GIMA
PROFESSIONAL MEDICAL PRODUCTS

ULTRA SINGLE PATIENT ANAESTHESIA FACE MASKS

REF **CM-65810 (57711) ULTRA SINGLE PATIENT ANAESTHESIA FACEMASK N. 1 - infant - pink**
REF **CM-65820 (57712) ULTRA SINGLE PATIENT ANAESTHESIA FACEMASK N. 2 - child - yellow**
REF **CM-65830 (57713) ULTRA SINGLE PATIENT ANAESTHESIA FACEMASK N. 3 - adult small - green**
REF **CM-65840 (57714) ULTRA SINGLE PATIENT ANAESTHESIA FACEMASK N. 4 - adult medium - red**
REF **CM-65850 (57715) ULTRA SINGLE PATIENT ANAESTHESIA FACEMASK N. 5 - adult large - blue**

Product Description:
Ultra single patient anaesthesia facemask is connected to manual resuscitator for use as an adjunct to artificial respiration and cardiopulmonary resuscitation. Five different sizes available, form infant to adult large. Injectable and non-injectable cushion types.

Specification of material: PP and TPR.

Usage instruction:
1. Check for clean airflow passage prior to use.
2. Connect to breathing circuit or resuscitation with appropriate fitting.
3. Make sure all connections are compatible and fit firmly.
4. Hand-hold secure with heat strap.

Warning:
1. The product should be placed in a dry and clean place and sealed conditions.
2. The operator should be trained and know the skill in CPR resuscitation.

Precaution:
1. Federal Law (USA) restricts this device to sale by or on the order of a physician.
2. Patient should be monitored constantly whenever the device is on use.
3. Store in clean dry conditions. Away from heat and light.
4. For single patient use only. Reuse may result in cross-infection. Do not soak, wash, rinse or sterilize this product. These procedures may leave harmful residues on the mask.

Destroy the product after use.

GIMA WARRANTY TERMS
The Gima 12-month standard B2B warranty applies.

	Keep in a cool, dry place	LOT	Lot number		Manufacturer
	Keep away from sunlight		Caution: read instructions (warnings) carefully		Date of manufacture
REF	Product code		Follow instructions for use		Disposable device, do not re-use
	Expiration date	CE	Medical Device complies with Directive 93/42/EEC	EC REP	Authorized representative in the European community

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