Recent advances in 4-aminoquinazoline based scaffold derivatives targeting EGFR kinases as anticancer agents

Nasser Saad ,Rania S.M. Ismail, Sahar Abuserii d, Dalal A. Abou El Ella c

Associate Professor of Pharmaceutical Chemistry

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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