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WARTA KERAJAAN PERSEKUTUAN

*FEDERAL GOVERNMENT
GAZETTE*

PERINTAH RACUN (PINDAAN SENARAI RACUN) 2014

POISONS (AMENDMENT OF POISONS LIST) ORDER 2014



DISIARKAN OLEH/
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AKTA RACUN 1952

PERINTAH RACUN (PINDAAN SENARAI RACUN) 2014

PADA menjalankan kuasa yang diberikan oleh seksyen 6 Akta Racun 1952 [*Akta 366*], Menteri, selepas berunding dengan Lembaga Racun, membuat perintah yang berikut:

Nama

1. Perintah ini bolehlah dinamakan **Perintah Racun (Pindaan Senarai Racun) 2014**.

Pindaan Jadual Pertama

2. Akta Racun 1952 dipinda dalam Jadual Pertama—

(1) dalam Senarai Racun—

(a) dengan memasukkan selepas butiran “Antimony; its chlorides, oxides, sulphides, antimonates, antimonites; organic compounds of antimony” dan butir-butir yang berhubungan dengannya butiran yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Apixaban	-	All preparations”;				

(b) dalam butiran “Arsenic; its organic and inorganic compounds”, dengan menggantikan butir-butir yang berhubungan dengannya dengan butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Arsenic; its organic and inorganic compounds	-	-	All medicinal preparations unless exempted	Preparations other than medicinal preparations unless exempted	-	(1) Cosmetic which is notified under the Control of Drugs and Cosmetics Regulations 1984 [P.U. (A) 223/1984] containing not more than 5 parts per million of arsenic calculated as the metal; (2) Product which is registered under the Control of Drugs and Cosmetic Regulations 1984 [P.U. (A) 223/1984] containing not more than 5 parts per million of arsenic calculated as the metal”;

(c) dengan memasukkan selepas butiran “Atracurium; its salts” dan butir-butir yang berhubungan dengannya butiran yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Avanafil	-	All preparations”;				

- (d) dengan memasukkan selepas butiran “Avoparcin” dan butir-butir yang berhubungan dengannya butiran yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Axitinib	-	All preparations”;				

- (e) dalam butiran “Mercury”, dengan menggantikan butir-butir yang berhubungan dengannya dengan butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Mercury	-	-	-	All preparations unless exempted	-	(1) Cosmetic which is notified under the Control of Drugs and Cosmetics Regulations 1984 [P.U. (A) 223/1984] containing not more than 1 part per million of mercury calculated as the metal; (2) Product which is registered under the Control of Drugs and Cosmetic Regulations 1984 [P.U. (A) 223/1984] containing not more than 0.5 part per million of mercury calculated as the metal”;

(f) dengan memasukkan selepas butiran “Narcotic substances, the following: their isomers (whenever the existence of such isomers is possible within the specific chemical designation); their salts; their esters and ethers (whenever the existence of such esters or ethers is possible); the salts of their esters and ethers” dan butir-butir yang berhubungan dengannya butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Natalizumab	-	All preparations”;				

(g) dalam butiran “Natamycin”, dengan menggantikan butir-butir yang berhubungan dengannya dengan butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Natamycin	-	-	All preparations unless exempted	-	-	Natamycin when use as food additive”;

- (h) dengan memasukkan selepas butiran “Perindopril” dan butir-butir yang berhubungan dengannya butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Pertuzumab	-	All preparations”;				

- (i) dalam butiran “Potassium hydroxide”, dengan menggantikan butir-butir yang berhubungan dengannya dengan butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Potassium hydroxide	-	-	-	-	12% and over unless exempted	Under 12%, accumulators, batteries or when use as food additive”;

- (j) dengan memasukkan selepas butiran “Practolol; its salts” dan butir-butir yang berhubungan dengannya butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Pralatrexate	-	All preparations”;				

- (k) dengan memasukkan selepas butiran “Recombinant-Methionyl Human Granulocyte-colony stimulating factor” dan butir-butir yang berhubungan dengannya butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Regorafenib	-	All preparations”;				

- (l) dalam butiran “Sodium hydroxide”, dengan menggantikan butir-butir yang berhubungan dengannya dengan butir-butir yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Sodium hydroxide	-	-	All preparations for therapeutic or diagnostic use	-	All preparations containing 12% and over other than those in Part I or exempted and subject to the provisions of Poisons (Sodium Hydroxide) Regulations, 1962 [L.N. 145/62]	Under 12%; Sodium hydroxide when use as food additive”; dan

- (m) dengan memasukkan selepas butiran “Vecuronium; its salts” dan butir-butir yang berhubungan dengannya butiran yang berikut:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>	
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>			
“Vemurafenib	-	All preparations”; dan					

- (2) dalam Lampiran Senarai Racun—

- (a) dalam senarai “**Antineoplastic agents and Immunosuppressants**”—

- (i) dengan memasukkan selepas butiran “L-Asparaginase” butiran “Axitinib”;
- (ii) dengan memasukkan selepas butiran “Mycophenolic acid” butiran “Natalizumab”;
- (iii) dengan memasukkan selepas butiran “Penciclovir” butiran berikut:
 - “Pertuzumab
 - Pralatrexate”;
- (iv) dengan memasukkan selepas butiran “Rasburicase” butiran “Regorafenib”; dan
- (v) dengan memasukkan selepas butiran “Trofosfamide” butiran “Vemurafenib”.

