

Supplementary material

Effects of pH value, chloride and sulfate concentrations on galvanic corrosion between lead and copper in drinking water

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Table S1. SEM-EDX results of scales collected after 7 days in completely mixed conditions at pH 7.0

Conditions	250 mg L ⁻¹ Cl ⁻		250 mg L ⁻¹ SO ₄ ²⁻		250 mg L ⁻¹ Cl ⁻ and 250 mg L ⁻¹ SO ₄ ²⁻		
	Element	Weight %	Atomic %	Weight %	Atomic %	Weight %	Atomic %
	C	13.83	49.35	21.87	52.53	18.57	55.60
	O	12.61	33.75	21.27	38.35	13.91	31.22
	S	0.01	0.01	0.73	0.66	0.16	0.18
	Cl	1.09	1.32	0.59	0.48	1.56	1.58
	Cu	1.25	0.85	0.57	0.26	0.01	0.01
	Pb	71.20	14.72	54.7	7.62	65.79	11.42

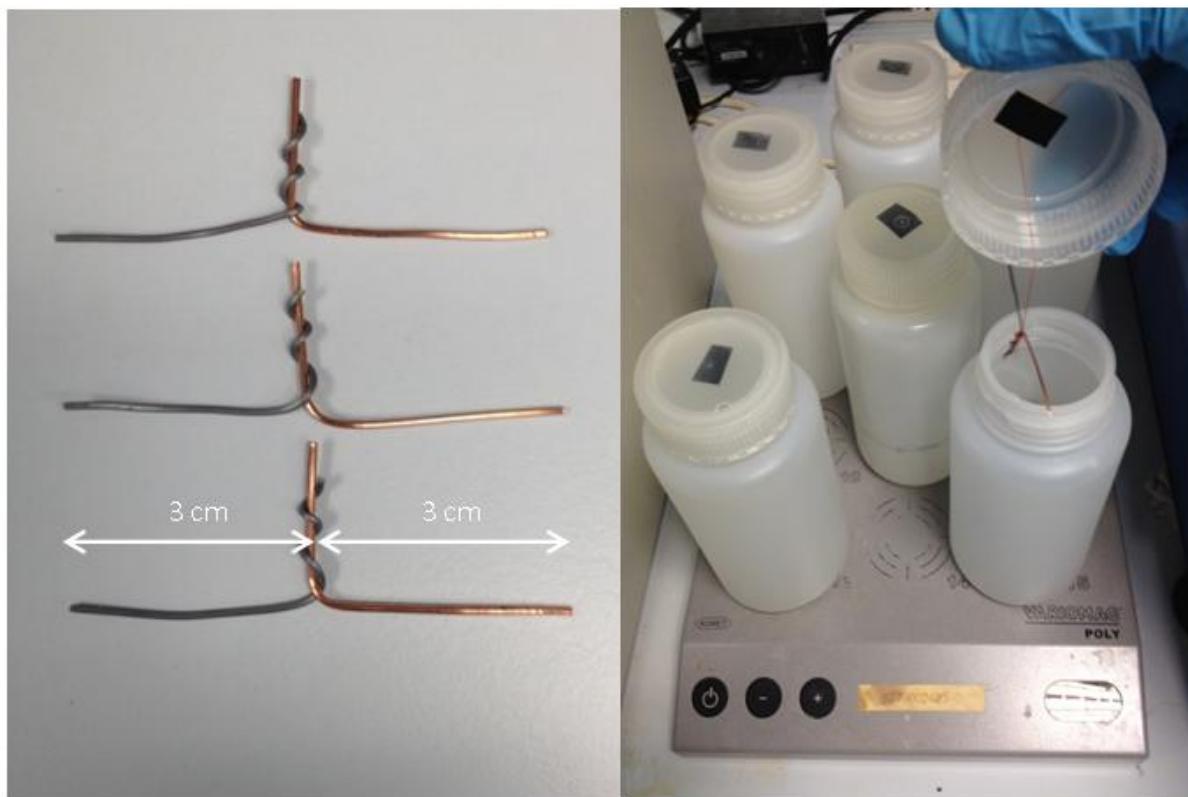


Fig. S1. Photographs of experimental setup. Lead wire tied round copper wire (left) and the wires are suspended in plastic bottle with a polyester thread (right).

