

Abstract

A compound represented by the following general formula (I) or a pharmacologically acceptable salt thereof, wherein A represents a C3 to C12 cycloalkyl group which may be substituted by one to three selected from a fluoro group, a hydroxy group, a C1 to C6 alkyl group, etc; R^1 , R^2 , and R^3 each independently represent a hydrogen atom, a fluoro group, or a C1 to C6 alkyl group; R^4 represents a hydrogen atom or a prodrug group; and Y represents $-CH_2-CHR^5-CH_2-NHR^6$ (wherein R^5 represents a hydrogen atom, a C1 to C6 alkyl group, or a C1 to C6 alkoxy group, and R^6 represents a hydrogen atom or a prodrug group), or the like exhibits excellent TAFIa inhibitory activity and is useful as a therapeutic drug for myocardial infarction, angina pectoris, acute coronary syndrome, cerebral infarction, deep vein thrombosis, pulmonary embolism, and the like.



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