



Medicines & Healthcare products
Regulatory Agency

**Medicines & Healthcare products
Regulatory Agency**

10 South Colonnade
Canary Wharf
London
E14 4PU
United Kingdom

gov.uk/mhra

**Viamed Limited
15 Station Road
Cross Hills
West Yorkshire
Keighley
BD20 7DT
England, United Kingdom**

14 May 2024

Dear **Stephen Nixon**

We are pleased to confirm that the application to register or update an existing registration for the following manufacturer, which you submitted on **30 April 2024** has been reviewed:

Application reference: **2024043001358644**

Manufacturer organisation: **Orantech Inc.**

Address:

Zone #A, 4F, 1st Bld.

7th Industrial Zone, Yulv Community,

Gongming,

Guangming, New District,

Guangdong

Shenzhen

518106

China

Manufacturer registration status: **Registered**

Device(s):

GMDN Code & Term	Status	Comment
31658 - Pulse oximeter probe, single-use	Registered	
37808 - Pulse oximeter probe, reusable	Registered	
37340 - General-purpose patient temperature probe, reusable	Registered	
35254 - General-purpose patient temperature probe, single-use	Registered	
46822 - Carbon dioxide sensor/cable	Registered	
41917 - Foetal Doppler system probe	Registered	
37326 - Blood pressure cuff, single-use	Registered	
35562 - Electrocardiographic lead set, reusable	Registered	
34978 - Blood pressure cuff, reusable	Registered	

Please note this letter **does not** represent any form of accreditation, certification or approval by the UK Competent Authority.

If you stop placing devices on the market or if you are not complying with the Regulations, you should inform us so that we can amend our records. You should be aware that it is an offence to place on the market UKCA or CE marked devices that do not comply with the regulations.

Please inform us of the following chargeable changes:

- 1. company/organisation information e.g. name and address**
- 2. additional devices (GMDN code or term)**

Please also use the Devices Online Registration Database (DORS) to tell us of the following changes e.g. removal/discontinuation of a device (GMDN) or product from your registration record, change of contact person, telephone number and/or email address, for which payment of our statutory fee does not apply.

Please note that the name and address of manufacturer, UK Responsible Person or Authorised Representative (Northern Ireland only) and devices that have been registered will be published on our [Public Access Registration Database](#) (PARD). In vitro diagnostic medical devices registered as undergoing performance evaluation study are not published on this database.

The account number for your company/organisation is **0000031562**.

Please do not respond directly to this email address. The originating email account is not monitored.

Yours sincerely,

A handwritten signature in black ink, appearing to read 'Ngozi Onyeukwu', with a stylized flourish at the end.

Ngozi Onyeukwu

Device registrations service

Devices division

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