F. No. 12-01/21-DC (Pt-306 – XXXVI- MDTL)
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organization
(New Drugs Division)

FDA Bhavan, Kotla Road,
New Delhi
Dated: 14 OCT 2021

Notice

Subject: Review of regulatory regime for drug approval – Reg

In order to enhance the testing capacity of medical devices in the country identification registration & empanelment of government private testing laboratories for medical device testing is under consideration as per medical devices Rules, 2017.

In this regards, laboratories which have capacity for testing of medical devices & are NABL accredited & interested in above process are requested to intimate the same to CDSCO at dci@cdsco.nic.in & ddcimd-cdsco@nic.in along with details of categories of such medical devices and capacities & capabilities of testing to CDSCO for initiating further examination and communication in this regard .

V.G.

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

To,

All stakeholders through CDSCO website

Copy to

PS to JS(R) , MoHFW , Nirman Bhawan, New Delhi