



## Skeletal muscle heat shock protein 60 increases after endurance training in mice and induces peroxisome proliferation-activated receptor- $\gamma$ coactivator-1 $\alpha$ 1 expression

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Heat shock protein (Hsp60) is a mitochondrial chaperonin whose unconventional cellular localizations and functions are discovered day by day.

In the present study, the levels of Hsp60 in fibres of the *soleus* muscle and its correlation to the expression of four isoforms of peroxisome proliferation-activated receptor- $\gamma$  (PPAR- $\gamma$ ) coactivator- $1\alpha$  (PGC1 $\alpha$ ) were investigated in 72 young (7-weeks old) healthy male mice (BALB/c AnNHsd) at baseline and after completing a 6-week endurance training program. The mice were assigned to one of the two experimental groups: SED (sedentary) or TR (trained). Short-term overexpression of hsp60, achieved by *in vitro* plasmid transfection, was then performed to determine whether this chaperonin could have a role in the activation of the expression levels of PGC- $1\alpha$  isoforms.

The levels of Hsp60 protein were fibre-type specific in the posterior muscles at baseline, and endurance training increased its content in type I muscle fibers. Concomitantly with the increased levels of Hsp60 released in the blood stream of trained mice, mitochondrial copy number and the expression of three isoforms of PGC-1 $\alpha$  increased. Overexpressing hsp60 in cultured myoblasts induced only the expression of PGC-1 $\alpha$ 1, letting us suppose a direct correlation between Hsp60 overexpression and PGC-1 $\alpha$ 1 activation.

Overall, these results suggest that during endurance training Hsp60 is upregulated and activates the mitochondrial biogenesis pathway, probably as a response to the oxidative stress induced by exercise. This study reveals a molecular response of skeletal muscle to a mechanical stress induced by training which involves the molecular chaperonin Hsp60 and the transcriptional co-activator PGC-1  $\alpha$ 1. The role of these proteins in aerobic adaptation and pathological conditions as cancer cachexia warrants further investigations.

## References

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