

HOSPITAL EPIDEMIOLOGY

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EDITORIAL

From Ritual to Reason and Back Again: OSHA and the Evolution of Infection Control

William M. Valenti, MD

SPECIAL ARTICLE

Infection Control in Latin America

Donald A. Goldmann, MD; Fernando Otaiza, MD; Samuel R. Ponce de Leon, MD, MSc; Idis Faingezicht Gutman, MD

ORIGINAL ARTICLES

Nosocomial Respiratory Tract Colonization and Infection with Aminoglycoside-Resistant Acinetobacter calcoaceticus var antitratus: Epidemiologic Characteristics and Clinical Significance

James E. Peacock, Jr., MD; LuAnne Sorrell, RN, CIC; Frank D. Sottile, MD; Loraine E. Price, BSN, CIC; William A. Rutala, PhD, MPH

Description of Case-Mix Adjusters by the Severity of Illness Working Group of The Society of Hospital Epidemiologists of America (SHEA)

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Introduction of a Plasmid Encoding the OHIO-1 b -Lactamase to an Intermediate Care Ward by Patient Transfer

David M. Shlaes, MD, PhD; Charlotte A. Currie-McCumber; Mary-Helen Lehman

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What is it?

Spread by blood or sexual contact.

300,000 new cases will occur in the U.S. this year.

Hemophiliacs, Asian immigrants, heterosexuals with multiple partners, male homosexuals, IV drug users, and health-care personnel are at highest risk.



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RECOMBIVAX HB*
[Hepatitis B Vaccine (Recombinant), MSD]

INDICATIONS AND USAGE

INDICATIONS AND USAGE
RECOMBINAX Hill is indicated for immunization against infection caused by all known suboypes of nepatitis Bivrus.
RECOMBINAX Hill will not prevent hepatitis Caused by other agents, such as hepatitis A vitus.
Caused by other agents, such as hepatitis A vitus on-A non-B hepatitis Writes, or other viruses known to infect the liver.
Vaccination is recommended in persons of all ager who are or will be at increased risk of infection with hepatitis B vitus in areas with high prevalence of infection, most of the population are arrisk of acquiring hepatitis B infection at ayoung age. Therefore, vaccination should be limited to those who are in groups identified as being at ingreased risk of infection.

CONTRAINDICATIONS

Hypersensitivity to yeast or any component of the vaccine.

WARNINGS

Patients who develop symptoms suggestive of hypersensitivity after an injection should not receive further injections of RECOMBIVAX HB (see CONTRAINDICATIONS).

RECOMBIVAX HB* [Hepatitis B Vaccine (Recombinant), MSD]

Because of the long incubation period for hepatitis B, it is possible for unrecognized infection to be present at the time RECOMBIVAX HB is given RECOMBIVAX HB may not prevent hepatitis B in such patients.

PRECAUTIONS

General

As with any percutaneous vaccine, epinephrine should be available for immediate use should an anaphylactoid reaction occur.

Any serious active infection is reason for delaying use of RECOMBIVAX HB except when, in the opinion of the physician, withholding the vaccine entails a greater risk.

Caution and appropriate care should be exercised in administering RECOMBINAY HB to individuals with severely compromised cardiopulmonary status of to others in whom a febrile or systemic reaction could pose a significant risk.

Pregnancy

<u>Pregnancy Category C.</u> Animal reproduction studies have not been conducted with RECOMBIVAX HB. It is also not known whether RECOMBIVAX HB can cause fetal narm when administered to a pregnant woman or can affect

RECOMBIVAX HB®
[Hepatitis B Vaccine [Recombinant], MSD]

reproduction capacity RECOMBIVAX HB should be given to a pregnant woman only if clearly needed.

Nursing Mothers

It is not known whether RECOMBIVAX HB is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when RECOMBIVAX HB is administered to a nursing woman.

Pediatric Use

RECOMBIVAX HB has been shown to be usually RECOMBIVAX HB has been shown to be usual well tolerated and highly immunogenic in infants and children of all ages. Newborns also respond well-maternally transferred antibodies do not interfere with the active immune response to the vaccine. See DOSAGE AND ADMINISTRATION for recommended pediatric dosage and for recommended obasge for infants born to HBsAg positive mothers.

ADVERSE REACTIONS

RECOMBINAX HB is generally well tolerated. No serious adverse reactions attributable to the vaccine have been reported during the course of clinical trials. No serious hypersensitivity reactions have been reported. No adverse experiences

RECOMBIVAX HB® (Hepatitis B Vaccine (Recombinant), MSD)

were reported during clinical trials which could be related to changes in the titers of antibodies to yeast. As with any vaccine, there is the possibility that broad use of the vaccine could reveal adverse reactions not observed in clinical trials. In a group of studies, 3258 doses of vaccine were administered to 1525 healthy adults who were monitored for 5 days after each dose ligication site and systemic complaints were reported following 17% and 15% of the injections, respectively.

