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### Article

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1 **Modeling transport and fate of heavy metals at the watershed**  
2 **scale: state-of-the-art and future directions**

3

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16

## 17 **Abstract**

18       A predictive understanding of the source-specific (e.g., point and diffuse sources)  
19 land-to-river heavy metal (HM) loads and HM dynamics in rivers is essential for  
20 mitigating river pollution and developing effective river basin management strategies.  
21 Developing such strategies requires adequate monitoring and comprehensive models  
22 based on a solid scientific understanding of the watershed system. However, a  
23 comprehensive review of existing studies on the watershed-scale HM fate and transport  
24 modeling is lacking. In this review, we synthesize the recent developments in the  
25 current generation of watershed-scale HM models, which cover a wide range of  
26 functionalities, capabilities, and spatial and temporal scales (resolutions). Existing  
27 models, constructed at various levels of complexity, have their strengths and  
28 weaknesses in supporting diverse intended uses. Additionally, current challenges in the  
29 application of watershed HM modeling are covered, including the representation of in-  
30 stream processes, organic matter/carbon dynamics and mitigation practices, the issues  
31 of model calibration and uncertainty analysis, and the balance between model  
32 complexity and available data. Finally, we outline future research requirements  
33 regarding modeling, strategic monitoring, and their combined use to enhance model  
34 capabilities. In particular, we envisage a flexible framework for future watershed-scale  
35 HM models with varying degrees of complexity to accommodate the available data and  
36 specific applications.

## 37 **Keywords**

38 Heavy metals, catchment-scale, fate and transport model, point and diffuse source  
39 pollution, heavy metal land process, in-stream heavy metal process

## 40 **1 Introduction**

41 Elevated concentrations of heavy metals (HMs) are a global threat to aquatic  
42 systems and human health owing to their potential accumulation, biomagnification, and  
43 toxicity. The 'heavy metals' is a collective term used here to represent a group of metals  
44 and metalloids such as cadmium (Cd), mercury (Hg), copper (Cu), arsenic (As), lead  
45 (Pb), chromium (Cr), zinc (Zn), and nickel (Ni) that cause toxicity and ecotoxicity.  
46 Although there still exist debates concerning the imprecise use of the term 'heavy  
47 metals' and its alternatives, including 'toxic metals', 'trace metals', 'trace elements', and  
48 'potentially toxic trace elements' (Duffus, 2002), we used the term 'heavy metals'  
49 consistently in this review for convenience. HMs are ubiquitous in the environment as  
50 naturally occurring elements. For example, As pollution from geogenic sources is a  
51 major problem in South Asia (e.g., Winkel et al., 2008). Meanwhile, HMs, such as Cd,  
52 Hg, Cu, As, Cr, and Pb, have contaminated river systems in many parts of the world as  
53 a by-product of industrialization (Johnson et al., 2018; Mason, 2013). For example, the  
54 signature of Pb mining and smelting activities by ancient Greeks and Romans have been  
55 documented in the Greenland ice cores (Hong et al., 1994). Moreover, over 100,000  
56 abandoned or inactive mining sites are spread over 2,000 km<sup>2</sup> in the United States

57 (U.S.EPA, 1997). England and Wales have over 3,000 abandoned metal mines (Jarvis  
58 et al., 2007). In China, a nationwide survey of soil contaminants during 2005–2013  
59 revealed that 16.1% of the soil samples exceeded the soil quality standards mainly  
60 because of high HM concentrations (MEPPRC and MLRPRC, 2014).

61 Knowledge of the HMs released from contaminated soils and sites in the upland  
62 and their subsequent migration in river networks is critical for assessing environmental  
63 risks, as well as developing effective pollution control and river basin management  
64 strategies. (Byrne et al., 2012; Le Roux et al., 2020). HMs can enter river systems via  
65 various diffusion pathways, including surface runoff, subsurface flow, groundwater  
66 flow, and soil erosion. Meanwhile, many biogeochemical processes greatly influence  
67 and regulate HM's mobility (Carrillo - González et al., 2006). These mechanisms  
68 involve sorption, complexation, precipitation, redox reactions as well as weathering  
69 processes influenced by environmental factors such as pH, redox potential, and  
70 temperature (Borch et al., 2010; Degryse et al., 2009). Extensive studies, including field  
71 measurements, have been performed on the mechanisms of different soil  
72 biogeochemical reactions at the plot and field scales (Selim and Kingery, 2003).  
73 However, when it turns to the watershed scales, a quantitative and complete description  
74 of metal migration may be hindered by the strong spatial heterogeneity and temporal  
75 variability of HM production, transformation, and transport processes (Li, 2019). To  
76 fully understand the complex mechanisms of HMs transport at a watershed scale, a  
77 large amount of time- and resource-consuming fieldwork is needed. Watershed-scale

78 HM models are increasingly used as essential tools for assessing and restoring surface  
79 waters because they can serve several aims: manifesting and examining pollutant  
80 transport mechanisms or hypotheses (Aim 1, improving scientific understanding);  
81 assessing the environmental risk (Aim 2, risk assessment); estimating pollutant fluxes  
82 and locating critical source areas (Aim 3 Identifying critical source areas); and  
83 evaluating the impacts of climate change and mitigation scenarios on water quality  
84 (Aim 4, scenario analysis).

85 Over the last few decades, water quality models concerning the quantification of  
86 nutrients (e.g., nitrogen and phosphorus) dynamics at the watershed-scale have been  
87 extensively reviewed ([Robson, 2014](#); [Rode et al., 2010](#); [Wellen et al., 2015](#)). In contrast,  
88 reviews on HM dynamics models remain scarce ([Ouyang et al., 2017](#); [Qiao et al., 2023](#)).  
89 Previous reviews have focused mainly on the spatiotemporal distribution of HMs in  
90 soil and river systems, their transformation and partitioning at solid-liquid interfaces  
91 (e.g., [Degryse et al., 2009](#)), and plot, field, and river-scale numerical transport models  
92 (e.g., [Carrillo-González et al., 2006](#); [Garneau et al., 2017](#)). To our knowledge, a holistic  
93 description of watershed HM processes accounting for natural and anthropogenic inputs,  
94 terrestrial delivers into streams, and in-stream dynamics is lacking. The last significant  
95 and most relevant review came from a USEPA workshop in 2007 ([Caruso et al., 2008](#)),  
96 which gathered experts from academia and the government to explore state-of-the-art  
97 models for simulating metal fate and transport from different scales and application  
98 domains, such as equilibrium, stream, and watershed models. [Caruso et al. \(2008\)](#) stated

99 that existing watershed models require further testing and evaluation using more field  
100 data, and a truly calibrated watershed HM model did not exist. In the last 15 years, new  
101 HM watershed models have been developed ([Meng et al., 2018](#); [Motovilov and](#)  
102 [Fashchevskaya, 2019](#); [Sui et al., 2022](#)), and monitoring methods have advanced ([Byrne](#)  
103 [et al., 2021](#); [Frau et al., 2018](#); [Hanhauser et al., 2020](#)). However, no follow-up review  
104 has been conducted till date. Therefore, we present a contemporary analysis of existing  
105 watershed pollution models by comparing the differences in functionality and  
106 underlying assumptions to highlight the challenges and opportunities for future model  
107 research and development.

108 Owing to the complexity of natural environmental systems, models are always  
109 necessarily simplified descriptions of reality. Moreover, models are often constructed  
110 at different levels of complexity in terms of structure, function, and processes  
111 depending on scientific knowledge, purposes, and data availability. Therefore, we  
112 evaluate models' strengths and weaknesses in this review according to their diverse  
113 intended uses. The remainder of this review is organized as follows. Section 2 outlines  
114 the primary sources, transformation mechanisms, and transport pathways of HMs in the  
115 natural environment. Section 3 reviews contemporary HM watershed models and  
116 highlights their basic assumptions, simplifications, components, and distinctions.  
117 Section 4 summarizes the research gaps and challenges in watershed HM modeling and  
118 applications. Finally, Section 5 concludes with future research needs in HMs  
119 monitoring, modeling, and their combined use.

