

Contact Officer: Janette Dunn

TGAIN: 151708
ENT ID: 31265

Medteq Innovations Pty Ltd
Level 2 Colonial State Bank Building
SPRINGWOOD QUEENSLAND 4127

ATTENTION: Phillip Stoneham

**NOTIFICATION OF APPROVAL OF A VARIATION TO A LISTING OF
THERAPEUTIC DEVICES**

AUST L: 73024

Manufacturer: MEDTEQ INNOVATIONS PTY LTD – ENT ID: 31265

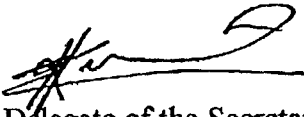
Variation: Change of brand name from
1. RES-Q-MASK TO CPR EZY MASK and
2. RES-Q-PAD TO CPR EZY PAD

Your application for a variation to the products within a listing has been approved and the ARTG listing record has been amended accordingly.

Further changes to this listing should be made in accordance with the TGA Publication, Device Requirements Version 4 (DR4), Appendix 3, 'Changes to Therapeutic Devices in the ARTG'. Information on the 'Conditions of Listing – Standard and Specific' are located at Appendix 4 of the DR4. The DR4 may be obtained from the TGA Publications Office, free call 1800 020 653 or via our web site at <http://www.health.gov.au/tga/docs/html/dr4.htm>.

Good Manufacturing Practice: Where certificates of Good Manufacturing Practice/Conformity Assessment are provided they must be either an original certificate or a notarised copy. (Refer page 3 of the "Guidelines on Standard of Overseas Manufacturers – 12th Edition" which is available from the Publications Office, TGA, Telephone (02) 6232 8610.)

As a replacement certificate will not be issued, you may wish to attach this letter to the original certificate for your records.



Delegate of the Secretary
Medical Devices Listing Section
Conformity Assessment Branch
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26th July 2001