(57) Abstract: According to the invention, derivatives of pyrazolyl[1,2,4]triazolo[4,3-α]quinoxaline according to formula I or the analog in the form of pyrazolyl-substituted tetrazolo[1,5-α]quinoxalines according to formula II are indicated. The substances according to formula 1 or 2 are suitable in this form or in the form of their pharmaceutically compatible salts as active ingredients, in particular as adenosine receptor ligands.



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