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Triheterocyclic Thiadiazepine Derivatives of Pharmacological Interest: Synthesis of 2-Aryl-4-[p-(2,3-dihydro-5-methyl-2-oxo-

 $1,\!3,\!4\text{-}oxadiazol\text{-}3\text{-}yl) phenyl]\text{-}6\text{-}(un) substituent\text{-}1\text{''},\!2\text{''},\!4\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{'},\!4\text{'-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{'},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-}b]\text{-}1\text{''},\!3\text{''-triazolo}[3\text{''},\!4\text{''-triazolo}[3\text{''},\,4\text{''-triazo$

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收稿日期 2004-12-8 修回日期 2005-8-1 网络版发布日期 接受日期

摘要 Synthesis of novel thiadiazepine derivatives based on the reactivity of 4-acetylphenylsydnone (1) was reported. The structures of the newly synthesized compounds have been proved by their physical, analytical, IR, ¹H NMR, ¹³C NMR and mass spectral data. 关键词 <u>4-acetylphenylsydnone</u> <u>1,3-dipolar cycloaddition</u> <u>1,3,4-thiadiazepine</u> <u>1,2,4-triazole</u>

1.3.4-oxadiazo-line triheterocycle ring closure condensation Michael addition cardiovascular agent

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1,3,4-oxadiazol-3-yl) phenyl]-6-(un) substituent-1",2",4"-triazolo[3",4"-b]-1',3',4'-thiadiaze pine

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Key words 4-acetylphenylsydnone 1 3-dipolar cycloaddition 1 3 4-thiadiazepine 1 2 triazole 1 3 4-oxadiazo-line triheterocycle ring closure condensation Michael addition cardiovascular agent

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