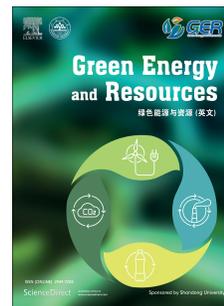


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Biochar for heavy metal remediation in aquatic and terrestrial environments:
Properties, influencing factors, and mechanisms

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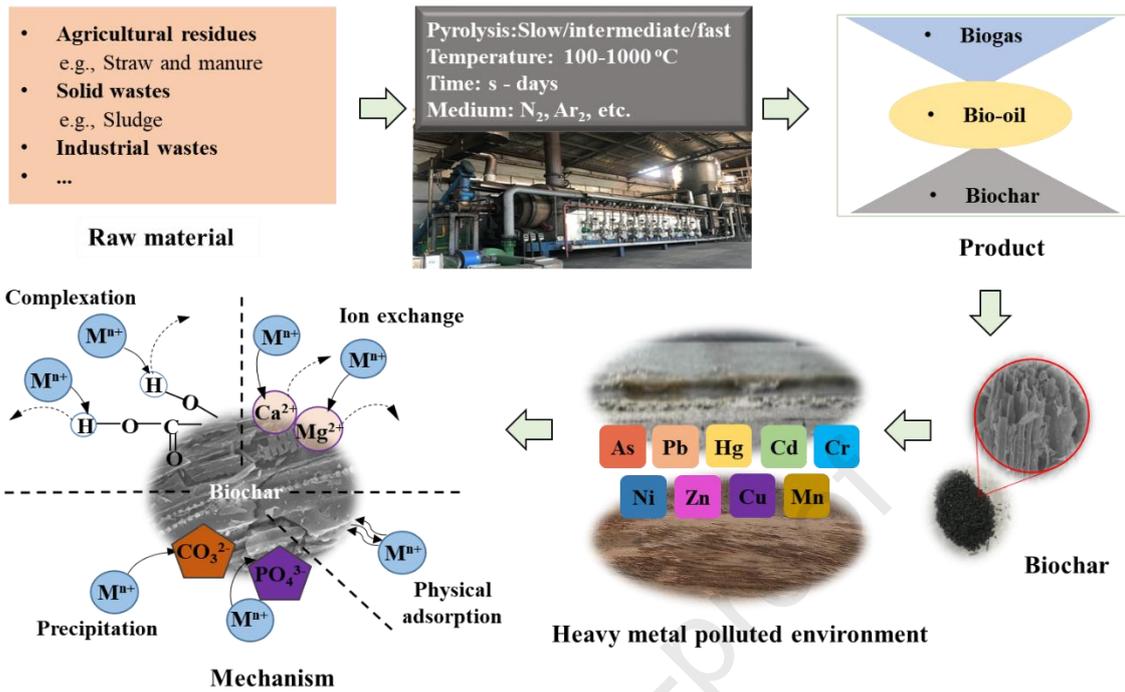
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Graphical Abstract

1 **Biochar for heavy metal remediation in aquatic and terrestrial environments:**
2 **Properties, influencing factors, and mechanisms**

3

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16 **Abstract:** With the rapid development of industrialization, heavy metal pollution in
17 water and soil has become increasingly serious. Biochar has attracted extensive
18 attention for its application in heavy metal remediation, which mainly depends on its
19 developed porous structure, large specific surface area, abundant functional groups and
20 minerals. This review outline the status of heavy metal pollution and summarize the
21 key physicochemical characteristics of biochar. Subsequently, a detailed analysis was
22 conducted on the critical influencing factors, encompassing biochar physicochemical
23 properties, environmental parameters, and heavy metal attributes. The core section
24 comprehensively examines the primary remediation mechanisms, such as physical
25 adsorption, ion exchange, precipitation, and complexation, highlighting how their
26 relative importance shifts with biochar type and environmental context. Furthermore,
27 remediation performance is not governed by a single mechanism but by the synergistic
28 interplay of biochar's evolving surface chemistry, solution conditions, and metal
29 speciation. Additionally, differences in dominant mechanisms and long-term stability
30 are critically compared between aquatic and soil systems. Finally, we identify current
31 knowledge gaps—particularly concerning short-term field performance, long-term
32 aging effects, and sustainability under real-world conditions—and propose targeted
33 future research directions to advance the practical, scalable, and safe application of
34 biochar in heavy metal remediation.

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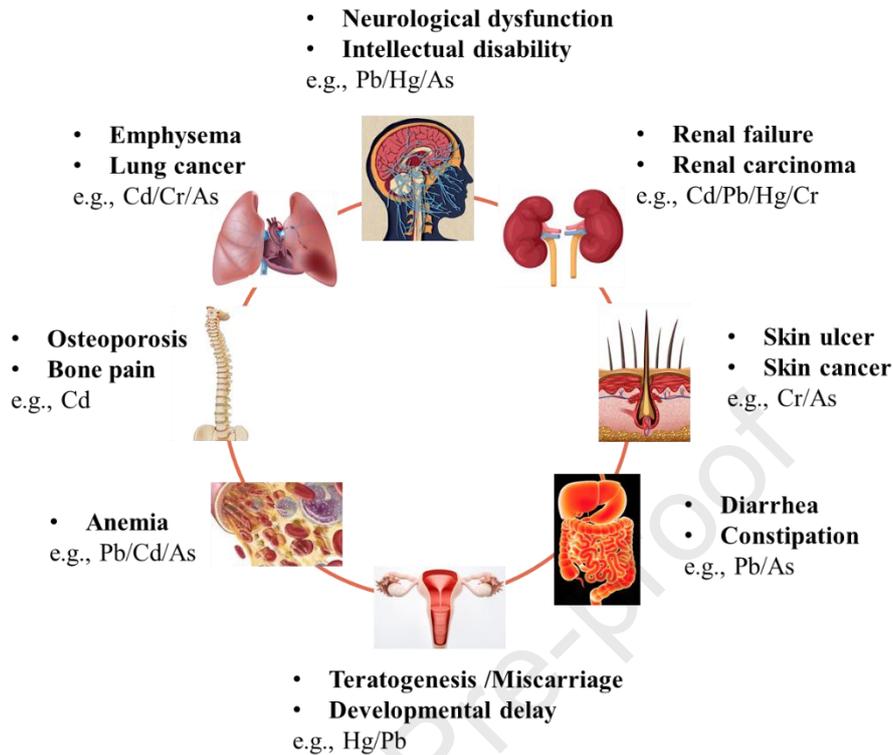
36 **Keywords:** Heavy metal; Biochar; Physicochemical property; Influence factor;
37 Remediation mechanism

38 1. Introduction

39 In recent years, the pollution of heavy metal in water and soil has become
40 increasingly serious with the rapid development of industrialization, which is mainly
41 caused by mining, smelting, fertilizers, and sewage irrigation (Ahmad et al., 2018; He
42 et al., 2019; Khan et al., 2022; Li et al., 2017; Lin et al., 2025; Meng et al., 2018).
43 According to the bulletin of soil contamination in China jointly issued by the Ministry
44 of Environmental Protection and the Ministry of Land and Resources, the percentages
45 of soil pollution sites exceeded 16.1%, and heavy metal was the main contaminant type,
46 its pollution sites exceeded 82.8% (He et al., 2019; O'Connor et al., 2018; Tan et al.,
47 2020; Wang et al., 2020). Also, heavy metal threatens human health through
48 contaminated food chains, leading to serious problems such as brain damage, and
49 kidney diseases, etc. (Fig.1). Therefore, it is a matter of people's health, environmental
50 protection, and ecological safety to strengthen the prevention and control of heavy
51 metal pollution in water and soil.

52 Heavy metal refers to the metal with a specific gravity greater than five, and
53 molecular weight between 63.5 and 200.6, mainly including Arsenic (As), lead (Pb),
54 mercury (Hg), cadmium (Cd), chromium (Cr), etc. (Chakma et al., 2025; Fu and Wang,
55 2011; Khan et al., 2022; Srivastava and Majumder, 2008). The top heavy metals and
56 their maximum allowable contaminant level are shown in Table 1. Heavy metals enter
57 aquatic and terrestrial systems through industrial effluents, mining and smelting
58 activities, agricultural inputs (fertilizers and pesticides), atmospheric deposition and
59 geological weathering. These pathways contribute to their widespread distribution and
60 long-term ecological risks. The heavy metals contained in the water and soil are difficult
61 to degrade, and easy to accumulate, which pose a serious threat to human health and
62 the environment (Aghababaei et al., 2017; Fu and Wang, 2011; Tepanosyan et al., 2018;

63 Wang and Chen, 2009).



64

65

Fig. 1 Human health risks from exposure to heavy metals.

66

Table 1 Top heavy metals and their maximum allowable contaminant level (Inyang et al., 2016; Srivastava and Majumder, 2008).

