Electronic Supplementary Material (ESI) for Metallomics. This journal is © The Royal Society of Chemistry 2020

- 1 Supplemental Data
- 2 Widhalm et al. "Human placental cell line HTR-8/SVneo accumulates cadmium by divalent metal
- 3 transporters DMT1 and ZIP14".
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- 6 Suppl. Table 1 Primers used in qPCR

Gene	Assay ID	Link
MT1A	Hs00831826_s1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs00831826_s1?CID=&ICID=&subtype=
MT1E	Hs01938284_g1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs01938284_g1?CID=&ICID=&subtype=
MT1X	Hs00745167_sH	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs00745167_sH?CID=&ICID=&subtype=
MT2A	Hs04194247_g1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs04194247_g1?CID=&ICID=&subtype=
SLC11A2	HS00167206_m1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs00167206_m1?CID=&ICID=&subtype=
SLC39A14	Hs00299262_m1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs00299262_m1?CID=&ICID=&subtype=
SLC39A8	Hs01061802_m1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs01061802_m1?CID=&ICID=&subtype=
UBC	Hs00824723_m1	https://www.thermofisher.com/taqman-gene-
		expression/product/Hs00824723_m1?CID=&ICID=&subtype=

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12 Suppl Fig.1



14 Suppl Fig. 2



- 17 Suppl Fig. 3

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- 24 Suppl Fig.4





33 TABLE and FIGURE LEGENDS

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35 Suppl. Table 1 Primers used in qPCR

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37 Suppl. Fig. 1 Validation of DMT1 antibodies for localisation studies (a) Detection of DMT1 by immunoblot 38 using either a rabbit anti-human DMT1 antibody from Biorbyt (#orb5976) or a rabbit anti-human DMT1 antibody 39 from Cell Signaling. Lysates of commercial negative (OriGene, #LY500001) and positive (OriGene, #LY433268) 40 controls (Neg. Con., Pos. Con.), lysates of BeWo and HTR-8/SVneo cells subjected to siRNA-mediated DMT1 41 gene knock down (si DMT1 B, si ctrl B; si DMT1 H, si ctrl H), whole placenta lysates 1-3, as well as human serum 42 albumin (HSA) were separated on reducing 7.5 % SDS-gels, transferred onto PVDF membranes and probed first with the DMT1 antibody from Biorbyt (1:200) followed by an HRP-labelled secondary antibody (1:10.000). 43 44 Afterwards, the membrane was stripped and incubated with the DMT1 antibody from Cell Signaling (1:1000) 45 followed by an HRP-labelled secondary antibody (1:10.000). Binding of antibodies was detected using enhanced chemiluminescence. Total proteins were detected with Amido black staining. Specific detection of DMT1 was 46 47 only observed for the employed Cell Signaling Antibody 48 49 Suppl. Fig. 2 Validation of ZIP14 antibody. Protein lysates (10 and 20 µg) from control and ZIP14 depleted cells were separated by SDS-PAGE and blotted on nitrocellulose. The membrane was cut and incubated over night 50

51 with antibodies for ZIP14 (Invitrogen, PA5-87880) and α -tubulin (A-tub; Sigma Aldrich, CP06) either 1:1000 or

52 1:2000 diluted and exposed to secondary antibody (IRDye® 800CW Goat anti-Rabbit IgG Secondary Antibody,

- 53 LI-COR, 926-32211) diluted 1:20.000 or 1:40.000 for 1h at room temperature and pictures taken with an Odyssey
- 54 imager (LI-COR).

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56 Suppl. Fig. 3 Levels of MT1A, MT1E and MT1X in Cd treated cells after MT2A depletion. Control and 57 MT2A depleted cells were treated with indicated Cd concentrations for 24h and expression of MT1A, MT1E and 58 MT1X evaluated. si ctrl treated with 0 μ M CdCl2 was set to 1. The data represent mean values \pm SD from three 59 independent experiments made in triplicate. The letters a-d denote homogeneous subgroups derived from one-way 60 ANOVA and S-N-K posthoc test (p<0.05)

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62 Suppl. Fig. 4 MT2A knockdown verification. Control and MT2A depleted cells were exposed to indicated Cd 63 concentrations for 24h (A), 48h (B) and 72h (C) and their MT2A expression analysed by qPCR. The data represent 64 mean values ± SD from three independent experiments made in six technical replicates. **p<0.01, *** p<0.001 65 from students t-test.

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67 Suppl. Fig. 5 Phase contrast images of MT2A depleted cells exposed to Cd. Control and MT2A depleted cells
68 were incubated with indicated Cd concentrations for 48h. Bar represents 500 μm.

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