Qiyun Wu^{1,2}, Aster H.Y. Fung², Miranda L. Xu^{1,2}, K.M. Poon², Etta Y.L. Liu^{1,2}, Xiang P. Kong¹, Ping Yao², Qing P. Xiong², Tina T.X. Dong^{1,2}, Karl W.K. Tsim^{1,2}

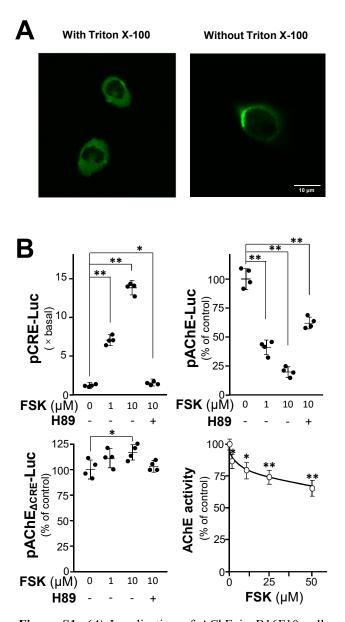


Figure S1. (A) Localization of AChE in B16F10 cells. B16F10 cells were fixed with 4% PFA for 15 min and stained with anti-AChE antibody, with or without 0.1% Triton X-100, followed with the Alexa 488 conjugated antibody. One representative result is shown, n = 4. (B) Cultured B16F10 cells transfected with pCRE-Luc, pAChE-Luc and AChE_{ACRE}-Luc constructs were treated with various concentrations of forskolin (FSK) for 12 hours. H89 (10 μ M) was used as a blocker of PKA. Cell lysates were collected for luciferase assay and Ellman assay.

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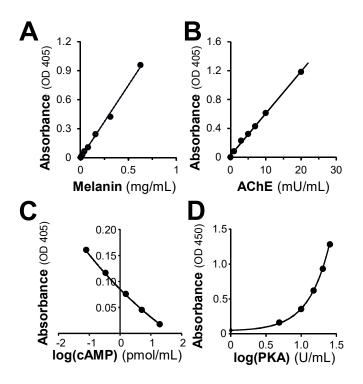


Figure S2. (A) Standard curve for synthetic melanin at concentration ranges between 0-1 mg/mL. (B) Standard curve for AChE activity ranges between 0-30 mU/mL for 30 min reaction time in Ellman assay. (C) Standard curve for cAMP ranges from 0-20 pmol/mL. (D) Standard curve for PKA activity ranges from 0-40 U/mL.