Inhibition of Herpes Simplex Virus Reactivation by Dipyridamole

RICHARD B. TENSER,* ANDREW GAYDOS, AND KATHLEEN A. HAY

Division of Neurology and Department of Microbiology and Immunology, The Pennsylvania State University College of Medicine, Hershey, Pennsylvania 17033

Received 16 April 2001/Returned for modification 12 July 2001/Accepted 27 August 2001

Herpes simplex virus (HSV) reactivation from latency was investigated. Reactivation of thymidine kinase-negative HSV, which is defective for reactivation, was greatly enhanced by thymidine (Tdr). The reactivation-enhancing effect of TdR was blocked by dipyridamole (DPM), a known nucleoside transport inhibitor. DPM also inhibited wild-type HSV reactivation, suggesting potential antiviral use.

In experimental models of herpes simplex virus (HSV) infection, expression of viral thymidine kinase (TK) has been shown to be important for viral latency. This was first suggested in studies of mice infected with TK-negative HSV. It was shown that in cases of acute infection, HSV replicated well in ocular tissues but not in trigeminal ganglia (TG) and that HSV reactivated poorly in ganglia during latency. Subsequently it was shown that establishment of latency was intact, i.e., latency-associated transcript (LAT) was readily detected in ganglion neurons, but reactivation was impaired (4, 5, 16). The defect of reactivation was explored in studies which showed that TK-negative HSV could in fact readily reactivate in ganglia if explant medium was supplemented with thymidine (Tdr) (17).

The present study extended this observation in three ways. First, it was shown that if a nucleoside transport inhibitor is added along with supplemental Tdr, the effect of Tdr on enhancing TK-negative HSV reactivation is blocked. This suggests the specific roles of Tdr and phosphorylation by TK in the reactivation process. Second, it is shown that the nucleoside transport inhibitor also blocks wild-type HSV reactivation from latency. Lastly, supplemental Tdr decreased the dipyridamole (DPM) block of wild-type HSV reactivation.

Latent infection of TG and lumbar dorsal root ganglia (DRG) was established in randomly bred CD-1 mice (Charles River Laboratories, Wilmington, Mass.) by standard methods. In brief, mice were anesthetized (methoxyflurane), and corneal inoculation (5 μl) or footpad inoculation (25 μl) was performed (17). Inoculation was performed with either TK-positive wild-type HSV type 1 (HSV-1; strain KOS, 5 × 10⁶ PFU/ml) or with mutant TK-negative HSV-1 (dlacTK, 4 × 10⁶ PFU/ml). The titers of the viruses were determined on Vero cell monolayers (17).

Inoculated mice were collected in accordance with institutional and federal guidelines. TG and DRG (from lumbar vertebrae 4 and 5) were removed and homogenized and tested for reactivated infectious HSV on Vero cell monolayers (17). Results are presented in terms of positive or negative for HSV reactivation.

Initially we investigated TK-negative HSV to supplement prior results which had demonstrated that although the dlsactk mutant virus reactivated poorly from latently infected ganglia, reactivation was greatly enhanced by the addition of Tdr to the explant medium. This was demonstrated for three different TK-negative mutants (17). In Table 1 it is shown that dlsactk reactivation did not occur when explant medium did not contain supplemental Tdr but that reactivation occurred at a frequency of 100% when explant medium contained supplemental Tdr (100 μM). In ganglia latently infected with TK-negative HSV, supplemental Tdr may have facilitated the synthesis of Tdr nucleotides by means of very low levels of viral TK that might have been present, but synthesis was more likely facilitated by means of cellular TK.

However, when the explant medium contained DPM in ad-

---

* Corresponding author. Mailing address: Division of Neurology, 500 University Dr., Hershey, PA 17033. Phone: (717) 531-8692. Fax: (717) 531-4694. E-mail: rtenser@psu.edu.