## 120 **2 Fate and transport of HMs in the real world**

### 121 **2.1 Sources of HMs in the environment**

122 There are two main categories of HM sources: natural and anthropogenic. Natural  
123 sources mainly include bedrock weathering, volcanic eruptions, and atmospheric  
124 deposition. Anthropogenic sources (e.g., industrial, agricultural, and municipal) have  
125 dramatically changed HM concentrations in the natural environment compared with  
126 pre-industrial times (Chen et al., 2008). For example, HM pollution in soil and water is  
127 often associated with metal mining and smelting, chemical production, dyeing and  
128 printing, and burning fossil fuels (Rauch and Pacyna, 2009). In addition, applying  
129 fertilizers, pesticides, manure, and wastewater irrigation has substantially increased  
130 HM fluxes to local water bodies near agricultural land (Shi et al., 2018). Domestic  
131 sewage discharge from treatment plants in urban areas can also contribute significant  
132 quantities of HMs to aquatic environments (Zhou et al., 2018). Vehicular exhaust and  
133 wear and tear of tires are other important HM sources (Werkenthin et al., 2014). For  
134 example, Zn isotopes in sediment cores of eight lakes across the United States indicated  
135 that vehicle emissions are the most significant source of Zn (Thapalia et al., 2015).  
136 Finally, it should be noted that human activities are deeply intertwined with natural  
137 processes, sometimes blurring the boundaries. For example, HMs emitted from  
138 industrial activities are transported through the atmosphere, deposited in the soil, and  
139 then transported to water bodies via runoff and erosion (Andronikov et al., 2021).



140 Water pollutants come from either point sources or dispersed sources. Thus, HM  
141 sources can also be divided into the point and nonpoint sources. A point source of  
142 pollution is discharged directly from a discernible source, such as a discharge pipe from  
143 a factory or sewage plant. For example, the Seine (Chen et al., 2008) and Rhine  
144 (Stigliani et al., 1993) river basins in Europe and Dongting Lake in China (Li et al.,  
145 2013) receive most of their HM pollution from point sources. Unlike point sources,  
146 diffuse (nonpoint) sources do not originate from a specified location. Instead, they are  
147 characterized by intermittent occurrences and spatial heterogeneity, such as runoff from  
148 agricultural land during rainfall, which makes them challenging to trace and control  
149 (Patterson et al., 2013).

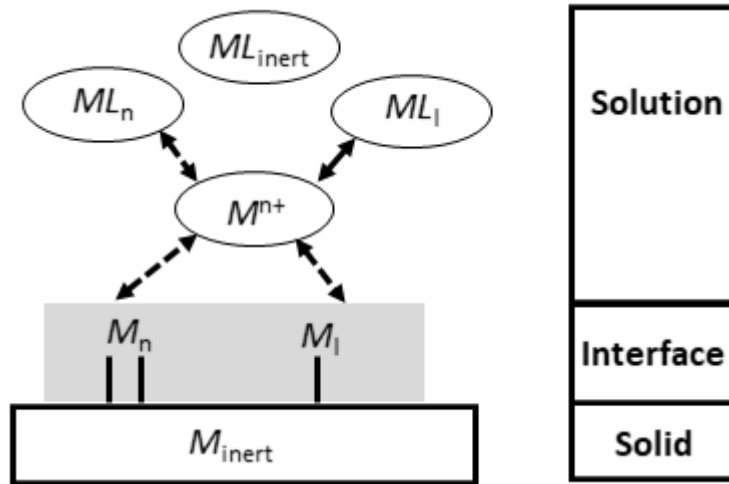
## 150 **2.2 Transformation mechanisms**

151 HM's mobility and bioavailability in terrestrial and aquatic systems depends on  
152 chemical speciation (van Leeuwen et al., 2005). In the most simple sense, chemical  
153 speciation refers to whether a metal exists in a dissolved/solute form or a solid/particle-  
154 contained form (Fig. 1). In the natural environment, several reactions can change HM  
155 speciation, such as sorption (adsorption/desorption), complexation  
156 (association/dissociation), precipitation/dissolution, diffusion into carbonates and  
157 oxyhydroxides (Degryse et al., 2009). In the solid phase, HMs are present as labile ( $M_l$ ),  
158 non-labile ( $M_n$ ), or inert metal ( $M_{inert}$ ) species. Labile metal can exchange rapidly with  
159 the solution phase. Furthermore, metals in the labile pool in the solid phase can slowly

160 transfer from/to a non-labile pool, which is a slow process taking years or longer  
161 (Buekers et al., 2008; Crout et al., 2006). In contrast, inert metals in parent minerals are  
162 unlikely to be released to a solution phase. In the solution phase, metals appear as free  
163 ions ( $M^{n+}$ ), complexes with inorganic or organic ligands ( $ML$ ), or associated with  
164 mineral colloids ( $ML_{\text{inert}}$ ) (Honeyman and Santschi, 1988). Generally, the free ions in  
165 the solution tend to react most actively with the solid phase. However, similar to inert  
166 solid metals, some metals in solution, such as those in colloidal minerals, may also be  
167 non-reactive. Metals complexes with inorganic or organic ligands could be divided into  
168 labile and non-labile  $ML$  ( $ML_{\text{labile}}$  and  $ML_{\text{non-labile}}$ ) according to the dissociation rate of  
169 these complexes.

170 Soil HM's solubility and mobility are governed by soil properties such as soluble  
171 ligands in soil pore water, soil matrix composition (e.g., oxides, clay, and organic  
172 matter), pH, temperature, and redox potential (Young, 2013). In soil solutions, HM  
173 species are significantly affected by the presence of different organic and inorganic  
174 ligands. The soil solid phase consists of various constituents (clay minerals, organic  
175 matter, iron, and aluminum oxides); HM species in the solution react with these  
176 constituents via different mechanisms. Soil pH is considered the most critical factor  
177 influencing metal speciation in soils, thus the master variable affecting metal behavior  
178 in soil systems. In addition, the increase/decrease of soil redox potential (Eh) could  
179 regulate a series of biogeochemical reactions. For example, some variable-valence HMs  
180 such as Hg, As, Cr, and Fe could undergo valence changes within the range of redox

181 potentials (Carrillo - González et al., 2006). Biomethylation could occur under  
182 anaerobic conditions, transforming the inorganic forms of Hg and As into methylated  
183 forms.



184  
185 Fig. 1 Schematic representation of metal species and reactions between the solid and  
186 solution phases. Broken and solid reversible arrows represent kinetically constrained  
187 and ‘instantaneous’ reactions, respectively Young (2013).

### 188 2.3 Transport pathways

189 HMs can enter river systems via various diffusion pathways, such as surface runoff,  
190 soil leaching, subsurface flow, groundwater flow, erosion, and atmospheric deposition  
191 (Fig. 2) (Foster and Charlesworth, 1996). Surface runoff occurs when the soil becomes  
192 saturated or rainfall intensity exceeds the infiltration rate (Yang et al., 2015). The  
193 amount and rate of metal fluxes in surface runoff depend on metal speciation and  
194 concentration in the soil, rainfall intensity, and watershed characteristics. Surface runoff  
195 moves quickly to the stream channel. Thus, the immediate response of the stream’s

196 metal concentrations to increased streamflow is an indicator of the surface runoff  
197 process because they have an immediate effect on the solutes of the stream ([Runkel et](#)  
198 [al., 2016](#)).

199 After water infiltrates the soil, it undergoes the processes of vertical leaching and  
200 lateral flow, with the concomitant of dissolved metals. Soil leaching is the downward  
201 movement of dissolved metals in the soil profile via percolating water. Subsurface  
202 lateral flow refers to soil water processes where infiltrating water accumulates and  
203 moves laterally downslope along the upper surface of a less permeable layer in the soil.  
204 Subsurface lateral flow is abundant at the interface between the soil and bedrock, where  
205 permeability changes dramatically. A range of terms are used to refer to subsurface  
206 lateral flow, including throughflow, subsurface runoff, and interflow. The relative  
207 contribution of soil leaching or subsurface lateral flow to HM transport varies largely  
208 depending on the soil attributes and topographic and meteorological conditions. [Xia et](#)  
209 [al. \(2014\)](#) found that soil leaching is the dominant export pathway of soil HM in the  
210 southern Song-nen Plain of Northeast China. The subsurface flow from acid mine  
211 drainage in Cement Creek, Colorado, is the largest contributor to Zn loads in the  
212 watershed ([Kimball et al. \(2002\)](#)). A national-scale modeling in the Netherlands reported  
213 that soil leaching was the most important contributor of Cd (20%), Zn (40%), and Pb  
214 (40%) to surface water ([Bonten et al., 2008](#)). Enhanced metal leaching is associated  
215 with acidic drainage due to high metal solubility and sulfide weathering rates under  
216 acidic conditions ([RoyChowdhury et al., 2015](#)).

