

	सीमाशुल्क आयुक्त का कार्यालय, एन.एस.-II, मुंबई अंचल-II OFFICE OF THE COMMISSIONER OF CUSTOMS, NS-II, MUMBAI ZONE-II जवाहरलाल नेहरू सीमाशुल्क भवन JAWAHARLAL NEHRU CUSTOM HOUSE पो: शेवा, ताल: उरण, नवीमुंबई - ४००७०७ PO: SHEVA, TAL: URAN, NAVI MUMBAI- 400707 Email: commr-ns2@gov.in	
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File No: GEN/AG/MTG/TMC/2/2026-A/M(E)-O/o-Commr-Cus-Nhava Sheva-II

DATE: As Signed

PUBLIC NOTICE NO.:121/2026

Subject: Extension of last date prescribed under Public Notice No. 88/2026 dated 20.07.2026-reg.

Attention of all concerned is invited to Public Notice No. 88/2026 dated 20.07.2026 regarding “*Procedures and documents required for Export Consignments of Drugs & Pharmaceuticals*”, wherein, vide Para 5.2, an interim facilitation measure was provided for export clearance up to 30.09.2026, subject to the conditions prescribed therein.

2. In continuation of the above, the aforesaid relaxation, is hereby extended up to **31.12.2026**. Thereafter, export clearance shall be strictly subject to compliance with the CDSCO Office Order dated 30.04.2024.
3. All other terms and conditions of Public Notice No. 88/2026 dated 20.07.2026 shall remain unchanged.
4. Difficulties, if any, in implementation of this Public Notice may be brought to the notice of the undersigned.
4. This Public Notice shall come into force with immediate effect.

Digitally signed by
Giridhar Gopalkrishna Pai
Date: 01-10-2026
18:01:04

(Giridhar G. Pai)

Commissioner of Customs, NS-II
JNCH, Nhava Sheva.

Encl: As above

To,

All Stakeholders,

Copy to:

- i. The Chief Commissioner of Customs, Mumbai Zone-II, Nhava Sheva.
- ii. The Commissioners of Customs, NS-I, NS-II, NS-III, NS-V, NS-General and NS Audit, Nhava Sheva.
- iii. All Additional/Joint/Dy./Asstt. Commissioners of Customs, Nhava Sheva.
- iv. AC/DC, EDI for uploading on JNCH website.
- v. Office copy.

	सीमाशुल्क आयुक्त का कार्यालय, एन.एस.-II, मुंबई अंचल-II OFFICE OF THE COMMISSIONER OF CUSTOMS, NS-II, MUMBAI ZONE-II जवाहरलाल नेहरू सीमाशुल्क भवन JAWAHARLAL NEHRU CUSTOM HOUSE पो: शेवा, ताल: उरण, नवीमुंबई - ४००७०७ PO: SHEVA, TAL: URAN, NAVI MUMBAI- 400707 Email: commr-ns2@gov.in	
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File No: GEN/AG/MTG/TMC/2/2026-A/M(E)-O/o-Commr-Cus-Nhava Sheva-II

DATE: As Signed

PUBLIC NOTICE NO.: ४४/2026

Subject: Procedures and documents required for export consignments of Drugs & Pharmaceuticals – reg.

Attention of all Exporters, members of the General Trade, Customs Brokers and other stakeholders is invited to Public Notice No. 51/2018 dated 28.03.2018 issued by Jawaharlal Nehru Custom House (JNCH); Public Notice No. 173 (RE-2008)/2004-2009 dated 13.04.2009 issued by the Directorate General of Foreign Trade (DGFT) and Office Order dated 30.04.2024 issued by the Central Drugs Standard Control Organization (CDSCO), Directorate General of Health Services, Ministry of Health & Family Welfare, Government of India, bearing F. No. IMP-12/1/2024-eOffice [copy enclosed].

2. Representations received from various trade associations and issues raised during stakeholder interactions have been examined in consultation with the office of the Assistant Drugs Controller (ADC), CDSCO. Accordingly, in continuation of and as a supplement to the aforesaid Public Notices, and with a view to facilitate exports of Drugs & Pharmaceuticals covered under the provisions of the Drugs and Cosmetics Act, 1940 and the Rules made thereunder, which are regulated by the Drugs Controller General of India (DCGI), Ministry of Health & Family Welfare, reduce procedural redundancies and promote Ease of Doing Business, the following procedures/guidelines are hereby prescribed.

3. In cases where the Shipping Bills are filed by the manufacturer for the export of drugs & pharmaceuticals covered under the provisions of the Drugs and Cosmetics Act, 1940 and the Rules made thereunder, other than unapproved/new/banned drugs, the documents prescribed under DGFT Public Notice No. 173 (RE-2008)/2004-2009 dated 13.04.2009 are required to be uploaded in e-Sanchit at the time of export.

4. In cases where the Shipping Bills are filed by Exporter other than manufacturer for the export of drugs & pharmaceuticals covered under the provisions of the Drugs and Cosmetics Act, 1940 and the Rules made thereunder, the exporter/CB shall produce all relevant documents before ADC/CDSCO to obtain ADC NOC as prescribed under DGFT Public Notice No. 173 (RE-2008)/2004-2009 dated 13.04.2009.

