.57 g kg^{-1} of soil, costing \$12.0 per 250 g, or $\$0.048 \text{ g}^{-1}$. With this cost and dose, the AMF cost kg^{-1} of soil was \$0.02736. There was no rent for the greenhouse, as it is provided free of charge to university faculty and research students. The cost of each plastic planter (length = 30 cm, diameter = 24.5 cm), which was locally fabricated and capable of retaining 15 kg of soil, was \$0.40. Therefore, the planter cost to hold each kg of soil was \$0.026. Whereas, the costs of three oat seeds ($\$0.01 \text{ pot}^{-1}$), fertilizer ($75 \text{ g pot}^{-1} = \0.35) and irrigation ($\$0.19 \text{ pot}^{-1}$) collectively were $\$0.55 \text{ pot}^{-1}$, containing 15 kg of soil. With this cost, oat seeds, fertilizer and irrigation were only $\$0.036 \text{ kg}^{-1}$ of soil.

Therefore, the combined cost for both amendments and experimental setup is:

- Cost of BDB: $50 \text{ g} \times \$0.10 = \0.005
- Cost of AMF: $0.57 \text{ g} \times \$0.048 = \0.02736
- Cost of planter kg^{-1} of soil = \$0.026
- Cost of oat seeds, fertilizer and irrigation kg^{-1} of soil = \$0.036
- Total cost kg^{-1} of soil: \$0.094 (9.4 cents)

Therefore, the price for remediating one kg of Pb-polluted soil was only 9.4 cents.

2.5. Crop harvesting and sample collection

A tapeline was used to measure plant height and ear length. Plant above-ground biomass was harvested carefully with a fine sickle. The number of tillers per plant was determined by counting. Further, the grain were collected from each pot, and the remaining aerial biomass of plants was denoted as straw. As for the collection of roots, the soil from each pot was carefully removed. The grain, straw, and roots were oven-dried (Memmert, Beschickung-loading, model 100–800, Schwabach, Germany) for 24 h at 70°C to measure their dry weight (DW). Later, the oven-dried samples were crushed in a mill, sieved, and stored for further analysis.

250 2.6. Analysis

251 2.6.1. Soil

252 2.6.1.1. Soil pH, Pb bioavailability and its immobilization index in soil

253 The soil pH was measured with pH meter (WTW 7110, Weilheim, Germany). Further, for
254 quantifying the labile soil Pb fraction, the soil was extracted with 5 mM diethylenetriaminepentaacetic acid
255 (DTPA) extractant (soil-DTPA solution = 1:2) (Lindsay and Norvell, 1978) and then measured on ICP–MS
256 (PerkinElmer's NexION® 2000). Likewise, to calculate the Pb immobilization index (IMDX–Pb) in soil,
257 Equation 1 was used (Naeem *et al.*, 2021).

$$258 \quad \text{IMDX – Pb (\%)} = \frac{\text{Pb extracted (control treatment)} - \text{Pb extracted (desired treatment)}}{\text{Pb extracted (control treatment)}} \times 100 \quad (1)$$

259 2.6.1.2. AMF root colonization, hyphal density, spore density, and easily extractable glomalin

260 For estimation of AMF root colonization and spore density, the standard methods were adopted,
261 already described in a previous publication (Farhad *et al.*, 2024). The grid-line intersection method was
262 used to calculate the hyphal length density after staining the hyphae with a trypan blue solution (0.05%,
263 w/v) (Jakobsen *et al.*, 1992). The easily extractable glomalin (EEG) was extracted from soil according to
264 the Wu *et al.* (2015) method. Briefly, a soil sample (0.5 g) was mixed with citrate buffer (4 mL, 20 mM,
265 pH = 7), extracted (103 kPa), heated (121 °C, 30 min), and later centrifuged (10,000× g min⁻¹, 5 min). The
266 EEG in the supernatant was measured by the Bradford method (1976).

267 2.6.1.3. Microbial numbers and microbial biomass carbon

268 For counting the microbial numbers and microbial biomass carbon (MBC) in the soil matrix, the
269 recommended methods were adopted (Iqbal *et al.*, 2023).

270 2.6.1.4. Soil enzymology

271 The wet soil was used to estimate the activities of chitinase, protease, catalase, urease, β-
272 glucosidase, and peroxidase by following various recommended protocols described in earlier research
273 (Zubair *et al.*, 2021).

274 **2.6.2. Plants**

275 *2.6.2.1. Pb concentrations in plant roots, straw, and grain*

276 The concentrations of Pb in straw, roots, and grain were measured after digesting them via the open
277 flask digestion method (Jones and Case, 1990). Briefly, samples (1 g) of each of them were digested in a
278 blend of HClO₄ and HNO₃ (2:1, v/v). Following, the ICP–MS was employed to measure Pb in these
279 digests.

280 *2.6.2.2. Nutrients and biochemical compounds in oat grain and straw*

281 The ICP–MS was used to check the Mg, Fe, Mn, Zn, and Ca concentrations in the grain and
282 straw digests. The phenol–sulfuric acid method was utilized to assess carbohydrate contents in the
283 grain and straw (Smith *et al.*, 1964). Further, for the quantification of N, P, and K concentrations and
284 the contents of protein, fat and fiber in grain and straw, the protocols can be seen in previous research
285 (Naeem *et al.*, 2021).

286 **2.7. Quality control and quality control in analysis**

287 To ensure the quality of analysis, certified reference materials, i.e., CTA-OTL-1 (plant material)
288 and DCI 7004 (soil material), as well as blank samples (simple solutions with no plant or soil samples),
289 were used. Pb recovery from these reference materials (both soil and plant materials) was 91-95%.
290 Before digesting the samples and extracting Pb, all laboratory wares were initially dipped in dilute
291 nitric acid and then washed several times with deionized water. Moreover, each chemical used in this
292 experiment was of analytical grade. All instruments were precise and calibrated before each analysis.

293 **2.8. Statistical analysis**

294 The data are represented as the mean of triplicates from each treatment with standard errors.
295 One-way ANOVA (through *Statistics 8.1*, Analytical Software, Tallahassee, FL, USA, Copyright
296 2005) and later least significant difference (LSD) test ($P < 0.05$) were carried out to equate significant
297 differences between treatment means of every treatment (Steel *et al.*, 1997) and were labelled by lower

case alphabet letters. Principal component analysis (PCA) was conducted to identify patterns and relationships among soil properties, enzymatic activities, plant biomass, and Pb accumulation across the treatments. Pearson correlation analysis was performed to quantify associations between variables and to assess the impact of soil and biological parameters on Pb mobility and plant performance. These statistical analyses were carried out using SPSS 26.0 and OriginPro 2024.