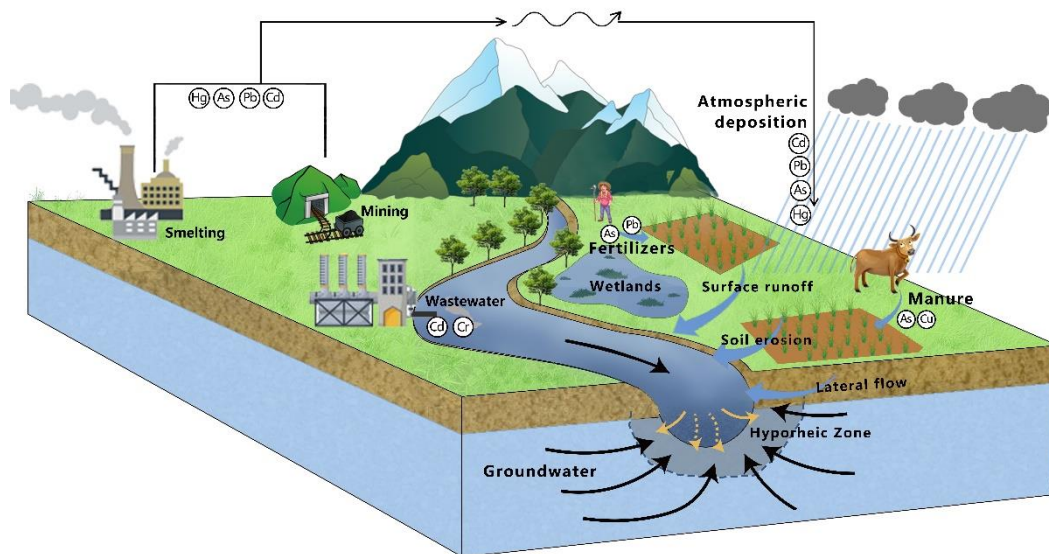
217 Groundwater discharge is another possible input of HMs into surface waters.  
218 Groundwater containing dissolved contaminants migrates from the soil into subjacent  
219 aquifers and finally enters adjacent streams. Several hydrological and biochemical  
220 factors determine the amount of metal transported from the groundwater to streams.  
221 Generally, HM fluxes via groundwater discharge are significant in mountainous mining  
222 areas. Polluted aquifers act as long-term pollution sources for the surrounding rivers,  
223 even after mining activities have stopped (Wang et al., 2019). For example, the inputs  
224 of Cd (43%) and Zn (28%) to the Riou Mort River in France are mainly through  
225 groundwater discharge (Coynel et al. (2007). In contrast, fieldwork in the Gilt Edge  
226 Mine, South Dakota, USA, showed that metal loads from bedrock fractures contributed  
227 <1% of the total load to the creek (Caruso and Dawson, 2008). Additionally, the slow  
228 groundwater movement and HM attenuation in aquifers due to sorption could reduce  
229 the metal loads to the streams. Thus, HM transport in subsurface environments can be  
230 significant within certain landscapes and geology.

231 Soil erosion plays an important role in the biogeochemical cycles of HMs.  
232 Scientometric analysis has revealed that erosion-induced transport is the most  
233 influential factor in HMs mitigation mechanisms (Ouyang et al., 2018). A recent  
234 national-scale study across China revealed the ubiquitous prominence of soil erosion  
235 contributions to land-to-river metal fluxes (e.g., As, Cd, Cr, Cu, Ni, Pb, Zn, and Hg)  
236 (Liu et al., 2019).

237 In the absence of local anthropogenic inputs, atmospheric deposition is the most

238 significant HM input to terrestrial environments (Lofts, 2007). For example, an  
239 inventory study of HM inputs to agricultural soils in China showed that atmospheric  
240 deposition accounted for 43–85 % of the total As, Cr, Hg, Ni, and Pb inputs (Luo et al.,  
241 2009). Atmospheric deposition contributes more Zn, Ni, and Pb to European  
242 agricultural soils than phosphate fertilizer application (Nziguheba and Smolders, 2008).

243 After HMs enter rivers, a number of physical (e.g., convection, diffusion, erosion  
244 and deposition of sediments), chemical (e.g., sorption, complexation, precipitation),  
245 and biological (e.g., bioturbation) processes could influence the fate of HMs in aquatic  
246 systems to some degree (e.g., Mason, 2013). The scavenging of HMs in river water  
247 depends largely on the solid-liquid distribution and the presence/properties of  
248 suspended/riverbed sediments (Honeyman and Santschi, 1988). Most HMs entering the  
249 rivers may be immobilized and stored by adsorption onto the riverbed sediments (Peng  
250 et al., 2009). A portion of HMs that adsorbed on fine suspended solids (e.g., hydrous  
251 oxides, clays) could transport downstream over long distances (Hochella et al., 2005).  
252 In addition, "big events", such as large storms and floods, have been shown to  
253 significantly affect HMs' remobilization and transportation (Ciszewski and Grygar,  
254 2016; Peraza-Castro et al., 2016).



255

256 Fig. 2 A schematic diagram of the heavy metal cycle at the watershed scale.

### 257 3 Overview of watershed-scale HM fate and transport models

#### 258 3.1 Existing representative models

259 Nine representative models were selected based on a thorough literature evaluation:

260 L-THIA (Park et al., 2013), METALPOL (Vink and Peters, 2003), WARMF (Chen et

261 al., 2000), ECOMAG-HM (Motovilov and Fashchevskaya, 2019), SWAT-HM (Meng

262 et al., 2018), INCA-Metals (Whitehead et al., 2009), CTT&F (Johnson et al., 2011),

263 TREX (Velleux et al., 2008), and TOPKAPI-ETH (Sui et al., 2022). A detailed

264 description of each model is provided in SI. It is not intended to be an exhaustive list

265 of all available watershed scale HM models. These representative models were selected

266 because they are capable of simulating metal fate and transport processes at the

267 watershed scale with various spatial and temporal resolutions. The manner of HM

268 fate/transport description varies among the models, ranging from simple export  
269 coefficient models (e.g., L-THIA and METALPOL) to more complex integrated  
270 watershed models (e.g., SWAT-HM and TOPKAPI-ETH). It should also be noted that  
271 urban watershed models, such as SWMM model, were not included in this review.  
272 Urban models focus more on the buildup and washoff of pollutants from impervious  
273 surfaces and subsequent transport and transformation processes in the urban water  
274 infrastructure systems. Though urban area is usually a significant part of a watershed,  
275 the prominent particularities of urban models render them incomparable with general  
276 watershed models which mainly target the natural to less urbanized environments.  
277 Useful reviews on urban water quantity and quality modeling can be found in [Zoppou](#)  
278 [\(2001\)](#).

279 As mentioned in Section 2, metal behavior in the natural environment is highly  
280 complex in terms of various pollution sources, transformation reactions, and transport  
281 pathways. Therefore, models are always simplified descriptions of environmental  
282 systems with different levels of complexity. Furthermore, models have evolved over  
283 time, becoming increasingly detailed in terms of spatiotemporal resolution and the  
284 number of components and processes included. The selected models have been  
285 developed for a range of different objectives. They reflect a broad spectrum of concepts  
286 and assumptions, which can be categorized according to various criteria, including  
287 representation of processes (e.g., empirical, conceptual, process-based, or physically-  
288 based), spatial scale (e.g., plot, field, or watershed), spatial discretization (e.g., lumped,



289 semi-distributed, and fully-distributed), and temporal scales (e.g., event-driven or long-  
290 term simulation). However, it is accepted that these categories are sometimes vague,  
291 and in practice, many models include elements of different categories. Generally, two  
292 main processes (contaminant partitioning and transformation and contaminant transport)  
293 are considered in watershed-scale metal models. The contaminant transport component  
294 commonly consists of three processes: (1) overland hydrological processes, (2) soil  
295 erosion and sediment transport, and (3) in-stream processes. The main features and  
296 components of the reviewed models and associated references are summarized in [Table](#)  
297 [1](#), with additional comparisons in [Sections 3.2](#) and [3.3](#).

Table 1 Existing metal models at different spatial and temporal scales considered in this review.

