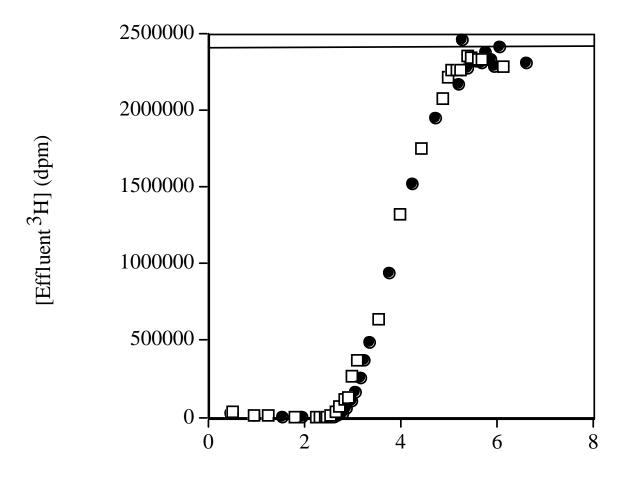
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Supporting Information

3 Hydrologic flow controls on biologic iron(III) reduction in natural sediments Morgan L. Minyard¹ and William D. Burgos^{1,*} 4 5 6 ¹ Department of Civil and Environmental Engineering, The Pennsylvania State University 7 * Corresponding author; address: 212 Sackett Building, University Park, PA, 16802-1408; telephone: 8 814-863-0578; fax: 814-863-7304; e-mail: wdb3@psu.edu 9 10 S2 – Supp. Info. Figure S1. ³H breakthrough curves from two of the columns operated at 11.1 pore 11 volumes (PV) d^{-1} . Open squares and closed circles represent the two replicate data sets, and solid line 12 represents the influent ³H concentration (disintegrations per minute). ³H breakthrough curves were used to define the PV as the volume displaced between the time when ³H was statistically greater than the 13 14 background concentration to the time when ³H was statistically inseparable from preceding and 15 proceeding values. The ³H breakthrough curves were analyzed to calculate a pore volume of 2.60 ± 0.13 16 mL (equivalent to a porosity of 0.44 v/v) and a dispersivity of 0.01 cm. 17 18 S3 - Supp. Info. Figure S2. Final biomass concentrations and distributions measured at the conclusion of

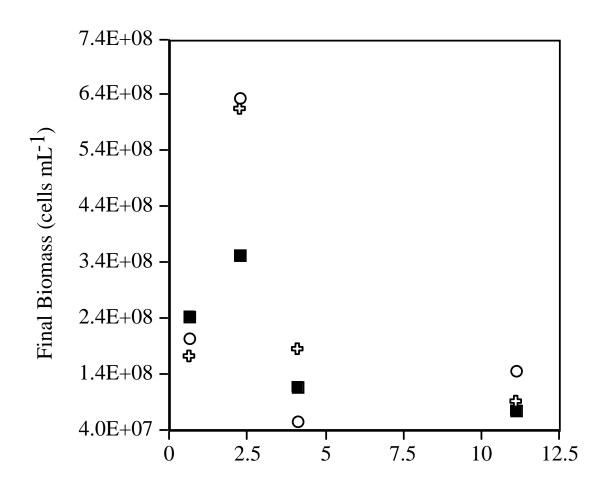
19 the column experiments. Phosphate-extractable cells were determined by combining $\sim 1/9$ of the column 20 contents with 5 mL of 20 mM Na₂HPO₄ (pH 7.0). The phosphate-sand suspension was vortex mixed for 21 one minute, allowed to settle overnight, and cells in the overlying water were enumerated by acridine 22 orange direct counts (AODC) using a Zeiss Axiophot microscope (Jena, Germany) based upon the average from 5 field counts (each field being $2.64 \times 10^{-4} \text{ cm}^2$) for each slide (three slides per column 23 24 region). Symbols represent mean values from influent (closed squares), middle (open squares), and 25 effluent (open crosses) regions of the column.



Cumulative Volume (mL)

26

Supporting Information Figure S1. ³H breakthrough curves from two of the columns operated at 11.1 pore volumes (PV) d⁻¹. Open squares and closed circles represent the two replicate data sets, and solid line represents the influent ³H concentration (disintegrations per minute). ³H breakthrough curves were used to define the PV as the volume displaced between the time when ³H was statistically greater than the background concentration to the time when ³H was statistically inseparable from preceding and proceeding values. The ³H breakthrough curves were analyzed to calculate a pore volume of 2.60 ± 0.13 mL (equivalent to a porosity of 0.44 v/v) and a dispersivity of 0.01 cm.



Flow Rate (pore volume da \bar{y}^1)

34

35 Supporting Information Figure S2. Final biomass concentrations and distributions measured at the 36 conclusion of the column experiments. Phosphate-extractable cells were determined by combining $\sim 1/9$ 37 of the column contents with 5 mL of 20 mM Na₂HPO₄ (pH 7.0). The phosphate-sand suspension was 38 vortex mixed for one minute, allowed to settle overnight, and cells in the overlying water were 39 enumerated by acridine orange direct counts (AODC) using a Zeiss Axiophot microscope (Jena, Germany) based upon the average from 5 field counts (each field being $2.64 \times 10^{-4} \text{ cm}^2$) for each slide 40 41 (three slides per column region). Symbols represent mean values from influent (closed squares), middle 42 (open squares), and effluent (open crosses) regions of the column.