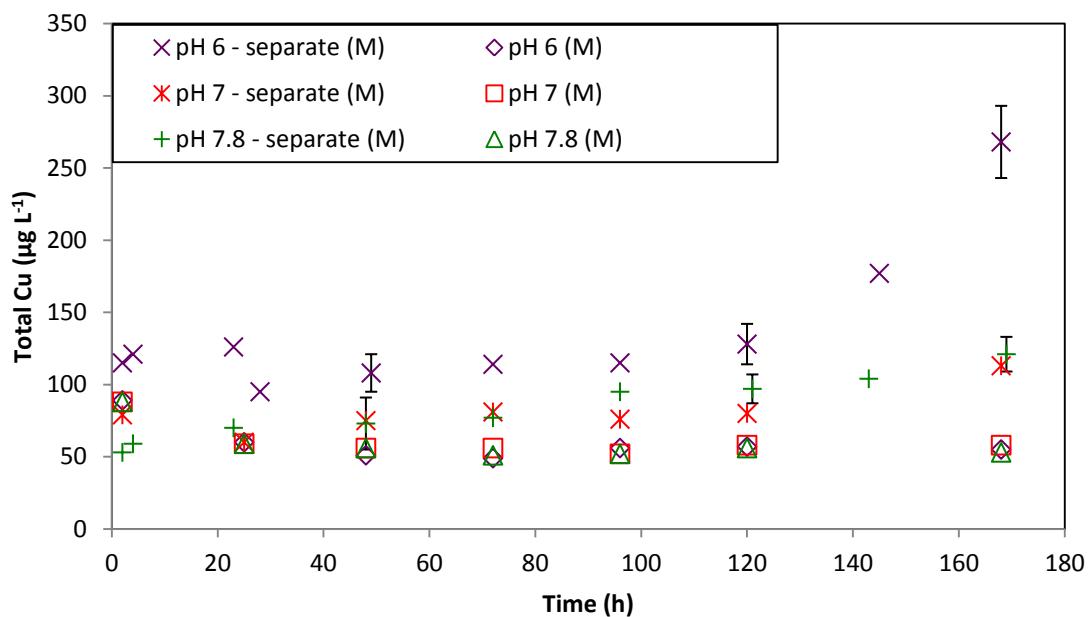


Fig. S2. Total copper as a function of time at different pH values under completely mixed conditions (M). Error bars represent standard deviation.

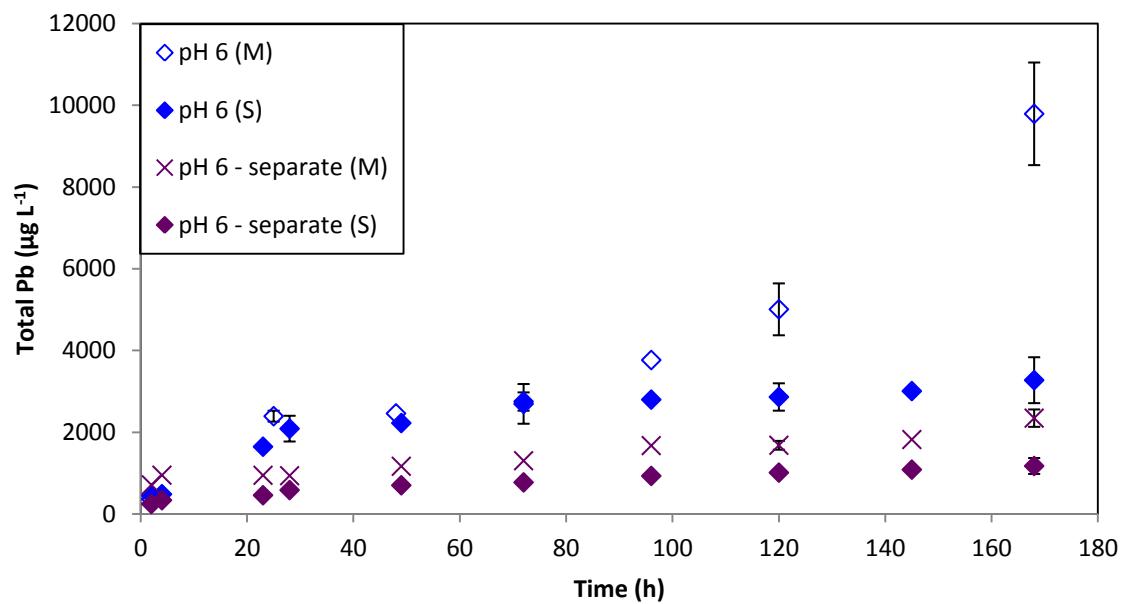


Fig. S3. Total lead as a function of time at pH 6.0. S, stagnant; M, completely mixed.

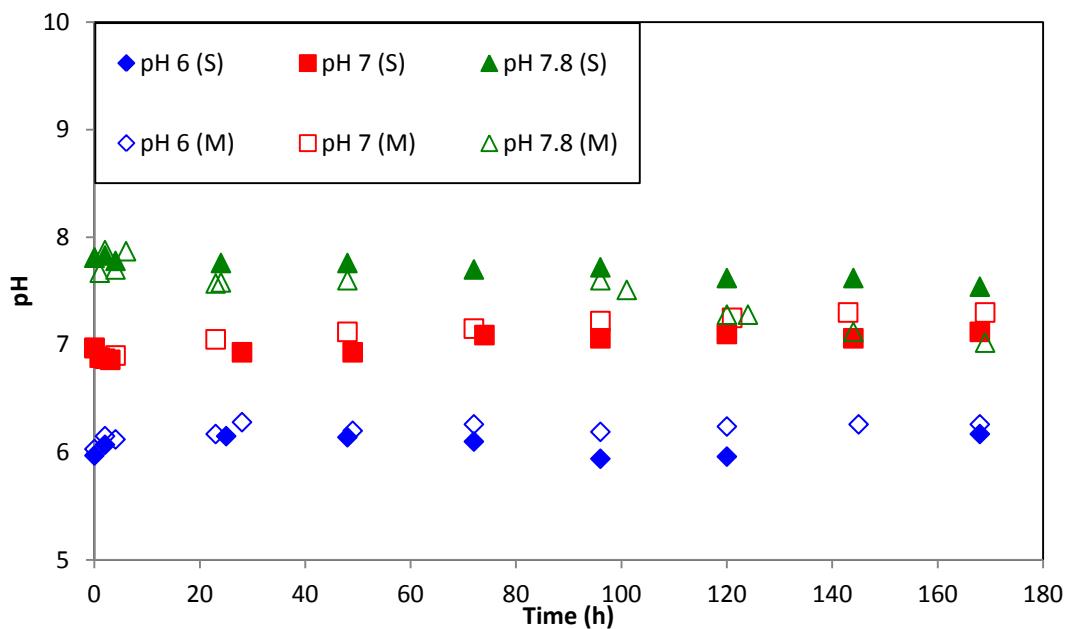


Fig. S4. Solution pH as a function of time at different solution pH. S, stagnant; M, completely mixed.