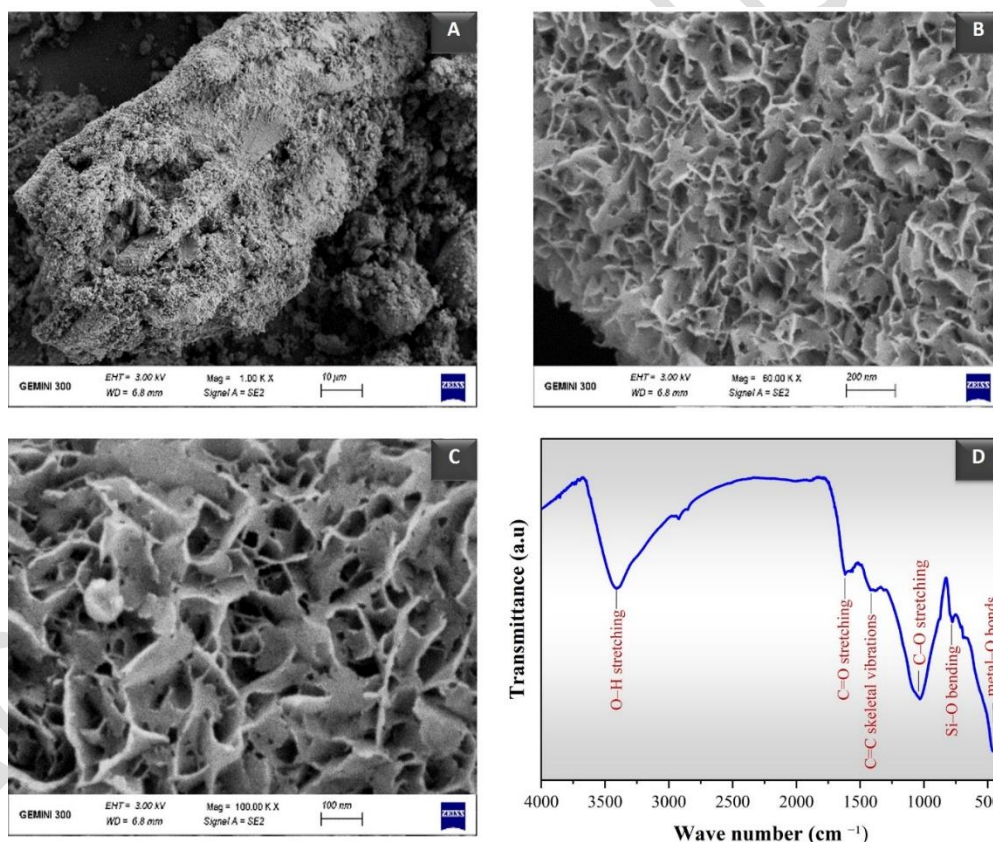
3. Results

3.1. SEM and FTIR analysis

The micromorphological features of digestate-derived biochar (BDB) were characterized using field-emission scanning electron microscopy (FE-SEM) at successive magnifications of 1,000 \times , 6,000 \times , and 100,000 \times (Figure 1A-C). At 1,000 \times , the BDB structure exhibited irregular macropores with diameters ranging from ~ 10 to $80\ \mu\text{m}$, likely resulting from the volatilization of organic fractions during pyrolysis at $600\ ^\circ\text{C}$. These large pore spaces contribute to aeration, increased water-holding capacity, and microbial ingress. At 6,000 \times magnification, the surface revealed a complex network of micro-pores and fissures with widths of approximately $1\text{--}5\ \mu\text{m}$ and depths exceeding $10\ \mu\text{m}$, increasing the specific surface area ($\sim 228.17\ \text{m}^2\ \text{g}^{-1}$, as previously measured). These features enhance the potential for root-fungal interface development, supporting colonization by AMF extraradical hyphae and soil bacterial consortia. At ultra-high magnification (100,000 \times), nanoscale surface irregularities became apparent, including ridge-like carbon lamellae and submicron particulate clusters ($\sim 200\text{--}500\ \text{nm}$). These nanostructures likely function as reactive sorption sites for Pb^{2+} ions via complexation, ion exchange, and surface precipitation mechanisms. The high surface charge density and morphological heterogeneity observed at this resolution affirm the suitability of BDB as both a Pb immobilizer and microbial habitat in Pb-contaminated soil.

The FTIR spectrum of BDB exhibited several distinct peaks corresponding to various surface functional groups (Figure 1D). A broad absorption band at $3409\ \text{cm}^{-1}$ is attributed to O–H stretching

322 vibrations, indicating the presence of hydroxyl groups. The peaks at 1619 cm^{-1} and 1420 cm^{-1}
 323 correspond to C=O stretching and C=C skeletal vibrations of aromatic rings, respectively, indicating
 324 partial retention of aromatic carbon structures after pyrolysis. A prominent peak at 1034 cm^{-1} is
 325 associated with C–O stretching vibrations, indicating the presence of oxygen-containing functional
 326 groups on the BDB surface. The absorption at 778 cm^{-1} is attributed to Si–O bending vibrations, likely
 327 originating from silica components in the digestate. The band at 453 cm^{-1} corresponds to metal–O
 328 bonds, indicating the presence of mineral components derived from the digestate. These functional
 329 groups indicate a chemically active BDB surface that can interact with HMs and enhance nutrient
 330 retention, thereby contributing to its effectiveness in soil remediation.



331
 332 Figure 1. SEM micrographs of digestate-derived biochar (BDB). (A) Low magnification (1,000×), (B)
 333 higher magnification (6,000×), (C) Ultra-high magnification (100,000×), and (D) FTIR spectrum of
 334 BDB, showing key functional groups.

335 3.2. *AMF-associated parameters, EEG, MBC, and microbial numbers*

336 AMF root colonization, contents of EEG and MBC, as well as numbers of fungi, bacteria, and
337 actinomycetes, ranged from 44.4 to 93.4%, 0.76–2.01 $\mu\text{g g}^{-1}$ soil, 261.4–453.3 mg C kg^{-1} soil,
338 14.9–25.8 $\text{CFU} \times 10^6 \text{ g}^{-1}$ soil, 34.6–115.8 $\text{CFU} \times 10^6 \text{ g}^{-1}$ soil and 14.1–33.2 $\text{CFU} \times 10^5 \text{ g}^{-1}$ soil,
339 correspondingly. Whereas, AMF hyphal length density and spore density in the soil were from 2.99 to
340 8.19 m g^{-1} soil and 124 to 381.3 number 100 g^{-1} soil, respectively (Figure 2). Compared to the control,
341 AMF root colonization, numbers of bacteria, MBC content and hyphal length density were
342 significantly improved with each treatment. Moreover, EEG content and the numbers of fungi,
343 actinomycetes and spore density were also enhanced in rest treatments, except BD, than control. The
344 BDB+AMF brought about the most remarkable improvements in the percentage of root colonization,
345 contents of EEG and MBC, numbers of fungi, bacteria, and actinomycetes, hyphal length density, and
346 spore density, by 110, 165, 73, 74, 235, 135, 173, and 207%, compared to the control.

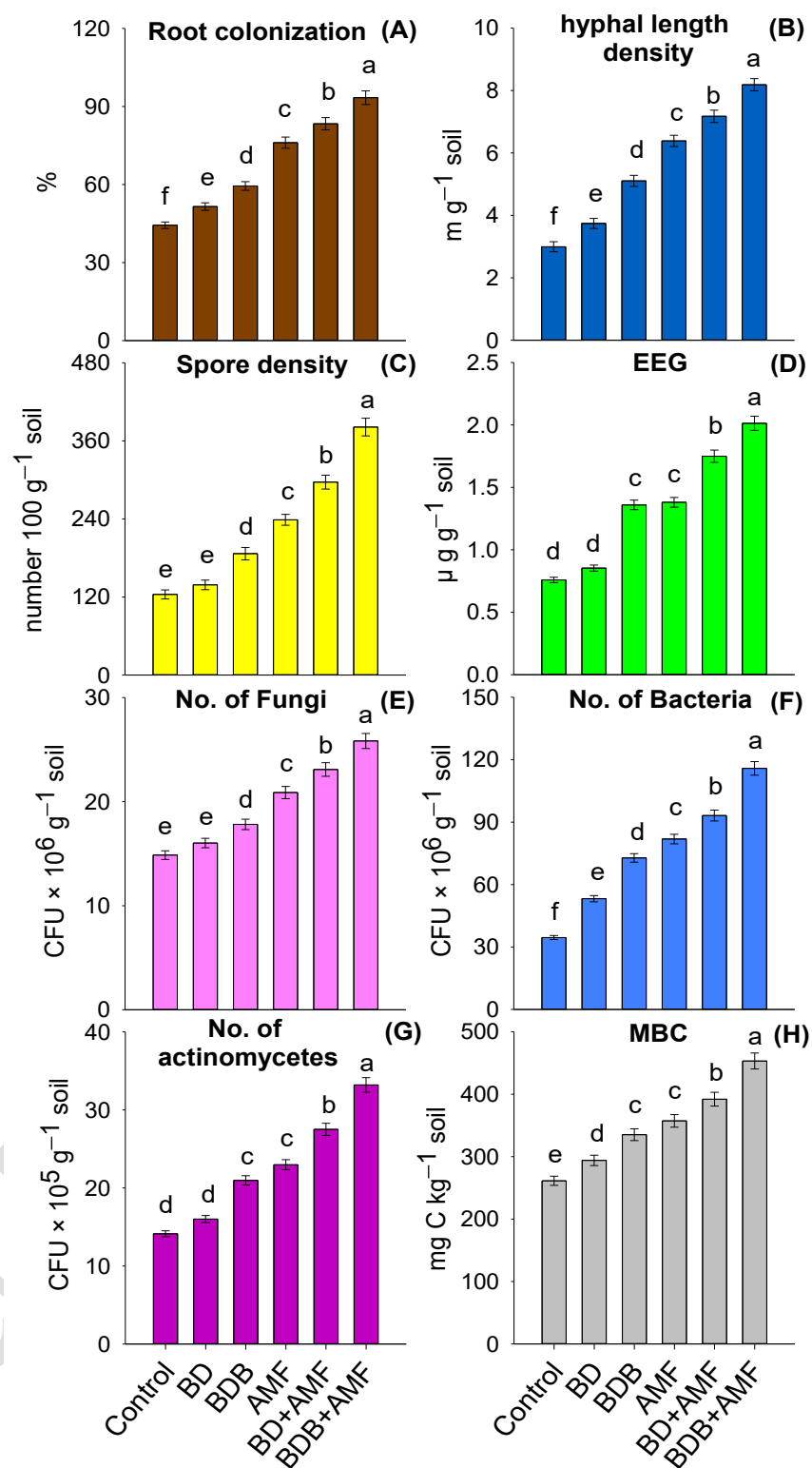


Figure 2. Effects of biogas digestate (BD), its biochar (BDB), AMF, and joint application of BD and BDB with AMF on the AMF root colonization (A), hyphal length density (B), spore density (C), easily extractable glomalin (EEG) (D), numbers of fungi (E), bacteria (F), actinomycetes (G) and microbial

351 biomass carbon (MBC) (H) in soil. The bars represent the mean of triplicate of each treatment, and the
352 error bars indicate the standard error (\pm SE). The different alphabets above the error bars differ
353 significantly ($P < 0.05$) from one another.

