



Medicines & Healthcare products
Regulatory Agency

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Regulatory Agency**

10 South Colonnade
Canary Wharf
London
E14 4PU
United Kingdom

gov.uk/mhra

**SHARE INFO LTD
3rd Floor, Office C, Townend House
Park Street
Walsall
West Midlands
WS1 1NS
England, United Kingdom**

22 June 2026

Dear **Simin Wang**

We are writing to inform you of the outcome of the application to register or update an existing registration for the following manufacturer, which you submitted on **16 June 2026**:

Application reference: **2026061601502069**

Manufacturer organisation: **Shenzhen Ruochuang Intelligent Manufacturing Co., Ltd.**

Address:

**Room 802, Building 1, Jia'an Science Park
No. 2 Longchang Road, Area 67, Xingdong Community
Xin'an Street, Bao'an District
Shenzhen
Guangdong
518101
China**

Manufacturer registration status: **Registered**

Device(s):

GMDN Code & Term	Status	Comment
41875 - Electric-motor-driven wheelchair, occupant-controlled	Registered	

Important Information:

Where new devices or device amendments within this application have been accepted and the status indicates 'Registered', this email confirmation does not represent any form of accreditation, certification or approval by the UK Competent Authority.

Where new devices within this application have been rejected, please review the reason/s for rejection. You will need to submit a new application. The [statutory fee](#) will be payable. Placing medical devices on the GB market that have not been registered with MHRA, or no longer comply with the regulations, constitutes a breach of the law. Different regulations apply to [Northern Ireland](#).

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

Keeping your Device Registration record up to date:

Please see [Making changes to your registration](#) for further information on changes you need to notify MHRA of, and the applicable [statutory fee](#). Full instructions on how to make changes are explained in our [Reference Guides](#) and [video tutorials](#).

Note:

The account number for your company/organisation is **0000038729**. Please keep a record of this.

Please do not reply directly to this email, as the originating email account is not monitored. Any queries must be sent to device.registrations@mhra.gov.uk.

Yours sincerely,



Ngozi Onyeukwu
MHRA Device Registration Service
 Data Assurance & Quality
 Healthcare, Quality & Access Group
 Medicines and Healthcare products Regulatory Agency
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