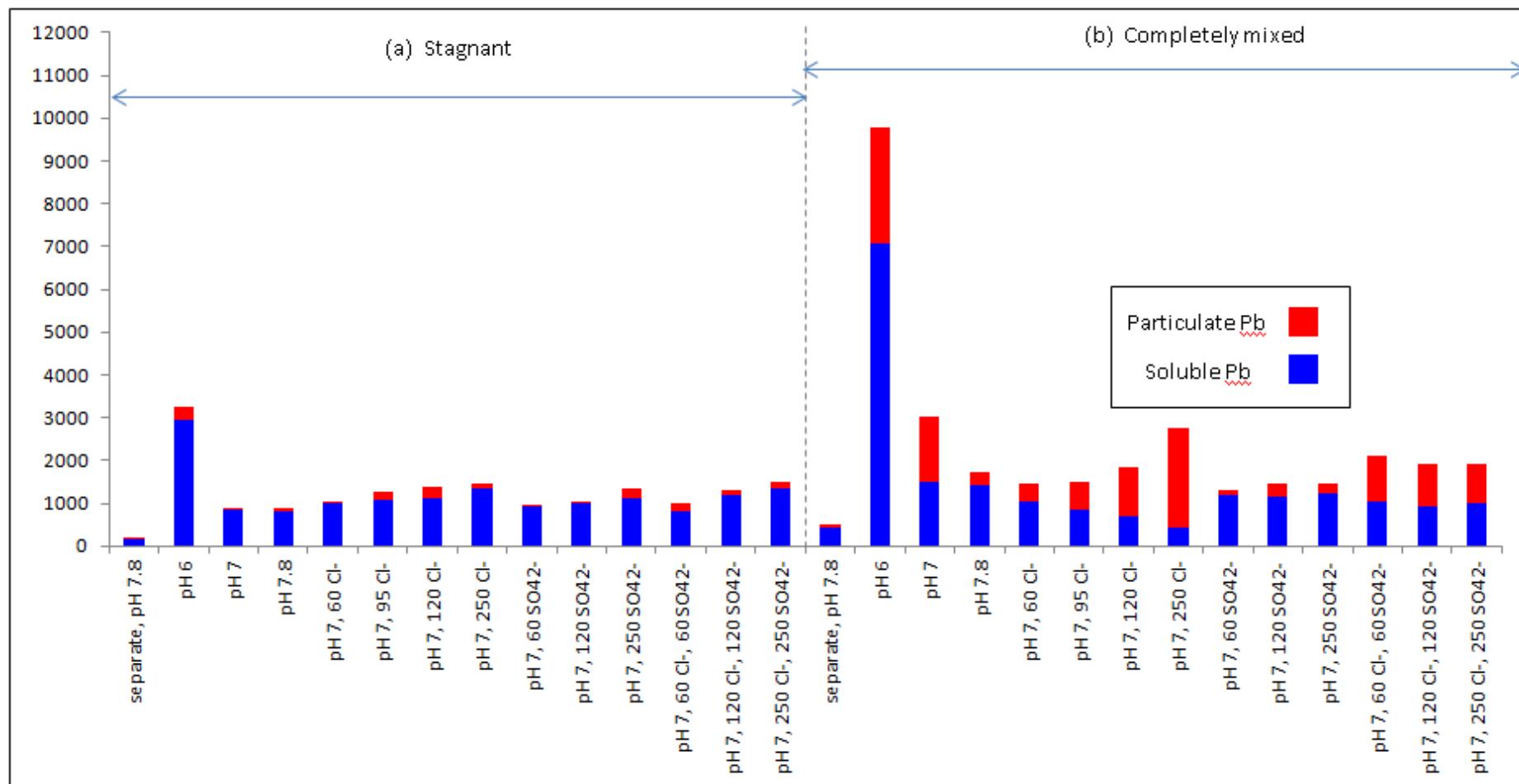


Fig. S5. Summary of total lead release under (a) stagnant and (b) completely mixed conditions after 7 days.

