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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 A_3 adenosine receptor subtype

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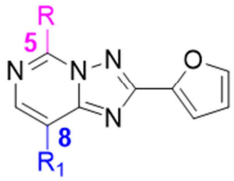
bifunctional moieties to create conjugates useful in theranostic



esters and amides to achieve A_3 potency and selectivity