

Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER : UK GMP 22500 Insp GMP/IMP 22500/1125023-0002 [H]

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER(1),(2)

Part 1

Issued following an inspection in accordance with :

Regulation 331A of The Human Medicines Regulations 2012 (SI 2012/1916)

The competent authority of United Kingdom confirms the following :

The Manufacturer : ANDERSEN CALEDONIA LIMITED

Site address : ANDERSEN CALEDONIA LIMITED, CALEDONIAN HOUSE, PHOENIX CRESCENT, STRATHCLYDE BUSINESS PARK, BELLSHILL, ML4 3NJ, UNITED KINGDOM

Other :

Is a contract laboratory that has been inspected in accordance with the Medicines Act as amended

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 30/04/2012, it is considered that it complies with

- The principles and guidelines of Good Manufacturing Practice laid down in Regulation B17 of the Human Medicines Regulations 2012 (as amended)

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. This certificate is valid only when presented with all pages and both Parts 1 and 2. The authenticity of this certificate may be verified in MHRA-GMDP. If it does not appear, please contact the issuing authority.

- (1) Guidance on the interpretation of this template can be found in the Help menu of MHRA-GMDP database.
- (2) These requirements fulfil the GMP recommendations of WHO.

Part 2

Human Medicinal Products

1. MANUFACTURING OPERATIONS

[1.6] Quality control testing

[1.6.2] Microbiological: non-sterility

Any restrictions related to the scope of this certificate:

Building Room Line/equipment QC Testing

Products

The scope of the microbiological testing at this site is limited to the handling of samples
from environmental monitoring

12/07/2013 Name and signature of the authorised person of the Competent Authority of United Kingdom
Confidential
Medicines and Healthcare products Regulatory Agency
Tel : Confidential