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No. S 1077

**HEALTH PRODUCTS ACT
(CHAPTER 122D)**

**HEALTH PRODUCTS (CLINICAL TRIALS)
(AMENDMENT NO. 3) REGULATIONS 2021**

In exercise of the powers conferred by section 72(1) of the Health Products Act, 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 (Clinical Trials) (Amendment No. 3) Regulations 2021 and come into operation on 3 January 2022.

Amendment of regulation 2

2. Regulation 2(1) of the Health Products (Clinical Trials) Regulations 2016 (G.N. No. S 331/2016) is amended by deleting the definition of “licensed healthcare institution” and substituting the following definition:

““licensed healthcare institution” means —

- (a) any premises or conveyance specified in a licence granted under the Healthcare Services Act 2020 for the provision of any licensable healthcare service; or
- (b) a healthcare institution that is licensed under the Private Hospitals and Medical Clinics Act 1980;”.

[G.N. Nos. S 107/2021; S 731/2021]