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## **No. S 817**

### **HEALTH PRODUCTS ACT 2007**

#### **HEALTH PRODUCTS (THERAPEUTIC PRODUCTS) (AMENDMENT NO. 3) REGULATIONS 2024**

In exercise of the powers conferred by section 72 of the Health Products Act 2007, the Health Sciences Authority, with the approval of the Minister for Health, makes the following Regulations:

#### **Citation and commencement**

1. These Regulations are the Health Products (Therapeutic Products) (Amendment No. 3) Regulations 2024 and come into operation on 1 November 2024.

#### **Amendment of First Schedule**

2. In the Health Products (Therapeutic Products) Regulations 2016 (G.N. No. S 329/2016), in the First Schedule, in paragraph 1, after the item “Bromazepam”, insert —

“Bromazolam”.

#### **Amendment of Second Schedule**

3. In the Health Products (Therapeutic Products) Regulations 2016, in the Second Schedule —

- (a) in Part 1, after each item specified in the first column of the following table, insert the item specified opposite it in the second column of the table:

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<i>First column</i>	<i>Second column</i>
<i>Item</i>	<i>Item</i>
Androsterone	Andusomeran
Auranofin	Avalglucosidase alfa
Bimatoprost	Binimetinib
Bromazepam	Bromazolam
Chlormethiazole	Chlormethine
Desvenlafaxine	Deucravacitinib
Eletriptan	Elobixibat
Encainide	Encorafenib
Etoxeridine	Etrasimod
Fexofenadine	Filgotinib
Fosinopril	Fosnetupitant
Gallium	Galsulfase
Gliquidone	Glofitamab
Goserelin	Gozetotide
Morphine	Mosunetuzumab
Niridazole	Nirmatrelvir
Obinutuzumab	Ocrelizumab
Regorafenib	Relatlimab
Romosozumab	Ropeginterferon
Selinexor	Selpercatinib
Somatostatin	Somatrogon
Spectinomycin	Spesolimab
Tazobactam	Teclistamab
Trazodone	Tremelimumab

; and

(b) in Part 3, delete the item “Policresulen”.

*[G.N. Nos. S 219/2017; S 119/2018; S 92/2019;  
S 969/2020; S 732/2021; S 1081/2021; S 458/2022;  
S 436/2023; S 681/2023; S 811/2023; S 610/2024;  
S 691/2024]*

Made on 12 October 2024.

BENJAMIN ONG  
*Chairperson,  
Health Sciences Authority,  
Singapore.*

[401:04/01-000; AG/LEGIS/SL/122D/2020/4]

(To be presented to Parliament under section 72(5) of the Health Products Act 2007).