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No. S 488

INTERPRETATION ACT 1965

INTERPRETATION (HEALTH SCIENCES AUTHORITY ACT — FEES) (AMENDMENT) ORDER 2024

In exercise of the powers conferred by section 46(1) of the Interpretation Act 1965, the Minister for Health makes the following Order:

Citation and commencement

1.—(1) This Order is the Interpretation (Health Sciences Authority Act — Fees) (Amendment) Order 2024 and, except for paragraph 3, comes into operation on 1 July 2024.

(2) Paragraph 3 is deemed to have come into operation on 31 December 2021.

Replacement of Schedule

2. In the Interpretation (Health Sciences Authority Act — Fees) Order 2017 (G.N. No. S 577/2017), replace the Schedule with —

"THE SCHEDULE

Paragraph 3

FEES

	First column	*Second column*
1.	Consultancy services on pre-registration requirements, to be provided to a person who is developing a medical device intended for registration under the Health Products Act 2007 —	
	(*a*) where the supply of services is a supply of prescribed international services under section 21(3)(*k*) of the Goods and Services Tax Act 1993	$486.24 per consultation, where each consultation is of a duration of 2 hours or shorter
	(*b*) where the supply of services is other than as described in paragraph (*a*)	$530 per consultation, where each consultation is of a duration of 2 hours or shorter
2.	Consultancy services on registration application requirements —	
	(*a*) where the supply of services is a supply of prescribed international services under section 21(3)(*k*) of the Goods and Services Tax Act 1993	$192.66 per consultation, where each consultation is of a duration of one hour or shorter
	(*b*) where the supply of services is other than as described in paragraph (*a*)	$210 per consultation, where each consultation is of a duration of one hour or shorter

Note:

(1) In this Schedule —

"pre-registration requirements" means the requirements imposed by or under the Health Products Act 2007 relating to the registration of medical devices, other than the registration application requirements;

"registration application requirements" means the particulars, documents, information and samples required for an application for registration of a medical device or cell, tissue or gene therapy product under section 30 of the Health Products Act 2007.".

Miscellaneous amendments

3. In the Interpretation (Health Sciences Authority Act — Fees) Order 2017 —

(*a*) in the following provisions, replace "(Cap. 122C)" with "2001":

Paragraph 2, definition of "Authority"

Paragraph 3; and

(*b*) in paragraph 2, in the definitions of "cell, tissue or gene therapy product" and "medical device", replace "(Cap. 122D)" with "2007".

[G.N. No. S 102/2021]

Made on 30 May 2024.

CHAN YENG KIT
Permanent Secretary,
Ministry of Health,
Singapore.

[401:04/05-000; AG/LEGIS/SL/1/2020/1]