354 ***3.3. Status of Pb in plant parts and its soil bioavailability***

355 The Pb concentrations in grain, straw, and roots ranged from 3.27 to 16.3, 54.5–157.1,
356 180.2–327.8 mg kg⁻¹ DW, correspondingly, while DTPA-extractable Pb from 1.16 to 3.06 mg kg⁻¹
357 soil. Similarly, the soil pH values varied between 6.60 and 7.71, whereas IMDX–Pb ranged from 6.89
358 to 62.1% (Figure 3). Compared to the control, every treatment markedly decreased Pb accumulation in
359 plant parts and DTPA extract. However, the maximum decline of 80, 66, 45, and 62% in Pb
360 concentrations of grain, straw, and roots and DTPA-extract, respectively, were noted with BDB+AMF
361 treatment. Moreover, the highest noteworthy increase in soil pH by 0.9 and 1.1 units was found in BDB
362 and BDB+AMF treatments, respectively, than control. Interestingly, the order of IMDX–Pb in post-
363 harvest soil was as follows: AMF (6.89%) < BD (20.7%) < BDB (31.0%) < BD+AMF (41.4%) <
364 BDB+AMF (62.1%).

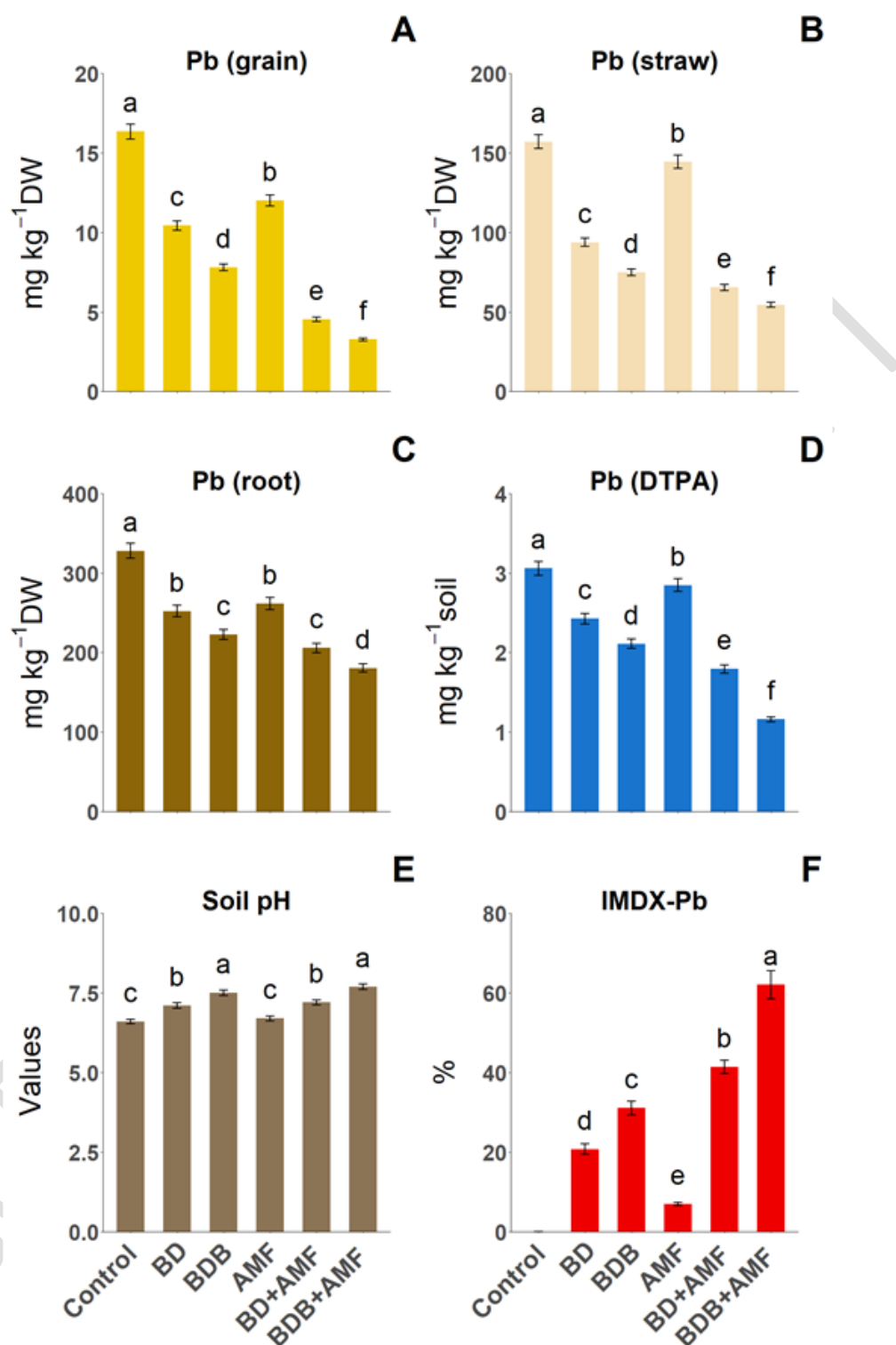


Figure 3. Effects of biogas digestate (BD), its biochar (BDB), AMF, and joint application of BD and BDB with AMF on Pb concentrations in oats grain (A), straw (B), roots (C) and DTPA extract (D), soil pH (E) and Pb immobilization index in soil (IMDX-Pb) (F). The bars represent the mean of triplicate of

each treatment, and the error bars indicate the standard error (\pm SE). The different alphabets above the error bars differ significantly ($P < 0.05$) from one another.

3.4. Plant growth and yield

Data of plant height, straw DW, root DW, grain DW, no. of tillers per plant, and ear length ranged as: 92.9 to 142.2 cm, 7.61–11.7, 1.01–1.65, 6.36–11.1 mg pot⁻¹, 1.67–4.33 number and 11.7–18.1 cm, respectively (Figure 4). Except for AMF, other treatments significantly increased plant height, straw DW, grain DW, and no. of tillers per plant than control. In contrast, root DW showed significant increases in all treatments over control. Soil addition of BDB+AMF resulted in maximum enhancement of plant height, straw DW, grain DW, and ear length by 53, 54, 74, and 47%, correspondingly, than unamended control. Interestingly, the topmost improvements in the no. of tillers per plant by 47, 32 and 21% were seen with BDB, BD+AMF, and BDB+AMF treatments, respectively, while by 57 and 65% in root DW with BD+AMF and BDB+AMF treatments, correspondingly, than control.

