File No.DCA-EST1/SEC/127/2024-Establishment Section

OFFICE OF THE DIRECTOR GENERAL::DRUGS AND COPYRIGHTS::DRUGS CONTROL ADMINISTRATION::VENGALRAONAGAR::HYDERABAD

Cir.Memo.Rc.No.2515312/JD/MISC/2024

Dated: .05.2024

Sub: Drugs & Cosmetics Act, 1940 & Rules made thereunder – NOC's for Manufacture of Unapproved / Banned / New Drugs Solely for Export Purpose – Certain instructions issued – Regarding.

Ref: Lr.No. IMP-12/1/2024-eoffice, dated 30.04.2024 of DCGI, New Delhi.

-x-x-x-

While enclosing a copy of the reference cited, all Drugs Inspectors in the State are instructed to direct all the manufacturers under your jurisdiction for ensuring the same and to upload the circular in Drugs Control Department website.

This may be treated as URGENT.

Sd/- V.B. KAMALASAN REDDY, IPS., DIRECTOR GENERAL

То

All the Assistant Directors in the State with a instructions to circular the same to all Drugs Inspectors working under their control.

Copy to: All the Deputy Directors in the State for information and necessary action.

//FORWARDED::BY ORDER// Signed by G.ramdhan Date: 16-05-2024 10:08:11 Reason: Approved JOINT DIRECTOR(I/c)

IMP-12/1/2024-eoffice

Government of India **Directorate General of Health Services Central** 0/0. DIRECTOR CINERALS Standard Control Organization Drugs Centrol Administration Division)

0 1 MAY 2024 No: Govt. of T.S. Vengalraonagar, Hyd

FDA Bhawan, Kotla Road, New Delhi-110002 Dated. ? 0 APR 2024

To

All States/UT Drugs Controllers

JD Section

Subject: NOC's for Manufacture of Unapproved/Banned/New Drugs Solely for Export Purpose -Reg

NOC's for manufacture of unapproved/banned/new drugs solely for export purpose are granted as per the Guidance Document issued by CDSCO.

> The above activity was delegated to State licensing Authorities w.e.f. 20th August, 2018 vide letter of F.No.7-5/2018/Misc./034(NOC) dated 2nd August, 2018

Now, it has been decided with the approval of Hon'ble HFM vide Ministry F. No. X.11035/210/2018-DR (Pt) dated 21st June, 2023 that Industry must be facilitated to file fresh applications for NOC for manufacture of unapproved/approved new drug/banned drugs solely for export purpose from15thMay, 2024 on online mode through CDSCO Zonal Offices. Accordingly, power delegated to State/UT Licensing Authority stands withdrawn w.e.f. 15 th May, 2024 and such NOC's shall be granted by the Head of respective CDSCO Zonal office w.e.f.15thMay 2024. Further All State/UT Drugs Controllers are required to handover all NOC's issued from 20th August, 2018 to14thMay, 2024 to respective Zonal Offices of CDSCO.

All manufacturers may be informed that they are required to obtain NOC from respective Zonal Offices of CDSCO through online mode (SUGAM Portal) w.e.f.15th May 2024 before issuing Manufacturing License from SLA for manufacture of Unapproved/Banned/New Drugs for export purpose.

Sh. Ranga Chandrashekar Rao Joint Drugs Controller (India) will be Nodal and designated person at CDSCO, HQ for said activity.

Yours faithfully

2100112 (Dr. Rajeev Singh Raghuvanshi) **Drugs Controller General (India)**

o. IMP-12/1/2024-eoffice (Computer No. 7581)

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Receipt No : 2515312/2024/INW-DCA

Copy

- 1. All Zonal/Sub-Zonal/Port Offices of CDSCO for necessary action.
- 2. All stake holders through CDSCO website.

Copy for information to:

- 1. Deputy Secretary (Drug Regulation), MoH F & W, Govt. of India
- 2. Joint Secretary, (Drug Regulation), MoH F & W, Govt. of India

F.No.X.11035/210/2018-DR (Pt) Government of India Ministry of Health and Family Welfare Department of Health and Family Welfare (DR Section)

Nirman Bhawan, New Delhi Dated the June, 2023

To DCGI, CDSCO, FDA Bhawan, New Delhi- 110002

Subject: NOC for manufacture of unapproved/banned/new drugs solely for export purpose - regarding.

Please refer to your e-file number 8200817 dated 03.01.2023 on subject cited above wherein you have proposed that in supersession of earlier CDSCO order issued vide F. No. 7-5/2018/Misc/034(NOC) dated 2nd August, 2018, a fresh order may be issued requiring that the manufacturer shall take NOC from respective zonal offices of CDSCO before taking manufacturing licenses from SLAs for the manufacture of Unapproved/Banned/New Drugs for export purpose.

2. In this regard, matter has been examined by this ministry and it is to inform you that industry must be facilitated to file fresh applications for NOC for manufacture of unapproved/banned/new drugs solely for export purpose from 15 May 2024 at a single designated person only on online mode to a point of contact at CDSCO headquarters and the applications can be disposed by CDSCO through their respective zonal offices.

3. Further, CDSCO may give necessary direction to all State/UT drugs controller to hand over details of all such export permissions given by them under the delegated provisions since 2018.

This is issued with approval of Hon'ble HFM.

Yours faithfully,

Signed by (Dr. Kiran Kurkina Karlen Karlapu Deputy Secretary to the Govi. of India Tele: 0 Data 2928023 14:27:43