Good Manufacturing Practice Certificate

MEDSAFE
NEW ZEALAND MEDICINES AND MEDICAL DEVICES SAFETY AUTHORITY
A BUSINESS UNIT OF THE MINISTRY OF HEALTH
www.medsafe.govt.nz

Ref: TT60-50-16-3

To whom it may concern

This is to certify that **Comvita New Zealand Limited** operating at **23 Wilson Road South, Paengaroa, Bay of Plenty 3189, New Zealand,** has been audited to the New Zealand Code of Good Manufacturing Practice for Manufacture and Distribution of Therapeutic Goods, Part 1: Manufacture of Pharmaceutical Products, and has been found to comply with the requirements for:

- Manufacture, packing and release for supply of non-sterile liquids, suspensions and sprays containing: herbal ingredients, apiary products, fish products, vitamins, minerals and enzymes.

- Packing and release for supply of tablets, lozenges and capsules containing: herbal ingredients, apiary products, fish products, dairy ingredients, vitamins, minerals and enzymes.

The following persons are currently nominated as the persons responsible for release for supply of finished product:

Liz Walker, Compliance Manager
Michel Eyskens, Quality Control Manager
David Jones, Process & Systems Manager

Note that this certificate only applies to products that:

- do not fall within the definitions of Medicine and Related Product in the New Zealand Medicines Act 1981
- are intended for export to Australia where they are categorised as 'listed medicines'

This certification is based on an audit carried out by an officer of the Ministry of Health at the Company's site on 15, 16 and 17 October 2013.

This certificate is valid until 24 July 2015.

Derek Fitzgerald
Manager, Compliance Management
24 January 2014

MINISTRY OF HEALTH 133 MOLESWORTH STREET WELLINGTON NEW ZEALAND