Model	Metal speciation and transformation	Hydrology	Soil erosion	In-stream	Space discretization	Temporal scale	Distinguishing features and applicability	Main developers and key references
L-THIA	Total metal concentration	SCS-CN; do not consider the spatial route of NPS pollution	None	None	Fully-distributed	Daily, yearly	Large-scale and long-term total metal loads where surface runoff pathway dominates.	Purdue University, USA ( <a href="#">Park et al., 2013</a> )
METALPOL	Total metal concentration	A water balance model based on Rhineflow	Modified USLE, sediment delivery ratio (SDR), and the specific enrichment ratio model	None	Fully-distributed	Yearly	Export coefficient method, large-scale and long-term total metal loads considering multiple pathways but ignoring in-stream processes.	Vrije University Amsterdam, Netherlands ( <a href="#">Vink and Peters, 2003</a> )
WARMF	Two-phase equilibrium partitioning	Overland Flow using Manning's equation Lateral flow using Darcy's Law	ANSWERS model	Continuously stirred tank reactor (CSTR) model	Lumped	Daily	Continuously stirred tank reactor (CSTR) model	Electric Power Research Institute (EPRI), USA ( <a href="#">Chen et al., 2000</a> )
TREX	Three-phase equilibrium partitioning	Diffusive wave approximation	CASC2D-SED model	Diffusive wave approximation	Fully-distributed	Variable time step	High-resolution and event-based simulation models for small-scale applications	Colorado State University, USA ( <a href="#">Velleux et al., 2008</a> )
SWAT-HM	Three-phase equilibrium partitioning and reaction	SCS-CN	MUSLE	Variable storage method or Muskingum method	Semi-distributed (HRU)	Daily	Long-term continuous models with reasonable model structure and computational efficiency suitable for meso-scale and large-scale watershed, easy to	Beijing Normal University, China ( <a href="#">Meng et al., 2018</a> )

							couple with other processes such as carbon cycle.	
CTT&F	Four-phase equilibrium partitioning	Designed for use within existing hydrological modeling systems	Designed for use within existing hydrological modeling systems	Designed for use within existing hydrological modeling systems	Fully-distributed	Variable time step	Metal fate and transport modeling at small watershed scale such as arms firing ranges	US Army Corps of Engineers ( <a href="#">Johnson et al., 2011</a> )
INCA-Metals	Two-phase equilibrium partitioning	Quick flow, soil water flow and groundwater flow.	Erosion is by splash detachment and flow erosion of bulk sediment; five grain size classes are considered for in-stream processes	Nonlinear reservoir model	Semi-distributed (landscape units)	Daily	Process-based representation, minimized data requirements and model structural complexity, easy to couple with other processes such as carbon cycle.	Reading university, UK ( <a href="#">Whitehead et al., 2009</a> )
ECOMAG-HM	Two-phase equilibrium partitioning	ECOMAG hydrological model	None	Kinematic wave equation	Semi-distributed	Daily	Accounts for process of dissolution by melt and rainwater, suitable for snow-dominated watershed	Russian Academy of Sciences, Russia ( <a href="#">Motovilov and Fashchevskaya, 2019</a> )
TOPKAPI-ETH	Four-phase equilibrium partitioning and reaction	Kinematic wave approximation	Overland flow erosion using transport capacity approach	Kinematic wave approximation	Fully-distributed	Variable time step	Physically explicit representation of the major hydrology-sediment-metal processes with a reasonable computational efficiency suitable for small-scale watersheds.	ETHZ, Switzerland ( <a href="#">Sui et al., 2022</a> )

## 300 **3.2 Representation of metal partitioning and transformation**

### 301 **3.2.1 Existing metal transformation schemes**

302 Existing models generally have four levels of complexity for reaction mechanisms  
303 (Fig. 3) (Degryse et al., 2009). The level 4 scheme (Fig. 3a) represents all the main  
304 metal species and transformation processes mentioned in Section 2.2. Although  
305 conceptually appealing, it requires extensive input information, often unavailable in  
306 routine research. The level 3 scheme does not consider the differences between non-  
307 labile and inert metals in the solid and solution phases. However, it requires  
308 simultaneous modeling or observation of dissolved organic matter (DOM) owing to its  
309 explicit modeling of metal complexes. The level 2 scheme consists of three metal pools:  
310 dissolved metal ( $M_d$ ), labile metal ( $M_l$ ), and non-labile metal ( $M_n$ ), in which free metal  
311 ions ( $M^{n+}$ ) and metal complexes ( $ML$ ) in solution are regarded as  $M_d$ . The last and most  
312 simplified level 1 scheme consists of two pools:  $M_d$  and particulate metal ( $M_p$ ), in which  
313  $M_l$  and  $M_n$  are further regarded together as  $M_p$ .

314 As discussed above, metal partitioning and transformation in the natural world are  
315 highly complex. Several mechanistic models (also called equilibrium and geochemical  
316 models), such as MINEQL (Westall et al., 1976), MINTEQA2 (Allison et al., 1991),  
317 WHAM (Tipping, 1994), and ORCHESTRA (Meeussen, 2003), have been developed  
318 to describe metal partitioning between solid and solution, or metal speciation in solution  
319 only. For example, the WHAM model could describe metal sorption on organic matter  
320 by nonspecific electrostatic sorption and specific competition sorption (protons and

321 metals compete for binding to two types of sites: carboxylic and phenolic groups)  
322 ([Tipping, 1998](#)). However, watershed-scale studies rarely provide such detailed input  
323 or validation information required for these mechanistic models. Most watershed-scale  
324 models employ the simpler level 2 and level 1 schemes that conceptually capture the  
325 dominant mechanism. For level 2, the HM transformation model considers two major  
326 reactions: (1) sorption and (2) slow reactions.

### 327 **Sorption**

328 Sorption refers to the adsorption-desorption processes between the dissolved metal  
329 in the solution phase and the labile metal in the solid phase. Several studies have  
330 demonstrated that adsorption-desorption is the most important process affecting the  
331 mobility and bioavailability of metals ([Degryse et al., 2009](#)). Dissolved metals (solution  
332 phase,  $[M_d]$ ) are reversibly adsorbed onto solids and become labile adsorbed metals  
333 (solid phase,  $M_l$ ). The solid-solution partition coefficient ( $K_d$ ) is defined as the ratio of  
334 labile metal concentration in the solid phase to dissolved metal concentration in the  
335 solution phase when equilibrium is attained:

$$336 \quad K_d = \frac{M_l}{[M_d]} \quad (1)$$

337 where  $K_d$  is the solid-solution partition coefficient ( $L\ kg^{-1}$ ), and  $M_l$  and  $[M_d]$  denote the  
338 labile metal concentration in the solid phase ( $mg\ kg^{-1}$ ) and the dissolved metal  
339 concentration in the solution phase ( $mg\ L^{-1}$ ), respectively.

340 It should be noted that  $K_d$  is an apparent (lumped) partition coefficient for  
341 describing the equilibrium speciation of metals between solid and solution phases, as it

342 describes both the sorption and complexation of free ions in solution rather than a single  
343 mechanism. Moreover, because  $K_d$  is a strict equilibrium concept, equilibrium is  
344 implicitly assumed when adopting the  $K_d$ -based model. Nevertheless, the  $K_d$  approach  
345 is often used in the existing watershed model for the following reasons. (1)  $K_d$  value  
346 can be determined from an extensive literature search (e.g., Allison and Allison, 2005)  
347 or estimated through laboratory adsorption/desorption batch tests. (2) Regression  
348 models can be incorporated with the  $K_d$  concept to reflect the spatial variability of soil  
349  $K_d$ . For example, most regression models involve a multivariate linear relationship  
350 between  $\log K_d$  and the routinely measured soil properties (e.g., pH and soil organic  
351 carbon) (De Groot et al., 1998).

### 352 **Slow reaction**

353 "Slow reaction" refers to all the slow chemical processes (between labile and non-  
354 labile phases) in the solid phase, such as the intra-particle diffusion of metals in  
355 carbonates and oxyhydroxides. It is modeled as a reversible conversion between labile  
356 adsorbed metals ( $M_l$ ) and their non-labile counterparts (solid phase,  $M_n$ ) with kinetic  
357 rates ( $k_1, k_{-1}$ ).

$$\begin{aligned} \frac{dM_l}{dt} &= -k_1 M_l + k_{-1} M_n \\ \frac{dM_n}{dt} &= k_1 M_l - k_{-1} M_n \end{aligned} \quad (2)$$

359 where  $k_1$  and  $k_{-1}$  are the forward and backward rates of the slow reaction ( $d^{-1}$ ).