67

Heavy metals	Rank No. (CERCLA)	Maximum contaminant level by EPA (mg/L)
As	1	0.01
Pb	2	0.015
Hg	3	0.002
Cd	7	0.005
Cr	17	0.1

68

Note: CERCLA, Comprehensive Environmental Response Compensation and Liability Act; EPA, Environmental Protection Agency.

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The conventional heavy metal remediation technologies include chemical precipitation, ion exchange, electrochemical and membrane separation for wastewater

71

72 (Aghababaei et al., 2017; Cui et al., 2016; Li et al., 2017; Li et al., 2023), and soil
73 replacement, heat treatment, electrodynamic repair, chemical leaching, and plant repair
74 for polluted soil (Kunkel et al., 2006; Robinson et al., 2003; Tognotti et al., 1991;
75 Wasay et al., 2001). The methods have the advantages such as good removal effects,
76 easy to recover heavy metals and disadvantages such as high cost, secondary pollution,
77 etc. (shown in Table 2). Notably, the biosorption is the technology utilizing biological
78 materials to removal the heavy metals, of which biochar is considered effective
79 approach for the remediation of heavy metal in water and soil because of its low cost,
80 simple methodology and environmentally friendly (Cui et al., 2016; He et al., 2019; Li
81 et al., 2017; Wang et al., 2015; Wang et al., 2018). Despite extensive literature, a unified
82 mechanistic framework linking biochar structure, environmental factors and metal
83 species across different environments is still lacking. This review was prepared by
84 systematically searching Web of Science, and Google Scholar for articles published up
85 to September 2025. Search terms included combinations of “biochar”, “heavy metal”,
86 “adsorption”, “remediation”, “influence”, and “mechanism”. Key studies were cross-
87 checked in cited references to ensure comprehensive coverage. This review integrates
88 the mechanisms of biochar–metal interactions across aquatic and terrestrial
89 environments and emphasizes how feedstock, pyrolysis conditions, environmental
90 factors and aging jointly determine remediation performance. Recent advances and
91 future research needs are critically discussed to provide a systematic and forward-
92 looking framework.

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94

Table 2 Technology for heavy metal remediation.

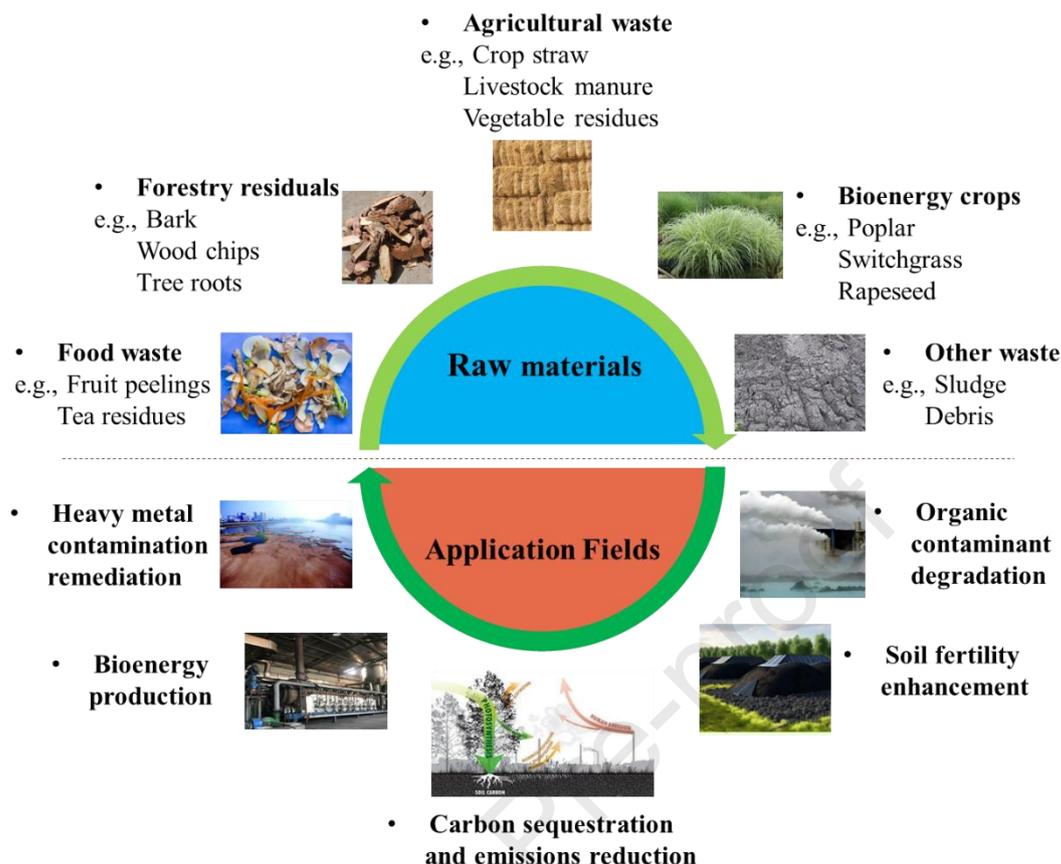
Matrix	Technology	Advantage	Disadvantage
Water	Chemical precipitation	Convenient, low cost, wide applicability	Sediment generated, difficult to handle
	Ion exchange	Materials can be reusable, high selectivity	High cost, low removal efficiency
	Electrochemical	Less chemical reagents, high selectivity and purity	High cost, strong current
	Membrane separation	Less solid waste and reagent, high removal efficiency	High cost, slow speed
Soil	Soil replacement	Efficiency	High cost, contaminated soil needs treatment
	Heat treatment	Efficiency	High cost
	Electrodynamic repair	Efficiency	High cost
	Chemical leaching	Long-lasting, simple operation	Secondary pollution
	Plant repair	Low cost, no secondary pollution	Long cycle, harsh conditions, slow effect

95 2. Biochar

96 Biochar, one carbon-rich product with highly aromatic and chemically stable, is
 97 derived by thermal degradation of biomass under anaerobic or anoxic conditions
 98 (Kambo and Dutta, 2015; Li and Li, 2023; Mimmo et al., 2014; Zhao et al., 2025). The
 99 history of biochar can be traced back to more than 2,500 years ago, the Native
 100 Americans of the Amazon basin improved the barren soil by adding a special fertilizer,
 101 and this Indian soil "Terra Preta" mainly contained a large amount of incompletely
 102 burned charcoal that was the so-called biochar (Laird et al., 2009; Lehmann et al., 2011;

103 Manyà, 2012). Besides, Lehmann pointed out that biochar could effectively reduce the
104 greenhouse effect of carbon dioxide and showed the considerable economic value
105 (Lehmann, 2007). Researches showed that the application of biochar was not only
106 reflected in the improvement of soil physicochemical properties, but also can be used
107 to repair pollution of heavy metals (Kan et al., 2016; Laird et al., 2009; Li et al., 2023;
108 Meyer et al., 2011; Zhu et al., 2017).

109 Biochar can be produced by pyrolysis of various feedstocks, including agricultural
110 residues (Cui et al., 2016; Jiang et al., 2012; Lin et al., 2025; Xu et al., 2023), industrial
111 wastes (Shen et al., 2018; Srivastava and Majumder, 2008), and solid wastes (Bogusz
112 and Oleszczuk, 2018; Chen et al., 2015; Li et al., 2022; Lu et al., 2012), and it is widely
113 applied in areas such as heavy metal contamination remediation, energy production,
114 and carbon sequestration and emissions reduction, etc. (shown in Fig.2). Pyrolysis, the
115 preparation technology of biochar, involves complex chemical reactions containing the
116 process that organic matter is continuously thermally decomposed to form gas-phase
117 and solid-phase (i.e. biochar) (Kan et al., 2016). With the temperature of the reaction
118 chamber decreasing, the compounds with higher polarity and molecular weight in the
119 gas phase are liquefied into bio-oil, and the compounds with lower molecular weight
120 are retained in syngas (Kan et al., 2016; Venderbosch and Prins, 2010). According to
121 the heating rate, pyrolysis can be divided into slow, medium and fast pyrolysis (Ahmad
122 et al., 2014; Manyà, 2012; Qambrani et al., 2017). The pyrolysis under different control
123 conditions and their products are shown in Table 3.



124

125

Fig. 2 Various sources and diverse applications of biochar.

126

Table 3 Typical products under different pyrolysis control conditions (Ahmad et al.,

127

2014; Manyà, 2012; Qambrani et al., 2017; Zhang et al., 2019).

