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WARTA KERAJAAN PERSEKUTUAN

*FEDERAL GOVERNMENT
GAZETTE*

PERINTAH RACUN (PINDAAN SENARAI RACUN) 2015

POISONS (AMENDMENT OF POISONS LIST) ORDER 2015



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AKTA RACUN 1952

PERINTAH RACUN (PINDAAN SENARAI RACUN) 2015

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) 2015**.

Pindaan Jadual Pertama

2. Akta Racun 1952 dipinda dalam Jadual Pertama—

(a) dalam Senarai Racun—

- (i) dengan menggantikan butiran “Abrisentan” dengan butiran “Ambrisentan”;
- (ii) dengan memasukkan selepas butiran “Alphadolone acetate” 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>		
“Alpha-Phenylacetoacetonitrile (APAAN)”	-	-	-	-	All preparations”;	

- (iii) dengan menggantikan butiran “Amilsulpride” dengan butiran “Amisulpride”;
- (iv) dengan menggantikan butiran “Anistreplace” dengan butiran “Anistreplase”;
- (v) dalam butiran “Antihistamines; the following:”—
 - (A) dengan menggantikan butiran “Dimenhydramine” dengan butiran “Dimenhydrinate”; dan
 - (B) dengan menggantikan butiran “Levocetirizine” dengan butiran “Levocetirizine”;
- (vi) dengan menggantikan butiran “Calcium polysterene” dengan butiran “Calcium polystyrene”;
- (vii) dengan memasukkan selepas butiran “Camylofine; 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>		
“Canakinumab	-	All preparations”;				

- (viii) dengan memasukkan selepas butiran “Cerivastatin; 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>		
“Certolizumab	-	All preparations”;				

- (ix) dengan memasukkan selepas butiran “2, 5-Dimethoxy-4-ethylamphetamine (DOET) (DD)” 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>		
“1,3-Dimethylamylamine (DMAA)	All preparations”;					

- (x) dengan memasukkan selepas butiran “Dinitrothymols” 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>		
“Diphenidol hydrochloride	-	-	All preparations”;			

- (xi) dengan memotong butiran “Fexofenadine; its salt” dan butir-butir yang berhubungan dengannya;

- (xii) dengan memasukkan selepas butiran “Gallamine; its salts; its quarternary compounds” 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>		
“Gamma Butyrolactone	-	-	-	-	All preparations unless exempted	When use as food additive in food”;

- (xiii) dengan memasukkan selepas butiran “Indomethacin; 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>		
“Infliximab	-	All preparations”;				

- (xiv) dengan menggantikan butiran “Lanthanum” 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>		
“Lanthanum carbonate	-	All preparations for therapeutic use”;				

- (xv) dengan memasukkan selepas butiran “Mipomersen sodium” 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>		
“Mirabegron	-	All preparations”;				

- (xvi) 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 in food”;

- (xvii) 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 in food”;

(xviii) 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	Under 12%; sodium hydroxide when use as food additive in food”;

(xix) dengan memotong butiran “Terfenadine” dan butir-butir yang berhubungan dengannya; dan

(xx) dengan memasukkan selepas butiran “Tofacitinib; 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>		
“Tofacitinib	-	All preparations”; dan				

(b) dalam Lampiran kepada Senarai Racun—

(i) dalam senarai “**Anticholinergics:**”—

(A) dengan memasukkan selepas butiran “Diethazine; its salts” butiran “Diphenidol hydrochloride”; dan

(B) dengan memasukkan selepas butiran “Methixene; its salts” butiran “Mirabegron”;

(ii) dalam senarai “**Antihistamines:**”—

(A) dengan memasukkan selepas butiran “3-Dibutylaminoethyl-4, 5, 6-trihydroxy-1-isobenzofuranone” butiran “Dimenhydrinate”; dan

(B) dengan menggantikan butiran “Levocetirizine” dengan butiran “Levocetirizine”;

(iii) dalam senarai “**Antineoplastic agents and Immunosuppressants:**”—

(A) dengan memasukkan selepas butiran “Cabazitaxel” butiran “Canakinumab”;

(B) dengan memasukkan selepas butiran “Carmustine; its salts” butiran “Certolizumab”;

(C) dengan memasukkan selepas butiran “Idarubicin” butiran “Infliximab”; dan

- (D) dengan memasukkan selepas butiran “Tocilizumab” butiran “Tofacitinib”;
- (iv) dalam senarai “**Cardio drugs:**”—
 - (A) dengan menggantikan butiran “Abrisentan” dengan butiran “Ambrisentan”; dan
 - (B) dengan menggantikan butiran “Anistreplase” dengan butiran “Anistreplase”;
- (v) dalam senarai “**Industrial and laboratory poisons:**”—
 - (A) dengan memasukkan selepas butiran “Acetyl chloride” butiran “Alpha-Phenylacetoacetonitrile (APAAN)”;
dan
 - (B) dengan memasukkan selepas butiran “Formaldehyde” butiran “Gamma Butyrolactone”;
- (vi) dalam senarai “**Sympathomimetic amines:**”, dengan memasukkan selepas butiran “Atomoxetine; its salts” butiran “1,3- Dimethylamylamine (DMAA)”;
- (vii) dalam senarai “**Tranquillisers:**”, dengan menggantikan butiran “Amisulpride” dengan butiran “Amisulpride”; dan
- (viii) dalam senarai “**Miscellaneous:**”—

- (A) dengan menggantikan butiran “Calcium polysterene” dengan butiran “Calcium polystyrene”; dan
- (B) dengan menggantikan butiran “Lanthanum” dengan butiran “Lanthanum carbonate”.

