檔號　　保存年限

台灣醫療暨生技器材工業同業公會 收 110. 05. 06 文 檔號: 110365

# 經濟部國際貿易局　函

機關地址：臺北市湖口街1號
承 辦 人：蔡濟鴻
聯絡電話：(02)23977250
傳　　真：(02)23584339
電子郵件：chris@trade.gov.tw

24159
新北市三重區重新路5段609巷6號3樓之3

受文者：台灣醫療暨生技器材工業同業公會

發文日期：中華民國110年5月5日
發文字號：貿雙一字第1100450504號
速別：最速件
密等及解密條件或保密期限：
附件：如文(1100450504-1.pdf)

主旨：有關印度政府簡化進口醫療用氧氣相關設備程序事，請查照並請轉知所屬會員。

說明：

一、依據駐印度代表處經濟組110年4月30日竺經字第1100001117號函辦理；本局110年4月28日貿雙一字第1107013074號函諒達。

二、印度糧食暨消費者事務部於110年4月28日宣布開放17項醫療器材進口3個月，包括噴霧器(Nebulizer)、氧氣濃縮機(Oxygen Concentrator)、氧氣瓶(Oxygen Canister)、氧氣鋼瓶(Oxygen Cylinder)、造氧機(Oxygen Generator)、呼吸器(Ventilator)、正壓器(Positive AirwayPressure Device)、氣壓錶(Flow Meter)、調節器(Regulator)及相關零組件等。醫療器材進口商原依印度度量衡規則(Legal Metrology Rules, 2011)須於通關前即完成申報，惟印度政府放寬廠商可於貨物通關後銷售前申報；另廠商於醫療器材進口後須立即向州政府通報進口數量，檢送印度政府公告(如附件)供參。

正本：台灣醫療暨生技器材工業同業公會、財團法人中華民國對外貿易發展協會
副本：駐印度代表處經濟組

局長 江文若

Speed Post

**I-10/22/2020-W&M**
Government of India
**Ministry of Consumer Affairs, Food and Public Distribution**
Department of Consumer Affairs
Legal Metrology Division

Krishi Bhawan, New Delhi
Dated:28.4.2021

To,

1. Director General, Foreign Trade,
Ministry of Commerce & Industry, Udyog Bhawan, New Delhi

2. Controllers of Legal Metrology,
All States/ UTs

**Subject:** Permission for making mandatory declarations required under the Legal Metrology (Packaged Commodities) Rules, 2011 after custom clearance and before sale to importers of medical devices due to prevalent situation of COVID-19 - reg.

Sir/ Madam,

The undersigned is directed to refer to the above-mentioned subject and to state that due to prevalent pandemic situation of COVID-19 there is a steep demand of medical devices in this critical condition on urgent basis in view of the emergent health concerns and immediate supply to the medical industry.

2. Therefore, considering the pandemic situation of COVID-19 and to meet the demand of medical devices, in exercise of the powers conferred by rule 33(1) and rule 6 of the Legal Metrology (Packaged Commodities) Rules, 2011, the Central Government hereby permits the importers of medical devices to import the following medical devices for three months from the date of this advisory, subject to the condition that the importers shall make all declarations required under these rules immediately after import/ custom clearance and before sale by way of stamping or putting sticker or online printing, as the case may be:

(i) Nebulizers,
(ii) Oxygen concentrators,
(iii) Continuous positive airway pressure (CPAP) devices,
(iv) Bilevel positive airway pressure (BIPAP) devices]
(v) Oxygen concentrator along with flow meter, regulator, connectors and tubing
(vi) Vacuum Pressure Swing Absorption (VPSA) and Pressure Swing Absorption (PSA) oxygen plants, Cryogenic Oxygen Air Separation Units (ASUs) producing liquid/ gaseous oxygen
(vii) Oxygen Cannister
(viii) Oxygen Filling Systems
(ix) Oxygen cylinders including cryogenic cylinders

(x) Oxygen Generators
(xi) Parts to be used for the manufacture of equipment for production, transportation, distribution or storage of Oxygen
(xii) Any other device from which Oxygen can be generated
(xiii) Ventilators (capable of functioning as high-flow devices) with nasal canula; Compressors including all accessories and tubing; humidifiers and Viral filters
(xiv) High flow nasal canula device with all attachments
(xv) Helmets for use with non-invasive ventilation
(xvi) Non-invasive ventilation oronasal masks for ICU ventilators
(xvii) Non-invasive ventilation nasal masks for ICU ventilators

3. The importers, importing the said medical devices under this permission shall inform all such imports with quantity imported to the Director (Legal Metrology) and the Controller (Legal Metrology) in the State, where the import is made, immediately after import.

Yours faithfully

(B. N. Dixit)
Director of Legal Metrology
Tel: 011-23389489/ Fax.-011-23385322
**Email: dirwm-ca@nic.in**

**Copy to**: All Industries/ Industry Associations/ Stake Holders