Classification	Pyrolysis temperature (°C)	Residence time	Production (%)		
			Solid	Gas	Liquid
Slow pyrolysis	100-1000	hour-days	35	35	30
Intermediate pyrolysis	~500	10-20 s	25	25	50
Fast pyrolysis	300-1000	< 2 s	12	13	75

128

The biochar mainly exhibits high carbon content, developed specific surface area

129

and porosity, alkaline pH, strong ion exchange capacity, rich functional groups and high

130

thermal stability (Feng et al., 2022; Kong et al., 2014; Novotny et al., 2015). Its

131

physicochemical properties not only depend on the parameters of preparation (pyrolysis

132 temperature, residence time, and heating rate, etc.), but also on the composition and
133 structural characteristics of biomass raw materials (type, particle size, and pretreatment
134 methods, etc.) (Akhtar and Amin, 2012; Jaffari et al., 2023; Kan et al., 2016; Manyà,
135 2012), which are described as follows:

136 *2.1 Elemental composition*

137 Biochar contains the main elements of its raw material, while the differences exists
138 in the productions under the different pyrolysis processes (Ahmed et al., 2016; Jaffari
139 et al., 2023; Qambrani et al., 2017). The C in biochar is mainly derived from the
140 degradation of carbon-containing organic substances (Ahmad et al., 2014; Kan et al.,
141 2016). The content of H or O decreases with increasing pyrolysis temperature, mainly
142 due to the decomposition of organic matter to form volatile gases (such as carbon
143 dioxide, and nitrogen oxides) (Zhang et al., 2020). The content of N is relatively low,
144 Si is mostly enriched in the form of silicon oxide crystals, and Metal elements (such as
145 Na, Mg, K, and Ca) often exist in the form of chloride, carbonate, phosphate, and sulfate
146 (Kan et al., 2016; Wang et al., 2016). In terms of feedstock, wood-based biochars led
147 to greater C content over other biochars, while manures-based biochars contained the
148 greatest N, S, P, Ca, and Mg (Ippolito et al., 2020).

149 *2.2 Specific surface area*

150 Biochar has a well-developed pore structure and a large specific surface area
151 (SSA), which is mainly due to the continuous degradation of organic matter during
152 pyrolysis (Chen et al., 2025; Novotny et al., 2015; Rio et al., 2005). With the increasing
153 pyrolysis temperature, the degradation of organic matter is violent, which is beneficial
154 to the formation of pores and the exposure of surface area. However, the SSA decreases
155 at excessively high temperature, leading to destruction of the pore structure (clogging

156 or collapse, etc.) (Clough et al., 2013; Li et al., 2017; Ma et al., 2025). In addition, the
157 differences in SSA exist between the different sources of biochar (Li et al., 2017; Tang
158 et al., 2013; Xie et al., 2016).

159 *2.3 pH*

160 The pH of biochar is alkaline, and pH value increases with increasing pyrolysis
161 temperature (Li et al., 2017; Suliman et al., 2016; Zhang et al., 2020). There is a
162 significant correlation between the pH value and ash content, which is mainly due to
163 the large amount of Na, Mg, K, Ca and carbonate generated during the preparation of
164 biochar. Meanwhile, the functional groups on the surface (such as -COOH and -OH)
165 are also one of the main reasons for its alkalinity (Li et al., 2017; Singh et al., 2010;
166 Yan et al., 2024; Zhao et al., 2015).

167 *2.4 Cation exchange capacity*

168 The cation exchange capacity (CEC) refers to the exchange capacity between the
169 cations in the biochar and the heavy metals in the water and soil, which can be
170 influenced by the minerals in biochar (Han et al., 2020; Xu et al., 2017; Zhao et al.,
171 2015). During the pyrolysis preparation of biochar, the CEC enhances as the metal
172 minerals enriched. The CEC of biochar mainly depends on the content and type of its
173 mineral components (Carrier et al., 2013; Leng et al., 2025; Uras et al., 2012), and has
174 a certain correlation with the soluble cation content (Xu et al., 2017). Generally, the ash
175 in crop straw is higher than that of wood, which is accordance with the fact that their
176 CEC values (Huijgen et al., 2014).

177 *2.5 Functional group*

178 Biochar contains a wide variety of functional groups, including oxygen-containing
179 functional groups (carboxyl, hydroxyl, ketone, and ester, etc.), and other active

180 functional groups (C=C and N-H with coordinated electron pairs, etc.) (Li et al., 2023;
181 Qambrani et al., 2017; Uchimiya et al., 2011; Várhegyi et al., 1998). With the increasing
182 pyrolysis temperature, the content of oxygen-containing functional groups decreases,
183 while the aromatic ring compounds and carbon skeleton structures are generated, which
184 is consistent with the decreasing atomic ratios of H/C, and O/C (Ahmed et al., 2016;
185 Mukherjee et al., 2011; Zhao et al.). This phenomenon is mainly due to the formation
186 of bio-oil by cellulose and hemicellulose, while lignin is retained as a carbon-based
187 solid during the pyrolysis process (Akhtar and Amin, 2012; Yang et al., 2007).

188 **3. Factors affecting biochar utilization for heavy metal remediation**

189 Biochar shows great potential in the remediation of heavy metal polluted water
190 and soil, because of its alkaline pH, strong ion exchange capacity, abundant functional
191 group and minerals, etc. (Ahmad et al., 2014; Kong et al., 2014; Li et al., 2022; Shahzad
192 et al., 2024; Yang et al., 2025). The biochar utilization for heavy metal remediation in
193 water and soil is summarized in Table 4.

194 There are many factors affecting the remediation of heavy metal by biochar
195 (Inyang et al., 2016; Shang and Chi, 2024; Uchimiya, 2014). Firstly, the removal
196 efficiencies of heavy metals are different between biochars derived from different raw
197 materials (Inyang et al., 2016; Xu et al., 2018). The composition, structure and pyrolysis
198 conditions of raw materials directly affect the removal efficiencies of heavy metals
199 (Uchimiya et al., 2011; Xiao et al., 2017; Xu et al., 2018). Furthermore, environmental
200 factors (pH, ionic strength, organic matter content, and redox potential, etc.) and heavy
201 metal properties (content, occurrence form, chemical valence state, etc.) will also have
202 a certain impact on the removal efficiencies of heavy metals (Uchimiya et al., 2011;
203 Uchimiya, 2014; Xiao et al., 2017; Xu et al., 2020; Xu and Zhao, 2013).

Table 4 Biochar utilization for the remediation of heavy metal in water and soil.

Matrix	Feedstock/pyrolysis temperature (°C)	Heavy metal	Condition			Efficiency (mg/g)	Reference
			Dosage (g/L)	pH	Time (h)		
	Date palm/700	Cd(II)	2	5.0	24	43.58	(Usman et al., 2016)
	Date palm/300	Cd(II)	2	5.0	24	26.96	(Usman et al., 2016)
	Empty fruit bunch/600	Cd(II)	5	6.0	24	85	(Sadegh-Zadeh et al., 2017)
	Rice husk/600	Cd(II)	5	6.0	24	36	(Sadegh-Zadeh et al., 2017)
	Water hyacinth/450	Cd(II)	5	-	24	70.3	(Zhang et al., 2015)
Water	Corn cob/550	Cd(II)	1	6.8	24	31.29	(Luo et al., 2018)
	Pine wood/400	Pb(II)	10	5.0	24	4.13	(Mohan et al., 2007)
	Wheat straw/600	Pb(II)	1	5.0	48	99.45	(Cao et al., 2019)
	Dairy manure/200	Pb(II)	5	-	4	109.4	(Cao et al., 2009)
	<i>Alternanthera philoxeroides</i> /600	Pb(II)	2	5.0	24	257.12	(Yang et al., 2014)
	Corn stalk/450	Pb(II)	2	5.5	24	49.7	(Liu et al., 2019)
	Tetra Pak/600	As(III)	2	-	24	24.2	(Ding et al., 2018)
Tetra Pak/600	As(V)	2	-	24	33.2	(Ding et al.,	

						2018)
Japanese oak wood/500	As(III)	1	7.0	2	3.16	(Niazi et al., 2018)
Japanese oak wood/500	As(V)	1	7.0	2	3.89	(Niazi et al., 2018)
Paper mill sludge/720	As(V)	1	-	24	22.8	(Yoon et al., 2017)
Banana peduncle/300	Cr(VI)	2	2.0	24	114	(Karim et al., 2015)
Wheat straw/500	Cr(VI)	2	3.0	24	12.23	(Zhu et al., 2016)
Sewage sludge/900	Cr(VI)	2	3.0	12	< 7	(Chen et al., 2015)
Sewage sludge/900	Cr(III)	2	3.0	12	25.27	(Chen et al., 2015)
Sewage sludge/300	Cr(VI)	12	-	3	64.10	(Agrafioti et al., 2014)
Sewage sludge/300	Cr(III)	12	-	3	94.34	(Agrafioti et al., 2014)
Sugarcane bagasse/600	Ni(II)	12	6.0	48	1949	(Lyu et al., 2018)
<i>Opuntia ficus-indica</i> cactus/600	Ni(II)	0.6	6.0	48	44	(Choudhary et al., 2020)
Dairy manure/350	Cu(II)	6	-	10	54.4	(Xu et al., 2013)
<i>Opuntia ficus-indica</i> cactus/600	Cu(II)	0.6	6.0	48	49	(Choudhary et al., 2020)
Rice straw/550	Zn(II)	1.25	5.0	48	39.46	(Zhao et al., 2020)
Dairy manure/350	Zn(II)	6	-	10	32.8	(Xu et al., 2013)

