## Safety Data Sheet according to the DIR. 2001/58/EEC CYBERSILICONE 380 PUTTY REGULAR SET

Revision date: 10/10/06 Print Date: 12/10/06 Page 1 of 2 REVISION DATE DESCRIPTION EMISSION APPROVAL 1 10/10/2006 First emission R&S RSGQA 1. IDENTIFICATION OF THE PREPARATION AND OF THE COMPANY 1.1 Identification of the preparation: *Trade name*: CYBERSILICONE 380 PUTTY REGULAR SET *Trade code*: 9002809 1.2 Utilization of the substance or of the preparation: Addition silicone for high precision impressions. 1.3 Identification of the company: *Supplier*.

DE Healthcare Products Gillingham ME8 0SB Denver PA 17517 / USA Fax.03 20 37 16 89

2. COMPOSITION/INFORMATION ON INGREDIENTS

Hazardous components in accordance with EEC directive 67/548 and successive amendments or components with a recognised exposure limit. None

3. IDENTIFICATION OF HAZARDS

PREPARATION NOT CLASSIFIED DANGEROUS

No specific hazards are encountered under normal use of the product. Contact with the eyes may cause slight irritation and reddening of the conjunctiva.

4. FIRST AID MEASURES

4.1 Contact with skin: The specific indications will not cause of irritation.

4.2 Contact with eyes: Wash immediately with water for at least 10 minutes.

4.3 Swallowing: If swallowed in a small quantity the product is harmless, otherwise consult a doctor.

4.4 Inhalation: Ventilate the premises.

5. FIRE-FIGHTING MEASURES

5.1 Recommended extinguishers: water, CO2, foam, or chemical powders, depending on

the materials involved in the fire.

5.2 Extinguishers not to be used: none in particular.

5.3 Risks arising from combustion: avoid inhaling the fumes.

5.4 Protective equipment: use protection for the respiratory tract.

6. ACCIDENTAL RELEASE MEASURES

6.1 Measures for personal safety: none in particular

6.2 Environmental measures: limit leakage with earth or sand. If the product has escaped into a water course, into the drainage system, or has contaminated the ground or vegetation, notify the competent authorities.

6.3 Cleaning methods: recover the product for re-use if possible, or for elimination. The product may, where appropriate, be absorbed by inert material. After the product has been recovered, rinse the area and materials involved with water.

7. HANDLING AND STORAGE

7.1 Handling precautions: refer to point 8 below. Do not eat or drink while working.

7.2 Incompatible materials: none in particular. Also see point 10 below.

7.3 Storage conditions: keep in a cool, dry place at a temperature of 5°C to 27°C (41 - 80°F).

7.4 Technical measures: adequately ventilated premises.

Cybertech DE Healthcare Prodcuts

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8. EXPOSURE CONTROLS / PERSONAL PROTECTION

8.1 Precautionary measures: ensure adequate ventilation of the premises where the

product is stored and/or handled.

8.2 Respiratory protection: not needed for normal use

8.3 Protection for hands: not needed for normal use

8.4 Eye protection: not needed for normal use. Always apply proper working practices.

8.5 Protection for skin: no special precaution required for normal use

8.6 Exposure limit(s) (ACGIH): none

9. PHÝSICAL AND CHEMICÁL PROPERTIES

9.1 Appearance and colour: Bicomponent paste, of which one part (base) is coloured (blue).

9.2 Odour: odourless

9.3 Relative density: 1.61 g/cc

9.4 Solubility in water: insoluble

10. STABILITY AND REACTIVITY

10.1 Conditions to avoid: stable under normal conditions.

10.2 Substances to avoid: the "base" component may produce hydrogen in contact with

alkaline and acid substances.

10.3 Hazardous decomposition products: none

11. TOXICOLOGICAL INFORMATION

The product itself has no cytotoxic effects on the L929 cellular line.

12. ÉCOLOGICAL INFORMATION

Adopt sound working practices, ensuring that the product is not released into the environment. 13. DISPOSAL CONSIDERATIONS

Recover if possible. In so doing, comply with the local and national regulations currently in force.

14. TRANSPORT INFORMATION

Not classified as dangerous under transport regulations.

**15. REGULATORY INFORMATION** 

The preparation is not considered hazardous under D.Lgs.65/03.

Medical Device CE marked according to the Directive 93/42/EEC.

**16. OTHER INFORMATION** 

The information contained herein is based on our state of knowledge on the date specified above. It refers solely to the product indicated and constitutes no guarantee of particular quality. It is the duty of the user to ensure that this information is appropriate and complete with respect to the specific use intended. This MSDS cancels and replaces any preceding release.