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Design and synthesis of novel 1*H*-tetrazol-5-amine based potent antimicrobial agents: DNA topoisomerase IV and gyrase affinity evaluation supported by molecular docking studies

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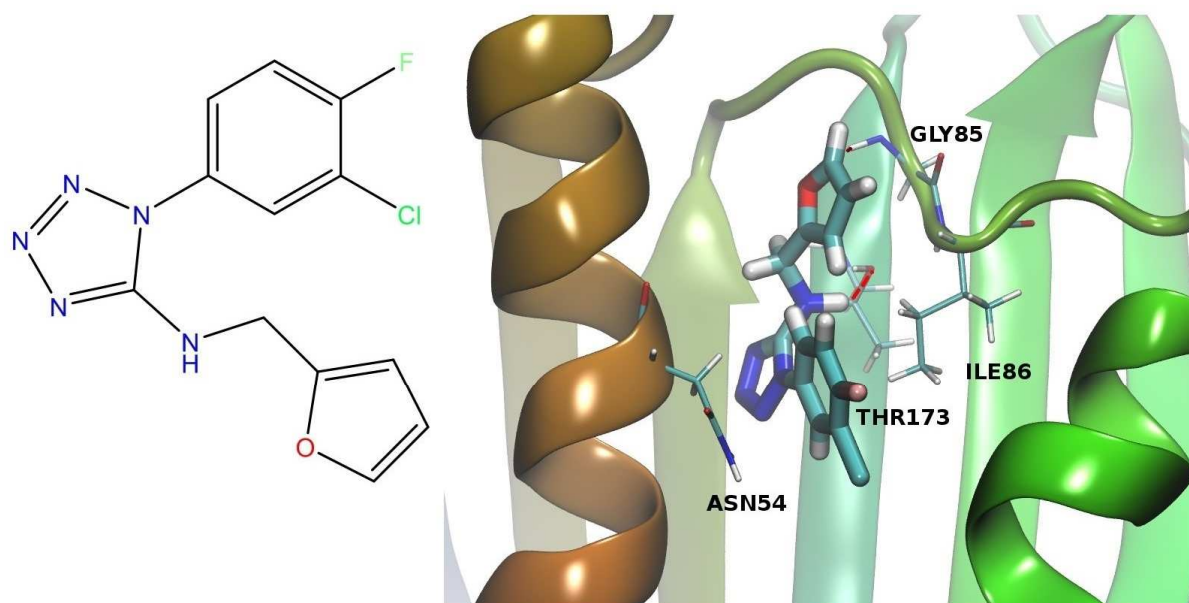
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Compound **11** was more potent to Ciprofloxacin against standard strains *E. faecalis*, *M. luteus*, *E. coli*, *P. vulgaris* (MIC 1 - 7 μ M).

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