

Supplementary Materials

Selenium Forms and Dosages Determined Their Biological Actions in Mouse Models of Parkinson's Disease

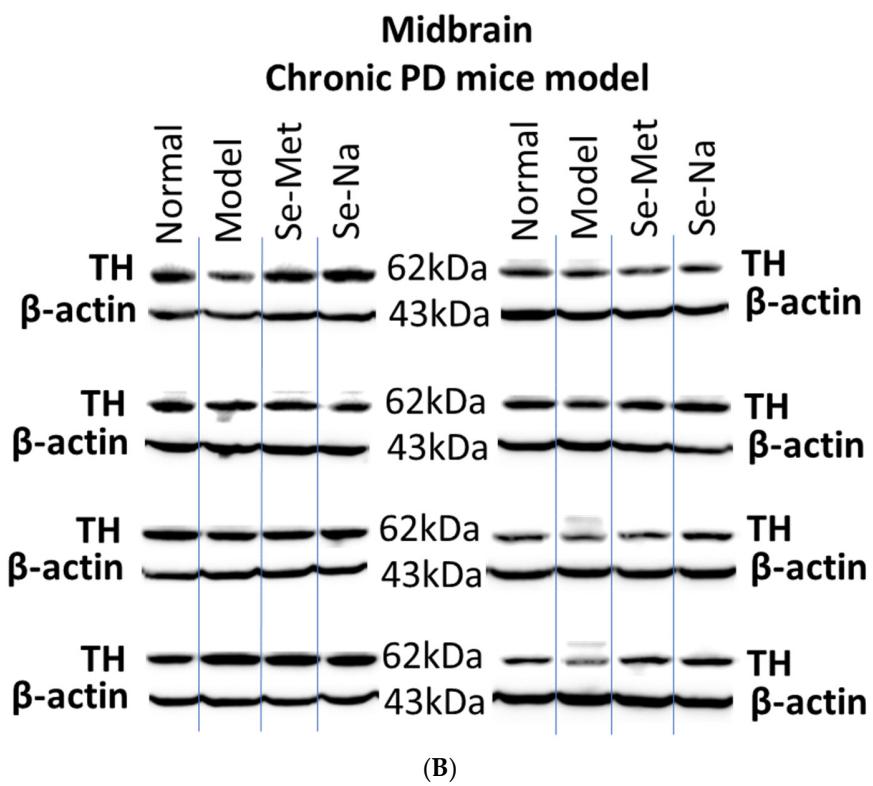
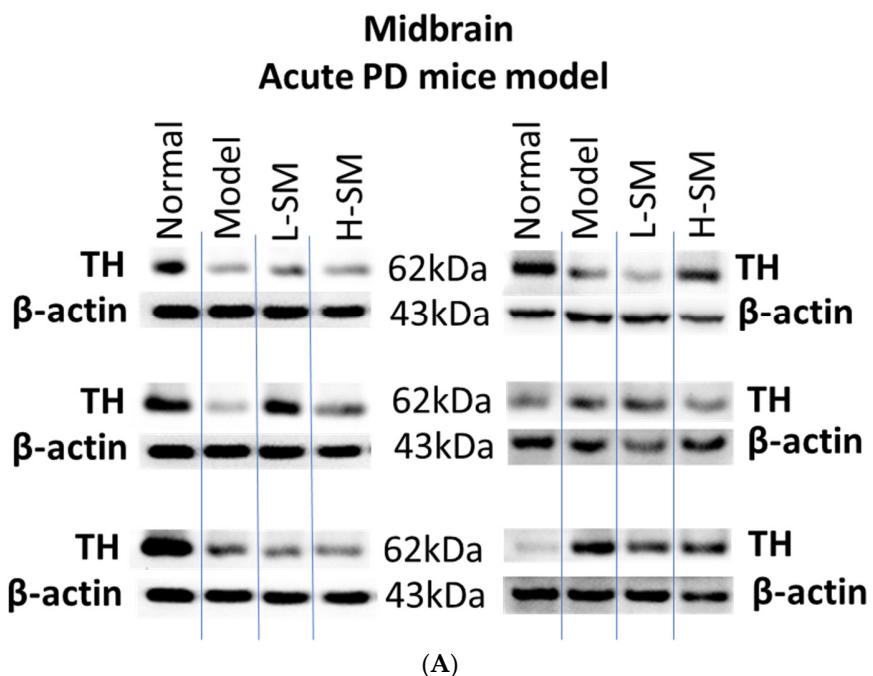
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Table S1. Primer list for real time RT-PCR analysis.