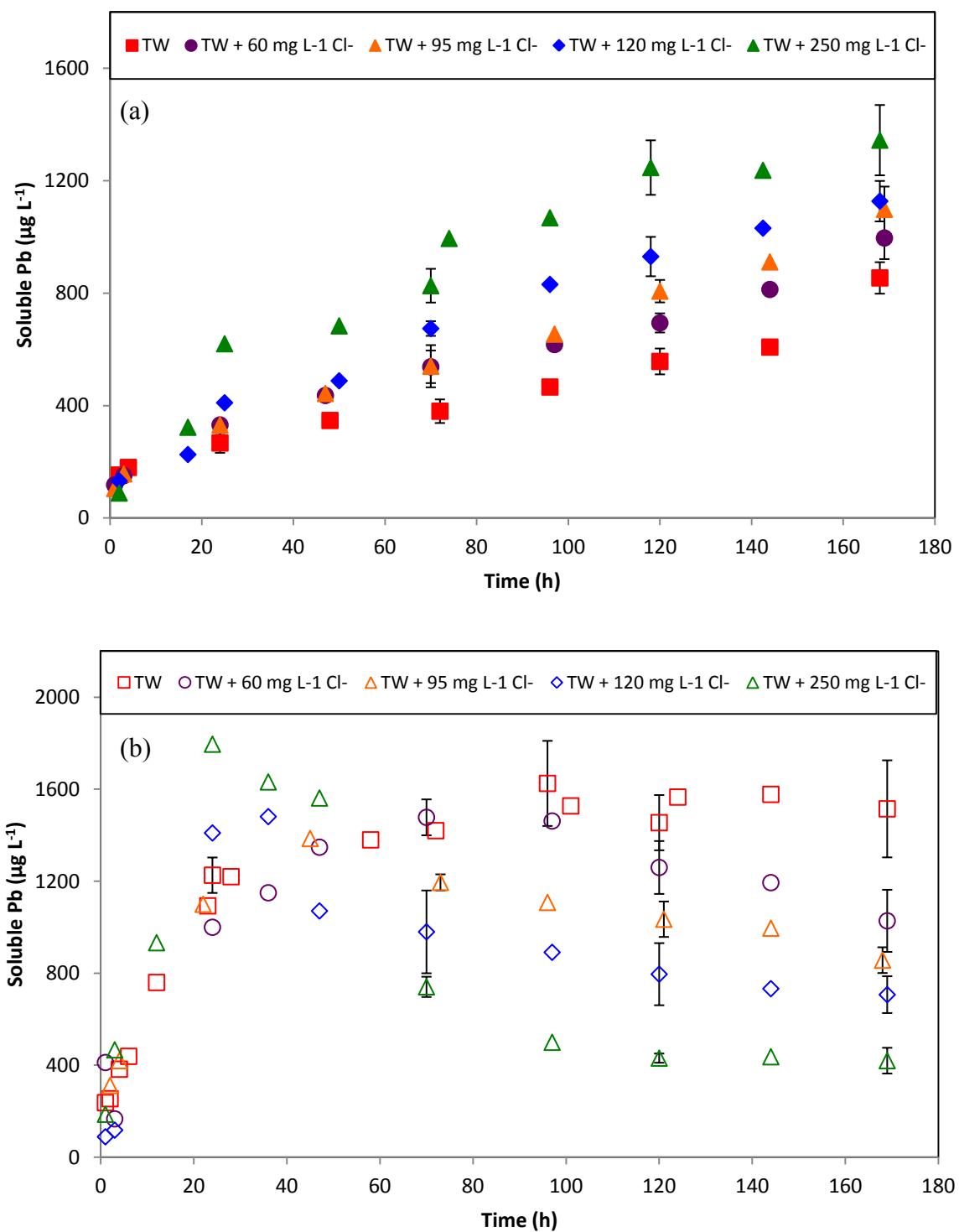


Fig. S6. Soluble lead under (a) stagnant and (b) completely mixed conditions as a function of time with different chloride dosages. TW, tap water ($\text{pH} = 7.0$, $35 \text{ mg L}^{-1} \text{Cl}^-$ and $18 \text{ mg L}^{-1} \text{SO}_4^{2-}$).

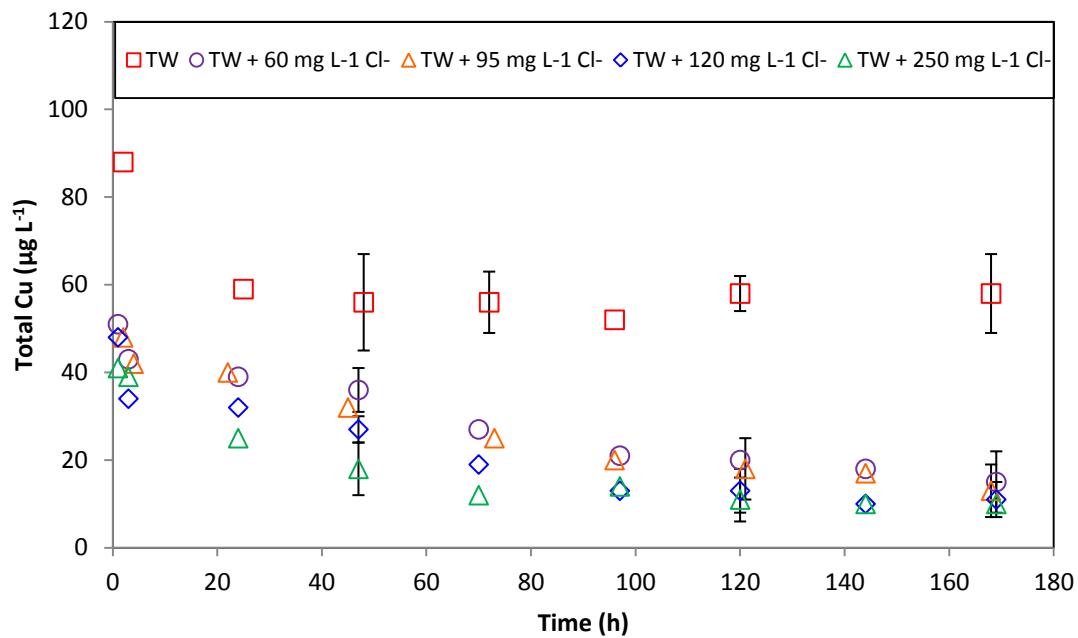


Fig. S7. Total copper as a function of time with different chloride dosages under completely mixed conditions.
TW: tap water ($\text{pH} = 7.0$, $35 \text{ mg L}^{-1} \text{Cl}^-$ and $18 \text{ mg L}^{-1} \text{SO}_4^{2-}$).

