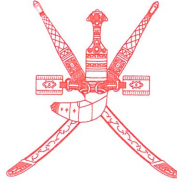


# Sultanate of Oman

Ministry of Health

Directorate General of Pharmaceutical Affairs  
and Drug Control  
MUSCAT



سِلْطَنَة عُومَان  
وَزَارَة الصِّحَّة  
المُتَّخِذَة الْعَامَّة لِالصِّيْدَة  
وَالرَّقَابَة الدَّوَاءِيَّة  
مَسْقَط

To:

**ALL MARKETING AUTHORISATION HOLDERS (MAHs)  
LOCAL PHARMACEUTICAL MANUFACTURERS  
LOCAL AGENTS**

**After Compliments,**

Enclosed please find our Circular No. <sup>circ 80</sup>..... dated  
10/12/18... regarding submission of CIOMS to Department of  
Pharmacovigilance & Drug Information.

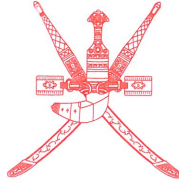
Cc:

**DG – for kind inf.  
SH (PV-Human Medicines)  
SH (PV-Herbal Medicines)  
Supdt. of Drug Information**

**Sultanate of Oman**

**Ministry of Health**

**Directorate General of Pharmaceutical Affairs  
and Drug Control  
MUSCAT**



سَلْطَنَةُ عُومَانِ  
وَزَارَةُ الصَّحَّةِ  
الْمُنِيرِيَّةِ الْعَامَّةِ لِلصِّيْدَانَةِ  
وَالرَّقَابَةِ الدَّرْوَائِيَّةِ  
مَسْقَط

**Circular No. ٨٥ / 2018**

٥2 -04-1440 H

١٥ -12-2018

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**Submission of CIOMS to Department of Pharmacovigilance & Drug Information**

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In order to simplify the reporting procedures for marketing authorisation holders (MAHs) and Pharmaceutical companies, please note that those who can create CIOMS in the ICH E2B-R2 (xml) format, can submit the CIOMS forms to the email ID: [xml-pv@moh.gov.om](mailto:xml-pv@moh.gov.om)

Kindly note that in each submission, there should not be more than 20 reports per mail.

For reporting timeframe, please refer to “Guideline on GVP in Oman for MAH/Pharmaceutical companies” Version 1/2017.

For companies who do not have this facility can continue using the MOH e-portal.

Effective Date: From the Date of this Circular

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**Ph. Hussain Al Ramimmy**  
**Director of Pharmacovigilance & Drug Information**

