Central Drugs Standard Control Organisation
Directorate General of Health Services
Ministry of Health & Family Welfare

Food and Drug Administration Bhawan
Kotla Road, New Delhi-110002

No. 7-5/2013/EU/WC-0169 Dated 28 JUN 2013

To

M/s. Avik Pharmaceutical Limited
A-1/7, 1st Phase GIDC,
Vapi-396 195, Valsad, Gujarat, India

SUB:- Written Confirmation of M/s. Avik Pharmaceutical Limited, A-1/7, 1st Phase GIDC, Vapi-396 195, Valsad, Gujarat, India, as per requirement of EU for import of active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC from India-Reg.

Sir,

Please refer to your application submitted to CDSCO, Ahmedabad Zone office and the recommendation received from DDC(I), Ahmedabad Zone, on the above noted subject.

Written Confirmation as required for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC from India is herewith granted subject to the following conditions:-

1. The Active Pharmaceutical Ingredients shall confirm to Good Manufacturing Practices mentioned in the EU directives or other equivalent (GMP of WHO/ICH Q7).

2. The manufacturer is subject to regular, strict and transparent controls and to the effective enforcement of Good Manufacturing Practice, including repeated and unannounced inspections, so as to ensure a protection of public health equivalent to that in the EU.

3. The manufacturer is required to follow the Guidance document for Issue of Written Confirmation as issued by CDSCO.

4. Written Confirmation shall be produced by the Authorized Exporter as and when required by the Drug Regulatory Authority.

5. The Written Confirmation will be withdrawn in the events of non compliance of Standards.

6. This Written Confirmation, unless it is sooner suspended o r cancelled, shall be valid for a period of three years.

7. In the event of any Non Compliance observed during inspections conducted by Local or International Drug Authorities, the same shall be forwarded to this office within 7 days of receipt of report.

8. In the event of any drug found not of standard quality, the same shall be reported to this office within 7 days of receipt of report.

Please note that Written Confirmation issued is liable to be suspended / cancelled, if any of the conditions stipulated above are not complied with or in case of violation of the provisions of the Drugs & Cosmetics Act, 1940 and the Rules thereunder as the case may be.

Please acknowledge the receipt.

Yours faithfully,

(Dr. G. N Singh)
Drugs Controller General (India)

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GOVERNMENT OF INDIA
MINISTRY OF HEALTH & FAMILY WELFARE
Central Drugs Standard Control Organization

CERTIFICATE. NO. : **WC-0169**

Written confirmation for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC

1. **Name and address of site:** M/s. Avik Pharmaceutical Limited
A-1/7, 1st Phase GIDC,
Vapi-396 195, Valsad, Gujarat, India

2. **Manufacturer's licence number**: G/424

Regarding the manufacturing plant under (1) of the following Active substance(s) exported to the EU for medicinal products for human use

As per List enclosed as Annexure-1

The issuing Regulatory Authority hereby confirms that:

The standards of good manufacturing practice applicable to this manufacturing plant are at least equivalent to those laid down in the EU(= GMP of WHO/ICH Q7);

The manufacturing plant is subject to regular, strict and transparent controls and to the effective enforcement of good manufacturing practice, including repeated and unannounced inspections, so as to ensure a protection of public health at least equivalent to that in the EU, and

In the event of findings relating to non-compliance, information on such findings is supplied by the exporting third country without delay to the EU

Date of inspection of the plant: 17th August, 2011

The Written Confirmation remains valid until: 02nd July, 2016

The authenticity of this written confirmation may be verified with the issuing regulatory authority

This written confirmation is without prejudice to the responsibilities of the manufacturer to ensure the quality of the medicinal product in accordance with Directive 2001/83/EC.

Address of the issuing regulatory authority: **Central Drugs Standard Control Organisation**
FDA Bhawan, Kotla Road,
New Delhi- 110 002, India

Name and function of responsible person: Dr. G.N. Singh,
Drugs Controller General (India)

E-mail: dcg@nic.in,
Telephone no.: +91-11-23236965
Fax no.: +91-11-23236973

Signature

O/C

Stamp of the authority and date

28 JUN 2013

GOVERNMENT OF INDIA
MINISTRY OF HEALTH & FAMILY WELFARE
Central Drugs Standard Control Organization

CERTIFICATE NO. :

Annexure-1
WC-0169

Written confirmation for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC

1. Name and address of site:

M/s. Avik Pharmaceutical Limited
A-1/7, 1st Phase GIDC,
Vapi-396 195, Valsad, Gujarat, India.