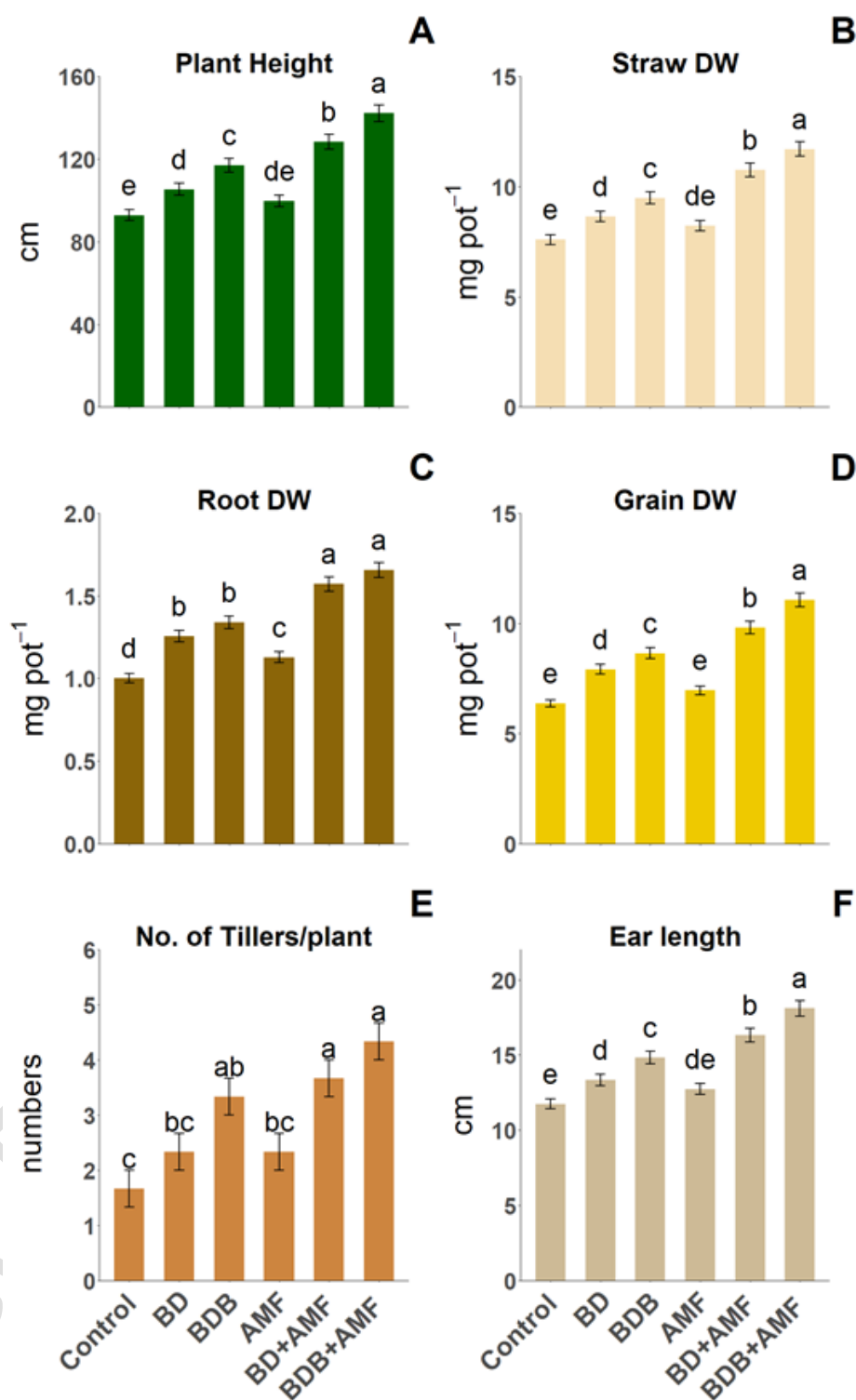


Figure 4. Effects of biogas digestate (BD), its biochar (BDB), AMF, and joint application of BD and BDB with AMF on plant height (A), straw DW (B), root DW (C), grain DW (D), number of tillers per plant (E) and ear length (F) of oats. The bars represent the mean of triplicate of each treatment, and the

error bars indicate the standard error (\pm SE). The different alphabets above the error bars differ significantly ($P < 0.05$) from one another.

3.5. Grain and straw status for nutrients and biochemical compounds

The protein, fat, fiber, and carbohydrate contents in grain ranged from 9.81 to 13.8, 3.27–5.06, 7.78–10.9, and 50.9–69.4%, while from 3.01 to 4.85, 1.50–2.27, 28.1–47.2, and 4.02–6.98% in straw, respectively (Table 2). Every treatment resulted in significant ($P < 0.05$) improvements in these compounds in both plant parts, than control. The topmost contents of protein, fat, and fiber were noted in the BDB+AMF treatment, depicting 41, 55, and 40, respectively, compared to control, in grain. Whereas, 62, 51, 68, and 74% improvements in straw protein, fat, fiber and carbohydrate were noted with BDB+AMF treatment than control. Improvements of 36% and 31% compared to the control were noted in grain carbohydrate content with BDB+AMF and BD+AMF, respectively. The N, P, K, Ca, and Mg concentrations were from 0.64 to 0.97, 0.51–0.83, 0.70–1.05, 0.34–0.57, and 0.22–0.43 g kg⁻¹ in straw, while from 2.01 to 3.47, 0.97–2.11, 2.06–4.41, 0.69–1.48, and 0.46–1.01 g kg⁻¹ in grain. While concentrations of Fe, Zn, and Mn in straw ranged from 41.4 to 75.7, 24.4–48.3, and 21.5–41.2 mg kg⁻¹, while from 62.8 to 114.9, 41.5–73.3, and 30.9–63.8 in grain, respectively (Figure 5). With the exception of N and Fe in grain and Mn in straw with AMF, other treatments brought significant improvements in the concentrations of all minerals in these plant parts. BDB+AMF brought the highest enhancements in N, P, K, Ca, Mg, Fe, Zn, and Mn up to 51, 63, 50, 66, 91, 82, 77, and 91% in straw, while 73, 117, 114, 112, 115, 83, 97 and 106% in grain, accordingly.

Table 2: Effects of biogas digestate (BD), its biochar (BDB), AMF, and joint application of BD and BDB with AMF on protein, fat, fiber and carbohydrate contents in oats grain and straw. The means of triplicate of each treatment with standard error (\pm SE), shown in each column, bearing dissimilar alphabets differ from one another significantly ($P < 0.05$).

Treatments	Protein		Fat		Fiber		Carbohydrate	
	<i>Grain</i>	<i>Straw</i>	<i>Grain</i>	<i>Straw</i>	<i>Grain</i>	<i>Straw</i>	<i>Grain</i>	<i>Straw</i>
<i>mg kg⁻¹ DW</i>								
Control	9.81±0.22 ^e	3.01±0.05 ^d	3.26±0.09 ^e	1.51±0.03 ^e	7.77±0.21 ^e	28.0±0.79 ^e	50.8±1.1 ^{5e}	4.02±0.1 ^{2e}
BD	11.4±0.26 ^{cd}	3.79±0.11 ^c	4.01±0.11 ^c	1.92±0.04 ^c	9.06±0.25 ^c	33.1±0.93 ^d	59.9±1.4 ^{2cd}	4.97±0.1 ^{5cd}
BDB	12.0±0.28 ^{bc}	4.22±0.12 ^b	4.22±0.12 ^b	2.06±0.04 ^b	9.69±0.27 ^b	36.3±1.02 ^c	62.9±1.4 ^{9bc}	5.39±0.1 ^{6c}
AMF	10.9±0.25 ^d	3.48±0.09 ^c	3.69±0.11 ^d	1.78±0.03 ^d	8.75±0.25 ^d	31.8±0.89 ^d	56.1±1.3 ^{1d}	4.55±0.1 ^{4d}
BD+AMF	12.7±0.29 ^b	4.42±0.12 ^b	4.53±0.13 ^b	2.14±0.04 ^b	10.0±0.28 ^b	41.7±1.17 ^b	66.9±1.6 ^{1ab}	6.13±0.1 ^{9b}
BDB+AMF	13.8±0.33 ^a	4.84±0.14 ^a	5.06±0.14 ^a	2.27±0.05 ^a	10.8±0.31 ^a	47.2±1.33 ^a	69.3±1.6 ^{7a}	6.98±0.2 ^{1a}

