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Electronic supplementary information

Contaminant removal by efficient separation of in-situ formed layered double hydroxide compounds from mine wastewaters

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23 **Table S1.** Ionic composition of acid (pH 4.5) mine wastewater.

Ion	mg/L	mM
Na ⁺	7.3×10 ²	3.2×10 ¹
K ⁺	3.2×10 ¹	8.2×10 ⁻¹
Mg ²⁺	3.5×10 ²	1.5×10 ¹
Ca ²⁺	3.0×10 ²	7.5
Al ³⁺	1.0×10 ¹	3.7×10 ⁻¹
Fe ³⁺	1.7×10 ¹	3.0×10 ⁻¹
Mn ²⁺	1.7×10 ¹	3.1×10 ⁻¹
Cl ⁻	1.5×10 ³	4.2×10 ¹
SO ₄ ²⁻	5.8×10 ²	6.0
Cd ²⁺	6.2×10 ⁻²	5.5×10 ⁻⁴
Ce ³⁺	1.5	1.0×10 ⁻²
Co ²⁺	3.4	5.8×10 ⁻²
Cu ²⁺	3.0×10 ¹	4.8×10 ⁻¹
La ³⁺	7.2×10 ⁻¹	5.2×10 ⁻³
Nd ³⁺	6.0×10 ⁻¹	4.2×10 ⁻³
Ni ²⁺	7.3	1.2×10 ⁻¹
Sr ²⁺	1.1	1.2×10 ⁻²
Y ³⁺	2.5×10 ⁻¹	2.8×10 ⁻³
Zn ²⁺	9.6	1.5×10 ⁻¹
CO ₃ ²⁻	2.1	3.4×10 ¹

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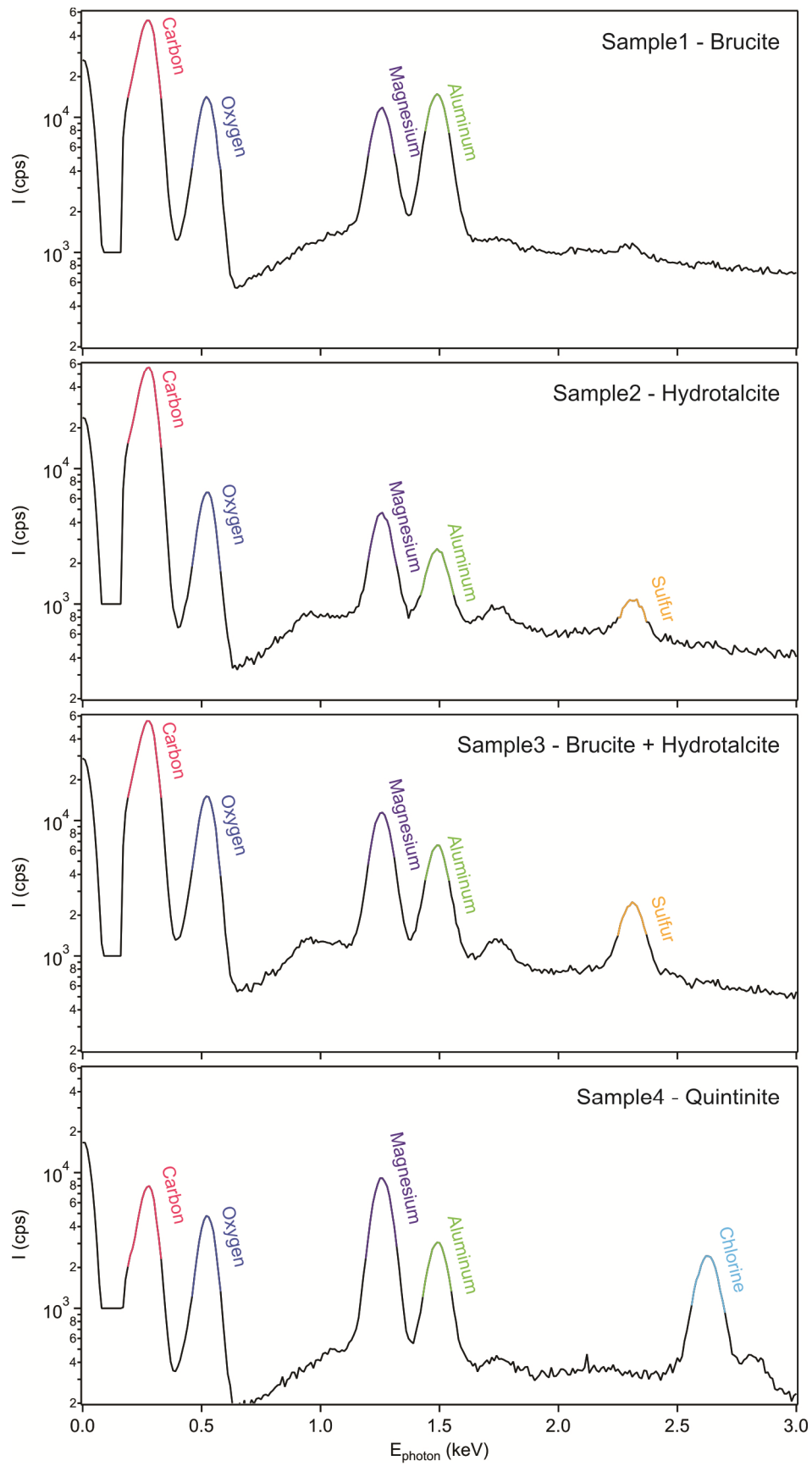
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26 **Table S2.** The concentration of PSS, DXS and phosphate at the isoelectric points.

