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INTERPRETATION ACT (CHAPTER 1)

INTERPRETATION (HEALTH SCIENCES AUTHORITY ACT — FEES) ORDER 2017

ARRANGEMENT OF PARAGRAPHS

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

Citation and commencement

1. This Order is the Interpretation (Health Sciences Authority Act — Fees) Order 2017 and comes into operation on 11 October 2017.

Definitions

2. In this Order —

"Authority" means the Health Sciences Authority established under section 3 of the Health Sciences Authority Act (Cap. 122C);

"medical device" means a health product categorised as a medical device in the First Schedule to the Health Products Act (Cap. 122D).

Fees for consultancy services

3. The fees specified in the second column of the Schedule are payable to the Authority for the consultancy services under section 12(*d*) of the Health Sciences Authority Act (Cap. 122C) specified opposite in the first column.

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 (Cap. 122D)	$500 per consultation, where each consultation is of a duration of 2 hours or shorter
2. Consultancy services on registration application requirements	$200 per consultation, where each consultation is of a duration of one hour or shorter

Note:

(1) Each fee specified in this Schedule is inclusive of goods and services tax chargeable under the Goods and Services Tax Act (Cap. 117A) on the supply of a service for which the fee is payable.

(2) Where the supply of services specified in this Schedule is a supply of prescribed international services under section 21(3)(*k*) of the Goods and Services Tax Act, no goods and services tax under that Act is chargeable on those services.

(3) In this Schedule —

"pre-registration requirements" means the requirements imposed by or under the Health Products Act (Cap. 122D) 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 under section 30 of the Health Products Act.

Made on 5 October 2017.

CHAN HENG KEE
Permanent Secretary,
Ministry of Health,
Singapore.

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