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[1,2,4]Triazolo[1,5-c]pyrimidines as adenosine receptor antagonists: Modifications at the 8 position to reach selectivity towards A₃ adenosine receptor subtype

Stephanie Federico, Enrico Margiotta, Veronica Salmaso, Giorgia Pastorin, Sonja Kachler, Karl-Norbert Klotz, Stefano Moro, Giampiero Spalluto

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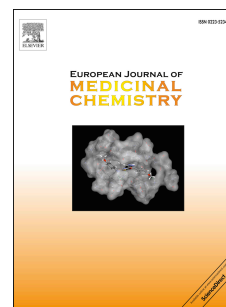
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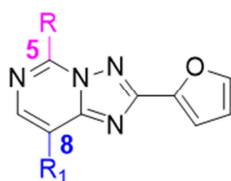
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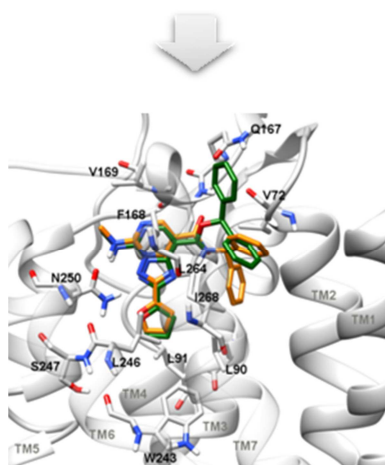
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bifunctional moieties to
create conjugates useful in
theranostic



esters and amides to
achieve A₃ potency and
selectivity



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