- (b) dalam senarai “**Vasodilators**”, dengan memasukkan selepas butiran “Amyl nitrite” butiran “Avanafil”; dan

- (c) dalam senarai “**Miscellaneous**”, dengan memasukkan selepas butiran “Anticoagulant substances including heparin and other substances structurally derived from coumarin and phenindione unless specified elsewhere in the List” butiran “Apixaban”.

Dibuat 11 Februari 2014
[KK(R)608(14); PN(PU2)172/XII]

DATUK SERI DR. S. SUBRAMANIAM
Menteri Kesihatan

POISONS ACT 1952

POISONS (AMENDMENT OF POISONS LIST) ORDER 2014

IN exercise of the powers conferred by section 6 of the Poisons Act 1952 [Act 366], the Minister, after consultation with the Poisons Board, makes the following order:

Citation

1. This order may be cited as the **Poisons (Amendment of Poisons List) Order 2014**.

Amendment of First Schedule

2. The Poisons Act 1952 is amended in the First Schedule—

(1) in the Poisons List—

(a) by inserting after the item “Antimony; its chlorides, oxides, sulphides, antimonates, antimonites; organic compounds of antimony” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Apixaban	-	All preparations”;				

- (b) in the item “Arsenic; its organic and inorganic compounds”, by substituting for the particulars relating to it the following particulars:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Arsenic; its organic and inorganic compounds	-	-	All medicinal preparations unless exempted	Preparations other than medicinal preparations unless exempted	-	(1) Cosmetic which is notified under the Control of Drugs and Cosmetics Regulations 1984 [P.U. (A) 223/1984] containing not more than 5 parts per million of arsenic calculated as the metal; (2) Product which is registered under the Control of Drugs and Cosmetic Regulations 1984 [P.U. (A) 223/1984] containing not more than 5 parts per million of arsenic calculated as the metal”;

- (c) by inserting after the item “Atracurium; its salts” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Avanafil	-	All preparations”;				

(d) by inserting after the item “Avoparcin” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Axitinib	-	All preparations”;				

(e) in the item “Mercury”, by substituting for the particulars relating to it the following particulars:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Mercury	-	-	-	All preparations unless exempted	-	(1) Cosmetic which is notified under the Control of Drugs and Cosmetics Regulations 1984 [P.U. (A) 223/1984] containing not more than 1 part per million of mercury calculated as the metal; (2) Product which is registered under the Control of Drugs and Cosmetic Regulations 1984 [P.U. (A) 223/1984] containing not more than 0.5 part per million of mercury calculated as the metal”;

- (f) by inserting after the item “Narcotic substances, the following: their isomers (whenever the existence of such isomers is possible within the specific chemical designation); their salts; their esters and ethers (whenever the existence of such esters or ethers is possible); the salts of their esters and ethers” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Natalizumab	-	All preparations”;				

- (g) in the item “Natamycin”, by substituting for the particulars relating to it the following particulars:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Natamycin	-	-	All preparations unless exempted	-	-	Natamycin when use as food additive”;

- (h) by inserting after the item “Perindopril” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Pertuzumab	-	All preparations”;				

(i) in the item “Potassium hydroxide”, by substituting for the particulars relating to it the following particulars:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Potassium hydroxide	-	-	-	-	12% and over unless exempted	Under 12%, accumulators, batteries or when use as food additive”;

(j) by inserting after the item “Practolol; its salts” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Pralatrexate	-	All preparations”;				

(k) by inserting after the item “Recombinant-Methionyl Human Granulocyte-colony stimulating factor” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Regorafenib	-	All preparations”;				

(l) in the item “Sodium hydroxide”, by substituting for the particulars relating to it the following particulars:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>		
“Sodium hydroxide	-	-	All preparations for therapeutic or diagnostic use	-	All preparations containing 12% and over other than those in Part I or exempted and subject to the provisions of Poisons (Sodium Hydroxide) Regulations, 1962 [L.N. 145/62]	Under 12%; Sodium hydroxide when use as food additive”; and

(m) by inserting after the item “Vecuronium; its salts” and the particulars relating to it the following item:

<i>Names</i>	<i>Part I</i>				<i>Part II</i>	<i>Exempt</i>	
	<i>Group A</i>	<i>Group B</i>	<i>Group C</i>	<i>Group D</i>			
“Vemurafenib	-	All preparations”; and					

(2) in the Appendix to the Poisons List—

(a) in the list of “**Antineoplastic agents and Immunosuppressants**”—

(i) by inserting after the item “L-Asparaginase” the item “Axitinib”;

(ii) by inserting after the item “Mycophenolic acid” the item “Natalizumab”;

(iii) by inserting after the item “Penciclovir” the following items:

“Pertuzumab
Pralatrexate”;

(iv) by inserting after the item “Rasburicase” the item “Regorafenib”; and

(v) by inserting after the item “Trofosfamide” the item “Vemurafenib”.

(b) in the list of “**Vasodilators**”, by inserting after the item “Amyl nitrite” the item “Avanafil”; and

(c) in the list of “**Miscellaneous**”, by inserting after the items “Anticoagulant substances including heparin and other substances structurally derived from coumarin and phenindione unless specified elsewhere in the List” the item “Apixaban”.

Made 11 February 2014
[KK(R)608(14); PN(PU2)172/XII]

DATUK SERI DR. S. SUBRAMANIAM
Minister of Health