Table 4 Biochar utilization for heavy metal remediation in water and soil. (*continued*)

Matrix	Feedstock/pyrolysis temperature (°c)	Heavy metals	Condition			Efficiency	Reference
			Dosage	pH	Time		
	Wheat straw/350-550	Cd	10–40 t/hm ²	5.36	3 y	CaCl ₂ extractable Cd was decreased by 54.5%–70.9%, 53.5%–64.8%, and 28.3%–60.9%.	(Bian et al., 2014)
	Wheat straw/350-550	Cd	20–40	5.01	2 y	CaCl ₂ extractable Cd was significantly decreased by 39%–92%.	(Bian et al., 2013)
	Green waste/550	Cd	5%	-	14 d	NH ₄ NO ₃ extractable Cd concentrations was reduced from 0.95 mg/kg to 0.67 mg/kg.	(Park et al., 2011)
Soil	Chicken manure/550	Cd	5%	-	14 d	NH ₄ NO ₃ extractable Cd concentrations was reduced from 0.95 mg/kg to 0.11 mg/kg.	(Park et al., 2011)
	Sugarcane bagasse/450	Cd	1.5–3 t/hm ²	5.8	30 d	Exchangeable Cd decreased from 8.5% (0.12 mg/kg) to 5.8% (0.08 mg/kg), 2.9% (0.04 mg/kg) and 2.9% (0.04 mg/kg).	(Nie et al., 2018)
	Wood/550	Cd	5–15 g/kg	6.38	10 w	The phosphate-extractable Cd was reduced from 0.28 mg/kg to 0.36 mg/kg, and 0.42 mg/kg.	(Namgay et al., 2010)

Water hyacinth/450	Cd	2%–5%	7.15	1 m	CaCl ₂ extractable Cd was decreased from 1.31 (Yin et al., 2016) mg/kg to 0.35 mg/kg, and 0.31 mg/kg.
Rice straw/450	Cd	2%–5%	7.15	1 m	CaCl ₂ extractable Cd was decreased from 1.31 (Yin et al., 2016) mg/kg to 0.81 mg/kg, and 0.49 mg/kg.
Wheat straw/450	Cd	10–40 t/hm ²	6.07	5 y	Exchangeable fractions of Cd were significantly decreased by 11.4%–44.6%, 19.8%–22.1%, 17.7%–22.1%, and 15.5%–24.2%. (Cui et al., 2016)
Wheat straw/350-550	Pb	10–40 t/hm ²	5.36	3 y	CaCl ₂ extractable Pb was reduced by 16.7%–33.3%, (Bian et al., 2014) 65.0%–79.6%, and 18.1%–59.1%.
Green waste/550	Pb	5%	-	14 d	NH ₄ NO ₃ extractable Pb concentrations was reduced (Park et al., 2011) from 11.3 mg/kg to 7.17 mg/kg.
Chicken manure/550	Pb	5%	-	14 d	NH ₄ NO ₃ extractable Pb concentrations was reduced (Park et al., 2011) from 11.3 mg/kg to 0.73 mg/kg.
Sugarcane bagasse/450	Pb	1.5–3 t/hm ²	5.8	30 d	Exchangeable and carbonate bound fractions decreased from 21.0% (73.5 mg/kg) to 15.7% (54.9 (Nie et al., 2018) mg/kg), 14.1% (48.7 mg/kg) and 12.6% (43.9 mg/kg).

Wheat straw/450	Pb	10–40 t/hm ²	6.07	5 y	Exchangeable fractions of Pb were significantly decreased by 14.2%–19.8%, 22.7%–45.5%, 40.8%–44.7%, and 28.2%–41.3%. (Cui et al., 2016)
Water hyacinth/450	Pb	2%–5%	7.15	1 m	CaCl ₂ extractable Pb was decreased from 2.81 mg/kg to 0.75 mg/kg, and 0.83 mg/kg. (Yin et al., 2016)
Rice straw/450	Pb	2%–5%	7.15	1 m	CaCl ₂ extractable Pb was decreased from 2.81 mg/kg to 1.59 mg/kg, and 1.09 mg/kg. (Yin et al., 2016)
Water hyacinth/450	As	2%–5%	7.15	1 m	KH ₂ PO ₄ extractable As was increased from 11.20 mg/kg to 11.97 mg/kg, and 12.52 mg/kg. (Yin et al., 2016)
Rice straw/450	As	2%–5%	7.15	1 m	KH ₂ PO ₄ extractable As was increased from 11.20 mg/kg to 11.92 mg/kg, and 13.07 mg/kg. (Yin et al., 2016)
Wood/550	As	5–15 g/kg	6.38	10 w	The phosphate-extractable As was increased from 0.12 mg/kg to 0.24 mg/kg, and 0.22 mg/kg. (Namgay et al., 2010)
Pine woodchip/450	As	3%	6.8	14 d	No significant difference in ammonium sulphate-extractable As between control and amended soil. (Brennan et al., 2014)
Olive tree pruning/450	As	3%	6.8	14 d	Ammonium sulphate-extractable As was increased by 2 folds. (Brennan et al., 2014)

Willow/350 & 550	As	10–20 g/kg	5.6	6 m	Compared to control soil, the concentration of water-soluble As had increased to the same value, the total As was increased by 14,000 mg/ha.	(Gregory et al., 2014)
Sugarcane bagasse/500	Cr	15 g/kg	7.54	4 m	Availability of Cr in the soil was reduced by 85%.	(Bashir et al., 2018)
Sugarcane bagasse/550	Cr	3%–7%	7.3	15 d	Bioavailable concentration of Cr was reduced from 23 mg/kg to 17.3 mg/kg, and 12.4 mg/kg.	(Khan et al., 2020)
Poplar wood/550	Cr	3%–7%	7.3	15 d	Bioavailable concentration of Cr was reduced from 23 mg/kg to 22.6 mg/kg, and 19.2 mg/kg.	(Khan et al., 2020)
Bagasse/600	Cr	2–10 g/kg	7.54	15 d	Immobilisation efficiency of Cr(VI) and Cr _{total} was 100% and 91.94%, respectively.	(Su et al., 2016)
Handwood/600	Ni	0.5%–2%	7.9	3 y	Residue fractions of Ni was increased from 51% to 61%–66%.	(Shen et al., 2016)
Willow/700	Ni	2.5%– 10%	6.53	18 m	Ni content in fractions (exchangeable, water soluble and bound to carbonates) decreased by 57–75%.	(Bogusz and Oleszczuk, 2018)
Green waste/550	Cu	5%	-	14 d	NH ₄ NO ₃ extractable Cu concentrations was reduced	(Park et al.,

					from 0.55 mg/kg to 0.42 mg/kg.	2011)
Sugarcane bagasse/450	Cu	1.5–3 t/hm ²	5.8	30 d	Two labile fractions (exchangeable and carbonate bound) decreased from 12.1% (33.9 mg/kg) to 8.7%, 6.3% and 6.1% (24.1, 17.4 and 16.9 mg/kg).	(Nie et al., 2018)
Handwood/600	Zn	0.5%–2%	7.9	3 y	Residue fractions of Zn was increased from 7% to 27%–35%.	(Shen et al., 2016)
Rice straw/500	Zn	0.75–22.5 t/hm ²	5.5	1 y	CaCl ₂ -extractable Zn concentrations were significantly decreased by 25.0%–83.6% for the paddy soil and 46.1%–90.3% for the wheat soil.	(Wu et al., 2018)
Holmoakwood/650	Zn	2%–4%	6.3	3 y	Acid-soluble fraction of Zn was reduced from 68%–81% to 65%–74%.	(Egene et al., 2018)

207

Note: y, year; m, month; w, week; d, day.

208 *3.1 Effect of physicochemical properties of biochar on heavy metal remediation*

209 As a pollution remediation agent, biochar's physicochemical properties (preparation
210 method, type of raw materials, pyrolysis temperature, cation exchange capacity and
211 surface charge, etc.) directly affect the removal efficiency of heavy metals (Inyang et al.,
212 2016; Qi et al., 2024; Qian et al., 2016; Uchimiya et al., 2011; Xiao et al., 2023). Biochar
213 preparation strongly influences its physicochemical properties and metal remediation
214 capacity. Pyrolysis temperature, heating rate, residence time and reaction atmosphere
215 determine aromaticity, surface area, porosity, functional groups, mineral phases and pH
216 (Mandal et al., 2021). For instance, increasing pyrolysis temperature enhances aromatic
217 structure and alkalinity but may reduce oxygen-containing groups that are crucial for
218 complexation with metal cations (Manyà, 2012; Xiao et al., 202).