360 Slow reaction is included in the SWAT-HM and TOPKAPI-ETH models because it  
361 plays an important role in long-term simulations (Buekers et al., 2008; Crout et al.,



### 371 **3.2.2 Equilibrium approach versus kinetic approach**

372 The equilibrium-based ( $K_d$ ) approach is commonly used for watershed-scale water  
373 quality models in [Caruso et al. \(2008\)](#) review. Recently developed models (e.g., SWAT-  
374 HM and TOPKAPI-ETH) have examined the equilibrium assumption adopted in the  
375 previous models to propose a new scheme (i.e., kinetic approach) to enhance their  
376 applicability. Specifically, the equilibrium assumption between the solution and solid  
377 phases without considering the slow reaction (transformation between labile and non-  
378 labile pools) may not reflect reality, especially in long-term metal simulations ([Sui et  
379 al., 2022](#)). The slow reaction is described as a reversible, first-order kinetic process with  
380 kinetic constants. However, it should be mentioned that mechanism-based principles  
381 have not yet been incorporated into the existing watershed HM models, although the  
382 development of reactive transport models (RTMs) in the subsurface geochemistry  
383 community has advanced rapidly since the 1980s ([Steeffel et al., 2015](#)). The reactions  
384 in RTMs models include both kinetically controlled (e.g., microbe-mediated redox  
385 reaction, mineral dissolution, and precipitation) and equilibrium-controlled ones (e.g.,  
386 ion exchange, surface complexation (sorption) and aqueous complexation) ([Li, 2019](#)).  
387 A few attempts (e.g., [Bao et al., 2017](#)) have been made to bring the RTMs from the  
388 "closed" groundwater systems into the "open" watersheds.

## 389 **3.3 Representation of watershed-scale transport processes**

### 390 **3.3.1 Spatial discretization and temporal scale**

391 A spatially distributed representation of the hydrology and contamination transport



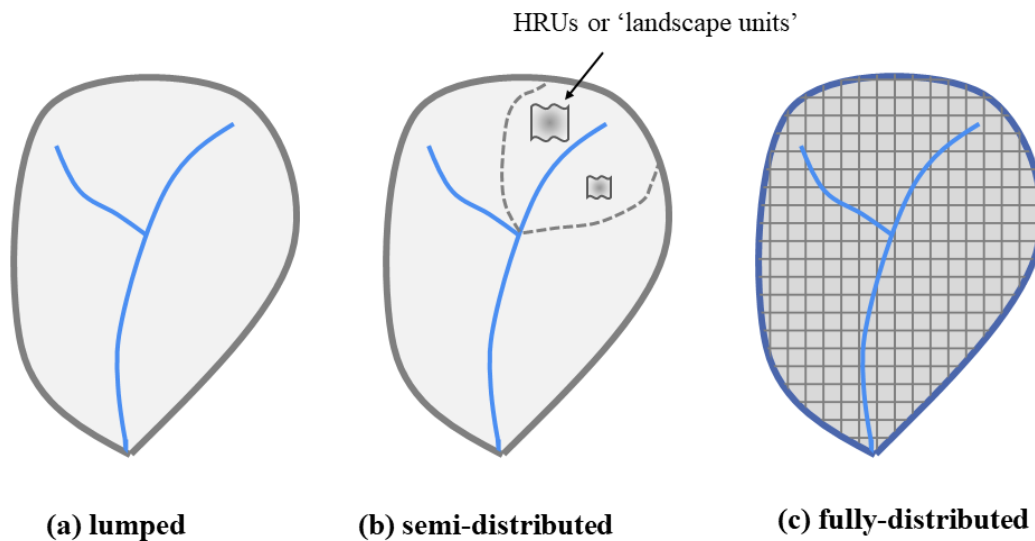
392 processes is necessary for watershed management. Thus, the watershed model  
393 simulates water flow and contaminant dynamics across discretized landscape units. The  
394 models differ in how they account for heterogeneity within each sub-basin. In  
395 watershed-scale models, spatial representation (discretization) is typically classified  
396 into three types: (1) lumped, (2) semi-distributed, and (3) fully-distributed models (Fig.  
397 4). The lumped spatial approach does not discretize the sub-basins and represents them  
398 using average lumped parameters to represent the physical processes within each sub-  
399 basin. The WARMF model follows this lumped approach (Chen et al., 2001). Semi-  
400 distributed approaches are based on properties of land use, soil type, and topography,  
401 such as slope. Examples of such models are the SWAT model, which uses Hydrologic  
402 Response Units (HRUs) (Arnold et al., 2010), and the 'landscape units' in INCA-Metals  
403 (Whitehead et al., 2009). However, as the semi-distributed model, SWAT-HM fails to  
404 show the interaction between the HRUs, as they are not internally linked within the sub-  
405 basin but are routed individually to the basin outlet. The fully-distributed approach  
406 divides the watershed and sub-basins into hydraulically connected elements, such as  
407 grid cells, to substantiate cell-to-cell transport. Each cell has unique properties, such as  
408 slope, land cover, and soil, in a fully-distributed model. Such an approach is  
409 implemented in TREX (Velleux et al., 2008) and CTT&F (Johnson et al., 2011). A  
410 principal advantage of a fully-distributed watershed model is the opportunity to identify  
411 the critical source areas (Aim 3) within the watershed and sub-basin, such as waste piles  
412 that contribute the most to HM transport. However, owing to the computational burden  
413 and high demand for data, applying a fully-distributed model is limited to the small and

414 mesoscale scale. The decreased spatial resolution of the semi-distributed model allows  
415 for a coarser calculation time step (e.g., daily), thus effectively reducing computational  
416 resources. The decreased spatial resolution of semi-distributed models allows for a  
417 coarser calculation time step (e.g., daily), thus effectively reducing computational  
418 resources. Semi-distributed models are still widely used for sub-basins prioritization  
419 when accurate location is not demanding. For instance, [Chen et al. \(2014\)](#) developed a  
420 framework integrating a watershed model with the Markov chain theory to pinpoint  
421 priority sub-basins.

422 For temporal scale, watershed-scale models can be divided into two categories:  
423 event and continuous models. Event models simulate watershed responses to a single  
424 rainfall event with a fine time resolution, like an hour or minutes, and are thus suit short-  
425 term simulating needs. In contrast, continuous models simulate the inter-rainfall  
426 environmental processes in the watershed as well as the rainfall events *alone*; thus, they  
427 usually simulate with daily timestep and suit long-term simulating needs. Among the 9  
428 selected models, TREX and TOPKAPI-ETH have the single-event simulation  
429 capability. Other models, such as INCA-Metals and SWAT-HM, are able to analyze the  
430 long-term effects of hydrological changes and water management practices. The daily  
431 time step is likely one of the best temporal resolutions for the ease of computation and  
432 availability of datasets while maintaining the capability to manifest temporal variation.  
433 The daily time resolution is probably adequate for larger creeks or rivers that are not  
434 "flashy" (i.e., the hydrograph peaks and falls back to normal flow quickly within 24  
435 hours). However, the daily time step may not be adequate to determine pollutant loads

436 from flashy systems. Concerning toxicity modeling (Aim 2, , risk assessment), if acute  
437 toxicity is of primary concern and the metal concentration dynamic fluctuates on an  
438 hour- or minute-scale, event models become necessary to perform reliable risk  
439 assessments. On the other hand, if chronic toxicity is an issue that does not appear until  
440 a longer exposure, such as several days, weeks, or even months, the daily time step is  
441 likely adequate. It should also be noted that although most applications of SWAT have  
442 been on a daily time step, recent modifications make the sub-daily calculation  
443 operational (Brighenti et al., 2019). These modifications include adding Green and  
444 Ampt infiltration equations using rainfall input at any time increment and channel  
445 routing at an hourly time step.

446



447

448 Fig. 4 Spatial discretization of the watershed model: (a) lumped, (b) semi-distributed,  
449 and (c) fully-distributed. HRUs are hydrological response units.

### 450 3.3.2 Overland hydrological processes

451 Modeling surface and subsurface hydrological processes is a prerequisite for  
452 process-based metal transport models because runoff drives soil HM deliveries into  
453 streams. Hydrological submodels in existing watershed-scale metal models can be  
454 divided into physically-based and conceptual. Every hydrological model requires two  
455 essential components: runoff generation and runoff routing. TREX and TOPKAPI are  
456 typical physically-based models that use the Green-Ampt equation to simulate  
457 infiltration and surface runoff generation. In contrast, the SCS-CN (curve number)  
458 method is widely used in models such as SWAT-HM and L-THIA to compute surface  
459 runoff. TREX and TOPKAPI route runoff using mass conservation-based continuity  
460 equations. The former uses diffusive wave approximation to simulate two-dimensional  
461 overland flow (i.e., surface runoff), while the latter uses kinematic wave approximation.  
462 They are both simpler forms of the St. Venant equations, also called dynamic wave  
463 equations.

