



## NOTIFICATION

### *Addendum*

The following communication, dated March 12<sup>th</sup>, 2020, is being circulated at the request of the delegation of Brazil.

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The Normative Instruction number 36, August 21<sup>st</sup>, 2019 – previously notified through [G/TBT/N/BRA/870/Add.3](#) – which adopts Complementary guidelines to the manufacture of biological medicinal substances and products for human use of the Pharmaceutical Inspection Convention and Pharmaceutical Inspection Co-operation Scheme (PIC/S) of July 1<sup>st</sup>, 2018, was rectified.

The final text is available only in Portuguese and can be downloaded at:

[http://portal.anvisa.gov.br/documents/10181/5389382/RET\\_IN\\_35\\_2019.pdf/f54076ca-e23f-4410-adab-b9b786bc670a](http://portal.anvisa.gov.br/documents/10181/5389382/RET_IN_35_2019.pdf/f54076ca-e23f-4410-adab-b9b786bc670a)

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