4.1. Upon due verification of the documents submitted by the Exporter other than manufacturer and being satisfied that the applicable regulatory requirements have been duly complied with, the ADC/CDSCO may issue a No Objection Certificate (NOC) for export of the consignment. The issuance of such NOC shall signify that the regulatory requirements falling within the jurisdiction of CDSCO have been examined by the office of the ADC.

4.2. Customs officers shall rely upon the verification carried out by the ADC/CDSCO in respect of the documents indicated in the ADC NOC. Ordinarily, customs officers shall not call for the same documents again from the exporter. In case any clarification or verification is considered necessary in respect of the regulatory aspects examined by CDSCO, the same may be referred to and obtained from ADC/CDSCO, through email at jnpt.mumbai@cdsco.nic.in.

5. In cases where the Shipping Bills are filed by the manufacturer for the export of unapproved/new/banned drugs, the provisions of the CDSCO Office Order dated 30.04.2024 (F. No. IMP-12/1/2024-eOffice) shall be applicable. As per the said Office Order, the powers delegated to the

State/UT Licensing Authorities for issuance of No Objection Certificates (NOCs) for manufacture of Unapproved/Banned/New Drugs solely for export purpose stand withdrawn with effect from 15.05.2024. All manufacturer exporters and other stakeholders are hereby informed that, with effect from 15.05.2024, it is mandatory to obtain a No Objection Certificate (NOC) from the respective CDSCO Zonal Office through the online SUGAM Portal prior to issuance of the Manufacturing Licence by the concerned State Licensing Authority (SLA) for manufacture of Unapproved/Banned/New Drugs intended solely for export purposes.

5.1. Accordingly, exporters seeking export clearance of Unapproved/Banned/New Drugs manufactured solely for export purpose are advised to ensure that:

- a. The requisite the CDSCO NOC has been obtained from the competent CDSCO Zonal Office through SUGAM Portal instead of State/UT Licencing Authorities;
- b. The prescribed chronology of regulatory approvals and manufacturing activities is duly complied with, i.e., the exporter obtains the CDSCO NOC prior to the Manufacturing Licence granted by the State Licensing Authority (SLA), and the exporter manufactures the drug(s) for export only after obtaining such Manufacturing Licence;
- c. In cases where the CDSCO NOC for manufacture of the drug(s) has been issued against a specific Purchase Order (PO) and for export to a specified overseas consignee/buyer, and the exporter is subsequently unable to fulfil such Purchase Order in part or full, the exporter can obtain revised/amended/new NOC, as applicable, from the competent CDSCO authority, incorporating the revised Purchase Order and/or overseas buyer, before export of the said already manufactured drug(s);
- d. The particulars declared in the Shipping Bill such as description, brand, grade, model, specification, unit quantity code, weight, country of origin etc. correspond with the details mentioned in the NOC.

5.2. As an interim facilitation measure, export clearance may be permitted in cases where the CDSCO NOC has been obtained post facto, provided the exporter holds a valid NOC issued by the State/UT Licensing Authority, valid Product Permission, Manufacturing Licence, and other prescribed regulatory approvals. This relaxation shall be applicable only up to **30.09.2026**, after which export clearance shall be strictly subject to compliance of the CDSCO Office Order dated 30.04.2024.

6. Difficulty, if any may also be brought to the notice of Deputy/Assistant Commissioner in charge of Appraising Main (Export) through email (email address: apmainexp@jawaharcustoms.gov.in).

7. Action to be taken in terms of decisions taken in this Public Notice should be considered as standing order for the purpose of officers and staff.

Encl: As above

Digitally signed by
Giridhar Gopalkrishna Pai
Date: 20-07-2026
16:29:40

(Giridhar G. Pai)
Commissioner of Customs, NS-II
JNCH, Nhava Sheva.

To,

All Stakeholders,

Copy to:

1. The Chief Commissioner of Customs, Mumbai Zone-II, Nhava Sheva.
2. The Commissioners of Customs, NS-I, NS-II, NS-III, NS-V, NS-General and NS Audit, Nhava Sheva.
3. All Additional/Joint/Dy./Asstt. Commissioners of Customs, Nhava Sheva.
4. AC/DC, EDI for uploading on JNCH website.
5. Assistant Drugs Controller (ADC), O/o cdsco; M/s JWR Logistics Pvt Ltd
6. Office copy.

IMP-12/1/2024-eoffice
Government of India
Directorate General of Health Services Central
Drugs Standard Control Organization
(Import & Registration Division)

FDA Bhawan, Kotla Road,
New Delhi-110002

Dated:

30 APR 2024

To

All States/UT Drugs Controllers

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 from 15th May, 2024 on online mode through CDSCO Zonal Offices. Accordingly, power delegated to State/UT Licensing Authority stands withdrawn w.e.f. 15th May, 2024 and such NOC's shall be granted by the Head of respective CDSCO Zonal office w.e.f. 15th May 2024. Further All State/UT Drugs Controllers are required to handover all NOC's issued from 20th August, 2018 to 14th May, 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



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

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

I/3597641/2023

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.

Sir,

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 Kumar Karlapu)
Deputy Secretary to the Govt. of India
Tele: 011-23062028
Date: 24.06.2023 14:27:43