464 In contrast, INCA-metals use storage-based (nonlinear reservoir) equations for  
465 flow routing. Subsurface hydrological processes (e.g., interflow, groundwater flow) are  
466 the most variable hydrologic components among the watershed models. Existing  
467 models use different approaches, ranging from ignoring all forms of subsurface fluxes  
468 (e.g., L-THIA) to empirical methods (e.g., INCA-Metals) to physically based equations  
469 (e.g., TOPKAPI-ETH). It should also be mentioned that all the reviewed models either  
470 lack groundwater modules or adopt simple ones. Regarding HMs, the convection–

471 diffusion–reaction equation (the conservation of solute mass) is the basic equation to  
472 simulate the movement of HMs driven by the hydrological flow and biogeochemical  
473 reactions (Steeffel et al., 2005). Some simplifications are made in the reviewed HM  
474 models. For example, the HM transport in grid-based TOPKAPI-ETH is approximated  
475 as an advection process, neglecting the diffusion process. For semi-distributed models,  
476 the HRU or 'landscape unit' is used as the basic calculation unit to calculate the mass  
477 balance equations assuming spatial uniformity.

### 478 3.3.3 Soil erosion and sediment transport

479 As a major part of soil HM is tightly adsorbed to mineral particles, bound with  
480 organic matter, or present in parent minerals, soil erosion is the primary pathway of soil  
481 HMs to water bodies and has been studied at various scales (plot, watershed, and  
482 national scales) under natural or simulated conditions (Huang et al., 2019a; Quinton  
483 and Catt, 2007; Zheng et al., 2016). More recent studies have highlighted the important  
484 role of the water erosion pathway in the movement of HM from soils to surface waters  
485 (Liu et al., 2019). Moreover, erosion is a highly selective process that enriches the  
486 detached material with small-sized silt, clay, and organic carbon. The enrichment ratio  
487 (ER) of HM is the key variable representing the mechanisms and is defined as

$$488 \quad ER = \frac{C_i}{C_0} \quad (3)$$

489 where  $C_0$  denotes the concentration in the original soil, and  $C_i$  represents the mean  
490 concentration in the eroded sediment.

491 Several soil erosion and sediment yield models are available in the existing

492 watershed-scale metal models with different process complexity and data requirements.  
493 The erosion and transport of particle HMs are directly coupled to corresponding fluxes  
494 of sediment particles. Soil erosion models quantify the amount of soil removed from a  
495 defined area over a given period. In contrast, the sediment yield models compute the  
496 amount of soil transported to water bodies over a defined area in a given period.  
497 Therefore, it is critical to account for all erosion and sediment transport processes  
498 within a basin to assess the sediment yield. The Universal Soil Loss Equation (USLE)  
499 and its descendants (e.g., RUSLE) are empirical models widely used for predicting soil  
500 erosion. However, USLE series models do not consider sediment deposition or route  
501 sediment in a spatial context; thus, they cannot be directly used to predict watershed  
502 sediment yield.

503 In many cases, USLE/RUSLE is applied to simulate hillslope erosion, along with  
504 sediment delivery ratios (SDRs), to determine the sediment delivered from the hillslope  
505 to water bodies. For example, METALPOL employed a modified USLE with the SDR  
506 to calculate the soil loss in river basins. The SWAT-HM uses the modified USLE  
507 (MUSLE) to simulate the sediment yield at the watershed scale. MUSLE replaces the  
508 rainfall energy factor in the USLE using a runoff rate factor, fulfilling the sediment  
509 yield prediction for a single storm event by considering the runoff characteristics.  
510 Physically-based models (e.g., TREX and TOPKAPI-ETH) integrate a spatially  
511 distributed soil erosion and suspended sediment transport module. This module  
512 simultaneously accounts for 4 main sediment transport processes, advection, dispersion,  
513 erosion and deposition on a two-dimensional overland plane; the latter two processes

514 are determined by the local transport capacity, which depends on the overland flow  
515 discharge and surface slope (Battista et al., 2020; Prosser and Rustomji, 2000).  
516 Additionally, ECOMAG-HM does not contain a sediment component; therefore, it is  
517 unsuitable for erosion-prone areas.

#### 518 **3.3.4 In-stream processes**

519 Current watershed metal models employ significantly different levels of  
520 complexity in modeling in-stream processes. Most models (e.g., TREX and TOPKAPI-  
521 ETH) assume a one-dimensional transport along a stream reach for simplicity. Models  
522 such as SWAT-HM and WARMF assume a well-mixed water column for each channel.  
523 Regarding channel flow, TREX and TOPKAPI-ETH use diffusive wave approximation  
524 and kinematic wave approximation under the overland flow modeling. SWAT-HM  
525 provides the variable storage and Muskingum methods for water routing in the channel  
526 network. Physically-based models commonly consist of two sediment transport  
527 processes in rivers/streams: (1) advection and dispersion and (2) erosion and deposition.  
528 A simplified stream power equation calculates the maximum sediment load in river  
529 channels in SWAT-HM. Concerning the metal module, the CTT&F model includes a  
530 four-phase equilibrium partitioning (dissolved, precipitated, sorbed to sediment  
531 particles, and complexed with the dissolved organic carbon (DOC)). In contrast, some  
532 models (e.g., METALPOL) do not explicitly consider the in-stream processes but  
533 quantify the "apparent" retention behavior of metals in river systems by computing the  
534 difference between the measured load of metals at a specific river station/section and

535 the sum of all point and diffuse (nonpoint) sources of metal emissions, which represent  
536 the inputs into a watershed.

### 537 **3.3.5 Point sources and atmospheric deposition**

538 There are generally two methods for quantifying point source emissions: the  
539 measurement method and the emission factor method when the source locations are  
540 identifiable and time series of effluent discharges and metal concentrations are available.  
541 Representing the point sources in the model is relatively easy. However, detailed  
542 observations are not available in most cases. Multiplying the estimated discharge  
543 (inhabitants connected to the point source  $\times$  water volume used per person per day) by  
544 the average HM concentration could be a reliable way to estimate the daily HM loads  
545 (Liu et al., 2018). Notably, this approach cannot account for short-term fluctuations in  
546 HM concentrations and loads. Moreover, detailed locations and point source emissions  
547 are unavailable in some large-scale applications. Therefore, a bottom-up approach  
548 combining the activity levels of various industry sectors with HM-specific emission  
549 factors have been preferred to evaluate the aqueous emissions of HMs (Huang et al.,  
550 2019b; Wu et al., 2018). In compliance with the spatial resolution of model watershed  
551 models, the technical implementation of point source modeling differs in semi-  
552 distributed model from that in fully distributed models. Semi-distributed models add  
553 point source loads to the inlets of the reaches receiving the discharge for the balance  
554 calculation, whereas fully-distributed models have the sink/source term of the river  
555 solute from the external point source appended to the transport continuity equation in a



556 river channel reach.

557       Atmosphere deposition is typically considered as the model input for watershed-  
558 scale water quality models. That is to say, most watershed models do not simulate  
559 atmospheric deposition processes, but instead use the measured fluxes of atmospheric  
560 deposition or simulated fluxes derived from air quality models. For example, modeling  
561 studies have mapped the global atmospheric concentrations and regional atmospheric  
562 deposition of priority heavy metals like Hg, Cd, and Pb under the United Nations'  
563 Convention on Long-Range Transboundary Air Pollution ([Ilyin et al., 2022](#)). Mosses  
564 have been used successfully as biomonitors to map the spatial patterns of HM  
565 deposition across Europe ([e.g., Harmens et al., 2010](#)).

