



26/05/2015 02:55 p.m.

cc:
bcc:

Subject: Fw: Pharmacovigilance Guideline



From: Devices/MOH
To: Mike Thompson <mike.thompson@paradise.net.nz>,
Cc: Devices <devices@moh.govt.nz>
Date: 26/05/2015 12:10 p.m.
Subject: Re: Pharmacovigilance Guideline
Sent by:

Dear Mike,

Thank you for your feedback on the guidance.

I have passed your comments on to the Clinical Risk Team.

Kind regards,

Medsafe
Clinical Leadership, Protection & Regulation
Ministry of Health
DDI: 04 819 6800
Fax: 04 819 6801

<http://www.medsafe.govt.nz>
<mailto:devices@moh.govt.nz>

Mike Thompson

26/05/2015 11:13:58 a.m.

From: Mike Thompson <mike.thompson@paradise.net.nz>
To: Devices <devices@moh.govt.nz>,
Date: 26/05/2015 11:13 a.m.
Subject: Pharmacovigilance Guideline

As mentioned on the phone, the draft PV guideline (consultation deadline was 15th May) is titled 'Therapeutic Products' but relates only to medicines, not medical devices. Either the title should be changed, or devices included as a sub-section. For completeness, the latter is preferable in my view.

Kind regards,

Mike

Mike Thompson | Ethos Consulting Group

ē' thōs n. distinctive character, spirit, attitude..

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Pharmacovigilance guideline - Medsafe consultation (draft 20Mar15).pdf
