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Are You Doing All You Can to Stop Methicillin-Resistant



BACTROBAN® NASAL (mupirocin calcium ointment), 2% Brief summary. For complete prescribing information, see package insert.

INDICATIONS AND USAGE

NDICATIONS AND USAGE Bactroban Nasal is indicated for eradication of nasal coloniza-tion with methicillin-resistant Staphylococcus aureus in adult patients and health care workers as part of a comprehensive infection control program to reduce the risk of infection among patients at high risk of methicillin-resistant *S. aureus* infection during institutional outbreaks of infections with this nathoraen pathogen.

- There are insufficient data at this time to establish that this product is safe and effective as part of an intervention program to prevent autoinfection of high-risk patients from their own nasal colonization with *S. aureus.*
- There are insufficient data at this time to recommend use (2)of Bactroban Nasal for general prophylaxis of any infec-tion in any patient population.
- tion in any patient population. Greater than 90% of subjects/patients in clinical trials had eradication of nasal colonization 2 to 4 days after therapy was completed. Approximately 30% recolonization was reported in one domestic study within 4 weeks after com-pletion of therapy. These eradication rates were clinically and statistically superior to those reported in subjects/ patients in the vehicle-treated arms of the adequate and well-controlled studies. Those treated with vehicle had eradication rates of 5% to 30% at 2 to 4 days post-thera-py with 85% to 100% recolonization within 4 weeks. INTERINDEATIONS (3)

CONTRAINDICATIONS

Bactroban Nasal is contraindicated in patients with known hypersensitivity to any of the constituents of the product WARNINGS

AVOID CONTACT WITH THE EYES. Application of Bactroban Nasal to the eye under testing conditions has caused severe symptoms such as burning and tearing. These symptoms resolved within days to weeks after discontinuation of the ointment.

In the event of a sensitization or severe local irritation from *Bactroban* Nasal, usage should be discontinued. PRECAUTIONS

PRECAULTIONS General: As with other antibacterial products, prolonged use may result in overgrowth of nonsusceptible microorganisms, including fungi. (See **DOSAGE AND ADMINISTRATION** in complete prescribing information.)

complete prescripting information.) Information for Patients: Patients should: apply approxi-mately one-half of the ointment from the single-use tube directly into one nostril and the other half into the other nos-tril; avoid contact of the medication with the eyes; discard the tube after using; press the sides of the nose together and gent-ly massage after application to spread the ointment through-out the inside of the nostrils; and discontinue using Bactroban Nasal and call a health care practitioner if sensitization or severa local irritation occurs. severe local irritation occurs.

Drug Interactions: The effect of the concurrent application of intranasal mupirocin calcium and other intranasal products has not been studied. Do not apply mupirocin calcium ointment, 2% concurrently with any other intranasal products.

2% concurrently with any other intranasal products. Carcinogenesis, Mutagenesis, Impairment of Fertility: Long-term studies in animals to evaluate carcinogenic poten-tial of mupirocin calcium have not been conducted. Results of the following studies performed with mupirocin calcium or mupirocin sodium in vitro and in vivo did not indicate a poten-tial for mutagenicity: rat primary hepatocyte unscheduled DNA synthesis, sediment analysis for DNA strand breaks, Salmo-nella reversion test (Ames), Escherichia coli mutation assay, metaphase analysis of human lymphocytes, mouse lym-phoma assay, and bone marrow micronuclei assay in mice. Aeproduction studies were performed in rats with mupirocin administered subcutaneously at doses up to 40 times the human intranasal dose (approximately 20 mg mupirocin per day) on a mg/m² basis and revealed no evidence of impaired fertility from mupirocin sediem.

fertility from mupirocin sodium. **Pregnancy: Teratogenic Effects. Pregnancy Category B.** Reproduction studies have been performed in rats and rabbits with mupirocin administered subcutaneously at doses up to 65 and 130 times, respectively, the human intranasal dose (approximately 20 mg mupirocin per day) on a mg/m² basis and revealed no evidence of harm to the fetus due to mupiro-cin. There are, however, no adequate and well-controlled stud-ies in pregnant women. Because animal reproduction studies are not always predictive of human response, this drug should be used during pregnancy only if clearly needed. **Nursing Mothers:** It is out hnown whether this drug storet.

Nursing Mothers: It is not known whether this drug is excret-ed in human milk. Because many drugs are excreted in human milk, exercise caution when Bactroban Nasal is administered to a nursing woman.

Pediatric Use: Safety in children under the age of 12 years has not been established. (See CLINICAL PHARMACOLOGY in complete prescribing information.)

ADVERSE REACTIONS

ADVERSE REACTIONS Clinical Trials: In clinical trials, 210 domestic and 2,130 foreign adult subjects/patients received Bactroban Nasal oint-ment. Less than 1% of domestic or foreign subjects and patients in clinical trials were withdrawn due to adverse events. In domestic clinical trials, 17% (36/210) of adults treat-ed with Bactroban Nasal ointmem reported adverse events thought to be at least possibly drug-related. The incidence of adverse events that were reported in at least 1% of adults enrolled in domestic clinical trials were as follows: headache, 9%; rhinitis, 6%; respiratory disorder; including upper respira-tory tract congestion, 5%; pharyngitis, 4%; taste perversion, 3%; burning/stinging, 2%; cough, 2%; and pruritus, 1%.

The following events though possibly drug-related were reported in less than 1% of adults enrolled in domestic clinical trials: blepharitis, diarthea, dry mouth, ear pain, epistaxis, nau-sea and rash. All adequate and well-controlled clinical trials have been performed using Bactroban Nasal ointment, 2% in one arm and the vehicle ointment in the other arm of the study.

OVERDOSAGE

Following single or repeated intranasal applications of Bactroban Nasal to adults, no evidence for systemic absorption of mupirocin was obtained.

Manufactured by DPT Laboratories, San Antonio, TX 78215 Distributed by SmithKline Beecham Pharmaceuticals, Philadelphia, PA 19101 BRS-BN:L3

INFECTIOUS DISEASES INFECTION CONTROL



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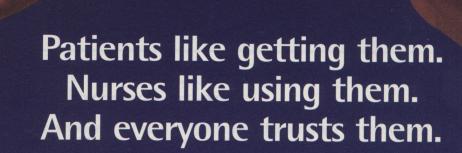
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Curriculum vitae and the names of three references should be sent to James C. Torner, Ph.D., Head, Department of Epidemiology, College of Public Health, 2800 Steindler Building, University of Iowa, Iowa City, Iowa 52242. Information on the College of Public Health and the Department of Epidemiology can be seen at www.publichealthmeh.uiowa.edu.

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INFECTION CONTROL AND HOSPITAL EPIDEMIOLOGY^{**}

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All The Facts For Your Immunization Program



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