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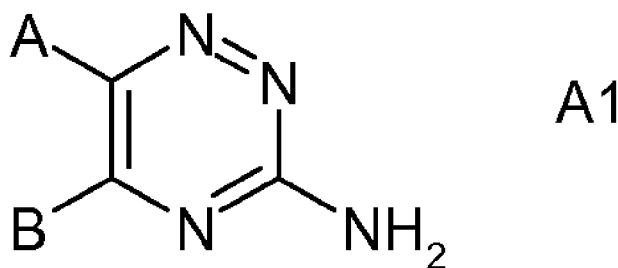
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(54) Title: 1,2,4-TRIAZINE-4-AMINE DERIVATIVES



(57) Abstract: According to the invention there is provided a compound of formula A1 which may be useful in the treatment of a condition or disorder ameliorated by the inhibition of the A<sub>1</sub>- A<sub>2b</sub> or, particularly, the A<sub>2a</sub> receptor wherein the compound of formula A1 has the structure, wherein, A represents Cy<sup>1</sup> or Het<sup>A</sup>; Cy<sup>1</sup> represents a 5- to 14-membered aromatic, fully saturated or partially unsaturated carbocyclic ring system comprising one, two or three rings, which Cy<sup>1</sup> group is optionally substituted by one or more R<sup>4a</sup> substituents; Het<sup>A</sup> represents a 5- to 14-membered heterocyclic group that may be aromatic, fully saturated or partially unsaturated, and which contains one or more heteroatoms selected from O, S and N, which heterocyclic group may comprise one, two or three rings and which Het<sup>A</sup> group is optionally substituted by one or more R<sup>4b</sup> substituents; B represents a Cy<sup>2</sup> or Het<sup>B</sup>; Cy<sup>2</sup> represents a 3- to 10-membered aromatic, fully saturated or partially unsaturated carbocyclic ring system comprising one or two rings, which Cy<sup>2</sup> group is optionally substituted by one or more R<sup>4c</sup> substituents; Het<sup>B</sup> represents a 3- to 10-membered heterocyclic group that may be aromatic, fully saturated or partially unsaturated, and which contains one or more heteroatoms selected from O, S and N, which heterocyclic group may comprise one or two rings and which Het<sup>B</sup> group is optionally substituted by one or more R<sup>4d</sup> substituents.



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