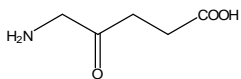
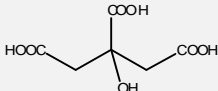
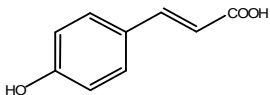
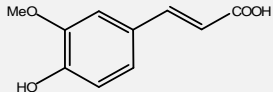
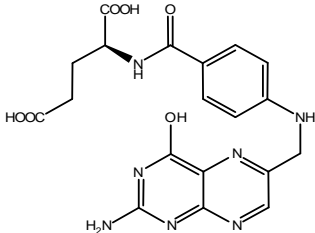
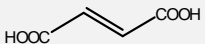
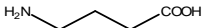
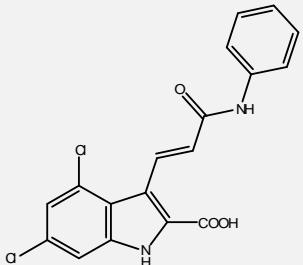
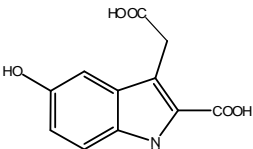
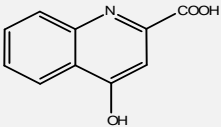
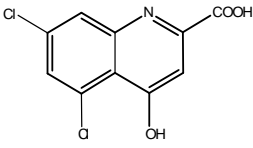
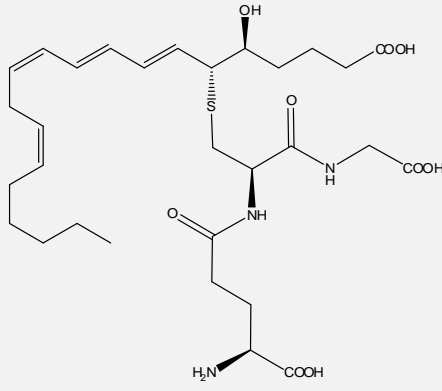
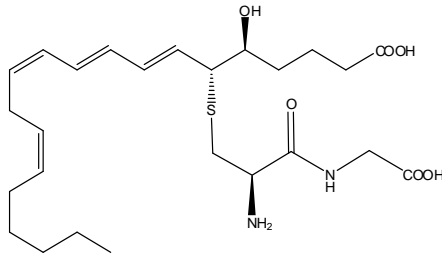
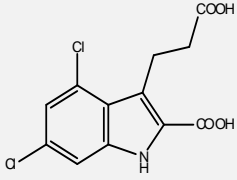
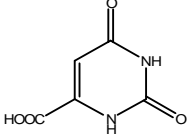
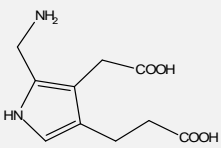


Supplemental Table 1

Test compound	Test concentration	Chemical structure
5-Aminolevulinic acid	100 μ M – 1 mM	
Citric acid	30 μ M	
p-Coumaric acid	30 μ M	
Ferulic acid	30 μ M	
Folic acid	30 μ M	
Fumaric acid	30 μ M	
γ -Aminobutyric acid	30 μ M	
Gavestinel	1-100 μ M	
5-Hydroxyindoleacetic acid	30 μ M	

Kynurenic acid	30 μ M	
5,7-Dichlorokynurenic acid	30 μ M	
LTC ₄	0.01 nM-1 μ M	
LTD ₄	0.1 nM – 1 μ M	
MDL29,951	1 nM – 100 μ M	
Orotic acid	30 μ M	
Porphobilinogen	10 μ M	

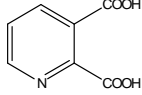
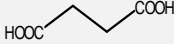
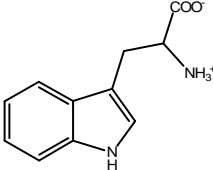
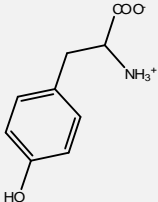
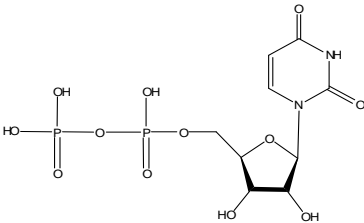
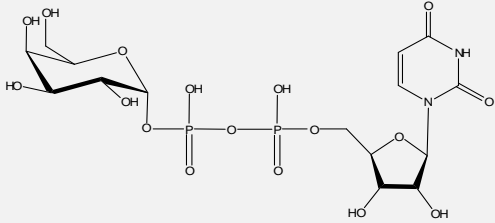
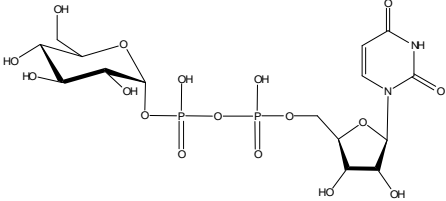
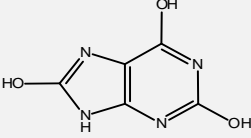
Quinolinic acid	30 μ M	
Succinic acid	30 μ M	
Tryptophane	30 μ M	
Tyrosine	30 μ M	
UDP	0.1 nM – 100 μ M	
UDP-galactose	0.1 nM – 100 μ M	
UDP-glucose	0.1 nM – 100 μ M	
Uric acid	30 μ M	

Table S1. Chemical structures of test compounds selected for biomolecular screening at GPR17. Molecules that qualified for the biomolecular screening contain at least one negative charge and/or represent precursors or intermediates of metabolic pathways with particular links to CNS activity. The concentration(range) applied in the biomolecular screening is indicated. UDP: uridine-diphosphate.