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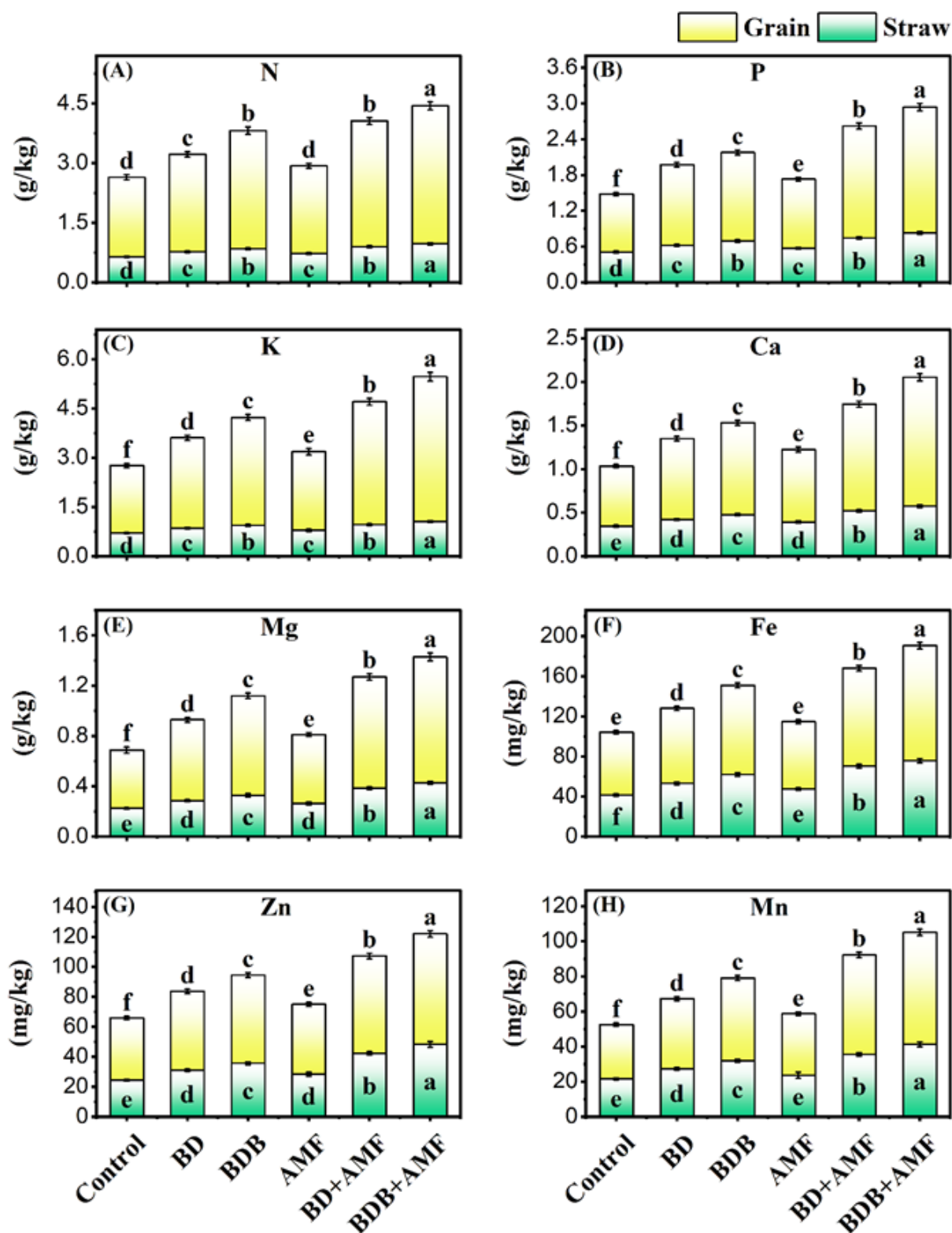


Figure 5. Effects of biogas digestate (BD), its biochar (BDB), AMF, and joint application of BD and BDB with AMF on N (A), P (B), K (C), Ca (D), Mg (E), Fe (F), Zn (G) and Mn (H) concentration in oat grain and straw. The bars represent the mean of triplicate of each treatment, and the error bars

418 indicate the standard error (\pm SE). The different alphabets above the error bars differ significantly ($P <$
419 0.05) from one another.

420 **3.6. Soil Enzymology**

421 The measured activities of soil enzymes, i.e., urease, protease, β -glucosidase, catalase,
422 chitinase, and peroxidase, were in the following ranges: 1.05–2.31 $\mu\text{g N-N (H}_4\text{ kg}^{-1}\text{ h}^{-1})$, 26.9–48.3 mg
423 $\text{kg}^{-1}\text{ 24 h}^{-1}$, 0.56–1.40 $\mu\text{g p-NP g}^{-1}\text{ 24 h}^{-1}$, 32.3–54.1 vol. of 0.1 M $\text{KMnO}_4\text{ g}^{-1}$ of soil, 3.90–8.52 mg
424 $\text{p-NP kg}^{-1}\text{ soil h}^{-1}$ and 2.11–4.64 $\text{mol g}^{-1}\text{ h}^{-1}$, respectively (Figure 6). Surprisingly, compared to the
425 control, each treatment led to significant improvements in soil enzymatic activities. In the case of
426 BDB+AMF treatment, the highest enhancements in urease, protease, β -glucosidase, catalase, chitinase,
427 and peroxidase activities by 119, 80, 151, 68, 118, and 121%, respectively, compared to control, were
428 noted.

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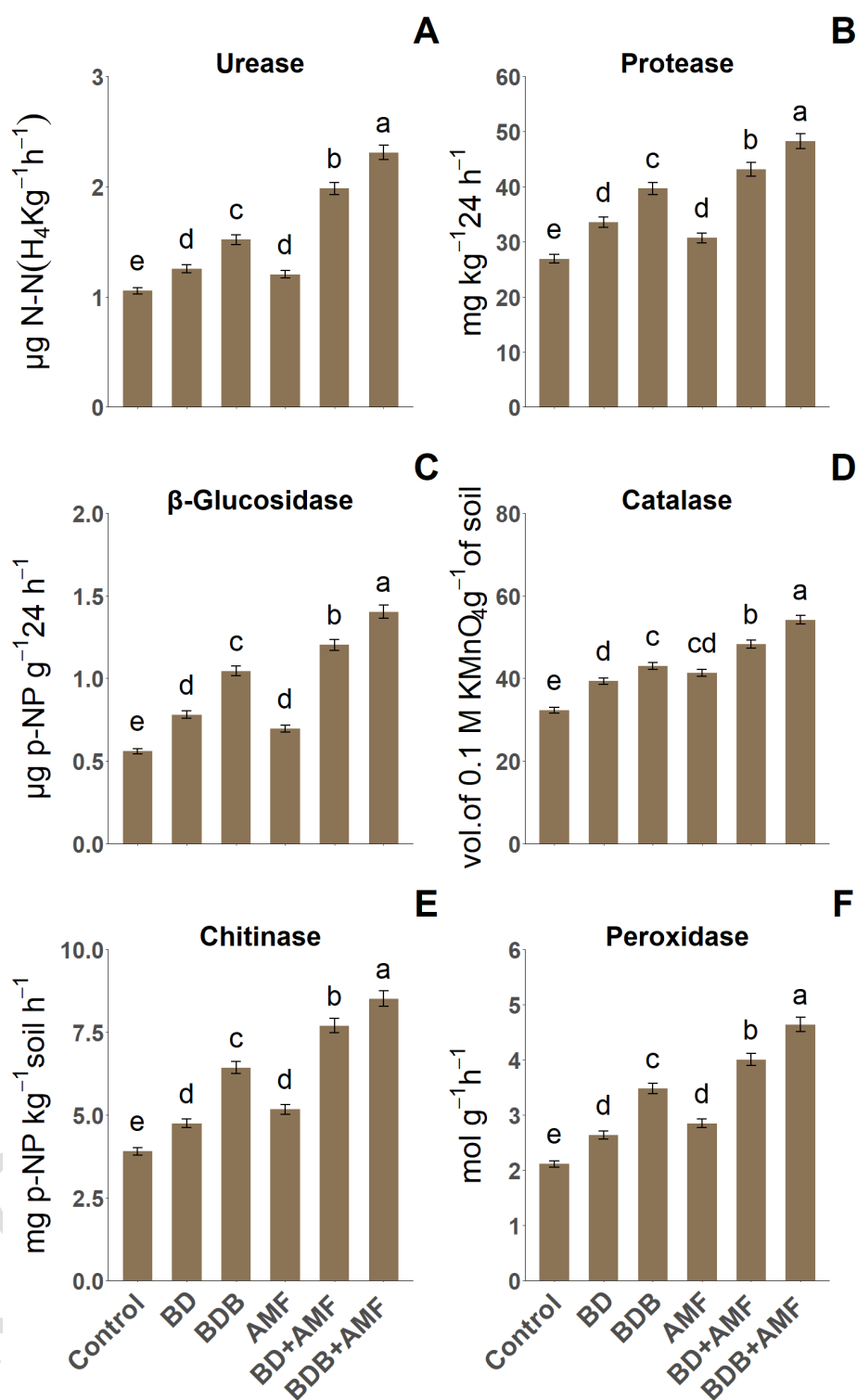


Figure 6. Effects of biogas digestate (BD), its biochar (BDB), AMF, and joint application of BD and BDB with AMF on urease (A), protease (B), β -glucosidase (C), catalase (D), chitinase (E) and peroxidase (F) in soil. The bars represent the mean of triplicate of each treatment, and the error bars

434 indicate the standard error (\pm SE). The different alphabets above the error bars differ significantly ($P <$
435 0.05) from one another.

