European Commission | EUDAMED

(https://ec.europa.eu)

Menu

- **Home**
- **Tasks**
- **Search & view**
- **Data Transfer**
- **News**
- **Help**

VIVEK NARANG

Logout

CURRENT ACTOR: Manufacturer, IN-MF-000018817, Narang Medical Limited [India]
Notifications

UDI-DI 08907097173612

Go back to the list

Manufacturer information

Organisation name: Narang Medical Limited

Actor ID/SRN: IN-MF-000018817

Address: Narang Tower New Delhi

Telephone number: -

Email: vivek@narang.com

Authorised Representative

Organisation name: CMC Medical Devices & Drugs SL

Eudamed Actor ID: ES-AR-000000293

Address: C/ Horacio Lengo N18 Málaga

Telephone number: -

Email: info@cmcmedicaldevices.com

Basic UDI-DI details

Version 1 [Current] | Last update date: -

Basic UDI-DI identification

Applicable regulation: MDR (REGULATION (EU) 2017/745 on medical devices)

Basic UDI-DI code: 8907097340.030XB

Issuing Entity: GS1

Is it a System or Procedure Pack which is a Device in itself?

System which is a device in itself

Risk class:	Class I
Implantable:	No
Measuring function:	No
Reusable surgical instruments:	Yes

Active device:	No
Device intended to administer and/or remove medicinal product:	No
Device model applicable:	Yes
Device model:	340.030
Name:	Self Centering Bone Holding Forceps, Size: 240mm

Tissues and cells

Presence of human tissues or cells, or their derivatives:	No
Presence of animal tissues or cells, or their derivatives:	No

Information on substances

Presence of a substance which, if used	No

separately, may be considered to be a medicinal product:	
Presence of a substance which, if used separately, may be considered to be a medicinal product derived from human blood or human plasma:	No

List of UDI-DIs for the Basic UDI-DI

UDI-DI details

Version 1 [Current] | Last update date: -

UDI-DI code:	08907097173612
Issuing Entity:	GS1

UDI-DI from another entity

UDI-DI from another entity (secondary) applicable:	No

Selected nomenclature codes

Code L091303 BONE REDUCTION FORCEPS, REUSABLE

Trade name

Trade name applicable:	Yes
Trade name:	NET [EN]
Reference/Catalogue number:	340.030

Is the device directly marked?

Is the device directly marked?:	Yes
*** Same as UDI-DI:**	Yes
*** Direct marking DI:**	08907097173612
*** Issuing Entity:**	GS1
Quantity of device:	1

Type of UDI-PI

Lot or Batch number:	Yes
Manufacturing date:	Yes

Additional product description:	Self Centering Bone Holding Forceps, Size: 240mm [EN]
URL for additional information (as electronic instructions for use):	-
UDI-DI status:	On the EU market

UDI-DI characteristics

Clinical size

Clinical size applicable:	No

Labelled as single use

Labelled as single use:	No
Maximum number of reuses applicable:	No
Maximum number of reuses:	-
Need for sterilisation	Yes

before use:	
Device labelled as sterile:	No
Containing latex:	No

CMR/Endocrine disruptor

Labelled for presence of Carcinogenic, Mutagenic and toxic to Reproduction (CMR) substances of category 1A or 1B:	No
Labelled for presence of substance(s) with endocrine-disrupting properties:	No

Storage/handling conditions

Storage/handling conditions, if applicable:	No

Critical warnings or contra-indications

Critical warnings or contra-indications, if applicable:	No
Reprocessed single use device:	No
Intended purpose other than medical (Annex XVI):	No

Information on substances

Presence of a substance which, if used separately, may be considered to be a medicinal product:	Not applicable
Presence of a substance which, if used separately, may be considered	Not applicable

to be a medicinal product derived from human blood or human plasma:	

Clinical Investigation(s)

Clinical Investigation

Clinical Investigation, if applicable:	No

Certificate information