Gene	Forward Primer (5'-3')	Reverse Primer (5'-3')
TH [1]	CCCAAGGGCTTCAGAAGAG	GGGCATCCTCGATGAGACT
GPX1 [2]	TGGACTGGTGGTGCTCG	CGTCACTGGGTGTTGGC
GPX4 [2]	ACGATGCCAACCCACT	CACGCAGCCGTTCTT
Sepp1 [2]	GGCCGTCTTGTGTATCACCT	TGGTGTGTTGTGGCTATG
18S rRNA [1]	TCGGAACTGAGGCCATGATT	TTTCGCTCTGGTCCGTCTTG
GAPDH [3]	CATGGCCTTCCGTGTTCTTA	CCTGCTTCACCACCTTCTTGAT

Note: the primer sequences for these genes refer to previously published papers as shown in this Table.
Abbreviations: TH, tyrosine hydroxylase; GPX1, glutathione peroxidase 1; GPX4, glutathione peroxidase 4; Sepp1, selenoprotein P



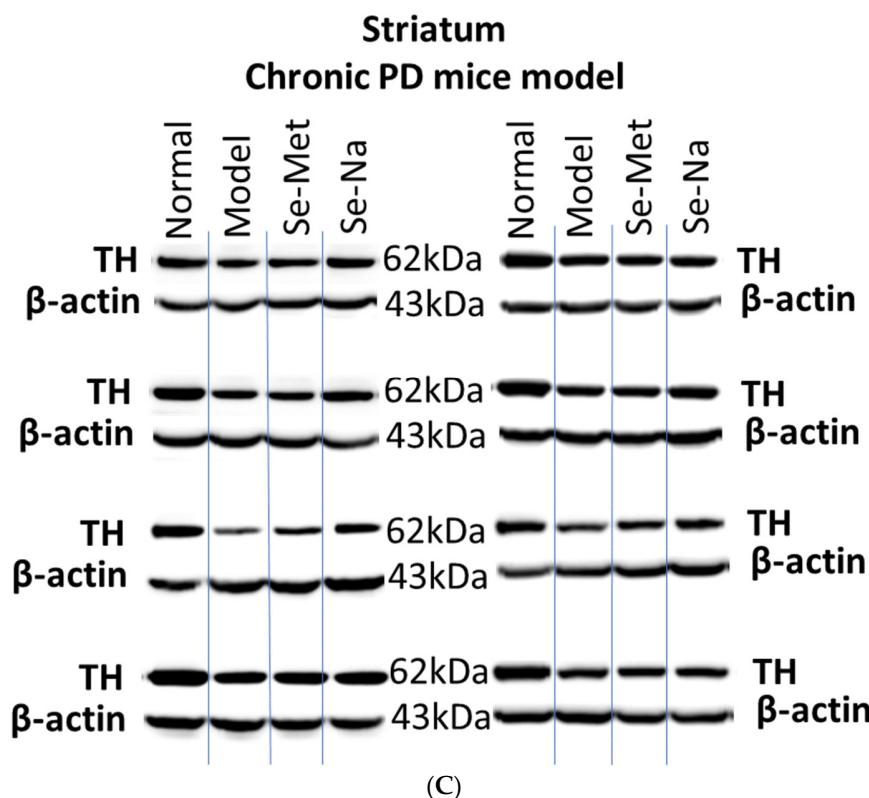


Figure S1. Original images of western blot (WB) bands to display protein expressions of tyrosine hydroxylase (TH) and its endogenous control (β -actin) in the midbrain of each mouse from acute PD model experiment (A); and the midbrain (B) or striatum (C) of each mouse from chronic PD model experiment. Grouping in acute PD mice model includes: Normal, Model, low (L-SM) and high (H-SM) dosage of seleno-L-methionine treated groups. Grouping in chronic PD mice model includes: Normal, Model, low dosage of seleno-L-methionine (Se-Met) and sodium selenite (Se-Na) treated groups.

References

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3. Yu, S; Luk, KH; Cheung, ST; Kwok, KW; Wong, KH; Chen, T: Polysaccharide-protein complex-decorated selenium nanosystem as an efficient bone-formation therapeutic. *J. Mater. Chem. B* **2018**, *6*, 5215–5219.