ADDITIONS AND CORRECTIONS

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ATP9B, a P4-ATPase (a putative aminophospholipid translocase), localizes to the *trans*-Golgi network in a CDC50 protein-independent manner.

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The evidence shown in Fig. 1 that ATP10A is expressed in HeLa cells is not correct. ATP10A expression was measured using RT-PCR with the sense (ccttatccccagtcacagctg) and anti-sense (ccgagtctgcctcttggtacc) primer pair. However, subsequent cloning and sequencing of the RT-PCR product showed that the original primer pair amplified glucosidase, beta, acid (GBA, XM006726211.1) by annealing to the mRNA of GBA (XM006726211.1) at positions 541 and 823 (marked with an *asterisk* in the revised Fig. 1) rather than ATP10A mRNA. When we examined ATP10A expression using two other primer pairs (pair 1, cacaatgttcgtgggcctcc (sense) and aaggacactgaagccacacg (anti-sense); pair 2, gtctccgaattcaatcctttgac (sense) and ccgagtctgcctcttggtacc (anti-sense)), we detected ATP10A expression in MDA-MB-231 cells but not in HeLa cells (Fig. 1 Addendum). Therefore, we conclude that the band identified as ATP10A in the original Fig. 1 was a product of GBA mRNA amplification, and ATP10A expression was not detected in HeLa cells. This correction does not affect the results or conclusions of this work.

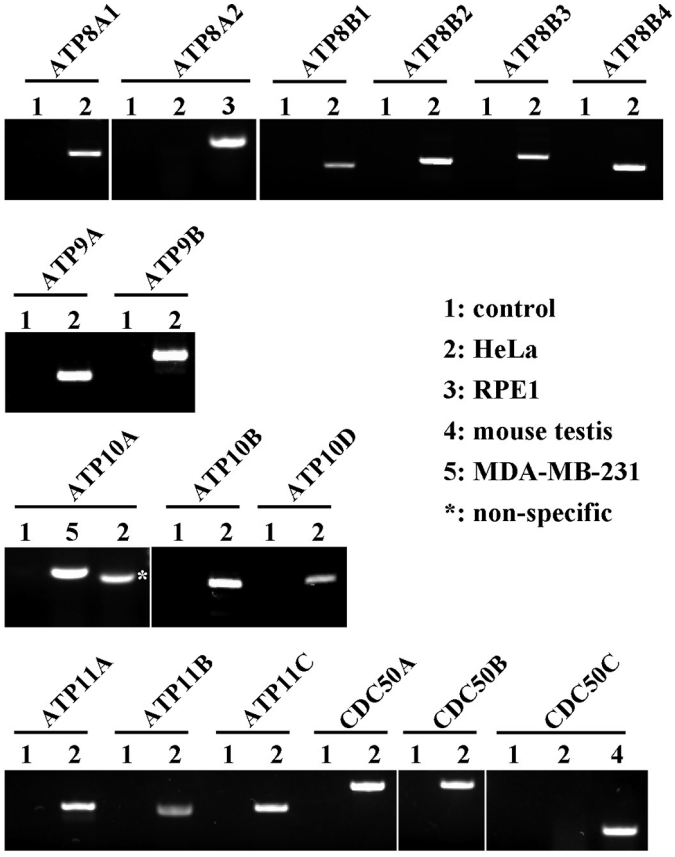

Fig. 1

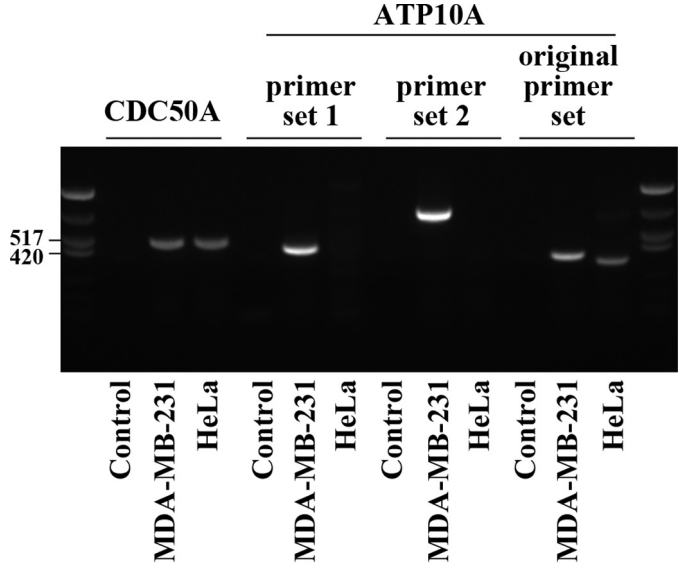

Fig. 1 Addendum

Authors are urged to introduce these corrections into any reprints they distribute. Secondary (abstract) services are urged to carry notice of these corrections as prominently as they carried the original abstracts.