

Supplementary Data

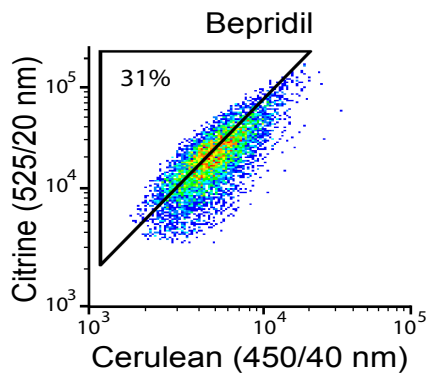
Identification of FDA-approved drugs targeting the Farnesoid X receptor

Sandra M.W. van de Wiel^{1,2}, Ingrid T.G.W. Bijsmans³, Saskia W.C. van Mil^{1,3} and Stan F.J. van de Graaf^{1,2}

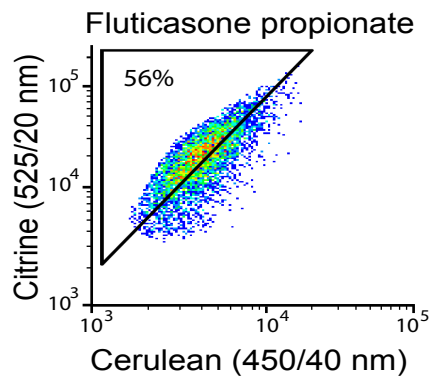
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Supplement Figure 1

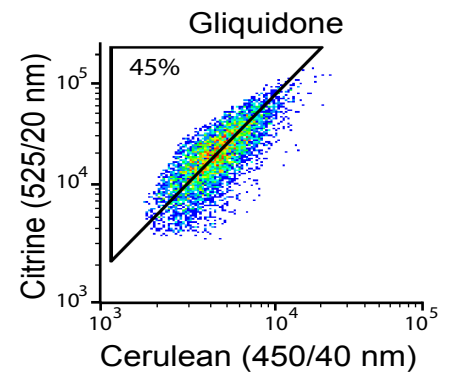
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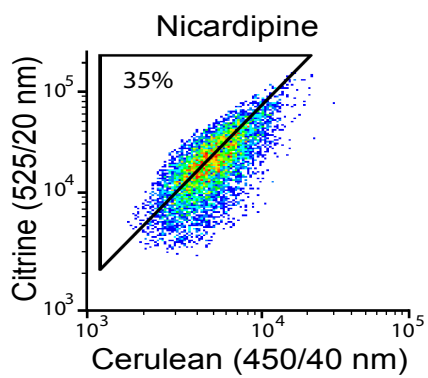
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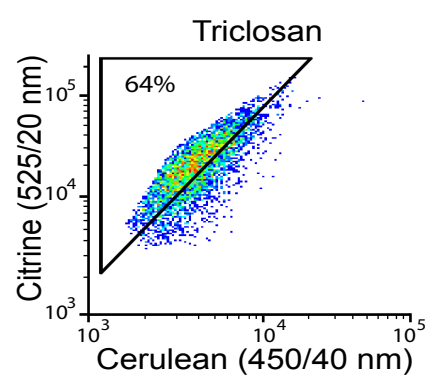
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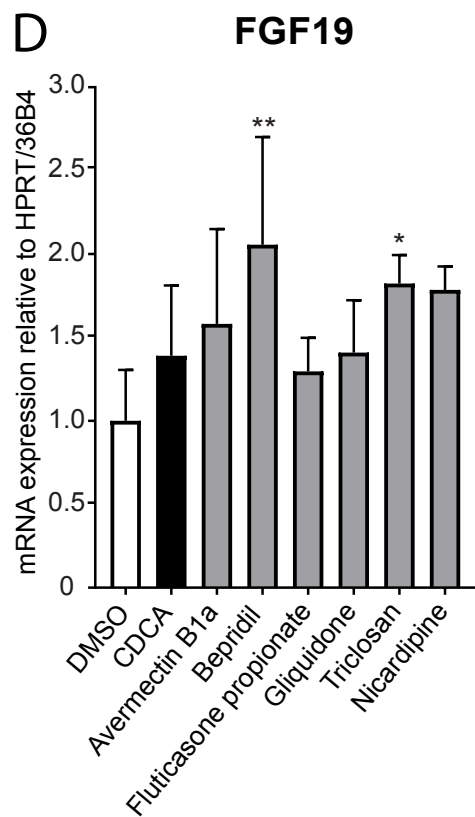
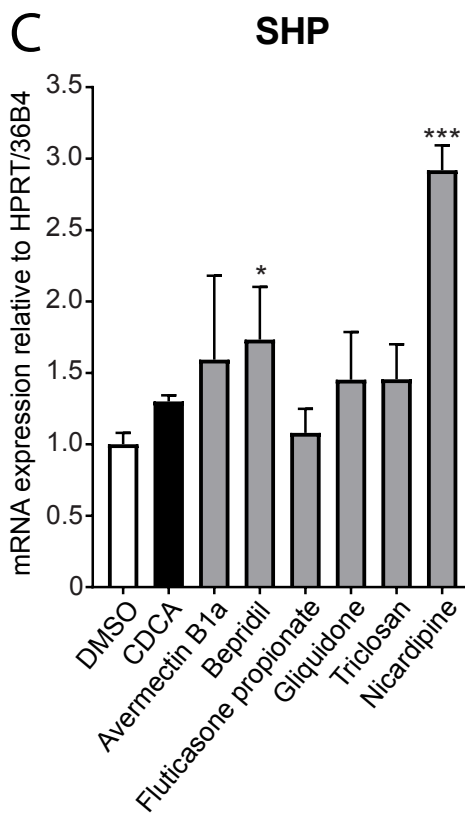
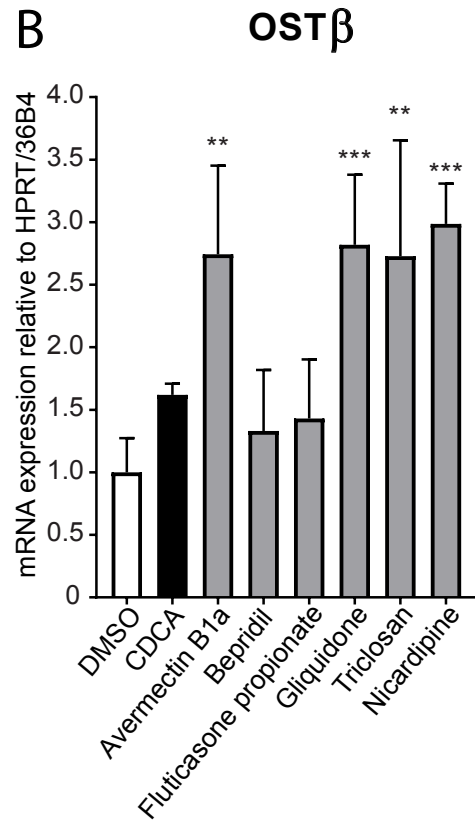
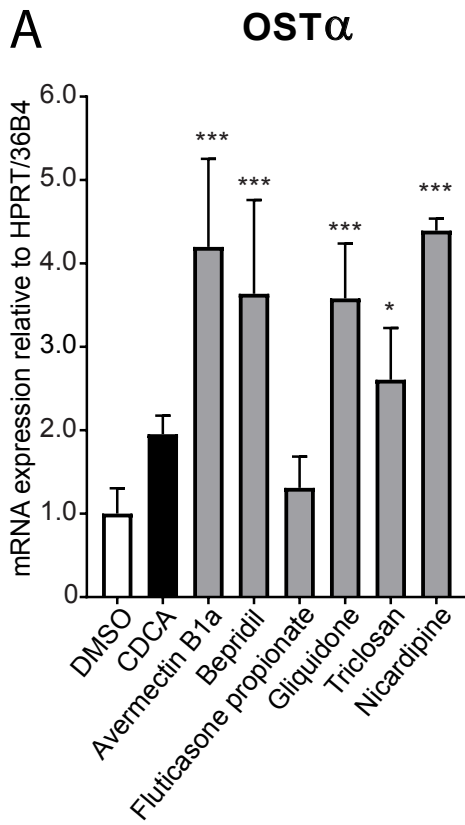
D