219 Usman et al. (2016) investigated the removal efficiency of Cd(II) onto different date
220 palm-derived biochars (BC-300 and BC-700) under pyrolysis temperatures of 300 and
221 700 °C, respectively, and found that the maximum adsorption capacity of 43.58 mg/g was
222 observed for BC-700 based on the Langmuir model, which was 1.6 times higher than that
223 of BC-300 (26.96 mg/g), indicating that pyrolysis temperature was the main factor
224 determining the adsorption efficiency of different biochars. Uchimiya et al. (2011) studied
225 the adsorption amounts of Pb, Cd, Ni, and Cd onto different cottonseed hull-based
226 biochars (CH200, CH350, CH500, CH650, and CH800) under 200, 350, 500, 650, and
227 800 °C for 4 h, respectively. The results suggested that CH350 was the most effective in
228 removing heavy metals, the phenomenon was due to the strong electrostatic interactions
229 between the negatively charged surfaces of biochar and heavy metal ions. Jiang et al.
230 (2013) prepared different biochars (derived from maize straw, rice straw, and bagasse,
231 respectively) at 400 °C for 4 h to remove As in the polluted soil. Compared with the raw

232 materials, the adsorption amounts of As onto biochars increased significantly, especially
233 the maize straw and rice straw-derived biochars, which were attributed to their larger
234 cation exchange capacity.

235 The biochars derived from different raw material have different removal efficiency
236 of different heavy metals, and the composition, structure and pyrolysis conditions of the
237 raw materials directly determine the properties of biochar, thus affecting the removal
238 efficiency of heavy metal (Park et al., 2011; Sadegh-Zadeh et al., 2017; Usman et al.,
239 2016; Ye et al., 2025; Yin et al., 2016). Generally, the adsorption amount of heavy metals
240 increases with the dosage increases to a certain extent and then reached equilibrium as
241 the dosage continues to increase, mainly because the adsorption sites provided for the
242 heavy metals are relatively saturated (Cao et al., 2019; Chen et al., 2015). In addition, the
243 aging phenomenon of biochar is mainly manifested in the oxidation of adding oxygen-
244 containing functional groups (such as carboxyl and hydroxyl groups, etc.), which causes
245 an increase in negative charge, enhanced CEC, and improves/affects the removal
246 efficiency of heavy metal (Kumar et al., 2018; Rechberger et al., 2019; Zhao et al., 2015).

247 *3.2 Effect of environmental factors on heavy metal remediation*

248 Environmental factors (pH, ionic strength, etc.) have varying degrees of influence
249 on the adsorption of heavy metals onto biochar (Feng et al., 2022; Uchimiya, 2014; Xiao
250 et al., 2023). Luo et al. (2018) investigated the removal efficiencies of Cd(II) in the
251 different type soils by rape straw and peanut shell-derived biochars. The results showed
252 that the Cd(II) removal efficiencies enhanced rapidly with the pH value increased from 2
253 to 4 and stabilized at pH 5–7, which was related to the fact that an increased pH decreases
254 the competitiveness of Cd(II) against protons for binding sites, and the precipitation,
255 hydroxide complexes of Cd(II) formed. Xu et al. (2020) studied the effects of FeCl₃ and

256 organic acids on remediation efficiency of Cr(VI) onto biochar, and found that FeCl₃
257 reduced the reduction rate of Cr(VI), mainly due to the covering effect between the iron
258 mineral salts and the iron-biochar composite and the oxidation of biochar by Fe(III); while
259 FeCl₃ and organic acid biochar can enhance the reduction rate of Cr(VI), the main reason
260 is that Fe(III) was reduced to Fe(II) by organic acid, which accelerated electron transfer.
261 Qian et al. (2019) analyzed the effects of ions on the remediation efficiency of Zn(II) and
262 Cd(II) onto wheat straw-derived biochar, the results showed that the ions in the acidic
263 environment weakens the remediation efficiency of heavy metals; while the ions in the
264 alkaline environment can enhance the remediation efficiency of heavy metals.

265 As an important environmental factor, pH directly affects the existence speciation of
266 heavy metal ions and the surface charge of biochar, thereby changing the way of
267 attraction/repulsion between biochar and heavy metals (Tan et al., 2020; Uchimiya, 2014;
268 Wang et al., 2018; Xiao et al., 2017; Yan et al., 2024). When the pH value is too low,
269 there are more H⁺ in the environment to form a strong competition with heavy metal ions,
270 reducing its adsorption capacity (Luo et al., 2019; Xiao et al., 2017). Under fluctuating
271 redox or pH conditions, dissolution of metal–biochar complexes or mineral phases may
272 occur. Acidic conditions enhance proton competition, whereas reducing conditions can
273 destabilize Fe/Mn oxide-bound metals, potentially causing desorption. Thus, long-term
274 stability should be evaluated under cyclic redox environments (Luo et al., 2025; Wu, et
275 al., 2025). On the other hand, ionic strength affects the amount of heavy metal onto
276 biochar, which is mainly attribute to the coexisting ions occupying the active sites on the
277 surface of the biochar and forming corresponding competition for heavy metals (Qian et
278 al., 2019; Zhang et al., 2017). Furthermore, temperature mainly affects the
279 thermodynamic process and adsorption heat capacity of biochar to adsorb heavy metals.

280 Generally, increasing the ambient temperature is beneficial to increase the amount of
281 heavy metal onto biochar, mainly because the adsorption process is an endothermic
282 reaction (Inyang et al., 2016; Luo et al., 2019; Xiao et al., 2017).

283 *3.3 Effect of heavy metal properties on heavy metal remediation*

284 The remediation efficiency of heavy metal is influenced by its properties (type of
285 heavy metal, ionic radius, and charge, etc.) (Li et al., 2023; Qian et al., 2016; Xiao et al.,
286 2023). Ko et al. (2004) investigated the ability of bone biochar to adsorb Cu(II), Zn(II)
287 and Cd(II), and the selectivity of metal ions follows the order $\text{Cu(II)} > \text{Zn(II)} > \text{Cd(II)}$,
288 the reversed order of hydrated ionic radii is $\text{Cu(II)} (4.19 \text{ \AA}) < \text{Cd(II)} (4.26 \text{ \AA}) < \text{Zn(II)}$
289 (4.30 \AA) . This result agrees with other studies (Lee and Moon, 2001; Wan Ngah and
290 Hanafiah, 2008). Beesley et al. (2011) employed the difference of As, Cd, and Zn from a
291 multi-element contaminated sediment-derived soil by a column leaching experiment, the
292 results showed that the capability of biochar to immobilize Cd and Zn assisted a 300- and
293 45-fold reduction in their leachate concentrations, respectively. While the As
294 concentration in eluate was not significantly reduced or increased during the leaching test,
295 which was due to the fact that As was so weakly water-soluble in this soil. Zhao et al.
296 (2020) indicated that the biochars had greater adsorption capacities for Pb(II) than for
297 Zn(II), which was due to the former's higher electronegativity, lower $\text{p}K_{\text{H}}$ (negative log
298 of hydrolysis constant), and lower binding energy.

299 In the adsorption process, the competition between heavy metals is induced
300 especially at high concentration, this is mainly attributable to differences in chemical
301 characteristics between the heavy metals (Park et al., 2016). The enhanced remediation
302 efficiency may be attributed to: (1) greater ionic radius (subsequently smaller hydrated
303 radius), (2) higher hydrolysis constant, (3) higher atomic weight, and (4) larger

304 electronegativity, all of which provide the heavy metal with improved efficacy for inner
305 sphere surface complexation or sorption reactions (Ko et al., 2004; Lee and Moon, 2001;
306 Park et al., 2016; Wan Ngah and Hanafiah, 2008).

307 Overall, in aquatic environments, the adsorption capacities of biochars toward heavy
308 metals vary substantially across feedstocks and pyrolysis temperatures (Usman et al.,
309 2016; Sadegh-Zadeh et al., 2017). For instance, the Cd adsorption capacity of date-palm
310 biochar increases from 26.96 mg g⁻¹ at 300 °C to 43.58 mg g⁻¹ at 700 °C (Usman et al.,
311 2016), while Pb adsorption rises dramatically from 4.13 mg g⁻¹ for pine-wood biochar
312 pyrolyzed at 400 °C to 257.12 mg g⁻¹ at 600 °C (Mohan et al., 2007; Yang et al., 2014).
313 Likewise, Ni exhibits the highest overall sorption capacity among the evaluated metals,
314 with high-temperature biochars achieving adsorption capacities exceeding 1900 mg g⁻¹
315 (Lyu et al., 2018). These trends are primarily attributed to the enlarged specific surface
316 area, increased mineral crystallinity, and enhanced aromaticity of biochars at elevated
317 pyrolysis temperatures, which favor metal retention through mineral precipitation and
318 surface complexation (Wang et al., 2024). However, the adsorption behaviors of anionic
319 species such as As and Cr(VI) are more complex; their sorption efficiencies are generally
320 lower and strongly influenced by solution pH (Chen et al., 2015; Ding et al., 2018; Niazi
321 et al., 2018).

