## Conjunctivitis Doesn't Always Need an Antibiotic

#### BY MICHELE G. SULLIVAN Mid-Atlantic Bureau

ost healthy children with conjunctivitis will get better by themselves and don't need an ophthalmic antibiotic, Peter W. Rose, M.B., and his colleagues reported.

"Parents should be encouraged to treat children themselves without medical consultation, unless their child develops unusual symptoms or the symptoms persist for more than a week," said Dr. Rose of Oxford (England) University, and his associates. They suggested that parents cleanse their children's eyes with lubricating eyedrops instead of rushing them off to the pediatrician at the first sign of

## Metapneumovirus Is Unseen Culprit In Bronchiolitis

Human metapneumovirus may be underreported as a pathogen in bronchiolitis and may lead to admittance to intensive care, especially when it infects infants in combination with human respiratory syncytial virus, reported Malcolm G. Semple, M.D., of the University of Liverpool (England), and his associates.

During the 2001-2002 winter season at one hospital, dual infection human metapneumovirus (hMPV) and human respiratory syncytial virus (hRSV) occurred at a significantly higher rate in infants with bronchiolitis who were admitted to the pediatric intensive care unit on mechanical ventilation (72%, 18 of 25) than in infants with bronchiolitis who were sent to the general wards (10%, 15 of 171). The investigators said that the temporal distribution of hMPV infections in infants in the pediatric ICU made it unlikely that the infections were nosocomial. In a subset of infants with complete clinical information, dual infection with hMPV and hRSV was not statistically significantly associated with disease severity in the retrospective study (J. Infect. Dis. 2005;191:382-6).

In nasopharyngeal aspirate and bronchoalveolar lavage samples that were taken at the same time from nine hMPV-infected infants on mechanical ventilation, hMPV was detected in only one nasopharyngeal aspirate and in all nine bronchoalveolar lavages. Of 18 infants who had mechanical ventilation, hMPV infection was found in bronchoalveolar lavages from 15 infants and in nasopharyngeal aspirates from 4 infants.

The discordance in the incidence of hMPV infection detected in bronchoalveolar lavages and nasopharyngeal aspirates raises the possibility that "hMPV infection during endemic seasons may be more common than is currently recognized and that it has been undetected because sampling from the lower respiratory tract is not possible on infants who do not require mechanical ventilation," Dr. Semple wrote. conjunctivitis (Lancet 2005;366:37-43).

The investigators randomized 326 children (mean age 3.3 years) with a clinical diagnosis of conjunctivitis to either chloramphenicol eyedrops (0.5%) or placebo (distilled water containing 1.5% boric acid and 0.3% borax). Parents applied the drops every 2 hours for the first 24 hours when the child was awake and four times a day until 48 hours after symptoms resolved.

After 7 days, 86% of those in the antibiotic group were clinically cured, compared with 83% of those in the placebo group. When 307 of the children were followed up at 6 weeks, fewer than 5% in each group had experienced a relapse or new infection. Only one reaction—a case of swollen eyelids and face—was attributed to antibiotic treatment.

Baseline cultures showed that 80% of the children had bacterial infections. Among this group, the clinical cure rate did not differ significantly between chloramphenicol and placebo (85% vs. 80%), but more of the chloramphenicol group than the placebo group experienced bacterial eradication (40% vs. 23%).

Although eradication is not necessary for a clinical cure, Dr. Rose and his associates said failure to achieve it could impact transmission. "Despite our results, antibiotic treatment might still reduce the absolute number, and, hence, transmissibility of pathogens, and further research might be necessary if antibiotics cease to be prescribed for this disorder."

# Change is in the Air

### Introducing ATROVENT<sup>®</sup> HFA:

The same medication, now in a different formulation

- ATROVENT\* (ipratropium bromide) Inhalation Aerosol (CFC) will not be available after December 2005
  ATROVENT\* HFA (ipratropium bromide HFA) Inhalation Aerosol has demonstrated comparable safety and efficacy to the CFC formulation
- Re-evaluate your patients on ATROVENT<sup>®</sup> Inhalation Aerosol (CFC) at their next office visit and consider the benefits of switching to SPIRIVA<sup>®</sup> HandiHaler<sup>®</sup>
- For patients switching from ATROVENT® Inhalation Aerosol (CFC) to ATROVENT® HFA, a new prescription specifying "ATROVENT® HFA" will be necessary
- ATROVENT® HFA (ipratropium bromide HFA) Inhalation Aerosol is indicated as a bronchodilator for maintenance treatment of bronchospasm associated with chronic obstructive pulmonary disease, including chronic bronchitis and emphysema, but is not indicated for the initial treatment of acute episodes of bronchospasm where rescue therapy is required for rapid response
- ATROVENT® HFA (ipratropium bromide HFA) Inhalation Aerosol is contraindicated in patients with a history of hypersensitivity to ipratropium bromide or other ATROVENT® HFA Inhalation Aerosol components. Immediate hypersensitivity reactions may occur after administration of ipratropium bromide, as demonstrated by rare cases of urticaria, angioedema, rash, bronchospasm, anaphylaxis and oropharyngeal edema. ATROVENT® HFA Inhalation Aerosol is also contraindicated in patients who are hypersensitive to atropine or its derivatives
- Inhaled medicines, including ATROVENT® HFA (ipratropium bromide HFA) Inhalation Aerosol, may cause paradoxical bronchospasm. If this occurs, treatment with ATROVENT® HFA (ipratropium bromide HFA) Inhalation Aerosol should be stopped and other treatments considered
- ATROVENT® HFA (ipratropium bromide HFA) Inhalation Aerosol should be used with caution in patients with narrow-angle glaucoma, prostatic hypertrophy or bladder-neck obstruction. Patients should avoid spraying the aerosol into their eyes. This may result in precipitation or worsening of narrow-angle glaucoma, mydriasis, eye pain or discomfort, temporary blurring of vision, visual halos or colored images in association with red eyes from conjunctival and corneal congestion. Should any combination of these symptoms develop, patients should consult their physician immediately
- The most common drug-related adverse events were dry mouth (1.6% of ATROVENT® HFA Inhalation Aerosol and 0.9% of ATROVENT® Inhalation Aerosol CFC patients), and taste perversion (bitter taste) (0.9% of ATROVENT® HFA Inhalation Aerosol and 0.3% of ATROVENT® Inhalation Aerosol CFC patients)
- SPIRIVA HandiHaler is intended as a once-daily maintenance treatment for COPD and is not indicated for the initial treatment of acute episodes of bronchospasm, i.e., rescue therapy. SPIRIVA HandiHaler is contraindicated in patients with a history of hypersensitivity to atropine or its derivatives, i.e., ipratropium or to any component of this product. The most commonly reported adverse drug reaction was dry mouth, which was usually mild and often resolved during continued treatment. Other reactions reported in individual patients and consistent with possible anticholinergic effects included constipation, increased heart rate, blurred vision, glaucoma, urinary difficulty and urinary retention. Inhaled medicines, including SPIRIVA, may cause paradoxical bronchospasm. If this occurs, treatment with SPIRIVA should be stopped and other treatments considered
  Please see Brief Summary of Prescribing Information for ATROVENT\* HFA Inhalation Aerosol and SPIRIVA HandiHaler on pages following this advertisement.





Boehringer Ingelheim

Copyright © 2005, Boehringer Ingelheim Pharmaceuticals, Inc. All rights reserved.

Printed in U.S.A.

(6/05)

AT-10286

—Jeff Evans