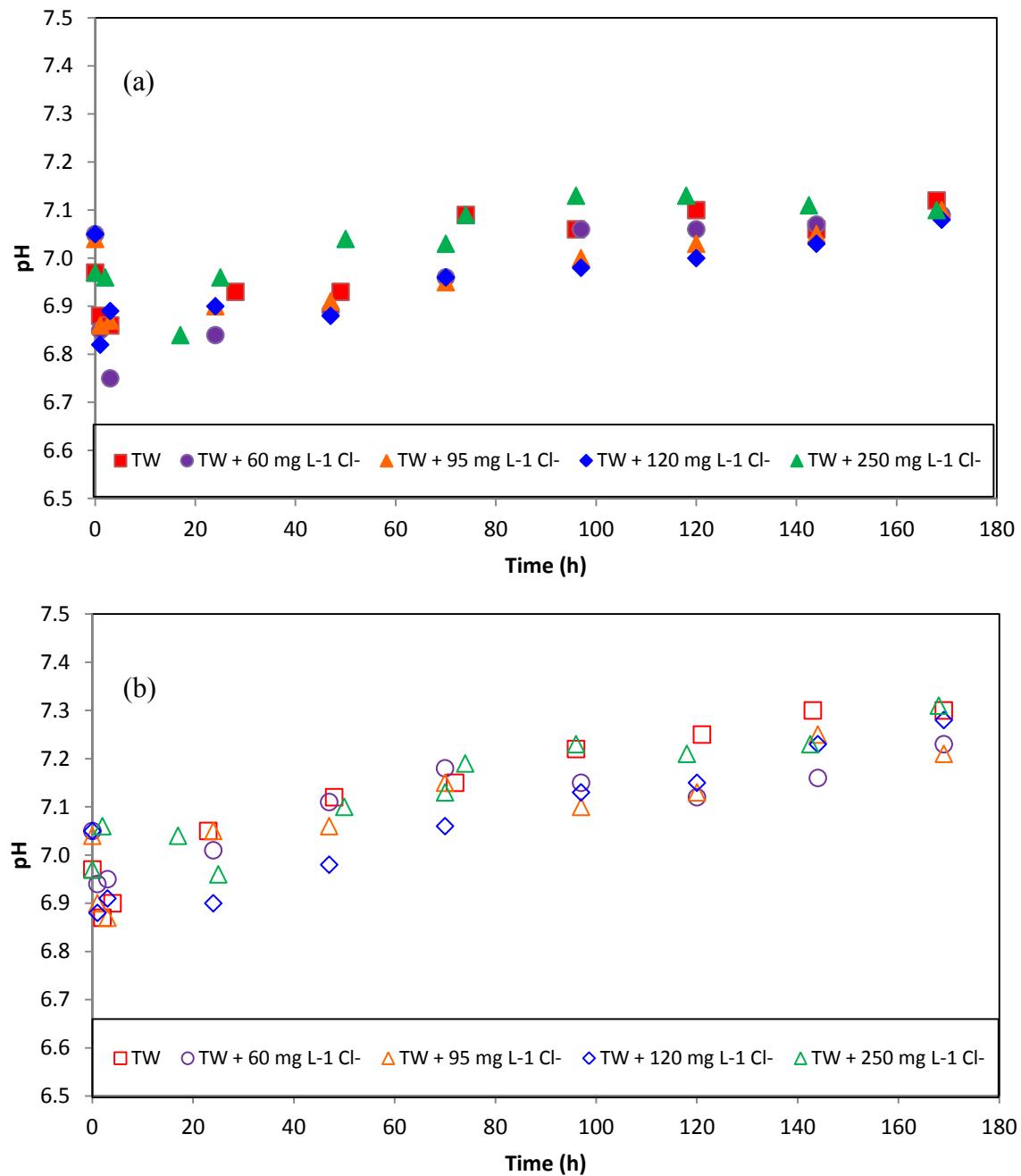


Fig. S8. Solution pH under (a) stagnant and (b) completely mixed conditions as a function of time with different chloride dosages. TW, tap water ($\text{pH} = 7.0$, $35 \text{ mg L}^{-1} \text{Cl}^-$ and $18 \text{ mg L}^{-1} \text{SO}_4^{2-}$).

