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Abstract
Oral and perioral herpes simplex virus (HSV) infections in healthy individuals often present with signs and symptoms that are clearly recognized by oral health care providers (OHCPs). Management of these infections is dependent upon a variety of factors and several agents may be used for treatment to accelerate healing and decrease symptoms associated with lesions. This article will review the pertinent aspects of topical and systemic therapies of HSV infections for the OHCP.

Disciplines
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Herpes Simplex Virus

Topical and Systemic Therapies for Oral and Perioral Herpes Simplex Virus Infections

ERIC T. STOOPLER, DMD, FDS RCS ED, AND RAMESH BALASUBRAMANIAM, BDSC, MS

Abstract

Oral and perioral herpes simplex virus (HSV) infections in healthy individuals often present with signs and symptoms that are clearly recognized by oral health care providers (OHCPs). Management of these infections is dependent upon a variety of factors and several agents may be used for treatment to accelerate healing and decrease symptoms associated with lesions. This article will review the pertinent aspects of topical and systemic therapies of HSV infections for the OHCP.

Authors

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Oral and perioral (herein referred to collectively as oral) herpes simplex virus (HSV) infections represent one of the most common oral soft tissue disease processes encountered in the general population. Oral HSV-1 serotype is the most common cause of orofacial infections, however, HSV-2 serotype has been implicated as a causative agent of these infections (and HSV-1 as the etiology for genital infections) due to sexual practices. Primary oral HSV infections usually occur in early childhood and while the majority are subclinical, clinical infections initially present with general symptoms, such as malaise, fever and lymphadenopathy (referred to as a prodrome) followed by vesicles and/or ulcers affecting a variety of intraoral surfaces. Most primary oral HSV infections are readily diagnosed based on clinical history, signs and symptoms and further laboratory investigation is generally not warranted. The majority of oral HSV infections are self-limiting with resolution usually within two weeks, often requiring only palliative treatment and supportive care as needed.

Following primary infection, the virus migrates to the trigeminal nerve ganglion where it can remain latent indefinitely but may be stimulated to reactivate under a variety of circumstances (environmental triggers, stress, illness, etc.) that results in clinical infection. The most common presentation of recrudescent HSV infection (development of clinical lesions) in healthy individuals is recurrent herpes labialis (RHL), observed as a lesion located...
Topical antiviral agents have demonstrated efficacy in accelerating the healing time of RHL lesions, especially if administered during the prodromal phase. The topical antiviral agents that are most commonly recommended to treat RHL include Acyclovir 5 percent cream, Penciclovir 1 percent cream and Docosanol 10 percent cream. Acyclovir is a nucleoside analogue of guanosine with a selective affinity for thymidine kinase (TK), which is necessary for activation of acyclovir, in virus-infected cells. Penciclovir is an acyclic guanine derivative with a similar antiviral spectrum as acyclovir. It is also phosphorylated by viral TK and inhibits viral DNA polymerase. Penciclovir has approximately 1/100th the potency of acyclovir, but is an effective antiviral agent due to its long half-life and high intracellular concentrations. Docosanol is a 22-carbon primary alcohol that blocks the virus from attaching to cells via interference of epithelial cell surface receptors and viral envelope proteins. Acyclovir 5 percent cream and Penciclovir 1 percent cream are available by prescription, while Docosanol is the only agent approved by the FDA as an OTC product for treatment of RHL.

Topical formuations of foscarnet, cidofovir and imiquimod are generally reserved for treatment of RHL lesions that are nonresponsive to typical antiviral agents and are rarely used in healthy individuals. In contrast to other antiviral agents dependent upon viral TK, foscarnet and cidofovir inhibit viral DNA synthesis independently of this mechanism. Foscarnet has demonstrated efficacy in treating acyclovir-resistant HSV infections, while cidofovir is generally reserved for both acyclovir and foscarnet-resistant HSV infections.
is a novel agent that enhances innate immunologic responses to viruses and topical formulations has shown to be effective in treating resistant HSV infection in the setting of HIV.15

Table 1 outlines the indications and usage recommendations for topical agents used for treatment of oral HSV infections.

