Supplementary data to:

Quantitative High-Throughput Identification of Drugs as Modulators of Human Constitutive Androstane Receptor

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Figure S1. Cytotoxicity of daunorubicin and bortezomib in HPH. Human primary hepatocytes (from donor HL-104) cultured in 96-well collagen coated plates were treated with 0.1% DMSO, daunorubicin, or bortezomib at indicated concentrations for 24 hours. MTT assay was carried out following the manufacturer's instruction. Data represent the mean \pm SD (n =3).



Figure S2. Effects of daunorubicin and bortezomib on the expression of CYP2B6 and CYP3A4 in HPH. HPH (from donor HL-104) were treated with the vehicle control (0.1% DMSO), daunorubicin (1, 2.5 and 5 μ M), or bortezomib (0.1, 0.5, and 1 μ M) for 24 h. Real-time PCR was used to analyze the expression of CYP2B6 (A) and CYP3A4 (B). Data represents the mean ± SD (n=3). #, P < 0.05.