	PSS (mg/g)	DXS (mg/g)	Phosphate (mM)
Sample 1	12	4.5	0.0084
Sample 2	0.4	6.8	0.0207
Sample 3	13.5	46.0	0.0096
Sample 4	0.9	1.1	0.0010

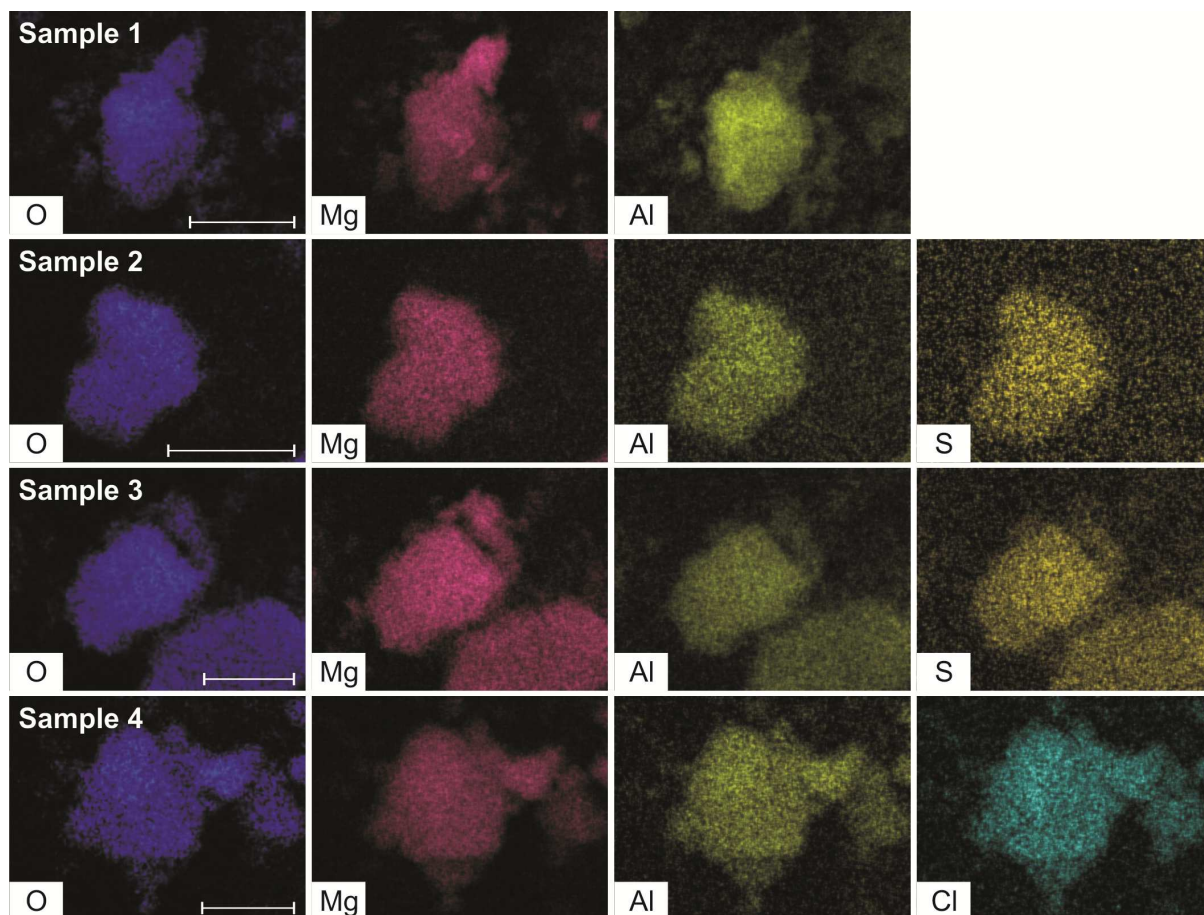
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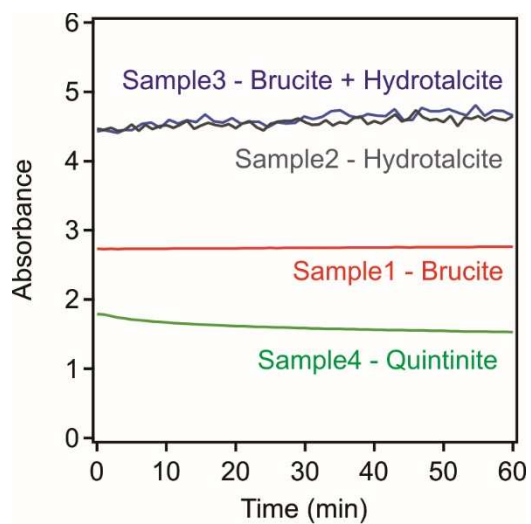
30 **Fig. S1** Elemental composition of the solid Samples 1-4. The presence of carbon is dominant
 31 due to the carbon tape sample holder.



