



Supplementary Material

A One-Step Extraction and Luminescence Assay for Quantifying Glucose and ATP Levels in Cultured HepG2 Cells

Rita Csepregi ^{1,3}, Viktória Temesfői ^{1,3}, Nikolett Sali ¹, Miklós Poór ^{2,3}, Paul W. Needs ⁴, Paul A. Kroon ⁴, and Tamás Kőszegi ^{1,3*}

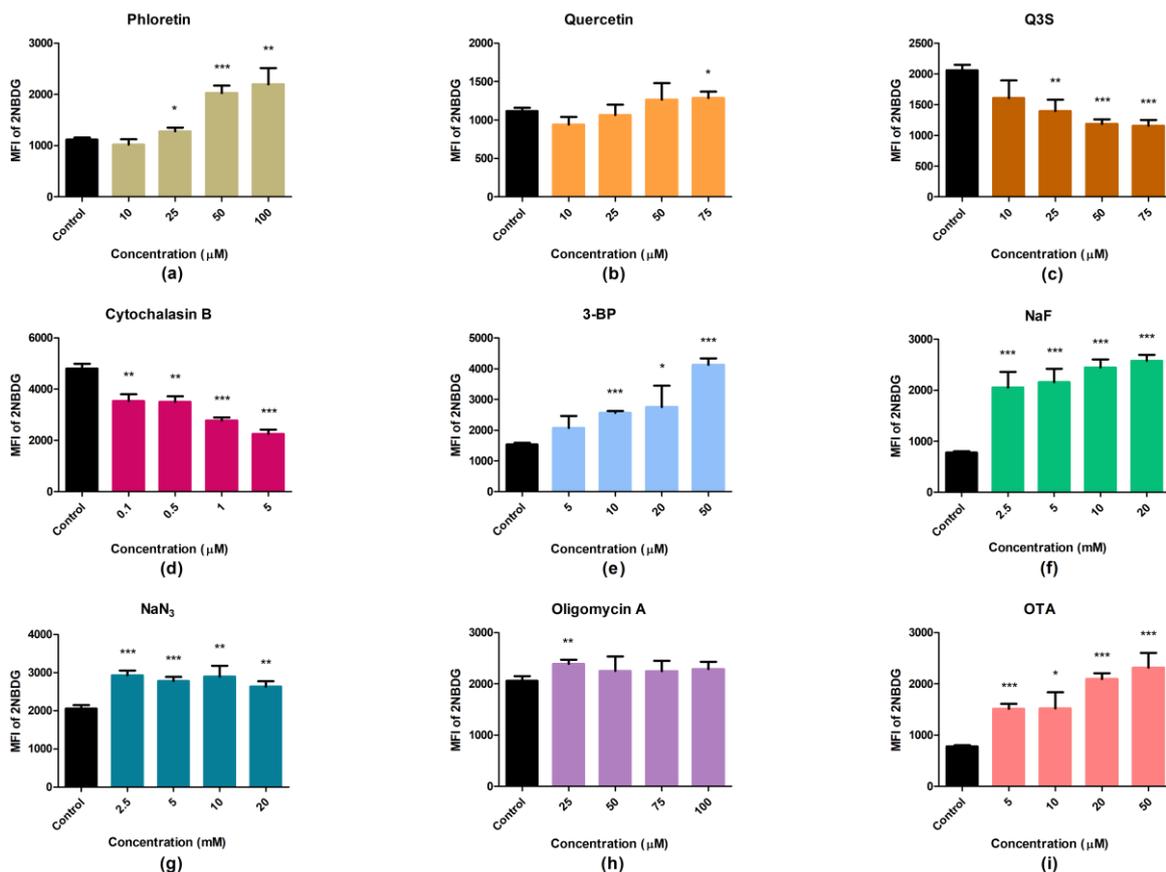
¹ Department of Laboratory Medicine, University of Pécs, Medical School, H-7624 Pécs, Ifjúság u. 13, Hungary; ritacsepregi93@gmail.com (R.C.); vtemesfoi@gmail.com (V.T.); niki26@gmail.hu (N.S.)

² Department of Pharmacology, University of Pécs, Faculty of Pharmacy, H-7624 Pécs, Szigeti u. 12, Hungary; poor.miklos@pte.hu (M.P.)

³ János Szentágotthai Research Center, H-7624 Pécs, Ifjúság u. 20, Hungary

⁴ Quadram Institute Bioscience, Norwich Research Park, Norwich NR4 7UA, UK; paul.needs@quadram.ac.uk (P.N.); paul.kroon@quadram.ac.uk (P.K.)

* Correspondence: tamas.koszegi@aok.pte.hu; Tel.: +36-30-491-7719; Fax: +36-72-536-121



Supplementary Figure 1. 2-NBDG uptake in HepG2 cells as results of (a) phloretin, (b) quercetin, (c) Q3'S, (d) cytochalasin B, (e) 3-BP, (f) NaF, (g) NaN₃, (h) oligomycin A, and (i) OTA treatments. Incubation time with metabolic inhibitors: 4 h, 2-NBDG: 1 h, MFI: mean fluorescence intensity. Columns represent the mean of medians, error bars show the interquartile range of fluorescence intensity of intracellular 2-NBDG. PI positive cells are included into the analysis. (* $p < 0.05$, ** $p < 0.01$ *** $p < 0.001$ compared with the controls).