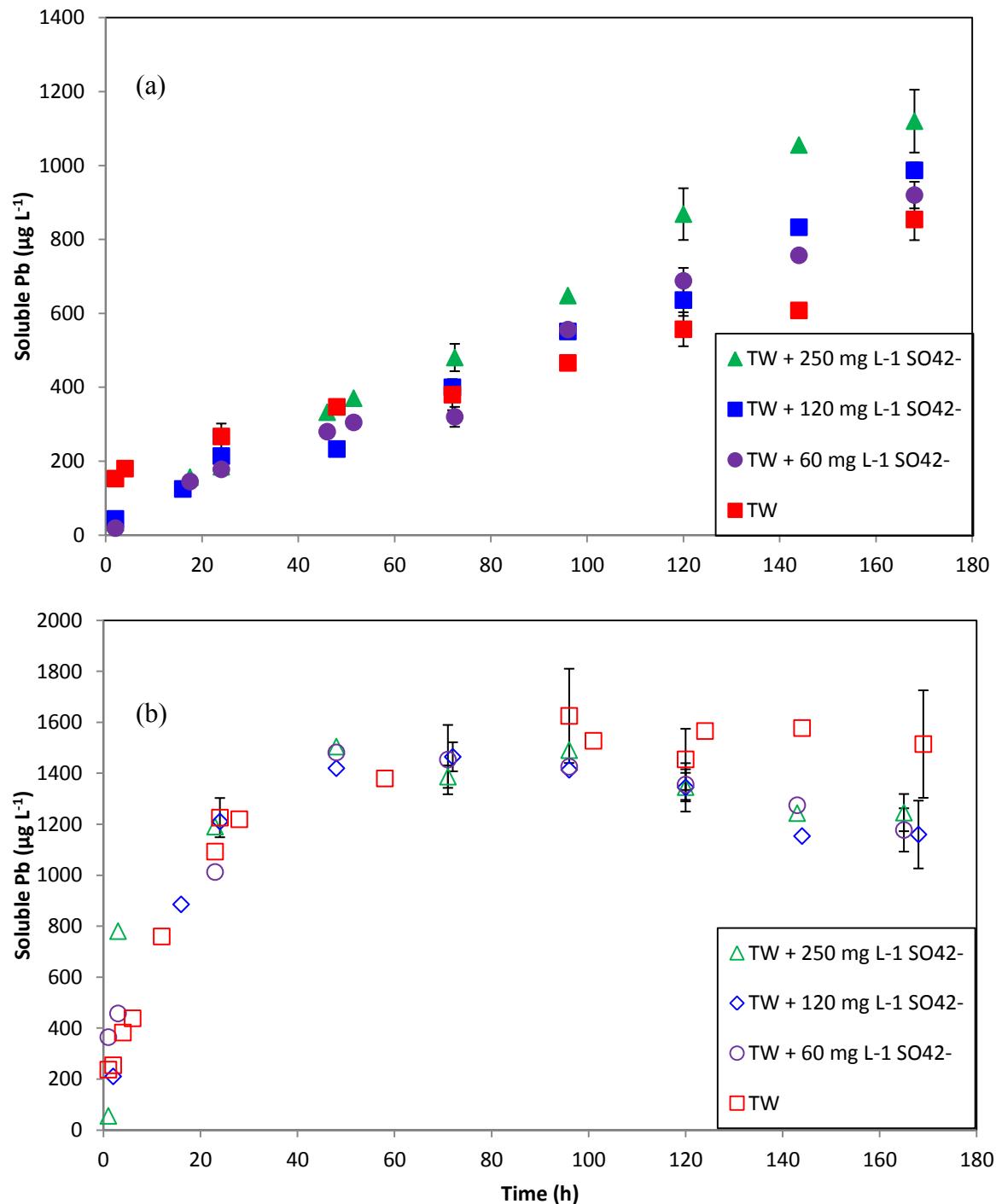


Fig. S9. Soluble lead under (a) stagnant and (b) completely mixed conditions as a function of time with different sulfate dosages. TW, tap water ($\text{pH} = 7.0$, $35 \text{ mg L}^{-1} \text{Cl}^-$ and $18 \text{ mg L}^{-1} \text{SO}_4^{2-}$).

