Australian Government
Department of Health and Ageing
Therapeutic Goods Administration

GMP CERTIFICATE OF MANUFACTURING FACILITY

No. MI-13102005-CE-000584-1

DXN Pharmaceutical SDN BHD, Lot 1109, Mukim Malau, Daerah Kubang Pasu, Kedah Darul Aman, Malaysia has been subject to audit by officers of the Manufacturer Assessment Branch, Therapeutic Goods Administration (TGA).

From the knowledge gained during the last audit, which was conducted on 26-27 June 2006, it is considered that the company complies with the current Australian Code of Good Manufacturing Practice for Medicinal Products.

The company manufactures the following traditional medicines: tablets, hard shell capsules and powders, which were included in the above audit for supply to Australia.

This certificate remains the property of the TGA and must be returned upon written demand. The validity of this certificate may be checked by contacting the undersigned.

This certificate remains valid for 3 years from the date of last audit, provided that re-audits are conducted when scheduled by the TGA.

Signed

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Date: 30/11/2006

Mr Andrew Muir
Medicines Audit Team Manager
Manufacturer Assessment Branch
PO Box 100, Woden ACT 2606, Australia
Tel: +61 (0)2 6232 8629
Fax: +61 (0)2 6232 8426

Commonwealth department of
Health and
Ageing
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