322 In soil systems, biochar functions predominantly by altering the speciation and
323 distribution of heavy metals, thereby reducing their bioavailability and mobility (Nie et
324 al., 2018; Khan et al., 2020). Numerous studies have shown that biochar application
325 significantly decreases the exchangeable or weakly bound fractions of Cd, Pb, Cu, and
326 Zn (e.g., CaCl₂-extractable or NH₄NO₃-extractable fractions). For example, wheat-straw
327 biochars (350–550 °C) reduce CaCl₂-extractable Cd by 28.3–70.9% over a three-year

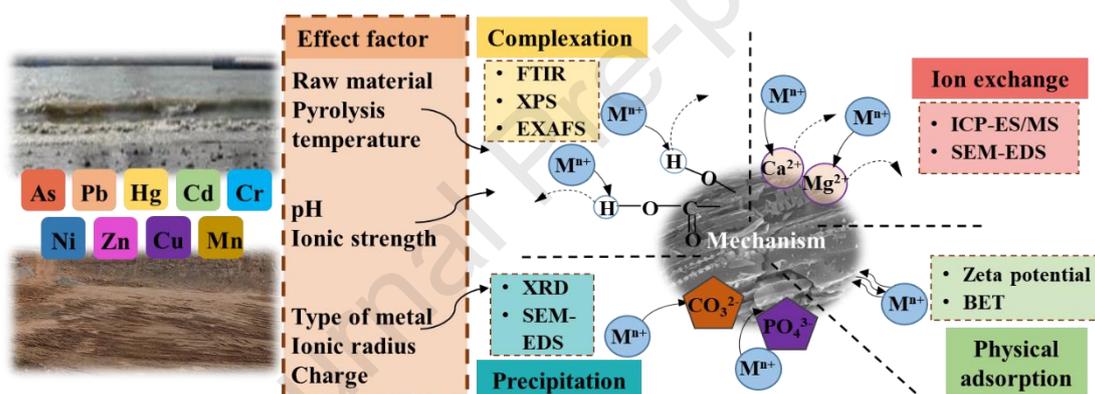
328 period (Bian et al., 2014). Similarly, poultry-manure biochar produced at 550 °C can
329 markedly decrease NH_4NO_3 -extractable Pb from 11.3 mg kg⁻¹ to 0.73 mg kg⁻¹ (Park et
330 al., 2011). Notably, for As, some studies have reported increased or unchanged
331 bioavailable concentrations following biochar addition, likely due to competitive
332 interactions between arsenate/arsenite and phosphate-like functional groups on biochar
333 surfaces, as well as elevated soil pH enhancing As solubility (Brennan et al., 2014; Yin et
334 al., 2016).

335 Importantly, the mechanisms governing remediation differ fundamentally between
336 aquatic and soil environments. In water, the process is dominated by rapid liquid–solid
337 interfacial adsorption kinetics, controlled mainly by diffusion and the competition
338 between H^+ and other cations (Park et al., 2016; Yoon et al., 2017). In contrast, soil
339 remediation involves complex multiphase interfacial processes that integrate interactions
340 among soil minerals, organic matter, plant roots, and microorganisms (Bashir et al., 2018;
341 Xiao et al., 2023). Therefore, the remediation performance of biochar is not dictated by
342 single variables such as pyrolysis temperature, metal speciation, or environmental pH,
343 but rather by the interplay among pollutant chemical characteristics, biochar structural
344 evolution during pyrolysis, and environmental application contexts (Suliman et al., 2016;
345 Xiao et al., 2023; Zhao et al., 2015).

346 **4. Main mechanism of heavy metal remediation by biochar**

347 The physical properties of biochar, including pore structure, specific surface area,
348 cation exchange capacity (CEC), and mineral content, directly influence adsorption and
349 ion-exchange processes. Macropores facilitate metal diffusion, while micropores enhance
350 surface interactions. CEC drives ion exchange between biochar-bound cations and target
351 metals. Mineral components (e.g., carbonates, phosphates, Fe/Mn oxides) promote

352 precipitation or complexation. Environmental conditions such as pH and ionic strength
 353 modulate these pathways. It is essential to reveal the remediation mechanism for
 354 exploring the role of biochar in removing heavy metals in water and soil. According to
 355 the interaction forces between the adsorbent and adsorbate, the remediation mechanism
 356 can be divided into physical adsorption, ion exchange, precipitation and complexation
 357 (shown in Fig. 3) (Chakma et al., 2025; Demirbas, 2008; Inyang et al., 2016; Li et al.,
 358 2023; Robalds et al., 2016). As mentioned above, the differences exist between the
 359 feedstock, environment, and heavy metal, therefore the remediation mechanism is
 360 required and compared shown is Table 5.



361
 362 **Fig. 3** The remediation mechanism of heavy metals by biochar (M, heavy metal).

363 **Table 5** Mechanism of heavy metals remediation by biochar.

Matrix	Feedstock/pyrolysis temperature (°C)	Heavy metal	Main mechanism	Reference
	Date palm/700	Cd(II)	Precipitation	(Usman et al., 2016)
Water	Date palm/300	Cd(II)	Ion exchange, complexation	(Usman et al., 2016)
	Empty fruit bunch/600	Cd(II)	Complexation	(Sadegh-Zadeh et al.,

			2017)
			(Sadegh-
Rice husk/600	Cd(II)	Complexation	Zadeh et al., 2017)
Water hyacinth/450	Cd(II)	Ion exchange, complexation, precipitation	(Zhang et al., 2015)
Corn cob/550	Cd(II)	Ion exchange, complexation	(Luo et al., 2018)
Pine wood/400	Pb(II)	Ion exchange	(Mohan et al., 2007)
Wheat straw/600	Pb(II)	Precipitation, cation exchange	(Cao et al., 2019)
Dairy manure/200	Pb(II)	Precipitation	(Cao et al., 2009)
<i>Alternanthera philoxeroides</i> /600	Pb(II)	Precipitation, complexation	(Yang et al., 2014)
Corn stalk/450	Pb(II)	Ion exchange, complexation, precipitation	(Liu et al., 2019)
Tetra Pak/600	As(III)	Complexation, precipitation	(Ding et al., 2018)
Tetra Pak/600	As(V)	Complexation	(Ding et al., 2018)
Japanese oak wood/500	As(III)	Complexation, precipitation, electrostatic interaction	(Niazi et al., 2018)
Japanese oak wood/500	As(V)	Complexation, precipitation, electrostatic interaction	(Niazi et al., 2018)
Banana peduncle/300	Cr(VI)	Reduction, complexation	(Karim et al., 2015)
Wheat straw/500	Cr(VI)	Electrostatic interaction, reduction	(Zhu et al., 2016)

	Sewage sludge/900	Cr(III)	Precipitation, cation exchange	(Chen et al., 2015)
	Sewage sludge/300	Cr(VI)	Electrostatic interaction, precipitation	(Agrafioti et al., 2014)
	Sewage sludge/300	Cr(III)	Electrostatic interaction, precipitation	(Agrafioti et al., 2014)
	Sugarcane bagasse/600	Ni(II)	Cation- π interaction, electrostatic interaction, complexation	(Lyu et al., 2018)
	<i>Opuntia ficus-indica</i> cactus/600	Ni(II)	Cation exchange, complexation, electrostatic interaction	(Choudhary et al., 2020)
	Dairy manure/350	Cu(II)	Precipitation, cation- π interaction, complexation	(Xu et al., 2013)
	<i>Opuntia ficus-indica</i> cactus/600	Cu(II)	Cation exchange, complexation, electrostatic interaction	(Choudhary et al., 2020)
	Rice straw/550	Zn(II)	Electrostatic interaction, cation exchange, complexation	(Zhao et al., 2020)
	Dairy manure/350	Zn(II)	Precipitation, cation- π interaction, complexation	(Xu et al., 2013)
Soil	Wheat straw/350-550	Cd	Precipitation, cation exchange, complexation	(Bian et al., 2014)
	Wheat straw/350-550	Cd	Precipitation	(Bian et al., 2013)
	Green waste/550	Cd	Precipitation, complexation	(Park et al., 2011)
	Chicken manure/550	Cd	Precipitation, complexation	(Park et al., 2011)
	Sugarcane bagasse/450	Cd	Complexation	(Nie et al., 2018)
	Wood/550	Cd	Complexation	(Namgay et al., 2010)

