

缬沙坦类似物合成中间产物的捕获(II)

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Intermediate Captured in the Reaction of Synthesizing Valsartan Analogue(II)

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Abstract: In this article we mainly discuss the effect of anion of Lewis acid on the structure of the intermediate in synthesis of valsartan analogue. The crystal belongs to monoclinic system with space group $P2_1/c$, and $a=1.0999(3)$ nm, $b=1.9134(5)$ nm, $c=0.9778(3)$ nm, $\beta=104.806(4)^\circ$, $V=1.9895(9)$ nm³, $D_c=1.708$ g·cm⁻³, $Z=18$. CCDC: 629931.

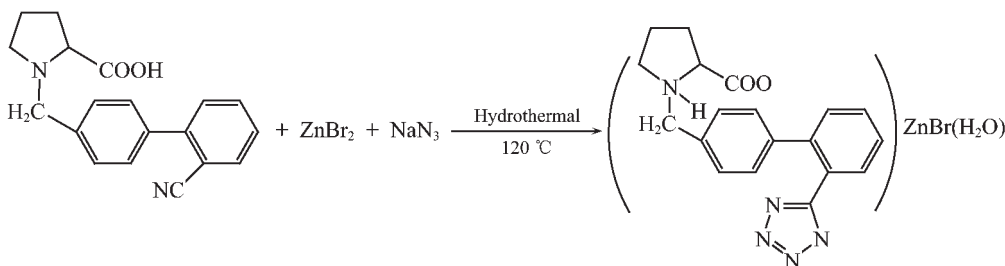
Key words: valsartan; Zn; tetrazole; hydrothermal synthesis

In order to investigate the effect of anion of Lewis acid on the structure and yield of intermediate^[1~10] produced by [2+3] cycloaddition reaction between cyano and azide, we have carried out the crystallographical characterization of the product obtained by *N*-(4-(2'-cyanophenyl)benzyl)proline reacting with ZnBr₂ as shown in Scheme I.

As depicted in Fig.1, the local coordination geometry around Zn center has a distorted tetrahedron

consisting of two N atoms, one O atom and a terminal Br. The *N*-(4-(2'-1H-tetrazol-5-yl-phenyl)benzyl)proline in-situ produced acts as a tridentate bridging binder to link two Zn centers to result in the formation of one-dimensional chain (Fig.2). The racemization occurred may be due to high temperature under hydrothermal conditions.

CCDC: 629931.



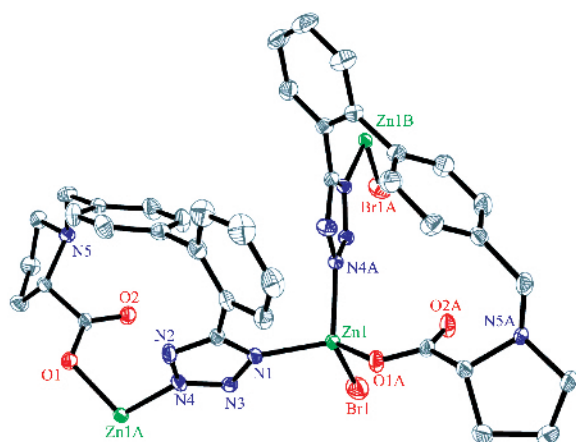
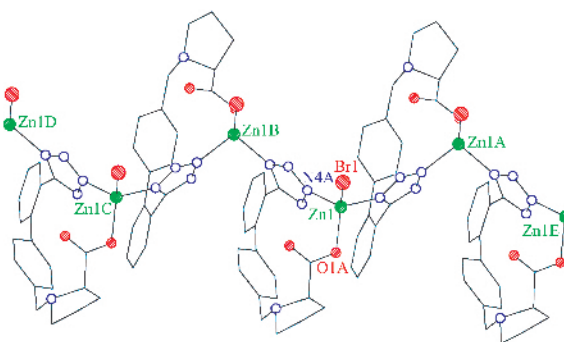
Scheme I

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Fig.1 Asymmetric unit of HTPBP-ZnBr(H₂O)Fig.2 1D chain structure of HTPBP-ZnBr(H₂O)

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