

10 August 2018

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## **Committee on Technical Barriers to Trade**

## Original: Spanish

## **NOTIFICATION**

## Addendum

The following communication, dated 9 August 2018, is being circulated at the request of the delegation of  $\underline{\text{Costa Rica}}$ .

<u>Draft Central American Technical Regulation (RTCA) No. 11.03.59:11 Pharmaceutical products.</u>
<u>Medicines for human use. Sanitary registration requirements</u>

On 13 July 2011, the Government of the Republic of Costa Rica notified in document G/TBT/N/CRI/118, draft Central American Technical Regulation (RTCA) No. 11.03.59:11 Pharmaceutical products. Medicines for human use. Sanitary registration requirements, with the aim of establishing the conditions and requirements for the registration of medicines for human use. It applies to medicines for human use that are manufactured or imported by natural and legal persons for marketing in the territory of the States Parties. It does not cover magistral preparations.

The purpose of the present Addendum is to notify WTO Members that this Technical Regulation has been revised and that there will be a comment period of 60 days as from the date of this notification.

The relevant document is available from:

http://reglatec.go.cr/reglatec/principal.jsp

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