

7-5/2016/EU/WC-0354
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan, Kotla Road
New Delhi-110002

Dated

19 7 OCT 2022

To

M/s. Elixir Pharma,
Plot No. 164/A, Phase-II, Opp. Lathia Industries GIDC,
Naroda, Ahmedabad-382330, Gujarat, India

SUB:- Written Confirmation of M/s Elixir Pharma, Plot No. 164/A, Phase-II, Opp. Lathia Industries GIDC, Naroda, Ahmedabad-382330, 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 online application no. WC/RE/2022/4909 submitted to CDSCO, Ahmedabad Zonal office and the recommendation received from DDC(I), Ahmedabad Zonal office 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 The Written Confirmation will be withdrawn in the events of non compliance of Standards.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 or 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
-	01	17 OCT 2022	08.07.2025

Yours faithfully,

(Dr. V. G. Somani)
Drugs Controller General (India)



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. Elixir Pharma,
Plot No. 164/A, Phase-II, Opp. Lathia Industries
GIDC, Naroda, Ahmedabad-382330, Gujarat, India

2. Manufacturer's licence number : G/25/1746

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

List of API(s):

S. No.	Active substance(s)	Activity(ies)
1.	Glibenclamide BP/EP	Manufacturing & Packing

ITEM(S) One (01) ONLY

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: 29.08.2022

The Written Confirmation remains valid until: **08.07.2025**

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. V. G. Somani,
Drugs Controller General (India)

E-mail:

Telephone no.:

Fax no.:

dcic@nic.in,

+91-11-23236965

+91-11-23236973

Signature

Stamp of the authority and date



7-5/2016/EU/WC-0354
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan, Kotla Road
New Delhi-110002

Dated

To

21 OCT 2022

M/s. Elixir Pharma,
Plot No. 164/A, Phase-II, Opp. Lathia Industries GIDC,
Naroda, Ahmedabad-382330, Gujarat, India

SUB:- Written Confirmation of M/s Elixir Pharma, Plot No. 164/A, Phase-II, Opp. Lathia Industries GIDC, Naroda, Ahmedabad-382330, 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 online application no. WC/RE/2022/4910 submitted to CDSCO, Ahmedabad Zonal office and the recommendation received from DDC(I), Ahmedabad Zonal office 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:-

9. The Active Pharmaceutical Ingredients shall confirm to Good Manufacturing Practices mentioned in the EU directives or other equivalent (GMP of WHO/ICH Q7).
10. 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.
11. The manufacturer is required to follow the Guidance document for Issue of Written Confirmation as issued by CDSCO.
12. Written Confirmation shall be produced by the Authorized Exporter as and when required by the Drug Regulatory Authority.
13. The Written Confirmation will be withdrawn in the events of non compliance of Standards.

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

15. 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.

16. 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
-	01	17.10.2022	08.07.2025
1	01	21 OCT 2022	08.07.2025

Yours faithfully,



(Dr. V. G. Somani)
Drugs Controller General (India)



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

CERTIFICATE NO. :

Annexure-1

WC-0354

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 Elixir Pharma,
Plot No. 164/A, Phase-II, Opp. Lathia Industries
GIDC, Naroda, Ahmedabad-382330, Gujarat, India

List of APIs:

Sr. No.	Active substance (s)	Activity(ies)
1.	Nalidixic Acid BP/EP/USP	Manufacturing & Packing

Item(s) One (01) Only

The Written Confirmation remains valid until: 08.07.2025

Signature

21 OCT 2022

Stamp of the authority and date



7-5/2016/EU/WC-0354
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation
(International Cell)

FDA Bhawan, Kotla Road,
New Delhi-110002

Dated: 10 JAN 2025

To,

**M/s. Elixir Pharma Chem Private Limited,
Plot No. 164/A, Phase-II, Opp. Lathia Industries,
G.I.D.C., Naroda, Ahmedabad-382330, Gujarat**

SUB:- Written Confirmation of **M/s. Elixir Pharma Chem Private Limited, Plot No. 164/A, Phase-II, Opp. Lathia Industries, G.I.D.C., Naroda, Ahmedabad-382330, Gujarat**, 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 online application no. WC/FR/2024/8224 dated 21.03.2024 submitted to CDSCO, DDC(I), Gujarat Zone, and the recommendation received from DDC(I), Gujarat 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 or 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.
9. The manufacturer is required to comply with the provisions of GSR 20(E), dated 18.01.2022.

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
--	01	17.10.2022	08.07.2025
01	01	21.10.2022	08.07.2025
02	02	10 JAN 2025	08.07.2025

Yours faithfully,

Chandrashekar
10/01/25
(Ranga Chandrashekar)
Joint Drugs Controller (India)

चंद्रशेखर रंगा/Chandrashekar Ranga
संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller (India)
केंद्रीय औषधि मानक नियंत्रक संगठन (मुख्यालय), स्वास्थ्य सेवा महानिदेशालय
C.D.S.C.O(HQ), Dte. General of Health Services
स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare
एन सी ० ५ धरम कोरना रोड, नई दिल्ली-110002 / FDA Bhaswan, Kotla Road, New Delhi-110002



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. Elixir Pharma Chem Private Limited,
Plot No. 164/A, Phase-II, Opp. Lathia Industries,
G.I.D.C., Naroda, Ahmedabad-382330, Gujarat

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1.	Glibenclamide BP/EP	Manufacturing & Packing
2.	Nalidixic Acid BP/EP/USP	Manufacturing & Packing

ITEM(S) Two (02) ONLY

The Written Confirmation remains valid until: 08.07.2025

Chandrashekar Ranga

Signature

Stamp of the authority and date



10 JAN 2025

चंद्रशेखर रंगा/Chandrashekar Ranga
संयुक्त औषधि नियंत्रक (भारत) / Joint Drugs Controller(India)
केंद्रीय औषधि यंत्रक नियंत्रण संगठन (मुख्यालय), स्वास्थ्य सेवा महानिदेशालय
C.D.S.C.O.(HQ), Dte. General of Health Services
स्वास्थ्य और परिवार कल्याण मंत्रालय / Ministry of Health and Family Welfare
एफ.डी.ए. भवन, कोटला रोड, नई दिल्ली-110002 / FDA Bhawan, Kotla Road, New Delhi-110002