### Systemic Therapies

Systemic therapies may be required for the treatment of primary oral HSV infection and treatment or prophylaxis of both RHL and RIH, especially in immunocompromised patients. Unlike topical agents, systemic medications enable greater drug exposure, rapid access to site of viral replication, better biocompatibility and less frequent dosing and improved compliance. Systemic medications are exclusively antiviral agents and may be administered orally or intravenously.8,27

As noted previously, treatment of primary oral HSV infection is typically based on supportive and symptomatic interventions.44 However, off-label use of systemic antiviral medications may accelerate healing time of primary oral HSV lesions by inhibiting DNA replication of infected cells if commenced when prodromal symptoms are recognized or within one day of vesicle eruption.4 Oral acyclovir 200 mg five times a day or 400 mg three times a day for 10 days may be used in severe cases of primary oral HSV infection in adults as currently prescribed in primary genital infection.4 In the pediatric patient, treatment with oral acyclovir suspension 15 mg/kg within three days of symptom onset and continued five times a day for one week was shown to accelerate healing, reduce viral shedding and improve oral intake.99 Contemporary antiviral medications such as famciclovir and valacyclovir may also be prescribed given their more convenient dosing and increased bioavailability.44 (Table 2) Famciclovir (prodrug of penciclovir) is a diacetyl-6-deoxy analogue that is rapidly absorbed and undergoes deacetylation in the gastrointestinal tract, blood and liver to its active form.1 Valacyclovir (prodrug of acyclovir) is an L-valine ester that is well absorbed and 99 percent converted to its active form in the gastrointestinal tract and liver.1 This results in a three- to five-times increase in bioavailability.44

Systemic antiviral medications may be used as prophylaxis or treatment in patients with severe, frequent, persistent and unsightly outbreaks.20 Oral valacyclovir has been shown to be effective and is approved by the FDA for the treatment of RHL.21 Oral acyclovir and famiciclovir are approved by the FDA specifically for the treatment and suppression of genital herpes, but have also been used for RHL therapy.13,21

In the immunocompromised individual, such as during chemotherapy or during
the use of immunosuppressive drugs, RHI may present as a severe outbreak.\textsuperscript{23} Oral or intravenous acyclovir has been shown to be effective in the prevention and treatment of RII in these patients.\textsuperscript{23} Similarly, valacyclovir and famciclovir may also be prescribed for the prevention and treatment of RII in immunocompromised patients. TABLE 3 summarizes the antiviral agents available, their dosages and duration of use based on the expert recommendations from the Fourth World Workshop in Oral Medicine.\textsuperscript{21} Never intravenous medications such as foscarnet and cidofovir may be necessary in acyclovir-resistant, severely immunocompromised patients. These medications are highly nephrotoxic and should be used with caution.\textsuperscript{21}

**TABLE 2**

<table>
<thead>
<tr>
<th>Systemic Antiviral Medications for the Treatment of Primary Herpes Simplex Virus Infection</th>
</tr>
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<tbody>
<tr>
<td>Acyclovir</td>
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<tr>
<td><strong>Dose</strong></td>
</tr>
<tr>
<td><strong>Frequency</strong></td>
</tr>
<tr>
<td><strong>Duration</strong></td>
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</tbody>
</table>

*Food and Drug Administration treatment recommendations for genital herpes
+ Recommendations from the Center for Disease Control and Prevention for genital herpes

**TABLE 3**

<table>
<thead>
<tr>
<th>Systemic Therapies for Treatment of Oral HSV Infections</th>
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</thead>
<tbody>
<tr>
<td><strong>Indication</strong></td>
</tr>
<tr>
<td>Treatment of RHL in the immunocompetent host</td>
</tr>
<tr>
<td>Prophylaxis of RHL in the immunocompetent host *</td>
</tr>
<tr>
<td>Treatment of recurrent HSV infections in the immunocompromised host</td>
</tr>
<tr>
<td>Prophylaxis of recurrent HSV infections in the immunocompromised host</td>
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</tbody>
</table>

*Duration of the prophylaxis is based on the extent and frequency of exposure to triggers of RHL episodes, such as sunlight, dental treatment, etc.

**Conclusions**

There is a variety of treatment modalities for oral HSV infections. OHCPs must be cognizant of the advantages and limitations of both topical and systemic therapies for this condition. It is imperative for OHCPs to determine the appropriate agents for treatment in the context of the patient’s disease presentation and overall medical status.

**REFERENCES**