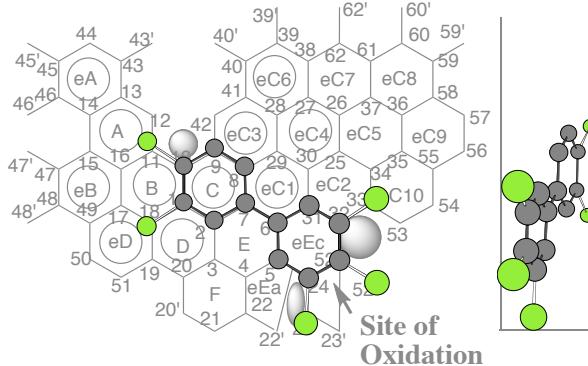
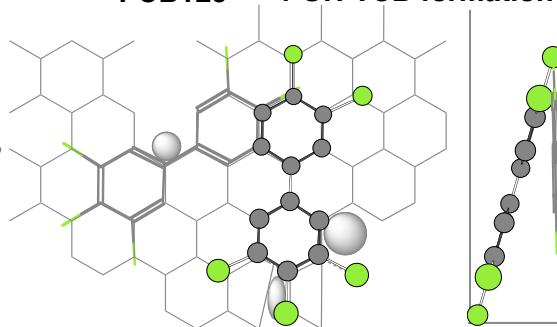
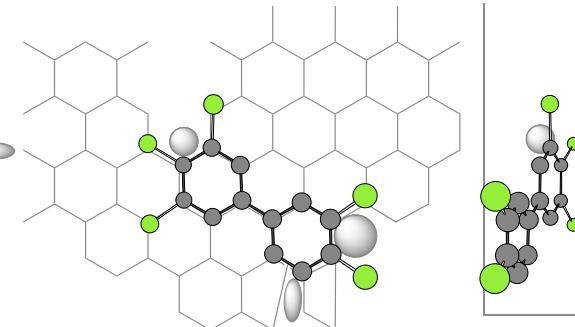
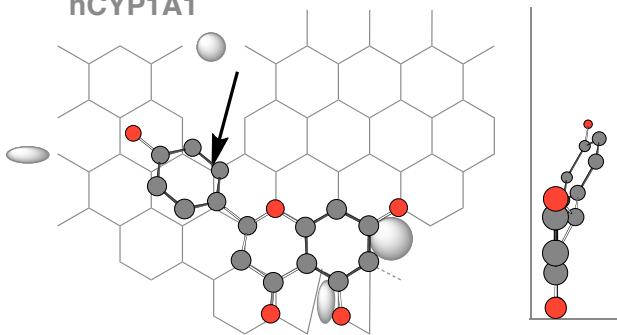
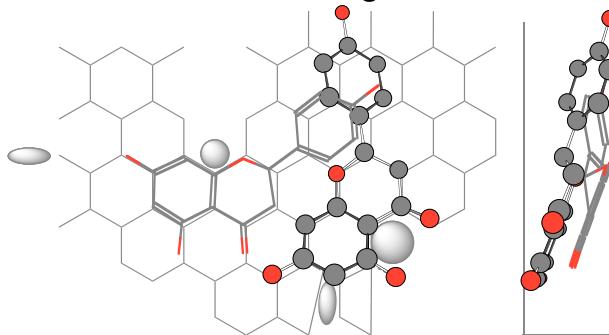
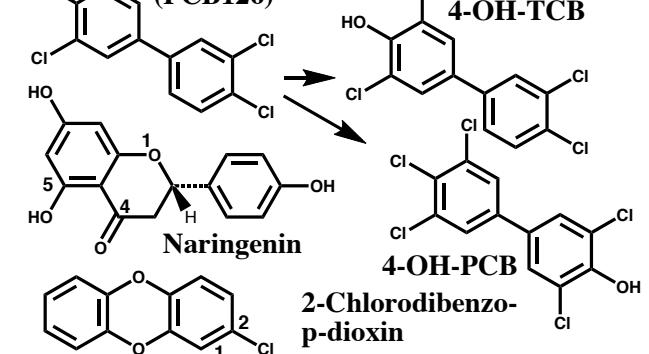
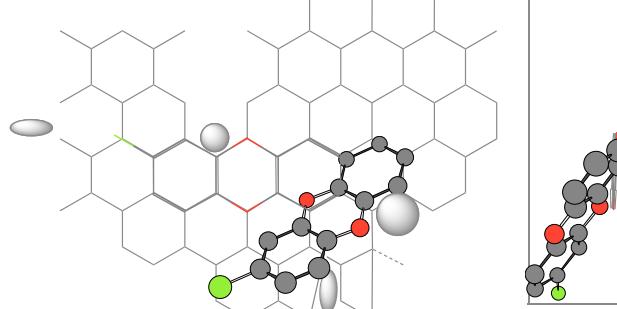
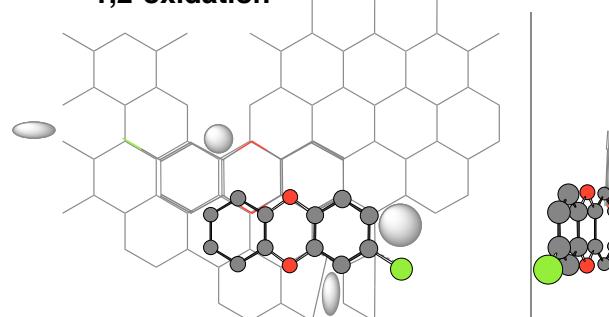
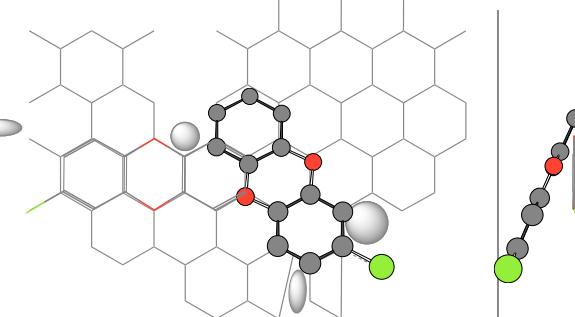


A rCYP1A1 PCB126 4-OH-TCB formation**B rCYP1A1 PCB126 bi-molecule binding 4-OH-TCB formation****C rCYP1A1 4-OH-PCB formation****D rCYP1A1 hCYP1A1 Naringenin inhibition****E rCYP1A1 Naringenin inhibition****3,3',4,4',5'-Hexachlorobiphenyl (PCB126)****F hCYP1A1 2-Chlorodibenzo-p-dioxin inhibition****G rCYP1A1 2-Chlorodibenzo-p-dioxin 1,2-oxidation****rCYP1A1 2-Chlorodibenzo-p-dioxin 2,3-oxidation**

Supplement Fig. 3 Rodent CYP1A1-selective Pier-sitting (Position 52')

Placements of PCB126 for the 4-OH-TCB (A and B) and 4-OH-PCB formations (C), of naringenin for the inhibition (D and E), and of 2-chlorodibenzo-p-dioxin (F, G and H) on CYP1A1 Template