

Erratum

Mitochondrial Transplantation Attenuates Airway Hyperresponsiveness by Inhibition of Cholinergic Hyperactivity: Erratum

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In our paper [1], the following corrections should be provided.

1. (corrected) page 1252, left column, line 4-7 from the bottom: status (7.87 ± 0.10 vs. 8.74 ± 0.10 Hz, $p < 0.05$). In addition, after 5 min administration of 30 mg/ml Ach, the CBF was significantly decreased in both CS/LPS (6.76 ± 0.12 Hz) and control groups (7.47 ± 0.22 Hz).

2. (corrected) Figure 3 legend:

Figure 3. Mitochondria in the regulation of airway responsiveness in rats. **A.** The airway responsiveness in control, CS/LPS exposed and CS/LPS exposed rats delivered with exogenous mitochondria prepared from control rats ($n=11-17$, * $p < 0.05$ vs. control, some data in Fig 1A were re-presented for comparison). **B-C** Electron micrographs showing long spindle mitochondria with clear cristae in ciliated airway epithelium in control rats (*left, B*, control) and in normal rats for mitochondrial preparation (*middle, B*, mito), round mitochondria with swelling, unclear cristae in airway epithelium in CS/LPS exposed rats (*middle, B*, CS/LPS), and mixture of two types of mitochondria with distinct morphology in ciliated airway epithelium in CS/LPS exposed rats after delivery of exogenous mitochondria prepared from control rats (*right, B*, CS/LPS+mito) and quantitative comparisons of two shapes of mitochondria (C). Quantitative analysis was obtained from 204, 187, 184 and 211 mitochondria of 11-17 separate ciliated airway epithelial cells from 5, 4, 6 and 7 rats, respectively for control, mito, CS/LPS and CS/LPS+mito rats respectively. The mitochondria in each cell were counted from six fields randomly chosen each with an area of $5.25 \mu\text{m}^2$. For comparison, the control group in Figure 2I was included here. **D-F** The tissue of tracheal epithelium was freshly stripped from the airway of rats after an administration of APEX-labeled mitochondria. The subsequent electronic microscopy examination identified the localization of the APEX-labeled, exogenous mitochondria mainly within the ciliated epithelial cells (**D**), occasionally within the goblet cells (**E**) and basal cells (**F**). The results were consistent in 4 tissues from 3 individual rats. endo mito:

endogenous mitochondria (white arrow); exo mito: exogenous mitochondria (black arrow). white star filled: ciliated epithelial cells; white star open: goblet cells; black star: basal cells.

3. (corrected) Figure 4 legend:

Figure 4. Mitochondria in the regulation of CBF. **A:** The effects of M receptor inhibitor, ipratropium bromide (IB) on CBF in rats without any treatment (blank) and acetylcholine (Ach)-induced decline in CBF in rat right inferior lobar bronchus in control and CS/LPS exposed rats (n=10-13, * $p < 0.05$). The waveform graph of cilia beating last for 1000 millisecond was drawn and columns represented Mean \pm SEM of CBF. **B:** The quantitative analysis of Ach level in BALF of CS/LPS-exposed rats (n=6 or 4 for control or CS/LPS, respectively, $p > 0.05$). **C:** The quantitative analysis of Ach in supernatant of cultured RAECs treated by CSE/LPS (n=5 for each, $p > 0.05$). **D:** Time-dependent decline in CBF of cultured RAECs exposed to LPS plus cigarette smoking extract (CSE/LPS) (n=5 for each, * $p < 0.05$). **E:** IB and mitochondrial transplantation on 3 hour exposure of CSE/LPS-induced decline in CBF of cultured RAECs and the effects of M receptor enhancer, win62577 (WIN) (n=4-8, * $p < 0.05$).

4. (corrected) Figure 5C and Figure 5 legend:

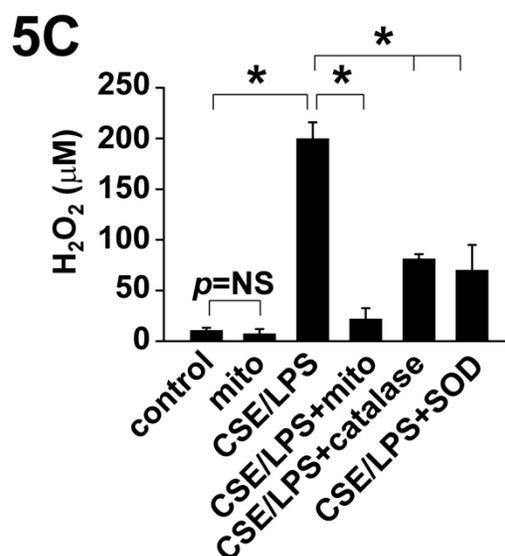


Figure 5. Mitochondria in the regulation cholinergic sensitivity and oxidant stress in cultured RAECs. **A:** Evaluation of cholinergic sensitivity by secreted level of LTB₄ from RAECs and the effects of CSE/LPS exposure. **B:** Mitochondria transplantation on CSE/LPS-enhanced cholinergic sensitivity in RAECs. For comparison, the control group and CSE/LPS group in **A** were included here. **C:** Mitochondrial transplantation on CSE/LPS-stimulated oxidant stress in RAECs. **D:** Oxidant stress on the regulation of CSE/LPS-enhanced cholinergic sensitivity in RAECs. **E-F:** Representative fluorescence of JC-1 showing CSE/LPS-induced alterations in mitochondria membrane potential and ratio of red against green fluorescence (n=4, * $p < 0.05$ for each).

5. (corrected) page 1255, right column, the bottom line:

the significantly increased mean linear intercept

6. (corrected) Figure S6.legend:

Figure S6. The dose-dependent effects of ipratropium bromide on CBF of airway epithelial cells. The effects of M receptor inhibitor, ipratropium bromide (IB) in 0, 25, 75, 125 and 250 µg/mL on the ciliary beating frequency (CBF) of cultured rat airway epithelial cells (n=4, R=0.9476, $p < 0.0001$).

References

- [1] Su Y, Zhu L, Yu X, Cai L, Lu Y, Zhang J, Li T, Li J, Xia J, Xu F, Hu Q. Mitochondrial Transplantation Attenuates Airway Hyperresponsiveness by Inhibition of Cholinergic Hyperactivity. *Theranostics*. 2016; 6(8):1244-60. doi: 10.7150/thno.13804