Water hyacinth/450	Cd	Complexation, precipitation, cation exchange	(Yin et al., 2016)
Rice straw/450	Cd	Complexation, precipitation, cation exchange	(Yin et al., 2016)
Wheat straw/450	Cd	Precipitation, complexation	(Cui et al., 2016)
Wheat straw/350-550	Pb	Precipitation, cation exchange, complexation	(Bian et al., 2014)
Green waste/550	Pb	Precipitation, complexation	(Park et al., 2011)
Chicken manure/550	Pb	Precipitation, complexation	(Park et al., 2011)
Sugarcane bagasse/450	Pb	Precipitation	(Nie et al., 2018)
Water hyacinth/450	Pb	Precipitation, complexation	(Yin et al., 2016)
Rice straw/450	Pb	Precipitation, complexation	(Yin et al., 2016)
Wheat straw/450	Pb	Precipitation, complexation	(Cui et al., 2016)
Wood/550	As	Complexation	(Namgay et al., 2010)
Water hyacinth/450	As	Complexation, precipitation	(Yin et al., 2016)
Rice straw/450	As	Complexation, precipitation	(Yin et al., 2016)
Willow/350 & 550	As	Precipitation	(Gregory et al., 2014)
Sugarcane bagasse/500	Cr	Precipitation	(Bashir et al., 2018)

Sugarcane bagasse/550	Cr	Precipitation	(Khan et al., 2020)
Poplar wood/550	Cr	Precipitation	(Khan et al., 2020)
Bagasse/600	Cr	Precipitation	(Su et al., 2016)
Handwood/600	Ni	Complexation, precipitation	(Shen et al., 2016)
Willow/700	Ni	Precipitation, electrostatic interaction, ion exchange, complexation	(Bogusz and Oleszczuk, 2018)
Green waste/550	Cu	Precipitation, complexation	(Park et al., 2011)
Sugarcane bagasse/450	Cu	Complexation	(Nie et al., 2018)
Handwood/600	Zn	Complexation, precipitation	(Shen et al., 2016)
Rice straw/500	Zn	Complexation, precipitation	(Wu et al., 2018)
Holmoakwood/650	Zn	Precipitation, complexation	(Egene et al., 2018)

364 4.1 Physical adsorption

365 The physical adsorption is electrostatic attraction or van der Waals force, and the
366 adsorption is relatively weak (Rahim and Kassim, 2008; Xiao et al., 2025). Physical
367 adsorption is mainly related to the specific surface area and porosity of the adsorbent
368 (Elaiwu et al., 2014). Zeta potential analysis determines the surface charge of biochar
369 under relevant pH conditions, clarifying its ability to attract or repel metal ions through
370 electrostatic interactions (Choudhary et al., 2020). Brunauer–Emmett–Teller (BET)

371 analysis provides information on specific surface area, pore size distribution, and
372 structural morphology, demonstrating their positive correlation with heavy-metal uptake
373 and highlighting the contribution of physical adsorption. While Cao et al. (2018) found
374 the specific surface areas and pore volumes, were not vital parameters for determining
375 the adsorption capacity of Pb(II) onto wheat straw samples, the reason was that the
376 influence of physical adsorption on Pb(II) adsorption amount was negligible. Also, Hi
377 gashikawa et al. (2016) showed the removal efficiencies of Ni by sawdust-derived
378 biochars under 350 °C (CM350) and 650 °C (CM650) were 8.45 and 10.94 mg/g,
379 respectively, which may be due to the greater surface area and better development of the
380 microstructures at higher temperatures. Therefore, the increased specific surface area and
381 porosity could contribute to the adsorption of heavy metal when the physical adsorption
382 occupies a certain proportion in the adsorption mechanism.

383 *4.2 Ion exchange*

384 Ion exchange is mainly combined with the adsorbent through the exchange of cations
385 (Na^+ , Mg^{2+} , K^+ , and Ca^{2+} , etc.) and heavy metal ions (Ahmad et al., 2014; Ahmad et al.,
386 2018; Bian et al., 2014; Cao et al., 2018; Ma et al., 2025). Inductively coupled plasma
387 (ICP) analysis detects changes in solution-phase concentrations of alkali and alkaline
388 earth metal cations (K^+ , Ca^{2+} , Mg^{2+} , Na^+) before and after adsorption, providing
389 quantitative evidence for cation exchange. Scanning electron microscopy coupled with
390 energy-dispersive X-ray spectroscopy (SEM–EDS) is employed to visualize the spatial
391 distribution of these exchangeable cations and their association with heavy metals at the
392 microscale (Niazi et al., 2018). Lu et al. (2012) studied the adsorption mechanism of Pb(II)
393 onto sludge-based biochar, the results showed that ion exchange accounted for the main
394 contribution (45.6% -60.2%), of which Ca^{2+} and Mg^{2+} play an important role instead of

395 K^+ and Na^+ . While Wang and Liu (2017) investigated the capacities of biochars to remove
396 the heavy metals (Cu, Pb, Cd, and Zn), and K^+ played an important role in ion exchange
397 with heavy metals. Through ion exchange, heavy metal ions can form complexes with
398 organic functional groups (e.g. $-COO^-$, $-O^-$) by covalent bond (Bian et al., 2014; Liu et al.,
399 2019; Luo et al., 2018; Park et al., 2011; Usman et al., 2016).

400 *4.3 Precipitation*

401 Precipitation is the combination of heavy metal ions and anions (OH^- , CO_3^{2-} , PO_4^{3-} ,
402 SO_4^{2-} , and SiO_4^{4-} , etc.) in the adsorbent to form precipitated salts (Ahmad et al., 2014;
403 Ma et al., 2025; Wang et al., 2015; Xu et al., 2017). X-ray diffraction (XRD) reveals
404 newly formed crystalline mineral phases by identifying characteristic diffraction peaks
405 corresponding to heavy-metal precipitates (Cao et al., 2019). SEM-EDS complements
406 XRD by mapping microregions enriched in heavy metals and demonstrating their co-
407 localization with elements such as C and Ca, supporting the occurrence of mineral
408 precipitation (Cao et al., 2019). Zhang et al. (2017) analyzed removal efficiencies of Pb(II)
409 onto celery-derived biochars under 350 and 500 °C, respectively. The results showed that
410 the precipitation contributed the most of adsorption mechanism, mainly in the form of
411 hydrocerussite ($2PbCO_3 \cdot Pb(OH)_2$) and hydro-pyromophite ($Pb_5(PO_4)_3OH$). Chen et al.
412 (2017) studied the precipitation mechanism of Pb onto wheat straw-derived biochar by
413 water-washing, and the contribution rate of anions (SO_4^{2-} , CO_3^{2-} , SiO_3^{2-} and PO_4^{3-})
414 followed the order: $SO_4^{2-} > CO_3^{2-} > SiO_3^{2-} > PO_4^{3-}$. Bian et al. (2014) pointed out that
415 precipitates ($Pb_5(PO_4)_3OH$ and $Cd(OH)_2$) were formed on the surface of biochar, which
416 was due to present of phosphate and hydroxide ions. Thus, the precipitation is directly
417 related to the cations or anions content, and is dominated by the mineral content (Cao et
418 al., 2019; Xu et al., 2017).

419 *4.4 Complexation*

420 Complexation mainly means the functional group on the surface of the adsorbent
421 and heavy metal ions to share the electronic equivalent to form complexes (He et al., 2025;
422 Wang et al., 2018; Zhang et al., 2017). Fourier-transform infrared spectroscopy (FTIR)
423 verifies the participation of oxygen-containing functional groups—such as C=O and O–
424 H—in metal coordination through characteristic peak shifts (Cao et al., 2019). X-ray
425 photoelectron spectroscopy (XPS) provides electronic-level evidence by revealing
426 changes in the O 1s high-resolution spectra (e.g., transformation of C=O to C–O–M
427 structures) and the emergence or binding-energy shifts of metal-specific peaks (e.g., Pb
428 4f) (Xiao et al., 2025). Extended X-ray absorption fine structure (EXAFS) further
429 confirms the formation of inner-sphere complexes by identifying atomic species (e.g.,
430 Pb–O), coordination numbers, and first-shell bond lengths (Lin et al., 2025). Uchimiya et
431 al. (2011) revealed that the stabilization ability of heavy metals (especially Pb and Cu) by
432 biochars directly correlated with their high containing oxygen functional groups, which
433 could be investigated in detail by the method of Fourier-transform infrared spectroscopy
434 (Song et al., 2014; Yuan et al., 2011). Yin et al. (2016) investigated that the varying effect
435 of biochar on Cd, Pb and As mobility in a multi-metal contaminated paddy soil, the results
436 revealed that the acidic functional groups and the hydroxylated surface of Fe/Mn oxides
437 could contribute to Cd immobilization by providing more complexing sites. Wu et al.
438 (2018) explored that the presence of biochar increased the proportion of Zn-organic
439 material in soils, which was mainly due to the fact that organic material could form
440 complexes with heavy metal, and surface oxygenated functional groups such as -OH and
441 -COOH could provide the binding sites. The O, N, and S, etc. in the functional groups (-
442 COOH, -OH, -NH, and -SH, etc.) can be used as coordinating atoms to provide electron

443 pair coordination with heavy metals (Qambrani et al., 2017; Rajapaksha et al., 2016).

