Recent advances in 4-aminoquinazoline based scaffold derivatives targeting EGFR

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Abstract

Quinazoline derivatives are fused heterocyclic ring systems which have been explored for
their inhibitory activity towards various protein kinase enzymes and their role as anticancer
agents. The present review to represent the most recent synthetic strategies and medicinal aspects
including structure activity relationships of substituted quinazolines as EGFR inhibitors reported
to date.

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