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## SELECTIVE INTRODUCTION OF TRIFLUOROMETHYL GROUPS INTO HETEROCYCLIC SYSTEMS

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Two new strategies for introduction of single trifluoromethyl groups into heterocyclic systems — starting from hexafluoro-acetone — are presented.

$$F_3C$$
  $\longrightarrow$   $\longrightarrow$   $CF_3$ 

Strategy I: A two-step procedure consisting of

- 1) ring synthesis with bis(trifluoromethyl) substituted hetero-1,3-dienes
- 2) HCF<sub>3</sub> elimination

offers versatile access to trifluoromethyl substituted ring systems.

Strategy II: A reaction sequence consisting of

- 1) reductive fluorine elimination
- 2) electrocyclic ring closure
- 3) elimination with aromatization

offers a new route to trifluoromethyl substituted 1,3-azoles. Scope and limitation of the two new procedures will be discussed.

$$F_{3}C = 0 \longrightarrow R \times SnCl_{2} F_{3}C \longrightarrow F_$$