444 The mechanistic evidence summarized in Table 5 supports the trends identified in
445 Table 4 and further explains the variability observed across studies. At lower pyrolysis
446 temperatures (<500 °C, particularly 300–450 °C), biochars retain abundant oxygen-
447 containing functional groups (e.g., –COOH, –OH), relatively high cation exchange
448 capacity (CEC), and a degree of amorphous carbon structure. Under these conditions, ion
449 exchange and surface complexation serve as the dominant mechanisms governing the
450 immobilization of cationic metals such as Cd and Cu (Choudhary et al., 2020; Usman et
451 al., 2016; Yin et al., 2016). For example, Cd(II) adsorption by water-hyacinth biochar
452 produced at 450 °C (70.3 mg g⁻¹) involves multiple mechanisms including ion exchange,
453 complexation, and precipitation (Zhang et al., 2015).

454 In contrast, high-temperature biochars (>500 °C, especially ≥600 °C) exhibit
455 markedly increased aromaticity, specific surface area, porosity, and surface alkalinity.
456 These structural characteristics enhance precipitation, complexation, and physical
457 adsorption interactions with heavy metals (Choudhary et al., 2020; Usman et al., 2016).
458 For certain metals such as Ni and Cu, cation– π interactions also become significant, which
459 refers to the electrostatic interaction between heavy metal cations and electron rich π
460 clouds on aromatic rings in the graphitized region of biochar (Lyu et al., 2018). For
461 instance, biochar produced from *Alternanthera philoxeroides* at 600 °C demonstrates a
462 Pb(II) adsorption capacity of 257.12 mg g⁻¹, primarily driven by precipitation and
463 complexation mechanisms (Yang et al., 2014). Notably, phosphorus-rich biochars (e.g.,
464 those derived from animal manure) tend to promote phosphate precipitation, whereas
465 Ca/Mg-rich biochars (e.g., those produced from crop residues) facilitate carbonate
466 precipitation (Cao et al., 2019; Xu et al., 2013).

467 Overall, cationic metals are mainly immobilized through ion exchange, electrostatic
468 attraction, and mineral precipitation—processes favored by the high pH and mineral-rich
469 surfaces produced at elevated pyrolysis temperatures (Yin et al., 2016; Bian et al., 2014).
470 In contrast, oxyanion species rely predominantly on ligand exchange and surface
471 complexation. These pathways diminish at higher temperatures as oxygen-containing
472 functional groups volatilize (Gregory et al., 2014; Bashir et al., 2018; Khan et al., 2020).
473 In summary, heavy-metal immobilization by biochar is governed by distinct dominant
474 mechanisms in different environmental systems, yet ultimately results from the
475 synergistic contributions of multiple interacting processes (Lin et al., 2025; Ma et al.,
476 2025).

477 **5. Challenge**

478 Short-term adsorption performance of biochar in field environments is consistently
479 lower than laboratory estimates. This discrepancy arises from competitive interactions
480 with coexisting ions, the presence of natural organic matter, and temporally variable redox
481 and pH conditions, which limit the effective availability of active sorption sites and
482 reduce the metal-binding efficiency observed under idealized laboratory conditions (Lin
483 et al., 2025; Ma et al., 2025).

484 Furthermore, Biochar exhibits declining regeneration efficiency during repeated
485 adsorption–desorption cycles, this decrease is primarily associated with pore clogging,
486 mineral precipitation, and the loss or transformation of surface functional groups (Guo et
487 al., 2025; Chakma et al., 2025). Such structural and chemical degradation reduces the
488 long-term stability of active sites and complicates efforts to maintain performance in
489 dynamic environmental settings.

490 Biochar aging presents a major challenge to sustained heavy metal remediation

491 performance. Abiotic oxidation, biotic weathering, wet–dry and freeze–thaw cycles, and
492 photo-oxidation collectively alter surface chemistry, pore architecture, and mineral
493 composition (Kumar et al., 2018; Long et al., 2024). Although aging can increase oxygen-
494 containing functional groups, enhancing complexation for some metals, it may also
495 induce physical degradation, pore collapse, and blockage (Kumar et al., 2018; Ma et al.,
496 2025). Therefore, aging effects are context-dependent; we recommend standardized aging
497 tests and multi-cycle regeneration studies to assess long-term stability.

498 In addition, scaling up biochar applications faces significant constraints related to
499 feedstock variability, technological bottlenecks, and economic constraints (Aziz, 2024).
500 Additional concerns arise from the potential release of polycyclic aromatic hydrocarbons
501 (PAHs), heavy metals, nanoparticles, and soluble organic fractions, particularly when
502 biochar is derived from contaminated or inadequately pyrolyzed feedstocks, posing
503 ecological risks, including adverse effects on soil biota and water quality (Guo et al., 2024;
504 Long et al., 2024).

505 **6. Conclusion and Prospects**

506 This review synthesizes current knowledge on biochar properties, influencing
507 factors and remediation mechanisms for heavy metals in aquatic and terrestrial
508 environments. The key conclusions are as follows: (1) The remediation performance of
509 biochar is highly dependent on feedstock type, pyrolysis parameters, and environment
510 conditions; (2) Complexation, ion exchange and mineral precipitation often co-occur and
511 their relative importance is system-specific; (3) Aging, redox/pH dynamics and natural
512 organic matter critically modulate stability and must be systematically tested.

513 Overall, Biochar has a great potential for application in the heavy metal remediation.
514 At present, the existing research has focused on the improvement of heavy metal

515 remediation, its influence factors and related mechanism, etc. However, the aging of
516 biochar, its component release, and the effects of complex environment systems during
517 heavy metal remediation are very limited. Considering the application of biochar, it is
518 very important to evaluate its stability, repeatability and reproducibility (Awual et al.,
519 2013; Aziz, 2024; Huang et al., 2017; Xu et al., 2018). The water-soluble components in
520 biochar are rich in carboxylic acid, humic acid, soluble mineral ions, and nanoparticles,
521 posing potential environmental risks, thus the relevant studies on pretreatment method is
522 necessary to minimize the negative impacts (Feng et al., 2022; Ghidotti et al., 2017; Guo
523 et al., 2024; Smith et al., 2016). At the same time, most of the experiments on heavy metal
524 remediation by biochar are performed in laboratory, but there is still lack of data for
525 natural conditions. Thus, the long-term monitoring research on heavy metal remediation
526 by biochar in the water and soil needs to be improved.

527 Future development directions focus on: (1) Adopt standardized reporting (feedstock,
528 pyrolysis conditions, full characterization, and experimental matrices); (2) Employ
529 spectroscopic/microscopic methods to link mechanisms to evidence; (3) Implement
530 standardized aging tests and multi-cycle regeneration studies to assess long-term stability;
531 (4) Perform pilot field trials with safety evaluation (soluble organics, PAHs and heavy
532 metal in nanoparticles) and life-cycle economic assessment before scale-up. These
533 targeted steps will accelerate safe, effective deployment of biochar for heavy-metal
534 remediation.

535 **CRedit authorship contribution statement**

536 **Yaoyao Cao:** Data curation, Funding acquisition, Visualization, Writing - original
537 draft. **Jian Chen:** Formal analysis, Writing - review & editing. **Hui Zhang:** Methodology,
538 Validation. **Hongjie Sheng:** Conceptualization, Investigation. **Kang Lv:** Methodology,

539 Validation. **Jinjin Cheng**: Formal analysis, Investigation. **Xiaolong Chen**: Formal
540 analysis, Writing - review and editing. **Dongfei Chen**: Data curation, Writing - review
541 and editing. **Xiangyang Yu**: Conceptualization, Funding acquisition, Supervision,
542 Writing - review & editing.

543 **Declaration of competing interest**

544 The authors declare that they have no known competing financial interests or
545 personal relationships that could have appeared to influence the work reported in this
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Highlights

- Biochar is widely used to the remediation of heavy metal in water and soil.
- Physicochemical properties and regularity of biochar are elaborated.
- Main factors affecting on the remediation of heavy metal by biochar are explored.
- Differences exist in the mechanisms of heavy metals remediation by biochar.
- The challenges and future studies on the application of biochar are prospected.

Declaration of competing interests

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.

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