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Structural optimization and structure-activity relationship studies of *N*-phenyl-7,8-dihydro-6*H*-pyrimido[5,4-b][1,4]oxazin-4-amine derivatives as a new class of inhibitors of RET and its drug resistance mutants

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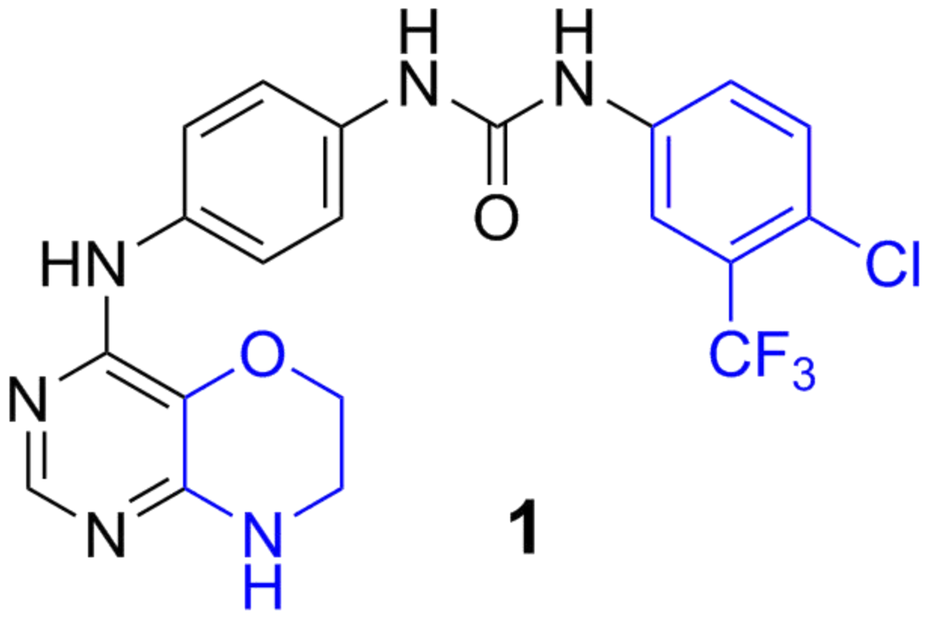
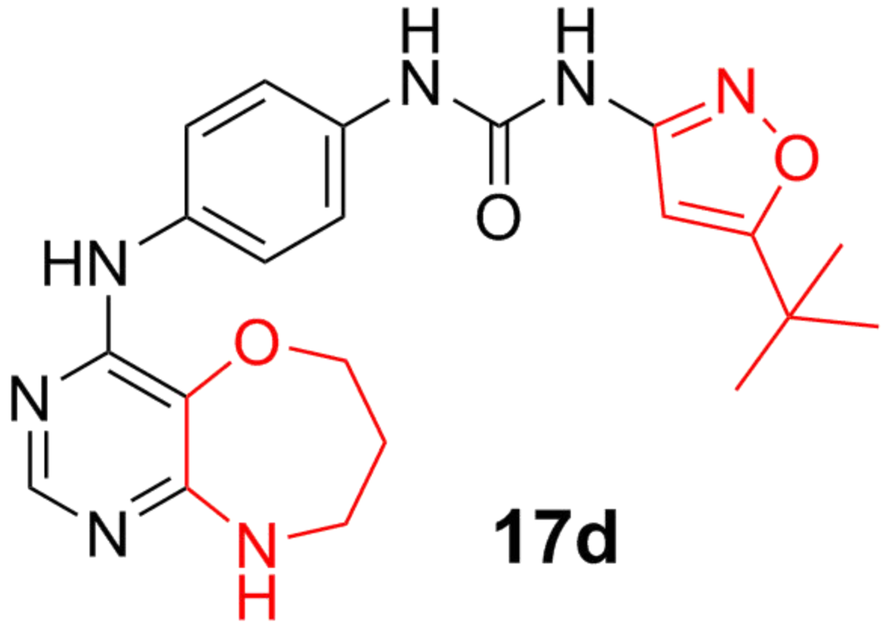


1

RET-wt: 0.177 μM
RET-V804M: 0.188 μM
RET-V804L: 0.285 μM

Structural optimization and SAR analyses



17d

RET-wt: 0.010 μM
RET-V804M: 0.015 μM
RET-V804L: 0.009 μM