436 ***3.7. Pearson correlation analysis and principal component analysis***

437 Pearson correlation analysis was conducted to investigate the relationships between soil
438 properties, enzyme activities, Pb accumulation, and plant growth traits (Figure 7). DTPA-extractable
439 Pb was significantly and negatively correlated with soil pH ($r = -0.92$, $p < 0.01$), microbial biomass
440 carbon (MBC, $r = -0.86$, $p < 0.01$), and enzymatic activities such as urease ($r = -0.86$, $p < 0.01$),
441 catalase ($r = -0.75$, $p < 0.01$), and dehydrogenase ($r = -0.70$, $p < 0.05$). This indicates that increased
442 biological activity and soil health are associated with reduced Pb bioavailability. Shoot and grain Pb
443 concentrations also exhibited strong negative correlations with plant biomass ($r = -0.85$ to -0.90 , $p <$
444 0.01), suggesting that Pb stress directly impairs growth. In contrast, shoot dry weight (SDW), root dry
445 weight (RDW), and grain dry weight (GDW) showed significant positive correlations with MBC and
446 antioxidant enzymes, highlighting the role of microbial and enzymatic enhancement in alleviating Pb
447 toxicity. Soil pH was positively associated with both biological activity and plant biomass, and
448 inversely related to Pb translocation. These findings collectively demonstrate that soil microbial and
449 enzymatic responses are strongly linked to reduced Pb mobility and improved plant performance,
450 affirming the effectiveness of the applied amendments in remediating Pb-contaminated soils.

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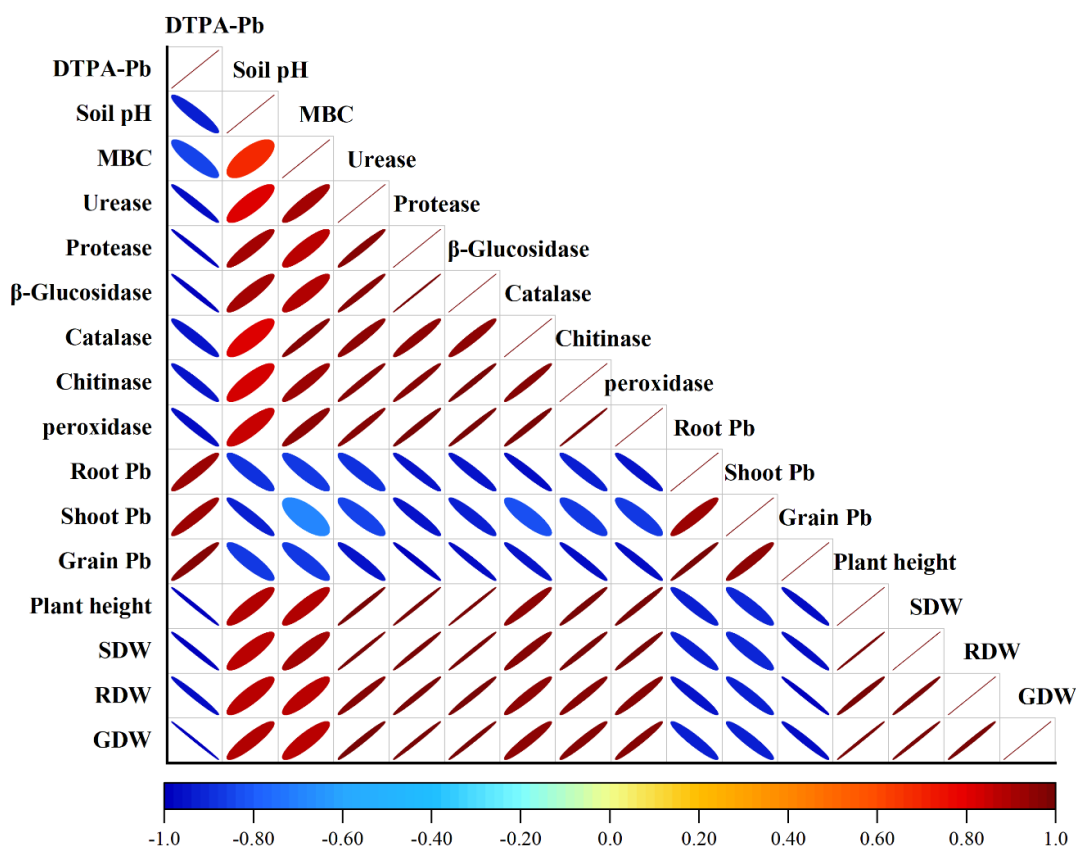
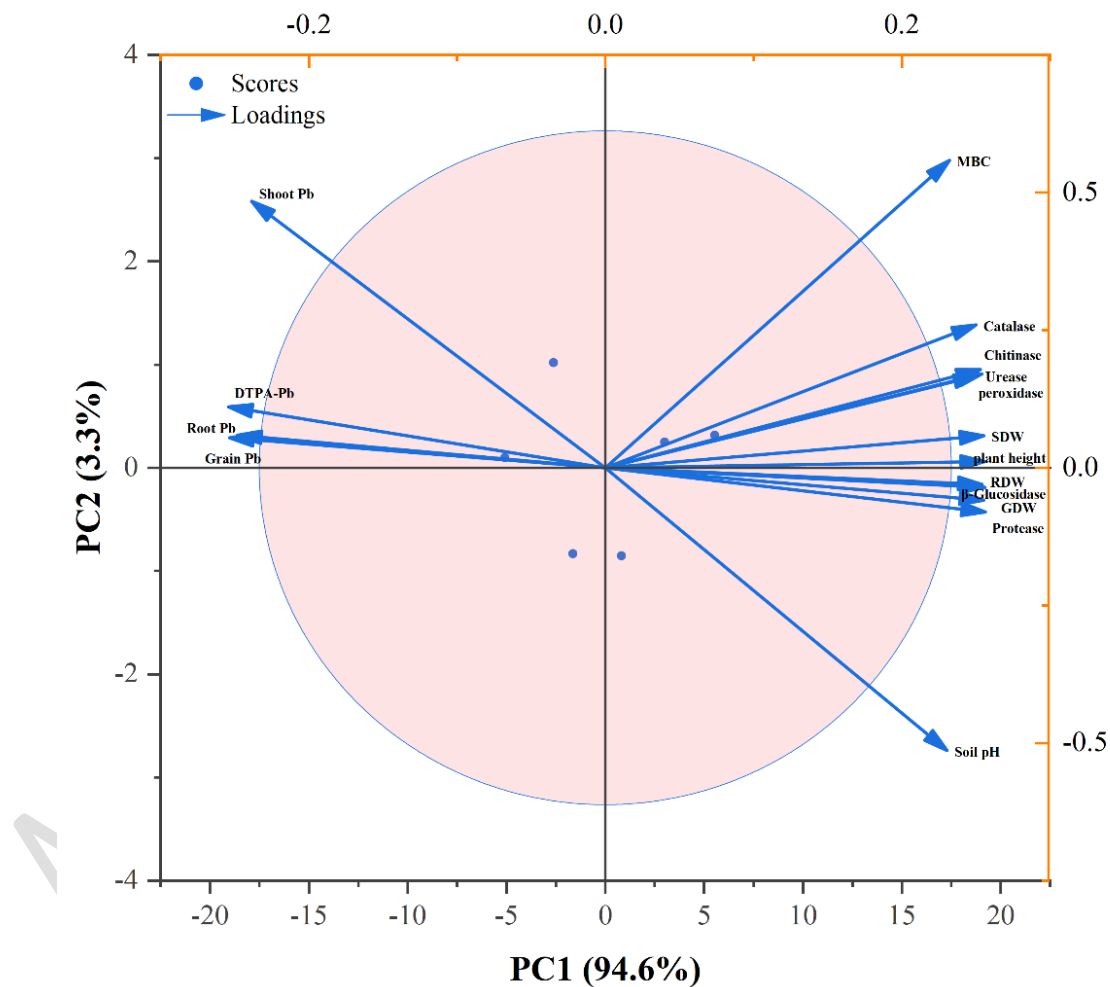


Figure 7. Correlation matrix showing relationships among soil biochemical properties, Pb accumulation in plant tissues, and plant growth parameters. Positive correlations are shown in red, negative correlations in blue, with the intensity of the colour reflecting the strength of the correlation (Pearson's r).

