

HC-PRE - pre-dip additive liquid
BouMatic, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).

HC-PRE

Pre-Dip Additive

Cleaning, foaming, skin conditioning

For use with HydroClean system

Mixed solution to be used within 72 hours

Use in a well ventilated area

Do not return unused product to original container

For Manufacturing Use Only

FIRST AID:

If in eyes: Hold eye open and rinse slowly and gently with water for 15-20 minutes. Remove contact lenses, if present after the first 5 minutes, then continue rinsing. Contact a physician immediately.

If swallowed: Have person sip a glass of water if able to swallow. Do not give anything to an unconscious person. Do not induce vomiting. Contact a physician immediately. If breathing difficulty occurs: Move person to fresh air. Contact a physician immediately.

If breathing difficulty occurs: Move person to fresh air. Contact a physician immediately.

If on skin: Take off contaminated clothing. Rinse skin immediately with plenty of water for 15-20 minutes. Contact a physician immediately.

Have the product container or label with you when going for treatment, calling a physician, the emergency number listed on this label or MSDS, or a poison control center.

PRECAUTION: Avoid eye and skin contact. Moderately irritating to the eyes. Contact with this product may cause severe eye damage. Contact with skin may cause irritation. May cause irritation of respiratory tract.

STORAGE: Store closed container in a cool, dry, well ventilated area. If product becomes frozen, thaw and mix well before use.

SEE MATERIAL SAFETY DATA SHEET



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Lot #:

Exp. Date:

Net Contents:

SKU:



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PRIMEROS AUXILIOS:

Contacto ocular: Mantenga los ojos abiertos y enjuague cuidadosamente durante 15-20 minutos. Qúitese los lentes de contacto, después de los primeros 5 minutos si los tiene puestos y continúe a enjuagar sus ojos. Contacte a un médico inmediatamente. **Si se ingiere:** Haga que la persona se tome un vaso de agua si puede. No le dé nada de tomar a una persona inconsciente. No induzca el vomito. Contacte a un médico inmediatamente. **Si se inhala:** Lleve a la persona al aire libre. Contacte a un médico inmediatamente. **Contacto con la piel:** Qúitese la ropa contaminada. Enjuague la piel inmediatamente con bastante agua durante 15-20 minutos. Contacte a un médico inmediatamente.

Asegúrese de tener el contenedor o la calcomanía del producto al momento de ir por atención médica, cuando hable con el médico, cuando hable al número de asistencia que se encuentra en la hoja de información del producto o el centro de control de productos venenosos.

PRECAUCION: Evite el contacto en los ojos y la piel. Este producto es moderadamente irritante para los ojos. Contacto con este producto puede causar daños graves en los ojos. Contacto con la piel puede causar irritación. Puede causar irritación de las vías respiratorias.

ALMACENAMIENTO: Almacene en contenedores cerrados en un lugar fresco y seco lejos de una fuente de calor. Si el producto se congela, descongélelo y agítelo antes de usarlo.

VEASE LA HOJA DE DATOS SOBRE MEDIDAS DE SEGURIDAD PARA EL MANEJO DE MATERIALES

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**Emergency No. Only/No. d'urgence Seulement/
No. de Emergencia Sólo:**
US/CANADA only (1-800-255-3924) (ChemTel, Inc)
International (+ 01-813-248-0585) (ChemTel, Inc)
Made in U.S.A.

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LN 9893000

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HC-PRE

pre-dip additive liquid

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:48106-1200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MYRISTAMINE OXIDE (UNII: J086PM3RRT) (MYRISTAMINE OXIDE - UNII:J086PM3RRT)	MYRISTAMINE OXIDE	300 g in 1 L

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48106-1200-2	56.8 L in 1 DRUM		
2	NDC:48106-1200-1	18.9 L in 1 DRUM		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/01/2013	

Labeler - BouMatic, LLC (124727400)

Registrant - BouMatic, LLC (124727400)

Revised: 7/2013

BouMatic, LLC