

IMMUNOSCAN (TM)
STAPH LATEX TEST KIT

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SECTION I IDENTITY

PRODUCT CLASS: DIAGNOSTIC TEST KIT.
PRODUCT CODE: B1045-1A (120 TESTS).

DESCRIPTION: RAPID SLIDE LATEX AGGLUTINATION TEST TO DETECT PROTEIN A AND/OR CLUMPING FACTOR COMMONLY USED AS AN AID IN THE IDENTIFICATION OF STAPHYLOCOCCUS AUREUS FROM MEDIA.

COMPONENTS: 1-(3-DIMETHYLAMINOPROPYL)-3-ETHYL-CARBODIMIDE HYDROCHLORIDE (EDAC), SODIUM PHOSPHATE BUFFER, 10% CARBOXYLATED LATEX, BORATE BUFFER, ETHANOLMINE BOVINE SERUM ALBUMIN, SODIUM AZIDE, DEXTRAN, HUMAN ALBUMIN.

HAZARDOUS INGREDIENTS: 1. HUMAN SOURCE MATERIAL (CONTAINED IN LATEX REAGENT).
2. SODIUM AZIDE (AS A PRESERVATIVE IN LATEX REAGENT). CAS #: 26628-22-8
CONTENT: 0.1%.

SECTION II PHYSICAL/CHEMICAL CHARACTERISTICS

CHEMICAL CHARACTERISTICS: NOT APPLICABLE. THIS KIT CONTAINS BIOLOGICAL MATERIAL AND BUFFERS.

APPEARANCE: TEST KIT CONSISTING OF CARD SLIDES, WOOD APPLICATORS, AND ONE PLASTIC BOTTLE CONTAINING STAPH LATEX REAGENT (MURKY WHITE LIQUID).

SECTION III FIRE AND EXPLOSION HAZARD DATA

CONDITIONS: NO FIRE OR EXPLOSION HAZARDS. PACKAGING MATERIAL WILL BURN IN A FIRE.

EXTINGUISHING MEDIA: USE STANDARD FIRE FIGHTING PROCEDURES DEPENDING ON THE SOURCE OF THE SURROUNDING FIRE.

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SECTION IV REACTIVITY DATA

INCOMPATIBILITIES: IF DISPOSED DOWN A DRAIN, THE SODIUM AZIDE IN THIS KIT MAY REACT WITH LEAD AND COPPER PLUMBING TO FORM HIGHLY EXPLOSIVE METAL AZIDES.

HAZARDOUS POLYMERIZATION: WILL NOT OCCUR.

CONDITIONS TO AVOID: NOT APPLICABLE.

HAZARDOUS DECOMPOSITION OR BYPRODUCTS: THIS PRODUCT IS STABLE. IF INVOLVED IN A FIRE, POISONOUS GAS MAY BE PRODUCED BY THE PACKAGING MATERIALS.

SECTION V TOXICOLOGY/HEALTH EFFECTS

WARNING - POTENTIAL BIOHAZARDOUS MATERIAL
SERUM USED IN THE PREPARATION OF THE REAGENT WAS TESTED BY AN FDA APPROVED
METHOD FOR THE PRESENCE OF THE ANTIBODY TO HIV AS WELL AS FOR HEPATITIS B
SURFACE ANTIGEN AND FOUND TO BE NEGATIVE. BECAUSE NO TEST METHOD CAN OFFER
COMPLETE ASSURANCE THAT HIV, HEPATITIS B VIRUS OR OTHER INFECTIOUS AGENTS ARE
ABSENT, THIS PRODUCT SHOULD BE HANDLED AT THE BIOSAFETY LEVEL 2 AS RECOMMENDED
FOR ANY POTENTIALLY INFECTIOUS HUMAN SPECIMEN IN THE CENTERS FOR DISEASE
CONTROL/NATIONAL INSTITUTES OF HEALTH MANUAL "BIOSAFETY IN MICROBIOLOGICAL AND
BIOMEDICAL LABORATORIES".

SECTION VI FIRST AID

ROUTE OF ENTRY: ACCIDENTAL INGESTION IS POSSIBLE.

MEDICAL CONDITIONS GENERALLY AGGRAVATED BY EXPOSURE: AS WITH ALL BIOLOGICAL
PRODUCTS, HYPERSENSITIVITY IS POSSIBLE. IF HYPERSENSITIVITY OCCURS, LIMIT
EXPOSURE.

EMERGENCY AND FIRST AID PROCEDURES:
FOR INGESTION, SEEK IMMEDIATE MEDICAL ATTENTION
FOR EYE CONTACT, FLUSH WITH PLENTY OF WATER AND SEEK MEDICAL ATTENTION.
FOR SKIN CONTACT, WASH WITH SOAP AND WATER.

SECTION VII PRECAUTIONS FOR SAFE HANDLING

GENERAL: READ THE PACKAGE INSERT. ALWAYS FOLLOW GOOD LABORATORY PRACTICES
WHEN USING THIS PRODUCT. HANDLE ALL CONTROLS AND TEST SPECIMENS AS IF CAPABLE
OF TRANSMITTING DISEASE. EMPLOYEE EXPOSURE TO HUMAN SOURCE MATERIAL IS
REGULATED UNDER THE CODE OF FEDERAL REGULATIONS 29 CFR 1910.1030.

STEPS TO BE TAKEN IN CASE MATERIAL IS SPILLED: DECONTAMINATE SPILL WITH A
BLEACH SOLUTION OR APPROPRIATE GERMICIDE PRIOR TO PICK UP. IF MATERIAL IS
SPILLED DOWN DRAIN, FLUSH WITH LARGE VOLUME OF WATER TO PREVENT AZIDE BUILDUP
IN COPPER OR LEAD PLUMBING. DECONTAMINATION PROCEDURES ARE AVAILABLE ON
REQUEST.

WASTE DISPOSAL METHOD: PLACE MATERIAL IN A SEALED CONTAINER AND DISPOSE OF AS
MEDICAL/INFECTIOUS WASTE IN ACCORDANCE WITH APPLICABLE ENVIRONMENTAL
REGULATIONS.

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SECTION VIII CONTROL MEASURES

PERSONAL PROTECTIVE EQUIPMENT: BARRIER GLOVES, EYE PROTECTION, AND LABORATORY
COAT MAY BE REQUIRED AS LABORATORY CONDITIONS INDICATE.

VENTILATION: A BIOSAFETY CABINET, AS RECOMMENDED IN THE CDC/NIH MANUAL, MAY BE
NECESSARY IF THERE IS A POSSIBILITY OF AEROSOLIZATION DURING HANDLING OF
CONTROLS OR TEST SPECIMENS.

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