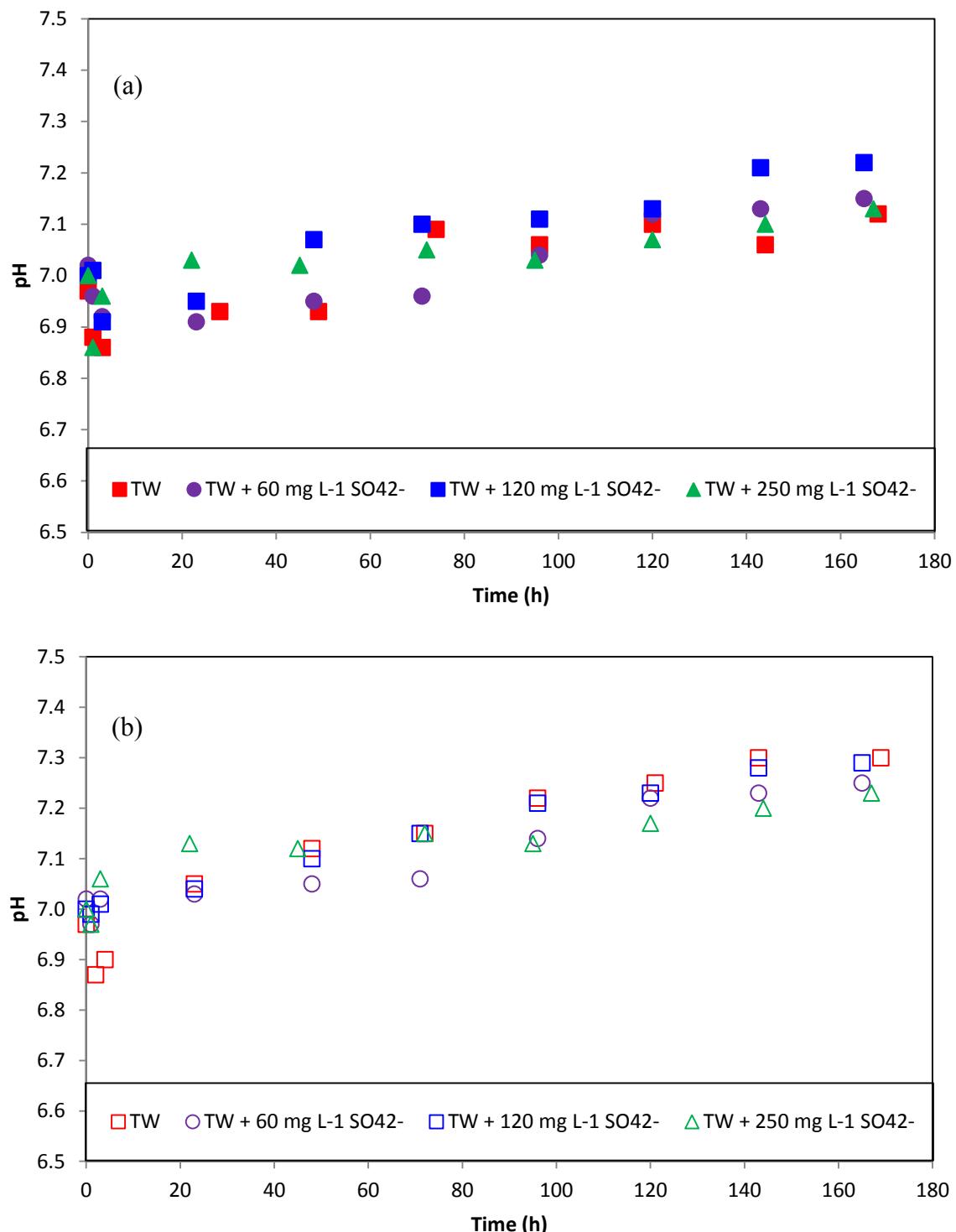


Fig. S10. Solution pH under (a) stagnant and (b) completely mixed conditions as a function of time with different sulfate dosages. TW, tap water ($\text{pH} = 7.0$, $35 \text{ mg L}^{-1} \text{ Cl}^-$ and $18 \text{ mg L}^{-1} \text{ SO}_4^{2-}$).

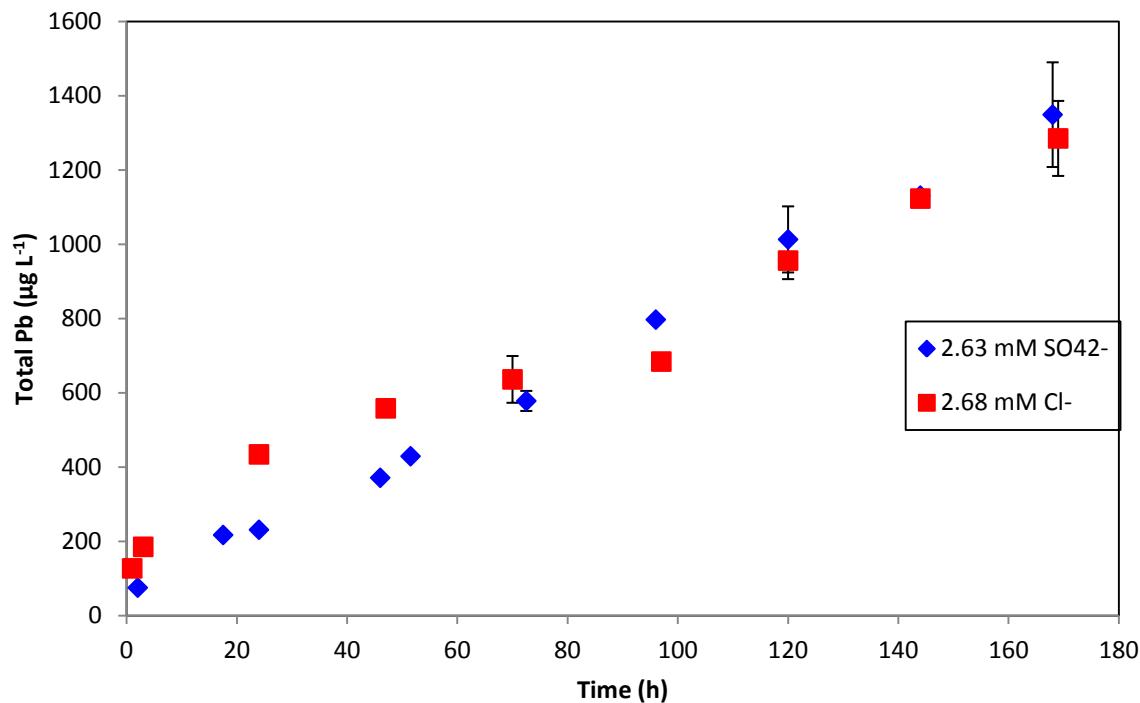


Fig. S11. Comparison between the effects of chloride and sulfate additions of about 2.65 mM on total lead concentration under stagnant conditions. Experimental conditions: pH = 7.0.

