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Green approach for the synthesis of thiophenyl pyrazoles and isoxazoles by adopting 1,3-dipolar cycloaddition methodology and their antimicrobial activity

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GRAPHICAL ABSTRACT

Green Approach for the Synthesis of Thiophenyl Pyrazoles And Isoxazoles By
Adopting 1,3-Dipolar Cycloaddition Methodology And Their Antimicrobial
Activity

$$S_{NO_2}^2$$
 $S_{NO_2}^2$
 $S_{NO_2}^2$

- Thiophenyl pyrazoles and isoxazoles were synthesized by 1,3-dipolar cycloaddition of nitrile imines and nitrile oxides to *N*-(arylethenesulfonyl)thiophene-2-carboxamides under green conditions followed by aromatization.
- The presence of electronwithdrawing substituents on the aromatic ring increased the activity.
- **4f**, **7e**, **7f**, **8e**, **8f**; MIC = $6.25 \mu g/well$ against *B. subtilis*.
- 8e, 8f; MIC = $6.25 \mu g/well$ against *A.niger*.

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