



## Corrigendum

## Corrigendum to “Decreased membrane cholesterol in liver mitochondria of the point mutation mouse model of juvenile Niemann-Pick C1, *Npc1<sup>nmf164</sup>*,” [Mitochondrion 51 (2019) 15–21]

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The authors regret that we neglected to say that the long lived mice were able to do so because of a GFAP-promoter, *Npc1* cDNA transgene, which allows long term survival (Marshall et al., 2018). We have only recently become aware that this transgene might be slightly expressed in the liver in stellate cells. Since the mice are hypomorphs with about 10–15% of normal *Npc1* activity already, this would be a very small addition since stellate cells are only 1/3 of the nonparenchymal cells (Baba et al., 2004). Also, in *Npc1*-deficient mice, cholesterol is not stored in perisinusoidal cells but in parenchymal cells and giant macrophages. The mice were not used for the mitochondrial imaging.

The authors would like to apologise for any inconvenience caused.

### References

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