https://www.wto.org/spanish/news_s/news18_s/tbt_20mar18_s.htm

4. EU laws, regulations, procedures and guidelines on marketing authorisation for medicinal products

India raised questions about EU processes for marketing authorizations for medicinal products, including for generic medicine, and the roles of various EU and member state authorities. The European Union said it welcomed further discussion on these matters in the EU-India Joint Working Group on Pharmaceuticals, Biotechnology and Medical Devices.