---

<table>
<thead>
<tr>
<th>Supplemental TdR in medium (concn)</th>
<th>No. of ganglia reactivated/no. tested (% positive) in medium with DPM at indicated concn (μM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>No</td>
<td>0/10 (0)</td>
</tr>
<tr>
<td>Yes (100 μM)</td>
<td>10/10 (100)</td>
</tr>
<tr>
<td></td>
<td>10/18 (56)</td>
</tr>
<tr>
<td></td>
<td>0/19 (0)</td>
</tr>
</tbody>
</table>

* ND, not done.
dition to 100 µM TdR, reactivation was inhibited (Table 1). DPM is a known nucleoside transport inhibitor (6, 11, 12). Blocking of the reactivation-enhancing effect of TdR, probably by inhibition of TdR transport into latently infected neurons, supports the existence of specific roles for TdR and phosphorylation by TK in facilitating HSV reactivation. This conclusion is supported by the observation that other nucleosides were minimally effective in enhancing reactivation of TK-negative HSV mutants (17).

With evidence that DPM blocked the effect of TdR on TK-defective HSV reactivation, we investigated the effect of DPM on reactivation of wild-type HSV in explant culture. Results with TG and DRG explanted in standard medium without supplemental TdR are shown in Table 2. Reactivation of HSV was inhibited in both tissues by DPM in a dose-dependent manner. Reactivation of HSV was inhibited in DRG somewhat more so than in TG. This was probably due to a greater HSV latency load in the latter, although it has been noted that HSV latency may otherwise differ somewhat between these tissues (13). Lastly, the effect of TdR on the DPM-mediated inhibition of wild-type HSV reactivation was evaluated. Supplemental TdR in the explant medium partially reversed the blocking effect of DPM (Table 3). Nucleoside transport in mammalian cells is mediated by multiple transport mechanisms, and some transporters are less sensitive to DPM than are others (2, 9). Excess TdR apparently circumvented the inhibition of reactivation by DPM, although specific mechanisms of inhibition remain to be determined.

DPM has been used occasionally in antiviral studies (18), including investigations of HSV (14). In the latter study, it was not shown to be a potent antiviral. However, that study investigated the effect of DPM on HSV replication in cell culture, a situation which differs markedly from reactivation from latency. First, the molecular state of the virus differs, and second, the amount of virus present probably differs. It is suggested that DPM may be effective in blocking HSV reactivation from latency because during reactivation the HSV genome is in a particularly vulnerable state or perhaps simply because only a small amount of virus is present.

The possibility of a toxic effect of DPM on latently infected neurons as a means of explaining inhibition of HSV reactiva-
tion cannot be completely excluded. This was investigated in part in a viral growth study (Fig. 1). Only a slight antiviral effect of DPM in a dose-dependent pattern was noted, and significant cellular toxicity was unlikely. In addition, the 25 μM DPM concentration used was similar to that used in other HSV studies, where there was no apparent cellular toxicity with 20 μM DPM (14). It does remain possible that DPM is particularly toxic to neurons and that destruction of neurons led to reactivation in ganglia.

However, the results in Table 3 show that reactivatable virus was present, albeit when supplemental TdR was added, indicating that at least some latently infected neurons survived DPM treatment. Lastly, DPM has been clinically used as an antiplatelet agent (6, 11, 12) and neurotoxicity has not been noted.

Although it is a nucleoside transport inhibitor, DPM has been shown to not inhibit transport into cells of nucleoside analogue antivirals such as acyclovir (8), zidovudine (1, 3), and lamivudine (3, 10). These observations and the reported DPM inhibition of transport of nucleosides such as TdR and deoxycytidine, which compete with the antivirals for kinase-mediated phosphorylation, have suggested mechanisms by which DPM may potentiate the antiviral effect of the dideoxynucleoside drugs (15).

There is widespread clinical experience with DPM as an antiplatelet agent; the mechanism of action may be inhibition of nucleoside transport, particularly of adenosine transport, into platelets (6, 11, 12). The antiviral results obtained in the present study suggest that DPM may also be clinically useful for the inhibition of HSV reactivation from latency.

This work was supported by National Institutes of Health grant NS20684 to Richard B. Tenser.

The secretarial assistance of Tracy Monette is gratefully acknowledged.

REFERENCES