

Mary Jordan

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**From:** Food Medicine Interface  
**Sent:** Thursday, 6 December 2018 4:28 PM  
**To:** s22  
**Subject:** RE: Grunbiotics Pty Ltd (Neurofolin) [DLM=Sensitive:Legal] [SEC=UNCLASSIFIED]

No problem, I've just added you now

s22

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**From:** s22  
**Sent:** Thursday, 6 December 2018 4:20 PM  
**To:** Food Medicine Interface <Food.MedicineInterface@foodstandards.gov.au>  
**Subject:** RE: Grunbiotics Pty Ltd (Neurofolin) [DLM=Sensitive:Legal] [SEC=UNCLASSIFIED]

Yes it would be useful to be copied into email distributions.

Thanks s22

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**From:** Food Medicine Interface  
**Sent:** Thursday, 6 December 2018 3:48 PM  
**To:** s22 <>  
**Subject:** RE: Grunbiotics Pty Ltd (Neurofolin) [DLM=Sensitive:Legal] [SEC=UNCLASSIFIED]

Hi

Yes, I would say the majority of goods fall into either of those general categories.

We don't meet regularly (either face to face or telecon), but mainly communicate via email. There has been one face to face meeting over the time the FMI Protocol has been in place and teleconference's are only organised if there's a dispute on the assessment of a product.

However it might be useful if I add you to the distribution list? You would then receive the FMI correspondence and regulators (and TGA) considerations when assessing products.

What do you think?

s22

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t s22

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From: s22  
Sent: Thursday, 6 December 2018 3:19 PM  
To: Food Medicine Interface <[Food.MedicineInterface@foodstandards.gov.au](mailto:Food.MedicineInterface@foodstandards.gov.au)>  
Subject: RE: Grunbiotics Pty Ltd (Neurofolin) [DLM=Sensitive:Legal] [SEC=UNCLASSIFIED]

s22 am I right in thinking that most products that are considered at the interface are either sports products or medical products?

I think it would be useful to attend FMI meetings so I get a sense of what regulators are thinking?

Possible??

Tx

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From: Food Medicine Interface  
Sent: Tuesday, 4 December 2018 10:40 AM  
To: s22 <[REDACTED]>  
Subject: FW: Grunbiotics Pty Ltd (Neurofolin) [DLM=Sensitive:Legal] [SEC=UNCLASSIFIED]

Hi [REDACTED]

I've sent you an invitation to share the letter mentioned below from the FMI folder (as I can forward an attachment), but here's the link again for reference:  
[http://teams/WG/FoodMed/Shared%20Documents/2018/Grunbiotics\\_non%20FMI/Grunbiotics%20Pty%20Ltd.pdf](http://teams/WG/FoodMed/Shared%20Documents/2018/Grunbiotics_non%20FMI/Grunbiotics%20Pty%20Ltd.pdf)

TGA and Victoria (home state) were both of the view that this product was a therapeutic and therefore 2.9.5 was not applicable.

Thanks

s22  
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t [REDACTED]

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From: s22  
Sent: Monday, 19 November 2018 3:55 PM  
To: Food Medicine Interface <[Food.MedicineInterface@foodstandards.gov.au](mailto:Food.MedicineInterface@foodstandards.gov.au)>  
Subject: Grunbiotics Pty Ltd (Neurofolin) [DLM=Sensitive:Legal]

Good afternoon,

I was advised by s22 to notify FSANZ about the communication the TGA have had with the company Grunbiotics Pty Ltd and its product Neurofolin.

I have attached a letter for your information only regarding the interpretation of 2.9.5. food for special medical purposes and the product Neurofolin.

Please don't hesitate to contact me should you have any queries.

Kind regards,

[REDACTED]

s22 [REDACTED]

Regulatory Compliance Section  
Regulatory, Education and Compliance Branch

Phone: [REDACTED]

Mobile: s22 [REDACTED]

Email: [REDACTED]

Therapeutic Goods Administration  
Department of Health  
PO Box 100  
Woden ACT 2606  
[www.tga.gov.au](http://www.tga.gov.au)



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