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33 **Fig. S2** Elemental distribution images in Sample1-4 determined by SEM-EDX
 34 measurements. The scale bars refer to 3 μm distance.

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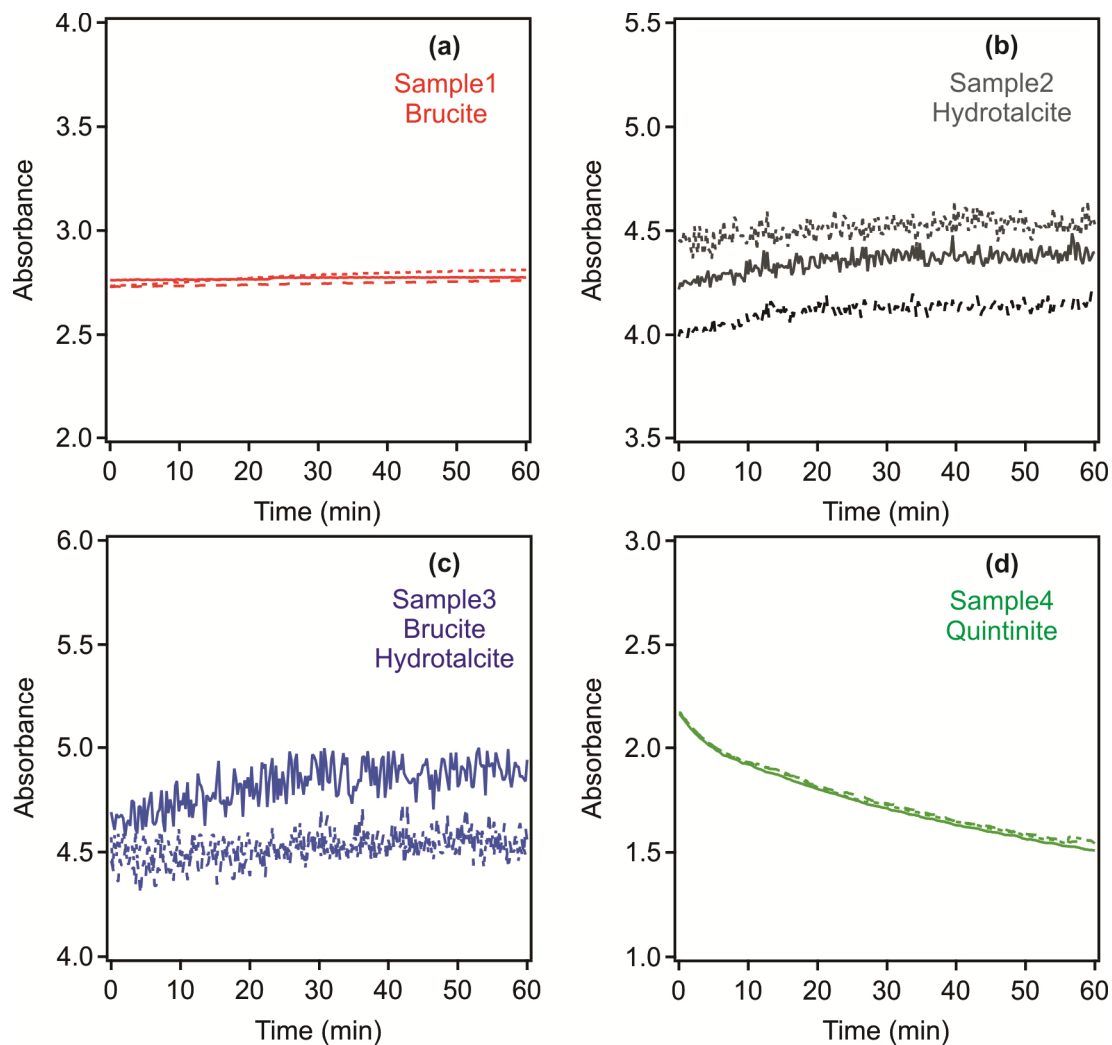


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37 **Fig. S3** Time-resolved absorbance measurements performed at 450 nm for the original

38 Samples 1-4.

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41 **Fig. S4** Time-resolved absorbance measurements performed at 450 nm for the original
 42 Samples 1-4 in the presence of PSS (solid lines), DXS (dotted lines) and phosphate (dotted
 43 line). The dose of the flocculants was set to the charge neutralization condition.

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