

THE DOHA/TRIPS PARAGRAPH 6 SYSTEM: REFLECTIONS FOR GLOBAL HEALTH

SOUTH CENTRE SIDE EVENT AT THE WTO TRIPS COUNCIL: PARAGRAPH 6 OF THE DOHA DECLARATION ON TRIPS AND PUBLIC HEALTH: AN EFFECTIVE SOLUTION? 8 NOVEMBER 2016

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- + Only WTO/TRIPS amendment is for global health
- System not workable reflected in data: 78/164 (48%)







TRIPS: ISSUES

TRIPS and public health

Events and briefing on Trade-Related Aspects of Intellectual Property Rights, patents, and pharmaceuticals and public health — including discussions in the WTO's TRIPS Council.

- 2005: Hong Kong Ministerial Declaration <u>Paragraph 40 on TRIPS and</u> Public Health
- 2007: <u>Decision to extend deadline for accepting TRIPS Agreement amendment</u>
- 2009: <u>Decision to extend deadline for accepting TRIPS Agreement</u> amendment
- 2011: <u>Decision to extend deadline for accepting TRIPS Agreement amendment</u>
- 2013: <u>Decision to extend deadline for accepting TRIPS Agreement</u> amendment
- 2015: <u>Decision to extend deadline for accepting TRIPS Agreement amendment</u>
- 2015: <u>Decision on the Extension of the Transition Period under Article 66.1 of the TRIPS Agreement for Least-Developed Country Members for Certain Obligations with Respect to Pharmaceutical Products adopted by the TRIPS Council on 6 November 2015. This Decision extends the earlier decision that was taken by the TRIPS Council in 2002 in order to implement paragraph 7 of the Doha Declaration on the TRIPS Agreement and Public Health. Least-developed countries will not have to protect pharmaceutical patents and test data until 1 January 2033.
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- Ironically, restricting trade
- Production capacity increasingly concentrated
 - WHO Pre-qualified products: 509
 - Indian manufacturing sites: 70% (358/509)
- Outbreaks and pandemics: from Anthrax to Zika



▲ 334.31 0.14%

Nikkei 7 17171.38 -0.03%

U.S. 10 Yr A 3/32 Yield 1.817%

Crude Oil A 45.10 0.47%

Euro 1.1055 0.14%

DJIA A 18259.60 2.08%

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CORPORATE INTELLIGENCE

Why More Indian Generic Drugs Will Make Their Way to the U.S.

By ED SILVERMAN

May 14, 2014 8:48 am ET

O COMMENTS

PEKKA SAKKI/REX USA/COURTESY EVERETT COLLECTION

Never mind the FDA crackdown on Indian drug makers for quality control problems. Salas of proceedintion drugs to the IIC by the Indian phermacoutical industry likely won't

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 - US FDA drug approvals = 39% Indian drugmakers
- UNSG High Level Panel on Access to Medicines



UNSG HIGH LEVEL PANEL ON ACCESS TO MEDICINES

"WTO Members should <u>revise</u> paragraph 6 decision in order to find a solution that enables a <u>swift and expedient</u> export of pharmaceutical products produced under compulsory license. WTO Members should, as necessary, adopt a waiver and permanent revision of the TRIPS Agreement to enable this reform."



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- UNSG High Level Panel on Access to Medicines
- Risk of "institutional stickiness" casting in stone undesirable rules
- 8y → 22+y of TRIPS; 2y → 13+y of Para 6





THANK YOU

Comments welcome: suerie.moon@graduateinstitute.ch smoon@hsph.harvard.edu

