F. No: 7-5/2016/EU/WC-0373
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organization
(International Cell)

FDA Bhawan, Kotla Road,
New Delhi- 110 002.
Dated: 18 MAR 2019

To,

M/s SMS Pharmaceuticals Ltd, (Unit-VII)
Kandivalasa (V),Poosapatirega(M),
Vizianagaram (Dist)-535 204
Andhra Pradesh, INDIA.

Sub: Written Confirmation of M/s SMS Pharmaceuticals Ltd,(Unit-VII) Kandivalasa(V), Poosapatirega(M), Vizianagaram (Dist)-535204, Andhra Pradesh, 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, Hyderabad Zonal office and the recommendation received from DDC (I), Hyderabad 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 The Written Confirmation will be withdrawn in the event of non-compliance of Standards.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 up to
01.	02	23.06.2016	22.06.2019
02	01	23.06.2016	22.06.2019
03	05	14.02.2017	22.06.2019
04	05	11.06.2018	22.06.2019
05	01	11.06.2018	22.06.2019
06	02	18 MAR 2019	22.06.2019
07	02	18 MAR 2019	22.06.2019

Yours faithfully,

(Dr.S.Eswara Reddy)
Drugs Controller General (India).

O/C
28.02.19
28-2-19
01/03/2019

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

Annexure – 06
WC-0373

CERTIFICATE NO. :

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 SMS Pharmaceuticals Ltd, (Unit-VII)
Kandivalasa (V),Poosapatirega(M),
Vizianagaram (Dist)- 535 204
Andhra Pradesh, INDIA.

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1	Solifenacin Succinate IH	Manufacturing & Packing
2	Ranolazine IH	Manufacturing &Packing

Item(s) Two (02) Only

The Written Confirmation remains valid until: 22.06.2019

Signature

Stamp of the authority and date

18 MAR 2019

O/C 28.02.19
28.2.19
01/03/2019

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

CERTIFICATE NO. : Annexure – 07
WC-0373

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 SMS Pharmaceuticals Ltd, (Unit-VII)
Kandivalasa (V),Poosapatirega(M),
Vizianagaram (Dist)- 535 204
Andhra Pradesh, INDIA.

List of APIs:

S. No.	Active substance(s)	Activity(ies)
1	Sitagliptin Hydrochloride IH	Manufacturing & Packing.
2	Paliperidone Palmitate IH	Manufacturing &Packing

ITEM(S) Two (02) ONLY

This certificate is being issued subject to condition that the firm shall obtain NOC from the Competent Authority, on case to case basis, to manufacture the above mentioned active substances for the purpose of export only, as the above mentioned active substances are not approved for manufacture or sale in India.

The **Written Confirmation remains valid until: 22.06.2019**

Signature

Stamp of the authority and date

18 MAR 2019

O/C
28.02.19.
28.2.19

01/03/19