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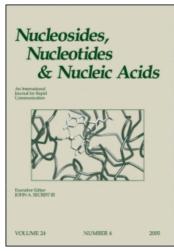
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Nucleosides, Nucleotides and Nucleic Acids

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Synthesis of Unusual Bicyclic Nucleosides Bearing an Unsaturated Side-Chain, as Potential Inhibitors of Varicella-Zoster Virus (VZV)

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Synthesis of Unusual Bicyclic Nucleosides Bearing an Unsaturated Side-Chain, as Potential Inhibitors of Varicella-Zoster Virus (VZV)

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We have previously reported a novel class of unusual bicyclic furano pyrimidine nucleosides bearing p-alkyl phenyl side-chains^[1] that showed an outstanding potency and highly selective activity against varicella-zoster virus (VZV). The p-pentyl phenyl analogue displayed an EC₅₀ value of 0.3 nM against the TK⁺ strain of VZV, representing an inhibitory activity 10,000 times more potent than that of acyclovir (ACV).

In an effort to further explore the structure-activity relationship in this series, we decided to introduce a more rigid side chain by inclusion of an alkene unsaturation

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to the α -phenyl unit. The substitution of the side-chain on the phenyl group at the *meta* position **1a**–**c** was carried out to probe the shape and size of the presumed lipophilic pocket of the target enzyme.

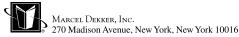
As previously reported by Robins and Barr,^[2] the target molecules were synthesized by the Pd(0) -catalyzed coupling of the alkenylphenyl iodide (5a–c) with 5-ethynyl-2'-deoxyuridine^[3] to yield 5-alkenylphenyl nucleosides which were finally cyclized in presence of CuI (1a–c). Compound 3 was prepared from the coupling of 5-iodo-2'-deoxyuridine with trimethylsilylacetylene, followed by desilylation. Alkenylphenyl iodide was the product of a two step reaction involving an Aldol-Grob^[3] reaction on the bromobenzaldehyde and a reverse halogen exchange.^[4]

i) HC=C-Si(Me)3, DIPEA, Pd(PPh3)4, Cul, DMF; ii) 10% NH3/H2O, MeOH; iii) CH3-(CH2)n-CO-(CH2)n-CH4, BF3-El2O, Hexane, r.t; iv) Kl, Cul, HMPA; v) DIPEA, Pd(PPh3)4, Cul, DMF; vi) Cul, Et3N, 80°C.

The target bicyclic compounds were evaluated for their ability to inhibit the replication of VZV in human embryonic lung (HEL) fibroblasts. The results showed a severe reduction of activity by more than 40,000-fold compared to the *p*-pentyl phenyl nucleoside. Thus, it is noticed the lipophilic pocket does not tolerate any alkenyl extension at the *meta* position of the aromatic system.

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