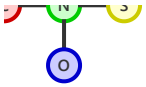
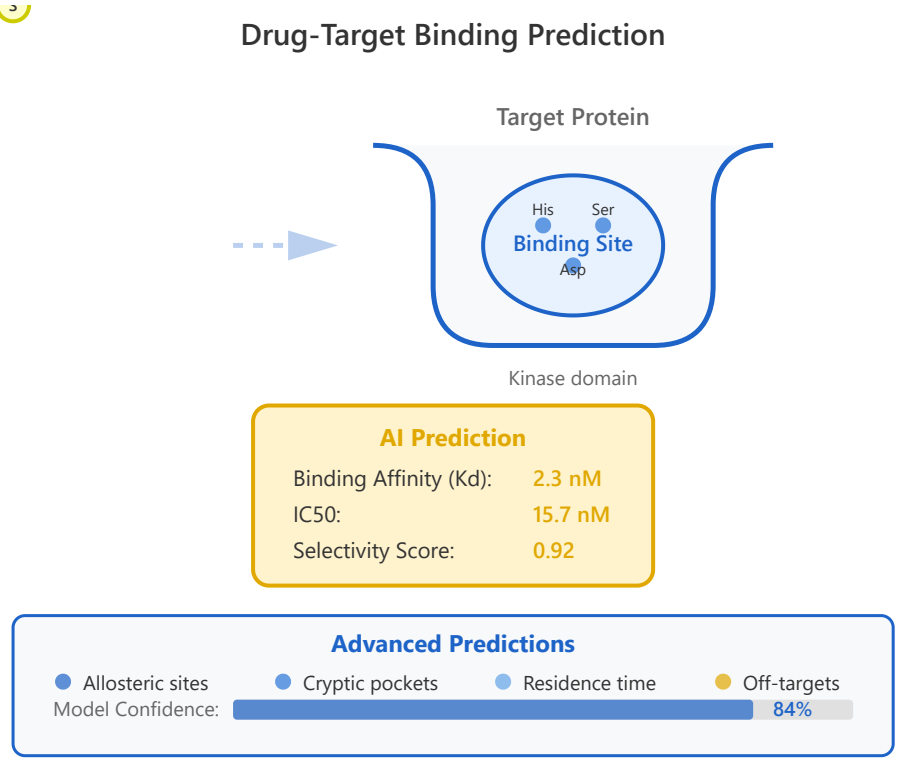
Drug-Target Affinity



MW: 350.4



Binding prediction

Kd, Ki, IC50 values

Kinase selectivity

Off-target profiling

Allosteric sites

Non-competitive binding

Cryptic pockets

Hidden binding sites

Residence time

Drug-target kinetics