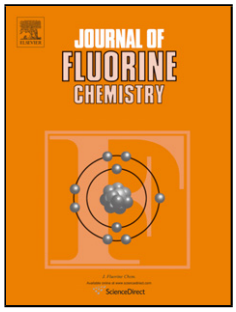

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SYNTHESIS AND ANTIMYCOBACTERIAL EVALUATION OF NEW (2-OXO-2*H*-CHROMEN-3-YL) SUBSTITUTED FLUOROQUINOLONES

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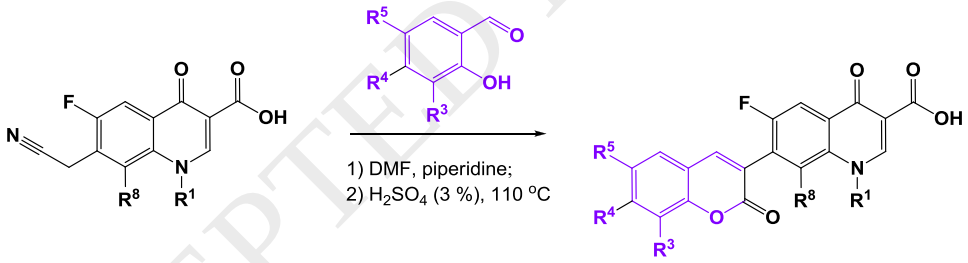
Graphical Abstract



Highlights

- The cyanomethyl fragment was incorporated into the benzene ring of fluoroquinolones
- Cyanomethyl fluoroquinolones was converted to (2-oxo-2*H*-chromen-3-yl) fluoroquinolones
- Substitution of another fluorine atom provides 1,4-dihydro[1]-benzoxepino[2,3-*g*]quinolones
- Some of new fluoroquinolones exhibit antimicobacterial activity

Abstract

An efficient method for incorporation of the cyanomethyl fragment into the benzene ring of bi- and tricyclic fluoroquinolones through the nucleophilic substitution of a fluorine atom with carbanions derived