BYOND HEALTHCARE

Byond Healthcare, a SAHPRA licenced distributor, importer and exporter of medical devices is excited to launch the following bioelectronic medicines (BEM): portable and convenient, non-invasive Vagus Nerve Stimulation (nVNS) devices.

Please see video for an overview on nVNS

gammaCore Sapphire™

gammaCore Sapphire™, an FDA, EU CE, TGA and Health Canada cleared medical device stimulating the cervical branch of the vagus nerve. GammaCore has been cleared by the FDA for the prevention and acute treatment of migraines (age 12 years and older), adjunctive use for preventative treatment of cluster headaches (adults), acute treatment of pain associated with episodic cluster headache (adults) and treatment of hemicrania continua and paroxysmal hemicrania (adults). UK and TGA have also cleared gammaCore for medication overuse headaches (adults).

The Parasym™

The Parasym™, a CE cleared device, is a non-invasive Vagus Nerve Stimulator (VNS) intended to provide non-invasive auricular VNS delivered through the tragus. The Parasym device is indicated for the management of symptoms associated with depression, pain, anxiety and insomnia.

Option 1 - To purchase devices or find out more, scan the QR code or visit our product page.

OUR PRODUCTS

Option 2 - To become a partner and one of a growing number of HCPs offering cutting edge technology in a clinical practice setting directly to their patients scan the QR code or email info@byondhealthcare.com

NAPPI codes can be requested from Byond Healthcare for the individual device options.

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CONTACT US

REFERENCE 1: SAHPRA CLASS B non-IVD Medical Device, gammaCoreTM Sapphire (cervical non-invasive Vagus Nerve Stimulator), US FDA licence No. K211856, EU/UK CA licence no. CE 571753/571753, TGA licence No. ARTG 355575, Health Canada licence No. 105996. See important safety information and frequently asked questions on https://www.gammacore.com/about/important-safety-information/ and https://www.byondhealthcare.com/device-faqs. For full information, refer to latest approved instruction for use (IFU) included with the marketed device and cleared by US FDA, EU CE Mark, TGA Australia and Health Canada.

REFERENCE 2: SAHPRA CLASS B non-IVD Medical Device, ParasymTM tVNS (non-invasive transcutaneous auricular Vagus Nerve Stimulator), licence No. CE Mark CE0197. See important safety information and frequently asked questions on https://www.byondhealthcare.com/device-faqs. For full information, refer to the latest approved user guide/instruction for use (IFU).

NAME AND BUSINESS ADDRESS OF THE LICENCED DISTRIBUTOR: BYOND HEALTHCARE (PTY) LTD. Reg. No. 2020/790377/07. 1011 Courson Street, Le Domaine, Val de Vie Estate, Paarl, Western Cape, 7646, Republic of South Africa. E-mail: info@byondhealthcare.com.