

[A0034]

Synthesis of Novel Acyclic Nucleoside and Acyclic Nucleoside Phosphonate Analogues of a Ring-Expanded ("Fat") Nucleobase Containing the Imidazo[4,5-*e*][1,3]diazepine Ring System

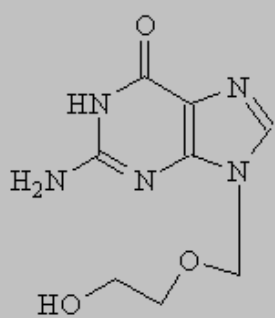
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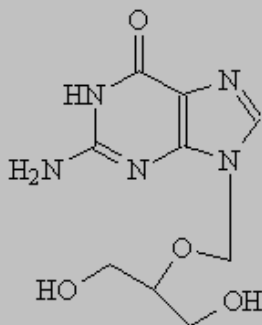
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Keywords: Organic Synthesis, Ring-Expanded ("Fat") Acyclic Nucleoside, Acyclic Nucleoside Phosphonate

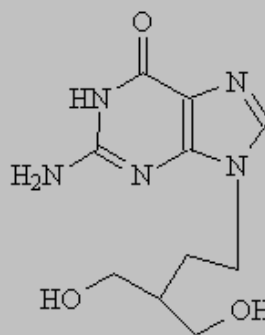
A number of acyclic nucleosides and nucleoside phosphonates have exhibited potent antiviral activities in recent years, and many of them have been approved by U.S. Food and Drug Administration (FDA) for a variety of viral infections. These include, but are not limited to, Acyclovir (**I**) for herpes simplex virus (HSV I and HSV II), Ganciclovir (**II**) for human cytomegalovirus (HCMV), Penciclovir (**III**) and Adefovir (PMEA) (**IV**) for hepatitis B virus (HBV). We have recently reported the synthesis and potent anti-HBV activity of a ring-expanded ("fat") nucleoside (**V**). We now report the synthesis of an acyclic nucleoside analogue (**VI**) as well as an acyclic nucleoside phosphonate analogue (**VII**) of the "fat" nucleoside **V**.



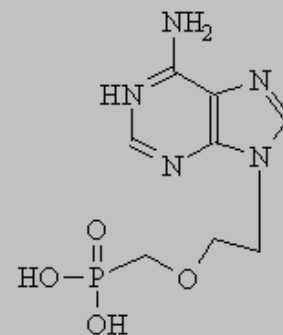
ACYCLOVIR (**I**)



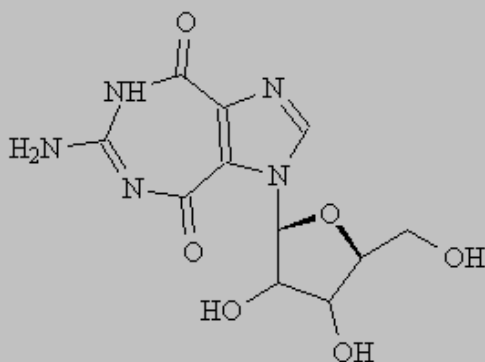
GANCICLOVIR (**II**)



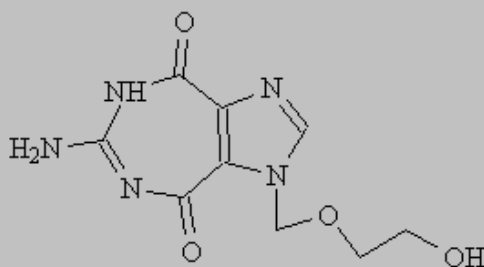
PENCICLOVIR (**III**)



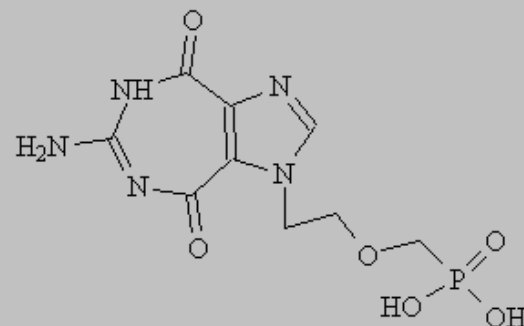
ADEFOVIR (**IV**)



(**V**)



(**VI**)



(**VII**)

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