ASAL
ORIGINAL

PIHAK BERKUASA
PERANTI PERUBATAN

MEDICAL DEVICE
AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran: **GA9442926-244979**
Registration No.:

Tarikh Sah Pendaftaran: **14/10/2026 - 13/10/2031**
Registration Validity Date:

Sijil ini adalah dengan ini diberi kepada: **GRZAN GROUP SDN. BHD.**
This certificate is hereby issued to:

yang beralamat di:
which is located at:

B39-7, THE SENTRAL RESIDENCES, JALAN STESEN SENTRAL 2,
50470 KUALA LUMPUR

Peranan establismen
Role of establishment

Wakil Diberi Kuasa
Authorized Representative

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

MURALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

LAMPIRAN 1
Attachment 1

No. Pendaftaran: **GA9442926-244979**
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Sijil ini adalah dengan ini diberi kepada: **GRZAN GROUP SDN. BHD.**
This certificate is hereby issued to:

Butir-butir peranti perubatan yang didaftarkan
Particulars of the registered medical device

Nama Peranti Perubatan *Medical Device Name*	**VR Cognitive Assessment and Rehab Training System**		
Kelas *Class*	**CLASS A**	Jenama *Brand*	**QuickMind**
Kelompok *Group*	**SINGLE**	Produk Identifikasi *Product Identifier*	**VIRTUAL BRAIN**
Nama dan alamat pembuat: *Name and address of manufacturer*	**YIWEI MEDICAL TECHNOLOGY CO., LTD ROOM 701-2, BLOCK B, SHENZHEN INTERNATIONAL INNOVATION CENTER, NO. 1006 SHENNAN BOULEVARD, XINTIAN COMMUNITY, HUAFU SUBDISTRICT, FUTIAN DISTRICT, SHENZHEN, GUANGDONG, 518000, CHINA 518000, CHINA**		

Medical Device AUTHORITY MALAYSIA