Principal component analysis (PCA) was performed to assess the relationships among soil properties, enzyme activities, biomass traits, and Pb accumulation under different treatments (Figure 8). The first two principal components (PC1 and PC2) explained 94.6% and 3.3% of the total variance, respectively, accounting for a cumulative 97.9% of the variation among the measured parameters. PC1 was primarily driven by positive loadings of soil microbial biomass carbon (MBC), catalase (CAT), chitinase, and plant growth parameters (shoot dry weight, root dry weight, and grain dry weight), while DTPA-extractable Pb, shoot Pb, and grain Pb were negatively associated with PC1. This indicates that

464 treatments enhancing microbial activity and plant biomass were negatively correlated with Pb
465 accumulation. PC2 contributed less variation but separated parameters such as soil pH and protease
466 activity. Treatments positioned on the positive side of PC1 were associated with improved soil
467 biological health and reduced Pb bioavailability, whereas those on the negative side were characterized
468 by high DTPA-Pb concentrations and Pb translocation to shoots and grains. The clear separation of
469 variables in the biplot highlights the contrasting effects of amendments on soil–plant interactions and
470 the effectiveness of treatments in our study in mitigating Pb stress and promoting plant performance.



471

472 Figure 8. PCA biplot showing the relationships among soil biochemical properties, Pb accumulation,
473 and plant growth parameters. Arrows represent variable loadings, and dots indicate treatment scores.

474 4. Discussion

475 4.1. AMF-associated parameters, EEG, MBC, and microbial numbers

476 Excessive Pb concentration in soil has detrimental effects on soil microbial survival and
477 activities, including declining microbial abundance, reduced AMF root colonization, and decreased
478 glomalin secretion by AMF (Siles *et al.*, 2022). Contrarily, BCR supports AMF root colonization, as
479 well as the contents of glomalin secreted by AMF and soil MBC content in Pb-polluted soil (Khan *et al.*,
480 2020). Additionally, it enhanced microbial diversity in soil with Cd and Zn pollution (Shi *et al.*,
481 2023). Inoculation of Pb-contaminated soil with AMF improved the ability of AMF to colonize roots,
482 expanded the diversity of beneficial microbes (*Rhodobacter*, *Archangium*, and *Longimicrobium*), and
483 soil glomalin content (Siles *et al.*, 2022; Li *et al.*, 2023; Adeyemi *et al.*, 2021). In our case, BDB
484 amplified AMF root colonization, the contents of EEG and MBC, and microbial numbers as well as
485 hyphal length density, and spore density (Figure 2) because 1) BDB improved soil C content (Tsai *et al.*,
486 2018), which was utilized by microbes including AMF, leading to their higher numbers, activities,
487 and perpetuation in soil (Shi *et al.*, 2023), and 2) a large surface area and porosity of BDB (Dutta *et al.*,
488 2021) provided habitat to soil microbes and AMF and protected them against predation and desiccation
489 (Hosseini *et al.*, 2021; Poveda *et al.*, 2021). Moreover, BDB immobilized Pb in soil and alleviated the
490 stress on microbes and AMF induced by Pb, leading to enhancements in their growth, perpetuation and
491 activities (Haider *et al.*, 2022). Interestingly, BCR improved soil conditions for AMF spores to
492 germinate, promoted the development of AMF mycelium, the ability to colonize roots, and secrete
493 EEG (Zhao *et al.*, 2024; Khan *et al.*, 2020). Furthermore, AMF inoculation raised glomalin and MBC
494 contents and microbial numbers by 1) solubilizing nutrients from soil and BCR (Zhao *et al.*, 2024;
495 Bisht and Garg, 2022) and 2) secreting EEG containing C, N, P, K, Ca, Mg, Fe, Zn, and Cu and organic
496 compounds required for boosting microbial activities, diversity, and structure (Singh *et al.*, 2022).
497 AMF accounts for 20 to 30% of soil microbial biomass (Sun *et al.*, 2022). Moreover, AMF improved

the aggregate stability of the soil, reduced wind and water erosion, and enhanced the infiltration of water and air, thereby boosting microbial growth and activities (Ai *et al.*, 2022).

4.2. Status of Pb in plant parts and its soil bioavailability

The higher bioavailability of Pb in soil led to increased Pb accumulation in the edible portions of crops (Pandey *et al.*, 2022). Applying BCR efficiently reduced soil Pb bioavailability and its concentrations in the aerial parts of *Prosopis laevigata* (Ramírez-Zamora *et al.*, 2022) and in the roots and shoots of *Zea mays* L. (Afzal *et al.*, 2024). Interestingly, *Funneliformis mosseae* inoculation in Pb-polluted soil declined Pb uptake (57.6%) in soybean (Adeyemi *et al.*, 2021). The lowest Pb concentrations in oat grain, straw, root and DTPA-extract, while the highest values of soil pH and IMDX-Pb were noted with BDB+AMF (Figure 3), with Pb concentration in grain (3.27 mg kg⁻¹ DW) below the permissible level of WHO/FAO (2007) with this treatment (Figure 3). With BDB+AMF, numerous mechanisms caused reduced Pb accumulation in oat roots, straw, and grain. When BD was pyrolyzed to BDB, basic cations (Mg, K, Na, and Ca) of BD were converted to their oxides, carbonates, and hydroxides (Tsai *et al.*, 2018), which raised soil pH upon their dissolution in the soil and precipitated Pb as hydroxyl-bound species of Pb, i.e., PbOH⁺, which reduced Pb bioavailability in the soil (Haider *et al.*, 2022). Interestingly, BDB had a spongy structure, a larger surface area, surface charges, aromatic functional groups, and negative sites, which reduced Pb bioavailability in the soil through immobilization, thereby reducing its plant uptake (Wang *et al.*, 2022a; Dutta *et al.*, 2021). Not only that, AMF also has additive roles in reducing Pb phytoavailability in soil and its uptake in oats. During colonizing plant roots, AMF reduced Pb entry into the roots by 1) acting as a physical barrier and 2) sequestering Pb into vesicles, spores, and mycelia (both intra and extraradical), which also minimized Pb concentrations in the parts of oat plants (Farhad *et al.*, 2024). AMF secrete EEG and EPS, which bind Pb in the soil, thus reducing its plant uptake (Singh *et al.*, 2022; Farhad *et al.*, 2024).

521 A higher production of PCs is reported in AMF-colonized roots, which chelate Pb and reduce its
522 mobility to plant aerial parts (Gu *et al.*, 2025).

523 **4.3. Plant growth and yield**

524 The growth, biomass and yield of cereals are reduced by Pb toxicity (Wang *et al.*, 2022b;
525 Pandey *et al.*, 2022). Conversely, improved growth, yield and 1000-grain weight of barley were
526 reported with soil conditioning with BCR (Hosseini *et al.*, 2021). BCR addition in Pb-polluted soil
527 augmented plant height, fresh and dry weights of roots and shoots of sedum (Shi *et al.*, 2023) and
528 mesquite (Ramírez-Zamora *et al.*, 2022). Interestingly, inoculating Pb-polluted soil with AMF
529 improved root and shoot biomass of Whipstick Wattle (Li *et al.*, 2023) and rosemary (Alinejad *et al.*,
530 2024). Various mechanisms of BDB+AMF improved plant height, straw DW, root DW, grain DW,
531 number of tillers per plant and ear length of oats (Figure 4). Being porous, nutrient-rich and having a
532 large surface area (Ayaz *et al.*, 2022; Dutta *et al.*, 2021; Tsai *et al.*, 2018), BDB improved soil WHC,
533 porosity, nutritional status, and plant water relations, which promoted plant growth (Kumar *et al.*,
534 2024; Ayaz *et al.*, 2022). Besides this, BDB improved oats growth and yield by reducing nutrient
535 leaching and enhancing organic matter content in soil (Ramírez-Zamora *et al.*, 2022; Dutta *et al.*,
536 2021). BDB immobilized Pb and reduced Pb-induced toxicity to oats plants, thereby increasing plant
537 growth and yield (Ayaz *et al.*, 2022). Moreover, AMF solubilized nutrients from the soil matrix,
538 improving the ability of roots to absorb these nutrients along with water, thereby enhancing plant
539 growth and yield. This concept of integrating cost-effective and localized strategies to address
540 environmental challenges aligns with broader innovation pathways, including digital finance,
541 ecological restoration, and sustainable agricultural mechanization, which promote environmental
542 recovery in resource-constrained settings (Wang and Ma, 2024; Ma *et al.*, 2025b; Ma *et al.*, 2025c).