Dibuat 26 Januari 2015
[KK(R)608(14); PN(PU2)172/XIII]

DATUK SERI DR. S. SUBRAMANIAM
Menteri Kesihatan

POISONS ACT 1952

POISONS (AMENDMENT OF POISONS LIST) ORDER 2015

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 2015**.

Amendment of First Schedule

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

(a) in the Poisons List—

- (i) by substituting for the item “Abrisentan” the item “Ambrisentan”;
- (ii) by inserting after the item “Alphadolone acetate” and the particulars relating to it the following item:

Names	Part I				Part II	Exempt
	Group A	Group B	Group C	Group D		
“Alpha-Phenylacetoacetonitrile (APAAN)”	-	-	-	-	All preparations”;	

- (iii) by substituting for the item “Amilsulpride” the item “Amisulpride”;
- (iv) by substituting for the item “Anistreplace” the item “Anistreplase”;
- (v) in the item “Antihistamines; the following:”—
- (A) by substituting for the item “Dimenhydramine” the item “Dimenhydrinate”; and
- (B) by substituting for the item “Levocetazine” the item “Levocetirizine”;
- (vi) by substituting for the item “Calcium polysterene” the item “Calcium polystyrene”;
- (vii) by inserting after the item “Camylofine; 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>		
“Canakinumab	-	All preparations”;				

- (viii) by inserting after the item “Cerivastatin; 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>		
“Certolizumab	-	All preparations”;				

- (ix) by inserting after the item “2, 5-Dimethoxy-4-ethylamphetamine (DOET) (DD)” 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>		
“1,3-Dimethylamylamine (DMAA)	All preparations”;					

- (x) by inserting after the item “Dinitrothymols” 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>		
“Diphenidol hydrochloride	-	-	All preparations”;			

- (xi) by deleting the item “Fexofenadine; its salt;” and the particulars relating to it;

- (xii) by inserting after the item “Gallamine; its salts; its quarternary compounds” 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>		
“Gamma Butyrolactone	-	-	-	-	All preparations unless exempted	When use as food additive in food”;

(xiii) by inserting after the item “Indomethacin; 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>		
“Infliximab	-	All preparations”;				

(xiv) by substituting for the item “Lanthanum” and 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>		
“Lanthanum carbonate	-	All preparations for therapeutic use”;				

(xv) by inserting after the item “Mipomersen sodium” 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>		
“Mirabegron	-	All preparations”;				

(xvi) 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 in food”;

(xvii) 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 in food”;

(xviii) 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	Under 12%; sodium hydroxide when use as food additive in food”;

(xix) by deleting the item “Terfenadine” and the particulars relating to it; and

(xx) by inserting after the item “Todrazoline; 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>		
“Tofacitinib	-	All preparations”; and				

(b) in the Appendix to the Poisons List—

(i) in the list of “**Anticholinergics:**”—

(A) by inserting after the item “Diethazine; its salts” the item “Diphenidol hydrochloride”; and

(B) by inserting after the item “Methixene; its salts” the item “Mirabegron”;

(ii) in the list of “**Antihistamines:**”—

(A) by inserting after the item “3-Dibutylaminomethyl-4, 5, 6-trihydroxy-1-isobenzofuranone” the item “Dimenhydrinate”; and

(B) by substituting for the item “Levocetizine” the item “Levocetirizine”;

- (iii) in the list of “**Antineoplastic agents and Immunosuppressants:**” —
 - (A) by inserting after the item “Cabazitaxel” the item “Canakinumab”;
 - (B) by inserting after the item “Carmustine; its salts” the item “Certolizumab”;
 - (C) by inserting after the item “Idarubicin” the item “Infliximab”; and
 - (D) by inserting after the item “Tocilizumab” the item “Tofacitinib”;
- (iv) in the list of “**Cardio drugs:**” —
 - (A) by substituting for the item “Abrisentan” the item “Ambrisentan”; and
 - (B) by substituting for the item “Anistreplase” the item “Anistreplase”;
- (v) in the list of “**Industrial and laboratory poisons:**” —
 - (A) by inserting after the item “Acetyl chloride” the item “Alpha-Phenylacetoacetonitrile (APAAN)”; and
 - (B) by inserting after the item “Formaldehyde”, the item “Gamma Butyrolactone”;

- (vi) in the list of “**Sympathomimetic amines:**”, by inserting after the item “Atomoxetine; its salts” the item “1,3- Dimethylamylamine (DMAA)”;
- (vii) in the list of “**Tranquillisers:**”, by substituting for the item “Amilsulpride” the item “Amisulpride”; and
- (viii) in the list of “**Miscellaneous:**”—
 - (A) by substituting for the item “Calcium polysterene” the item “Calcium polystyrene”; and
 - (B) by substituting for the item “Lanthanum” the item “Lanthanum carbonate”.

Made 26 January 2015
[KK(R)608(14); PN(PU2)172/XIII]

DATUK SERI DR. S. SUBRAMANIAM
Minister of Health