The following adverse reactions were reported

Incidence Equal to or Greater than 1% of Injections

LOCAL REACTION (INJECTION SITE)

Injection site reactions consisting principally of soreness and including pain, tenderness, pruritus, erythema, ecchymosis, swelling, warmth, and nodule formation

BODY AS A WHOLE

The most frequent systemic complaints include fatigue/weakness; headache, fever (≥100°F); malaise.

DIGESTIVE SYSTEM Nausea: diarrhea

HepatitisB

It can be prevented.

A vaccine is available.

Help eliminate the risk.... Recombivax HB (Hepatitis B Vaccine [Recombinant] | MSD)

RECOMBIVAX HB is contraindicated in the presence of hypersensitivity to yeast or any other component of the vaccine.

See below for a Brief Summary of Prescribing Information for RECOMBIVAX HB.

RECOMBIVAX HB® [Hepatitis B Vaccine [Recombinant], MSD]

RESPIRATORY SYSTEM

Pharyngitis; upper respiratory infection. Incidence Less than 1% of Injections

BODY AS A WHOLE Sweating; achiness; sensation of warmth; lightheadedness, chills; flushing.

DIGESTIVE SYSTEM Vomiting; abdomina. pains/cramps; dyspepsia; diminished appetite.

RESPIRATORY SYSTEM Rhinitis, influenza: cough

NERVOUS SYSTEM Vertigo/dizziness; paresthesia

INTEGUMENTARY SYSTEM Pruntus; rash (non-specified); angioedema; urticana.

MUSCULOSKELETAL SYSTEM Arthralgia including monoarticular; myalgia; back pain; neck pain; shoulder pain; neck stiffness. HEMIC/LYMPHATIC SYSTEM

PSYCHIATRIC/BEHAVIORAL

RECOMBIVAX HB® [Hepatitis 8 Vaccine (Recombina inanti. MSD1

SPECIAL SENSES Earache

UROGENITAL SYSTEM Dysuna

CARDIOVASCULAR SYSTEM

Potential ADVERSE EFFECTS

Potential ADVERSE EFFECTS
In addition, a variety of adverse effects, not observed inclinicatinals with RECOMBIVAXHB, have been reported with HEPTWAXH 8* (Hepatitis B Vaccine). MSD) (plasma derived nepatitis B vaccine). Those listed below are to serve as alerting information to physicians:

Hypersensitivity. An apparent hypersensitivity syndrome of delayed onset has been reported days to weeks after vaccination. This has included the following findings: arthritis (iusually transent), fever, and demarkologic reactions such as utilicanal erythems multiforme, or ecohymoses.

Nervous System: Neurological disorders such as optic neuritis, myelitis including transverse myellis, acute radiculoneuropathy including Guillain-Barré syndrome; penpheral neuropathy including Bell's palsy and herpes zoster.

RECOMBIVAX HB® [Hepatitis 8 Vaccine (Recombinant), MSD]

Special Senses; Tinnitus; visual disturbances.

DOSAGE AND ADMINISTRATION

Do not inject intravenously or intradermally.

Bo not inject intravenously of intradermany, RECOMBIVAX HB is for intramuscular injection. The <u>detaid muscle</u> is the preferred site for intramuscular injection in adults. Data suggest that injections given in the buttock's frequently are given into fatty issue instead of into muscle. Such injections have resulted in a lower seroconversion rate than was expected. The <u>anterolateral thigh</u> is the recommended site for intramuscular injection in infants and young children.

RECOMBIVAX HB may be administered subcutaneously to persons at risk of hermorrhage following intramuscular injections. However, when other aluminum-adsorbed vaccines have been administered subcutaneously, an incread incidence of local reactions including subcutane-us nodules has been observed. Therefore, subcutaneous administration should be used only in persons [e.g., hermophilacs] at risk of hemorrhage following intramuscular injections.

The immunization regimen consists of 3 doses of vaccine. The volume of vaccine to be given on each occasion is as follows:

RECOMBIVAX HB® [Hepatitis 8 Vaccine [Recombinant], MSD]

Formulation Initial I month 6 month: 05mL 0.5mL 0.5ml Adults and Older Children Adult 10mL 10mL 10mL 10mL 10mL

Whenever revaccination or administration of a booster dose is appropriate, RECOMBIVAX HB

booster dose is appropriate, RECOMBIVAX HB may be used. For dosage for infants born of HBsAg positive montrers and for dosage for known or presumed exposure to HBsAg, see the Prescribing Information. The vaccine should be used as supplied; no dilution or reconstitution is necessary. The full recommended dose of the vaccine should be used.

Store vials at 2–8°C [35.6–46.4°F]. Storage above or below the recommended temperature may reduce potency.

Do not freeze since freezing destroys potency.

For more detailed information, consult your MSD Representative or see Prescribing Information. Merck Sharp & Dohme, Division of Merck & Co., INC., West Point, PA 19486. J7RX04 (201)