Medicines & Healthcare products Regulatory Agency

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10 South Colonnade
Canary Wharf
London
E14 4PU
United Kingdom

gov.uk/mhra

HDJ LIMITED
16
Churchfields Avenue
Feltham
England
Tw135pb
England, United Kingdom

08 October 2025

Dear **Abdul Razzaq**

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 **04 October 2025:**

Application reference: **2025100401439672**

Manufacturer organisation: **SCISSORS TOWN**
Address:
Sadra Badra Opposite Old Saga Yard Daska Road
Sialkot
Punjab
51310
Pakistan

Manufacturer registration status: **Registered**

Device(s):

GMDN Code & Term	Status	Comment
12726 - Suturing needle holder, reusable	Registered	

GMDN Code & Term	Status	Comment
38727 - General-purpose surgical scissors, reusable	Registered	
12235 - Scalpel handle, reusable	Registered	
10455 - Bone cutter	Registered	
42500 - Dressing/utility forceps, tweezers-like, reusable	Registered	
35048 - Bone lever/elevator, reusable	Registered	
33542 - Bone hook	Registered	
45918 - Hand-held surgical retractor, reusable	Registered	
35352 - Vaginal speculum, reusable	Registered	
33209 - Orthodontic pliers	Registered	
35350 - Nasal speculum, reusable	Registered	
31335 - Bone curette, reusable	Registered	
62466 - Surgical soft-tissue manipulation forceps, tweezers-like, reusable	Registered	

GMDN Code & Term	Status	Comment
61424 - Laparoscopic trocar blade, reusable	Registered	
11775 - Open-surgery biopsy forceps, reusable	Registered	
32853 - Orthopaedic joint/limb rongeur	Registered	
32312 - Surgical mallet	Registered	
12844 - Orthopaedic osteotome	Registered	
12245 - Ear knife	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 (PARD). 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 **0000036313.** 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
10 South Colonnade, Canary Wharf, London, E14 4PU
device.registrations@mhra.gov.uk