## 566 **4 Current challenges and research needs**

### 567 **4.1 Improving the model representation of real-world processes**

#### 568 **4.1.1 Integrating with a carbon cycle module**

569       The terrestrial and aquatic carbon cycles play a critical role in the biogeochemical  
570 cycling of HMs in natural environments ([Warren and Haack, 2001](#)). For example, the  
571 decomposition of soil organic matter (SOM) and its associated processes directly  
572 influence soil carbon, oxygen, and nitrogen cycles, and indirectly regulate soil HM  
573 cycles by modulating Eh and pH. SOM decomposition reactions have long been  
574 recognized as complex and mostly microbe-mediated. Among the development and  
575 applications of 9 reviewed models, only a few works has explicitly considered the  
576 dynamic linking between SOM and HM ([Du et al., 2019](#)). In soil solutions, Metals such

577 as Cu and Pb may mostly be present as metal complexes, while Cd and Zn are usually  
578 present as free ions or labile complexes (Nolan et al., 2003). From the perspective of  
579 transport modeling, the total solution concentration, and not the metal speciation (e.g.,  
580 free-ion and metal complex concentrations) in the solution, must be known. When the  
581 objective is to predict metal risk and toxicity (Aim 2, risk assessment), it is crucial to  
582 identify which metal species are taken up since bioavailability is influenced by the  
583 activity of free metal ions and "labile" metal complexes pool (Parker et al., 2001).  
584 However, DOM/DOC is an often-overlooked variable in data collection and modeling  
585 despite its importance in metal risk and toxicity assessments (Caruso et al., 2008).  
586 SWAT and INCA are capable of characterizing metal complex dynamics because these  
587 model families have been developed for a range of water quality variables, including  
588 organic carbon. For example, Futter et al. (2007) presented a process-based model for  
589 simulating DOC in soil and river water called the Integrated Catchments Model for  
590 Carbon (INCA-C), and Zhang et al. (2013) developed the SWAT-C model to simulate  
591 the mass balance of soil organic carbon. In addition, recent studies have enhanced the  
592 SWAT model by introducing microbe-mediated SOM turnover processes based on  
593 Microbial Kinetics and Thermodynamics model using dual Michaelis-Menten kinetics  
594 (Bhanja et al., 2019a; 2019b). Thus, SWAT-HM and INCA-metals are the most  
595 prospective model among the 9 models due to their amenability to integrate the  
596 SOM/DOM-HM interaction by taking the advantage of the existing carbon modules.

#### 597 4.1.2 Reflecting on the role of the hyporheic zone

598 The hyporheic zone is defined as the portion of a streambed that contains a mixture  
599 of surface water and groundwater (Fig. 2). All of the nine models mentioned above  
600 ignore the metal attenuation (sink) and release (source) processes in the hyporheic zone.  
601 Thus, the metal attenuation and release processes in the hyporheic zone should be  
602 incorporated to understand the transport mechanisms in the river (Aim 1, improving  
603 understanding). Biogeochemical processes (both chemical and microbial) occurring  
604 within the hyporheic zone can significantly influence the fate and transport of HMs  
605 (Boano et al., 2014; Gandy et al., 2007). For example, the Mn oxide formation in the  
606 hyporheic zone could uptake the HMs and decrease the metal loads (Fuller and Harvey,  
607 2000). In contrast, microorganisms' respiration may deplete the hyporheic zone's  
608 dissolved oxygen (Bourg and Bertin, 1993). The induced redox change could promote  
609 the dissolution of iron and manganese oxides and the adsorbed metals, causing their  
610 release into the solution and making the hyporheic zone a metal source (Coynel et al.,  
611 2007). Hyporheic zones are closely dependent on riverbed morphology and hence vary  
612 with higher spatial resolution. Meanwhile, the river morphology could affect the  
613 physicochemical properties of river sediment and floodplain soil and thus contribute to  
614 the further redistribution of HMs (Wei et al., 2022). Integrating hyporheic zones into  
615 the watershed model may be cumbersome for semi-distributed models because of their  
616 predefined flow direction and usually an inadequate resolution of the reach channel. On  
617 the other hand, fully-distributed models with grid cell discretization have better

618 potential to model hyporheic exchange in detail. Integrating the transient storage reach-  
619 scale models (e.g., OTIS) within the fully-distributed watershed framework could be a  
620 solution. However, it is still challenging because of the high data requirements to reflect  
621 the remarkable heterogeneity of hyporheic exchange across the river network.

#### 622 **4.1.3 Refining the solid/liquid distribution ( $K_d$ ) in rivers**

623 In rivers, the fate of HMs depends on their solid/liquid distribution between water  
624 and suspended solid (SS) phases and the behavior of these two phases according to  
625 hydro-sedimentary processes. The solid/liquid fractionation is usually modeled with the  
626 fixed " $K_d$ " using the recommended screening values or finite field measurements.  
627 However, the  $K_d$  variability is in the order of magnitudes depending on several  
628 environmental factors, such as SS, DOM/DOC content, and pH (Lu and Allen, 2006).  
629 None of the existing HM models has the capability to reflect the dynamic changes in  
630  $K_d$  values under different hydrological conditions. Recently, Tomczak et al. (2019)  
631 compiled a database containing 8564  $K_d$  values from 50 elements with their potential  
632 explanatory factors and determined the  $K_d$  distributions for *in situ* water/SS conditions  
633 as a function of SS, DOC, and pH. For example, assuming a log-normal distribution,  
634 the changes of geometric mean ( $K_d$ ) and geometric standard deviation ( $K_d$ ) were  
635 identified as power laws of  $m/V$  (ratio of solid mass to water volume) for Cd, Cu, Hg  
636 and Pb (Fig. 4 in Tomczak et al. 2019). Adding such relations to existing models could  
637 reduce the global variability of  $K_d$  values.

## 638 **4.2 Gearing toward mitigation practices and climate change**

639 Measuring pollution loads from all pathways within a watershed and evaluating the  
640 effectiveness of mitigation practices through actual implementation in the field is time-  
641 consuming and resource intensive. Numerical models can be valuable tools for  
642 developing targeted remediation strategies and assessing the impact of climate change  
643 on water quality. However, only a few watershed-scale HM models have considered  
644 mitigation practices. For example, [Whitehead et al. \(2009\)](#) investigated a range of  
645 cleanup scenarios considering point source reductions to evaluate the effects of mine  
646 restoration on water quality. Nonpoint source control measures should be evaluated in  
647 future watershed management plans because numerous studies suggest they play an  
648 important role in metal transport at the watershed scale ([Liu et al., 2019](#); [Zhou et al.,](#)  
649 [2023](#)). For example, soil conservation measures (e.g., terracing, contour farming, and  
650 strip cropping) are common management operations for soil pollution control.  
651 Additionally, the extensive use of nitrogen fertilizers has resulted in significant soil  
652 acidification in several areas of China over the last three decades. This has contributed  
653 to increased metal availability and metal loss in soil ([Guo et al., 2010](#)). Field  
654 experiments have shown that liming can effectively reduce Cd accumulation in rice  
655 grains, as it increases soil pH and reduces Cd's bioavailability (CaCl<sub>2</sub>-extractable) in  
656 the rhizosphere ([Chen et al., 2018](#)). Therefore, further investigations are required to  
657 represent mitigation practices in watershed-scale HM models for effective management  
658 decisions.

659 Climate change is modifying metal deliveries from soil to water due to changes in  
660 hydrological processes (Byrne et al., 2012). For example, in the Snake River watershed  
661 in the Rocky Mountains, USA, intensified generation of Acid Mine Drainage has been  
662 linked to warmer summer air temperatures and earlier drying of shallow soils, which  
663 expands weathering fronts and promotes oxidation (Rue and McKnight, 2021). In UK  
664 watersheds where low river flows are expected to occur more frequently and severely  
665 due to climate change, metal-rich groundwater may significantly influence stream  
666 metal concentrations (Byrne et al., 2020). Thus, assessing metal transport dynamics and  
667 responses to climate change scenarios is essential to develop effective watershed  
668 management strategies.

### 669 **4.3 Reforming model calibration and uncertainty analysis**

670 In the watershed-scale models, except for the physical parameters, some  
671 parameters are difficult to measure directly; therefore, they must be estimated from the  
672 collected data through a calibration procedure (Herrera et al., 2022). Calibration can be  
673 conducted manually by trial and error or automatically through a computer-based  
674 procedure or a combination of the two. Currently, the general sequence for calibrating  
675 a watershed-scale HM model is as follows: 1) calibration of hydrological parameters,  
676 focusing on the proportion of surface runoff and baseflow; 2) calibration of sediment  
677 parameters, focusing on the ratio of upland and channel erosions; and 3) calibration of  
678 metal parameters, focusing on the land-to-river metal fluxes and riverine metal  
679 concentrations and loads (Meng et al., 2018; Sui et al., 2022).