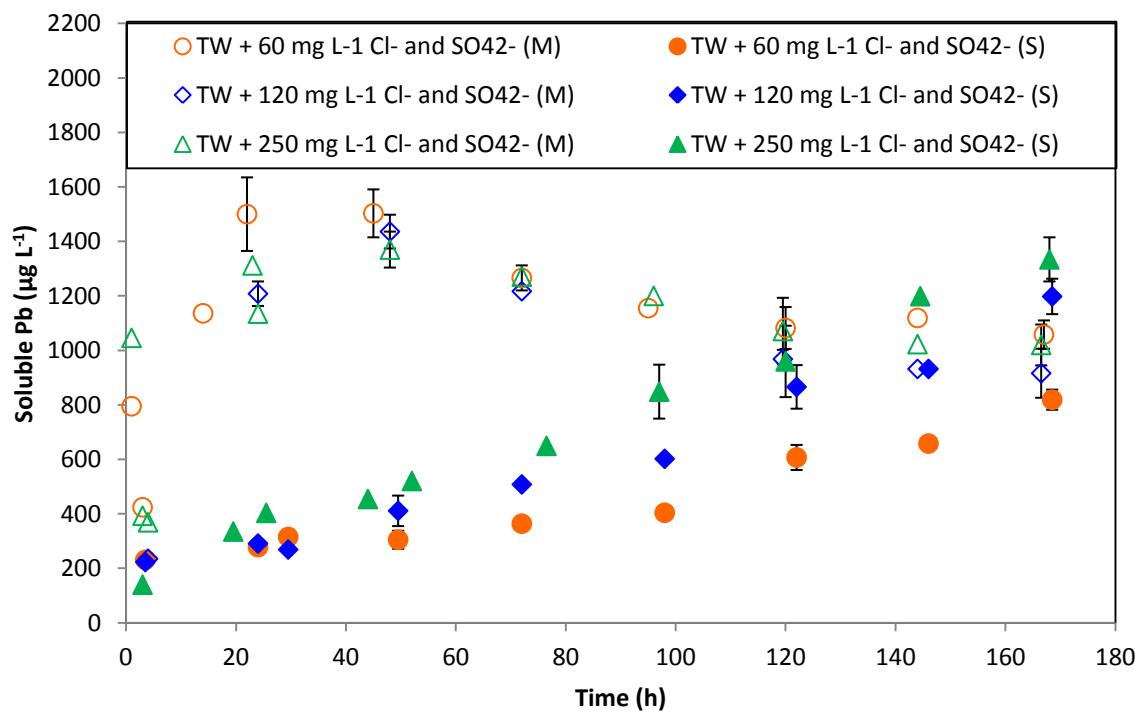


Fig. S12. Soluble lead as a function of time with different chloride and sulfate additions at a constant CSMR of 1 in stagnant and completely mixed conditions. TW, tap water (pH = 7.0, $35 \text{ mg L}^{-1} \text{Cl}^-$ and $18 \text{ mg L}^{-1} \text{SO}_4^{2-}$); S, stagnant; M, completely mixed.

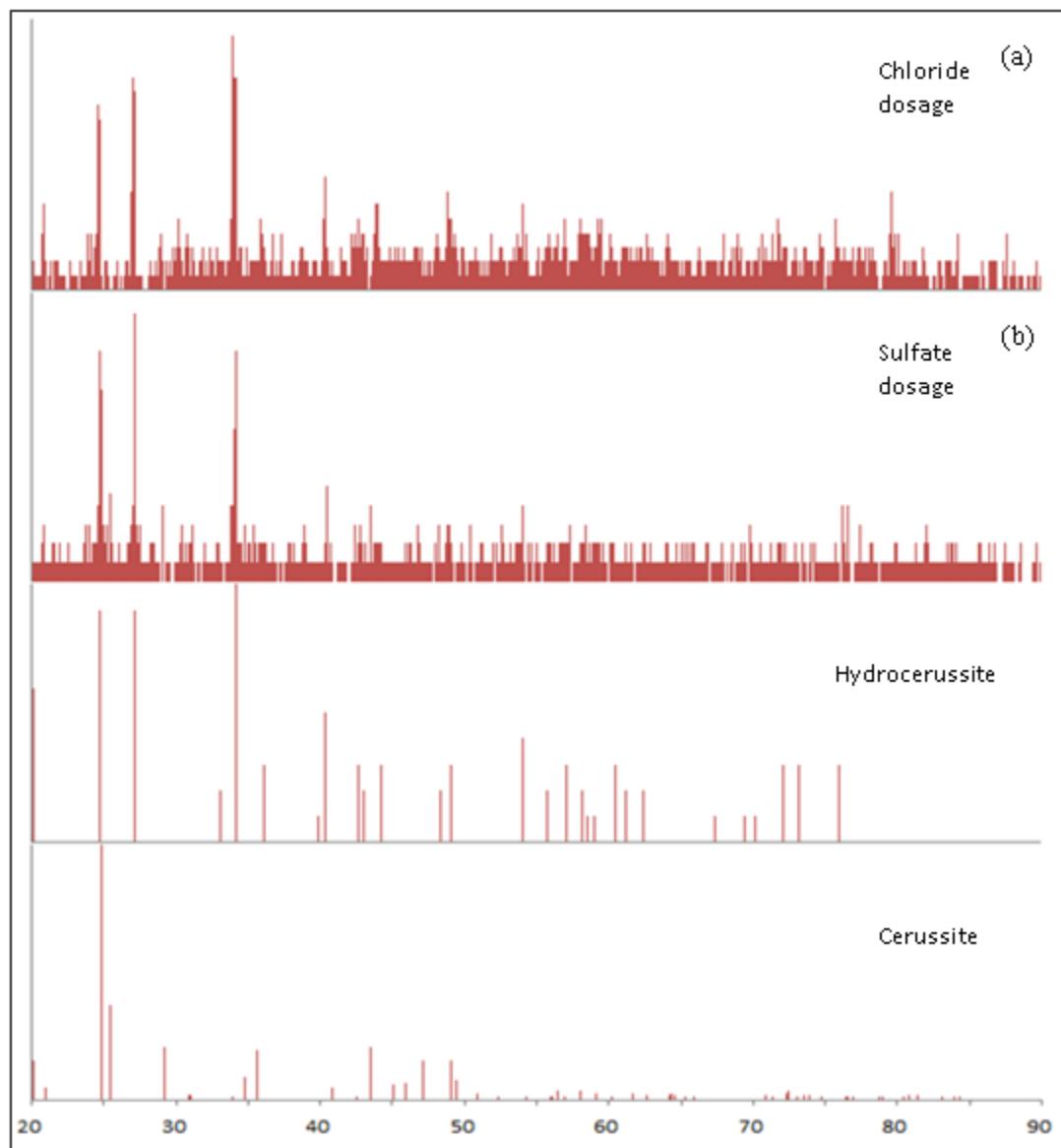


Fig. S13. X-Ray diffraction (XRD) patterns of scaling collected after 7 d under completely mixed conditions at pH 7.0. (a) Chloride dosage of 250 mg L^{-1} , (b) sulfate dosage of 250 mg L^{-1} . Reference standards for hydrocerussite and cerussite obtained from PDF cards: 00-013-0131 and 00-047-1734 respectively.