TO ALL TO WHOM these presents shall come

I IAN ALASTAIR FISHER of the City of Liverpool in England Notary Public by Royal Authority duly authorised admitted and sworn and practising in the said City **DO HEREBY CERTIFY AND ATTEST** that the attached document consisting of six pages is a true and complete copy of the original **CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER** of Thompson & Capper Limited whose site address is 1-13 Hardwick Road Astmoor Industrial Estate Runcorn Cheshire WA7 1PH United Kingdom I having carefully compared the said copy with the original and found the same to agree therewith

IN FAITH AND TESTIMONY whereof I have hereunto subscribed my hand and affixed my Notarial Seal of Office at the said City this Twenty-eighth day of October in the year of Our Lord Two Thousand and Fifteen

Notary Public
26 Exchange Street East,
Liverpool, L2 3PH
England

MHRA
Regulating Medicines and Medical Devices

MHRA
151 Buckingham Palace Road
London SW1W 9SZ
United Kingdom

mhra.gov.uk

RESTRICTED – COMMERCIAL
Mrs AL Lambert
THOMPSON & CAPPER LIMITED
1-13 HARDWICK ROAD
ASTMOOR INDUSTRIAL ESTATE
RUNCORN
WA7 1PH
UNITED KINGDOM

UK MIA 1359 Insp GMP 1359/700383-0005

MHRA
Regulating Medicines and Medical Devices

Medicines and Healthcare products Regulatory Agency

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with Art. 111(5) of Directive 2001/83/EC.

The competent authority of the United Kingdom confirms the following:

The manufacturer	THOMPSON & CAPPER LIMITED
Site address	1-13 HARDWICK ROAD ASTMOOR INDUSTRIAL ESTATE RUNCORN WA7 1PH UNITED KINGDOM

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. MIA 1359 in accordance with Art. 40 of Directive 2001/83/EC transposed in the following national legislation: The Human Medicines Regulations 2012 (SI 2012/1916).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 01/06/2015, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC.

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 only valid when presented with all pages and both parts 1 and 2.

The authenticity of this certificate may be verified in EudraGMDP. If it does not appear please contact the issuing authority.

MHRA
Regulating Medicines and Medical Devices

art 2

Human Medicinal Products

1. MANUFACTURING OPERATIONS

1.1 Sterile products

Not Authorised

1.2 Non-sterile products

1.2.1 Non-sterile products (processing operations for the following dosage forms)

1.2.1.1 Capsules, hard shell
1.2.1.8 Other solid dosage forms
1.2.1.13 Tablets
1.2.1.17 Other non-sterile medicinal products
Other solid dosage forms, Sachets and sticks.

1.3 Biological medicinal products

Not Authorised

1.4 Other products or manufacturing activity

1.4.1 Manufacture of

1.4.1.1 Herbal products
1.4.1.2 Homeopathic products
1.4.1.3 Other
Anthroposophic remedy.

1.5 Packaging

1.5.1 Primary packaging

1.5.1.2 Capsules, soft shell
1.5.2 Secondary packaging

1.6 Quality control testing

1.6.3 Chemical/physical

2. IMPORTATION OF MEDICINAL PRODUCTS

2.1 Quality control testing of imported medicinal products

Not Authorised

2.2 Batch certification of imported medicinal products

Not Authorised

2.3 Other importation activities

Not Authorised

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3. MANUFACTURING OPERATIONS

3.1 Manufacture of Active Substance by Chemical Synthesis
Not Authorised

3.2 Processing Activities of Active Substance from Natural Sources
Not Authorised

3.3 Manufacture of Active Substance using Biological Processes
Not Authorised

3.4 Manufacture of sterile active substance
Not Authorised

3.5 General Finishing Steps
Not Authorised

3.6 Quality Control Testing
Not Authorised

4 Other Activities
Not Authorised

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Any restrictions or clarifying remarks related to the scope of this certificate:

N/A

1. Building(s)/Area(s)

   N/A

2. Room(s)

   N/A

3. Line(s) Equipment(s)

   N/A

4. QC testing

   N/A

5. Medicinal Product(s)/IMP(s)

   N/A

Name of the authorised person of the Competent Authority of the United Kingdom

Ian Holloway
GMP Inspector
Ian.Holloway@mhra.gsi.gov.uk

Date: 13/10/2015