680        Although numerous algorithms and tools exist for parameter calibration, the non-  
681 uniqueness feature (equifinality) in calibration makes the estimation of a unique or best  
682 parameter set meaningless (Abbaspour, 2022; Beven, 2006). Given the various types  
683 and sources of uncertainty, no calibration problem in the hydrology and water quality  
684 domain is uniquely solvable. In other words, if a single model (a best parameter set) fits  
685 the measurements well, many models (multiple parameter sets) exist that can achieve  
686 statistically similar model accuracy. For example, a model may exhibit satisfactory  
687 statistical agreement with measured streamflow data while misrepresenting flow  
688 pathways (e.g., surface runoff, lateral flow, and baseflow).

689        Mitigation of the non-uniqueness problem may be partially achieved by: (1)  
690 improving model representation and developing more robust models (discussed in  
691 Section 4.1), (2) implementing thorough parameter sensitivity analyses prior to  
692 calibration, (3) collecting more and better quality data (e.g., incorporating soft data)  
693 and extracting more information from the collected data, and (4) obtaining  
694 unconditional parameters through multi-variable, multi-objective calibration  
695 procedures. For example, sensitivity analysis can reduce the number of parameters for  
696 calibration by identifying the parameters with negligible influence on the output,  
697 thereby reducing the combinations of parameter sets that produce similar results. Some  
698 studies have separately calibrated surface runoff and baseflow (Zhang et al., 2011).  
699 Calibrating metal parameters using both dissolved and total metal concentrations could  
700 also improve parameter identifiability. For example, a long-term and high-frequency  
701 estimation of metal concentrations and fluxes in rivers could be obtained by analyzing

702 the relationship between sporadic sampled (dissolved/particulate/total) metal  
703 concentrations and continuously monitored proxy indicators (e.g.,  
704 discharge/SS/turbidity) (Nasrabadi et al., 2016).

705 Moreover, tracer and isotope data may assist watershed-scale model calibration  
706 because they provide information on both hydrologic pathways and specific  
707 contaminants (Jensen et al., 2018). Multi-objective (e.g., multi-criteria and multi-  
708 variable) calibration routines have been proposed to estimate model parameters using  
709 multiple types of data (Bennett et al., 2013). The use of soft data in automated  
710 calibration procedures is also recommended to constrain the model equifinality and  
711 improve the model's realism (Arnold et al., 2015). Typical soft data for metal modeling  
712 could be long-term average sediment yield, event mean concentration, and source  
713 apportionment using isotope analysis.

714 Recognizing and quantifying model uncertainties are vital for applying watershed  
715 HM models. As mentioned above, the uncertainties could be introduced by model  
716 validation data (e.g., limited data) and model structures (e.g., different representations  
717 of metal transformation). It should also be noted that model input (e.g., precipitation,  
718 DEM, digital elevation model) is another non-negligible source (Shen et al., 2012). The  
719 precision of the DEM is expected to affect the delineation of watersheds and the  
720 calculation of terrain factors (e.g., slope, slope length), which will further influence  
721 models' simulations of flow, sediment, and metal transport processes (Wechsler, 2007).  
722 Most HM model studies have been calibrated against HM concentrations in river water  
723 but occasionally report model uncertainty. Zhou et al. (2020) applied the SUFI-2



724 algorithm ([Abbaspour et al., 2007](#)) and behavioral simulations (e.g., model  
725 performance above a threshold of one/multiple objective functions) to represent the  
726 metal flux uncertainty. Thus, further research is needed to quantify uncertainty and its  
727 relative source strengths to support robust decision-making. Some existing long-term  
728 and high-frequency datasets in experimental watersheds (e.g., [Neal et al., 2013](#)) provide  
729 a valuable opportunity for model comparison, testing, and reducing model uncertainty.

#### 730 **4.4 Balancing model complexity and data requirements**

731 In Section 3, the problems of model complexity and data requirements have been  
732 raised. Well-informed modeling relies on a well-designed monitoring network.  
733 However, an enormous amount of input and calibration data is required to build and  
734 calibrate complex water quality models. Moreover, the data requirements of water  
735 quality models vary with model complexity and purpose. Thus, in addition to collecting  
736 more and better quality monitoring data, it is necessary to determine the amount of data  
737 ([Ledergerber et al., 2019](#)) and the value of the data ([Abbaspour et al., 1996](#)) to be  
738 collected. The optimal experimental design (OED) technique is a useful tool for  
739 identifying data shortages and maximizing available information behind the  
740 measurement data. For example, suppose a new dataset is to be collected; model-based  
741 OED can evaluate the informational gain from the perspective of model calibration  
742 from a bundle of proposed experiments and help reach a cost-effective experimental  
743 design ([Vanrolleghem et al., 1995](#)).

744 Regarding better quality monitoring data, existing monitoring protocols by

745 environmental regulators often collect grab samples, which cannot detect dynamic  
746 changes in HM concentrations, particularly during high-flow periods (Neal et al., 2012).  
747 Future research should incorporate the new advancements in sampling and analytical  
748 technologies which enable more representative and efficient measurement of HMs in  
749 soil and water. For instance, Frau et al. (2018) investigated the possibility of using novel  
750 electromagnetic wave sensors for the real-time and continuous monitoring of HMs in  
751 surface water. Moreover, several studies have applied the tracer injection and synoptic  
752 sampling (TISS) method to quantify the spatially detailed point and nonpoint metal  
753 fluxes to streams, which could be crucial data for multi-objective calibration in  
754 watershed-scale models (Byrne et al., 2021; Kimball et al., 2002). TISS uses  
755 conservative tracers (typically bromide or chloride) and synchronous (or synoptic)  
756 water sampling to measure the flow and pollutants concentrations. For example, the  
757 TISS method revealed the spatial profiles of dissolved, colloidal, and total Zn loads in  
758 Cement Creek, Colorado, USA (Fig. 9 in Kimball et al., 2002). Furthermore, a new  
759 paradigm called SEPSTAT has been proposed for effective heavy metal monitoring  
760 based on dry preservation: solid-phase extraction, preservation, storage, transport, and  
761 analysis of trace contaminants. It was designed to overcome the logistical challenge and  
762 could create the required large-scale data in resource-limited settings. (Hanhauser et al.,  
763 2020). In addition, hyperspectral remote sensing offers a non-destructive and real-time  
764 method for retrieving soil metal concentrations, which could facilitate the simulation of  
765 ubiquitous spatial heterogeneity in watershed-scale HM models (Nawar et al., 2020).

## 766 **5 Conclusions and recommendations**

767 This study provided an overview of the current generation of watershed-scale HM  
768 models in terms of their simplifications and assumptions, components and functions,  
769 strengths and weaknesses. The primary conclusions and recommendations, including  
770 model improvement, data acquisition, and their combined use, are as follows:

### 771 ● **Model development in compliance with the intended use**

772 The future directions of model development were identified for four different aims.  
773 For Aim 1 (improving understanding), the metal attenuation and release processes in  
774 the hyporheic zone, as well as the dynamic  $K_d$  should be incorporated to understand the  
775 transport mechanisms in the river. SOM/DOM module and sub-daily schemes should  
776 be incorporated when the objective is to predict better metal toxicity and risk (Aim 2,  
777 risk assessment). For Aim 3 (identifying critical source areas), fully-distributed models  
778 would be suitable for small-to mesoscale watersheds, while semi-distributed models  
779 with the Markov chains theory could be applied on large scale watersheds. Aim 4  
780 (scenario analysis) requires an improved representation of mitigation practices in  
781 existing models to inform effective management decisions.

### 782 ● **Data acquisition through numerical and field experiments**

783 Watershed HM studies as a whole suffer from a lack of spatiotemporal  
784 measurements. Model developers and experimental researchers should work together  
785 to ensure that our models reflect the best available systems understanding and to ensure  
786 that monitoring programs provide the maximum information models require. Both

787 virtual (e.g., OED technique) and actual (e.g., TISS method) experiments could be  
788 helpful to optimize experimental schemes and obtain better quality data.

789 ● **'Benchmark' models/codes and data sets for model application**

790 The comparison and evaluation of available metal models using real data sets need  
791 to be done for developing benchmark models/data and pre-and post-processing tools to  
792 facilitate better visualization, accessibility, and commutations. The existing  
793 experimental watersheds with long-term and high-frequency data provide a valuable  
794 opportunity. Ultimately, the goal is to establish a flexible framework that combines  
795 hydrology, sediment, and chemical sub-models at various levels of complexity to match  
796 the available data and support the corresponding model purpose.

797

798 **Data availability**

799 No data was used for the research described in the article.

800

801 **Declaration of Competing Interest**

802 The authors declare no competing financial interest.

803

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