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Betamethasone Dipropionate(BP/USP/EP)	Manufacturing & Packing
2.	Clobetasol Propionate (BP/USP)	Manufacturing & Packing
3.	Mometasone Furoate (BP/USP/EP)	Manufacturing & Packing
4	Triamcinolone Acetonide (EP)	Manufacturing & Packing

ITEM(S) Four (04) ONLY

The Written Confirmation remains valid until 02nd July, 2016

Signature

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Stamp of the authority and date

28 JUN 2013

Central Drugs Standard Control Organisation
Directorate General of Health Services
Ministry of Health & Family Welfare

Food and Drug Administration Bhawan
Kotla Road, New Delhi-110002

No. 7-5/2013/EU/WC-0169 Dated 10.6 AUG 2013

To

M/s. Avik Pharmaceutical Limited,
A-1/7, 1st Phase GIDC,
Vapi-396 195, Valsad, Gujarat, India

SUB:- Written Confirmation of M/s. Avik Pharmaceutical Limited, A-1/7, 1st Phase GIDC, Vapi-396 195, Valsad, Gujarat, India, as per requirement of EU for import of active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC from India-Reg.

Sir,

Please refer to your application submitted to CDSCO, Ahmedabad Zone office and the recommendation received from DDC(I), Ahmedabad Zone, on the above noted subject.

Written Confirmation as required for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC from India is herewith granted subject to the following conditions :-

1. The Active Pharmaceutical Ingredients shall confirm to Good Manufacturing Practices mentioned in the EU directives or other equivalent (GMP of WHO/ICH Q7).
2. The manufacturer is subject to regular, strict and transparent controls and to the effective enforcement of Good Manufacturing Practice, including repeated and unannounced inspections, so as to ensure a protection of public health equivalent to that in the EU.
3. The manufacturer is required to follow the Guidance document for Issue of Written Confirmation as issued by CDSCO.
4. Written Confirmation shall be produced by the Authorized Exporter as and when required by the Drug Regulatory Authority.
5. The Written Confirmation will be withdrawn in the events of non compliance of Standards.
6. This Written Confirmation, unless it is sooner suspended o r cancelled, shall be valid for a period of three years.

7. In the event of any Non Compliance observed during inspections conducted by Local or International Drug Authorities, the same shall be forwarded to this office within 7 days of receipt of report.

8. In the event of any drug found not of standard quality, the same shall be reported to this office within 7 days of receipt of report.

Please note that Written Confirmation issued is liable to be suspended / cancelled if any of the conditions stipulated above are not complied with or in case of violation of the provisions of the Drugs & Cosmetics Act, 1940 and the Rules thereunder as the case may be.

Please acknowledge the receipt.

Annexure No.	No. of Products	Date of issue	Valid Upto
1	04	28/06/2013	02/07/2016
2	05	10 AUG 20	02/07/2016

Yours faithfully,

(Dr. G. N Singh)
Drugs Controller General (India)

Annexure-2

GOVERNMENT OF INDIA
MINISTRY OF HEALTH & FAMILY WELFARE
Central Drugs Standard Control Organization

CERTIFICATE. NO. : WC-0168

Written confirmation for active substances imported into the European Union (EU) for medicinal products for human use, in accordance with Article 46b(2)(b) of Directive 2001/83/EC

1. Name and address of site: M/s. Avik Pharmaceutical Limited
A-1/7, 1st Phase GIDC,
Vapi-396 195, Valsad, Gujarat, India

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1	Beclomethasone Dipropionate (BP/EP/USP)	Manufacturing & Packing
2	Betamethasone Valerate (BP/EP/USP)	Manufacturing & Packing
3	Clobetasol Propionate (EP)	Manufacturing & Packing
4	Prednisolone Sodium Phosphate (BP/EP/USP)	Manufacturing & Packing
5	Triamcinolone (BP/EP/USP)	Manufacturing & Packing

ITEM(S) Five (05) ONLY

The Written Confirmation remains valid until: 02nd July, 2018

Signature

Stamp of the authority and date

08 AUG 2013