543 AMF also supported plant growth and yield by secreting glomalin, which not only provides
544 essential nutrients to plants but also encourages soil microbes to secrete growth hormones, such as

indole-3-acetic acid (IAA) and 1-aminocyclopropane-1-carboxylic acid deaminase (ACC deaminase). These growth hormones also promoted growth, yield and biomass of oats (Singh *et al.*, 2022; Liu *et al.*, 2022). Upsurging levels of plant endogenous growth hormones, reducing Pb toxicity, Pb-induced oxidative stress, and nematode attack are other key roles of AMF by which the growth, biomass, and yield of oats were improved (Gu *et al.*, 2025; Ai *et al.*, 2022; Pires *et al.*, 2022).

4.4. Grain and straw status for nutrients and biochemical compounds

Reduced nutrient uptake and synthesis of biochemical compounds in various plant species are noted with Pb stress (Adeyemi *et al.*, 2021; Boorboori *et al.*, 2022). BCR incorporation in Pb-contaminated soil raised protein, fiber, and fat contents, while N, P, K, Ca, Mg, Mn, Fe, and Zn concentrations in spinach (Turan, 2020). Soil inoculation with *Diversispora eburnea* uplifted P, K, Ca, Mg, Mn, Fe and Zn contents in maize (Sun *et al.*, 2022). The rise in contents of nutrients and biochemical compounds (protein, fat, fiber and carbohydrate) of oat grain and straw with BDB+AMF is due to the versatile mechanisms of these amendments (Table 2, Figure 5). BDB was nutrient-rich, supplying essential nutrients to plants and thereby increasing the nutrient content of grain and straw, which is positively linked to enhanced synthesis of biochemical compounds (Tsai *et al.*, 2018; Ayaz *et al.*, 2022). Moreover, BDB improved soil moisture content, porosity, and health. Such ambient soil conditions enabled the plants to exhibit better growth and improved metabolism, ultimately enhancing the synthesis of biochemical compounds in them (Dutta *et al.*, 2021; Turan, 2020). BCR improves nutrient availability to plants as it minimizes 1) nutrient leaching from soil (Ramírez-Zamora *et al.*, 2022) and 2) enhances the numbers of soil microbes to solubilize nutrients (Kumar *et al.*, 2024). Moreover, BDB enhanced the synthesis of biochemical compounds in oat plants by mitigating Pb-induced oxidative stress, which they faced in the Pb-polluted soil (Afzal *et al.*, 2024; Haider *et al.*, 2022). Furthermore, AMF raised nutrient uptake by oats through 1) expanding root adsorptive area through tweaking its architecture (shape, branches, and volume) (Sun *et al.*, 2022; Zhou *et al.*, 2022;

569 Gu *et al.*, 2025), 2) minimizing nutrient leaching from the soil, and 3) fetching nutrients from distant
570 zones (Yang *et al.*, 2023). In BDB+AMF, the highest EG content was observed (Figure 2). Glomalin
571 contains significant concentrations of C, N, P, Fe, Cu, Zn, Mg, Ca, S, and K. These nutrients were
572 readily available to oat plants throughout their entire growth phase, resulting in enhanced nutritional
573 status of both straw and grain (Singh *et al.*, 2022). Interestingly, this rise in nutrient uptake by oats
574 enhanced the ability of plants to synthesize various biochemical compounds (Bisht and Garg, 2022;
575 Pirsarandi *et al.*, 2022).

576 **4.5. Soil Enzymology**

577 Soil health is crucial for improved plant growth, yield, and quality of the harvestable part.
578 Compared to traditional soil health monitoring methods, the recurrent neural networks (RNN)-IoT
579 methodology is efficient, accurate, and minimizes ecological impacts by judiciously using water and
580 fertilizer (Selvanarayanan *et al.*, 2024). Soil health is closely linked to nutrient cycling, which is driven
581 by the enzymes secreted by soil microbes (Siles *et al.*, 2022). BCR addition enhanced urease,
582 dehydrogenase, and β -glucosidase activities in Pb-polluted soil (Pandey *et al.*, 2022), while urease,
583 phosphatase, and protease activities in the soil with Cd and Zn pollution (Shi *et al.*, 2023). Moreover,
584 phosphatase, invertase, glucosidase, and protease activities were improved in Cd-polluted soil upon
585 inoculation with AMF mixture (*Rhizoglossus intraradices*, *Claroideoglossus etunicatum*,
586 *Claroideoglossus claroideum* and *Funneliformis mosseae*) (Bisht and Garg, 2022). The highest
587 activities of urease, protease, β -glucosidase, catalase, chitinase and peroxidase with BDB+AMF are
588 due to several mechanisms (Figure 6). BCR addition in soil improves microbial secretion of enzymes
589 by 1) alleviating Pb toxicity to microbes (Zhao *et al.*, 2024) and 2) improving the overall soil
590 environment (Shi *et al.*, 2023; Kumar *et al.*, 2024). The nutrient richness of BDB (Ayaz *et al.*, 2022;
591 Dutta *et al.*, 2021) and porosity (Dutta *et al.*, 2021) also raised microbial enzyme secretion by
592 providing nutrition and shelter to them (Kumar *et al.*, 2024). Surface adsorption of extracellular

enzymes and/or substrates or inhibiting the reaction site of enzymes in a way that affects their affinity for substrates with BCR also positively impacts soil enzymatic activities (Siles *et al.*, 2022). Interestingly, AMF boost activities and enzyme secretion by microbes by providing them with 1) glomalin, a rich source of nutrition, and 2) alleviating HMs toxicity (Singh *et al.*, 2022). Apart from the secretion of several extracellular enzymes on its own, AMF improve the contents and composition of root exudates, thus positively affecting microbial enzyme secretion (Singh *et al.*, 2022; Zhou *et al.*, 2022).

Conclusions

Lead can enter the food chain from the release of untreated effluents from Pb acid battery repairing workshops. The BD is a waste of biogas plants available the whole year. It is cheap and rich in nutrients, but has severe ecological issues at its disposal. Moreover, AMF inoculum is inexpensive, boosts plant growth and mineral nutrition, and reduces Pb uptake in plants. Thus, preparing BDB from BD is practical for such waste material to use as a source of plant nutrition and remediation of Pb-polluted soils. Charring of BD to prepare BDB increases its surface area and mineral nutrients, thus making it a suitable candidate for enhancing plant growth and reducing Pb mobility in soil. In the current pot experiment, BDB+AMF showed premium results in reducing Pb accumulation in grain (below the WHO/FAO limit [5 mg kg^{-1} of DW]), straw, and roots. Other outcomes of this treatment are 1) enhanced mineral nutrition and biochemical worth of grain and straw, and 2) improved plant growth and yield. Regarding soil health, this treatment also boosted the enzyme activities and microbial numbers in the soil. Conclusively, applying BDB with AMF has the suitability to remediate Pb pollution of soil but also boost the nutritive value of oat grain and straw. Due to the richness of BD with nutrients and its availability throughout the year, the conversion of BD into BDB can be very cost-effective and overcome the shortage issue of feedstock to prepare BCR for agricultural and environmental usage. However, certain limitations, such as the availability of inexpensive equipment to

617 produce BDB at low costs, the properties of BDB, the AMF inoculum with different types of AMF
618 strains, and local climatic conditions, including precipitation, humidity, and temperature, may
619 necessitate further management to utilize this amendment by farmers effectively. Since the
620 immobilization technique does not remove Pb from the soil but firmly fixes it in the soil matrix, long-
621 term monitoring of the amended soils is also crucial, which requires expertise from skilled persons or
622 scientists. Compared with existing research outcomes like AMF+BCR effects on wheat for reducing Pb
623 concentrations in grain (58%), shoots (16%) and roots (47%), while reduction of maize shoot Pb
624 (32.94%) and soil bioavailable Pb (30.94%), our findings are superior in terms of Pb reductions in soil
625 bioavailable fraction (62%), roots (45%), shoots (66%) and grain (80%), than control. Thus, our results
626 highlight the significance of BDB+AMF as a superior soil amendment. Future directions could involve
627 the use of advanced modelling tools to monitor Pb immobilization in soils with BDB+AMF, having
628 variable pollution levels, and their effects on soil health and Pb distribution in different edible parts of
629 food crops in pot experiments. Later, the efficacy of this amendment should also be observed in
630 contaminated agricultural fields, where soil, microbial communities, and climatic conditions vary
631 significantly from pots. Government research institutes should support the farming communities by
632 providing consultancy services and supplying them with BDB and AMF at cheaper rates to remediate
633 soil polluted with Pb.

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637 **Declaration of competing interest**

638 Authors declare no competing interests.

639 **Availability of data and material**

640 The data of this manuscript is present in this manuscript and can be made available on a reasonable
641 request.

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