E



Supplement Figure 2



Supplementary figure 1: Representative FACS-plots of five hits showing the amount of FRET+ cells after 30 minute treatment with negative control (DMSO), positive control (GW4064) or 10 μ M of compound dissolved in DMSO. Numbers indicate the percentages of cells within the FRET gate.

Supplementary figure 2: Compounds directly increase FXR target gene expression. (A-E) Huh7 cells were treated for 6 hours with either DMSO, 10 μ M CDCA or 10 μ M compound and were analysed for gene expression of FXR target genes *OST α* , *OST β* , *SHP* and *FGF19*. This experiment (n=4) was performed in duplicate. Data is given as means \pm SD. Significance is measured relative to DMSO controls. *p < .05 (One-way ANOVA; post hoc: Dunnett's multiple comparison).

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Supplementary table 1

Primer list for RT-qPCR analysis

Gene	Forward primer (5'-3')	Reverse primer (5'-3')
<i>HPRT</i>	TGACCTTGATTTATTTGCATACC	CGAGCAAGACGTTTCAGTCCT
<i>36B4</i>	TCATCAACGGTACAAACGA	GCCTTGACCTTTTCAGCAAG
<i>SLC51A</i> (OST α)	AGATAACGCTGACCCTGGTG	AATTTGGCTCCCATGTTCTG
<i>SLC51B</i> (OST β)	GCAGCTGTGGTGGTCATTAT	TAGGCTGTTGTGATCCTTGG
<i>ABCC2</i> (MRP2)	ACAGAGGCTGGTGGCAACC	ACCATTACCTTGTCACTGTCCATGA
<i>NR0B2</i> (SHP)	CGCCCTATCATTGGAGATGT	TGTCTATACAGGCTTGCCCC
<i>FGF19</i>	TTTCTCATCACTTCCCCAGG	AGGCTTCCCCTACTCCTGAA
<i>G6Pase</i>	CCTTGCTGCTCATTTTCCTC	GGCTGGCATTATAGATGCTGT
<i>CYP7A1</i>	AATCCATACCTGGGCTGTG	AGGCAGCGGTCTTTGAGTTA