

(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 08907097173636

≮ 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: 8907097340.050XH

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 No

function:

Reusable Yes

surgical

instruments:

Active No

device:

Device No

intended to administer and/or remove medicinal product:

Device model Yes

applicable:

Device 340.050

model:

Name: Self Centering Bone Holding

Forceps, Size: 280mm

Tissues and cells

Presence of No

human

tissues or

cells, or their

derivatives:

Presence of No

animal

tissues or

cells, or their

derivatives:

Information on substances

No

Presence of

a substance

which, if

used

separately,

may be

considered

to be a

medicinal

product:

Presence of

No

a substance

which, if

used

separately,

may be

considered

to be a

medicinal

product

derived from

human blood

or human

plasma:

List of UDI-DIs for the Basic UDI-DI

UDI-DI details

Version 1 [Current] Last update date: iii -

UDI-DI code: 08907097173636

Issuing GS1

Entity:

UDI-DI from another entity

UDI-DI from No

another

entity

(secondary)

applicable:

Selected nomenclature codes

Code L091303 BONE REDUCTION FORCEPS, REUSABLE

Trade name

Trade name

applicable:

Trade name: NET [EN]

Reference/Catalogue 0.050

number:

Is the device directly marked?

Yes

Is the device Yes

directly

marked?:

* Same as

Yes

UDI-DI:

* Direct

08907097173636

marking DI:

* Issuing

GS1

Entity:

Quantity of

1

device:

Type of UDI-PI

Lot or Batch

Yes

number:

Manufacturing

Yes

date:

Additional Self Centering Bone Holding product Forceps, Size: 280mm [EN]

description:

URL for -

additional information (as electronic instructions

for use):

UDI-DI On the EU market

status:

UDI-DI characteristics

Clinical size

Clinical size No

applicable:

Labelled as single use

Labelled as No

single use:

Maximum No

number of

reuses

applicable:

Maximum -

number of

reuses:

Need for Yes

sterilisation

before use:

Device No

labelled as

sterile:

Containing No

latex:

CMR/Endocrine disruptor

Labelled for No

presence of

Carcinogenic,

Mutagenic

and toxic to

Reproduction

(CMR)

substances

of category

1A or 1B:

Labelled for

No

presence of

substance(s)

with

endocrine-

disrupting

properties:

Storage/handling conditions

Storage/handling No conditions, if applicable:

Critical warnings or contra-indications

Critical No

warnings or

contra-

indications, if

applicable:

Reprocessed No

single use

device:

Intended

purpose

other than

medical

(Annex XVI):

Information on substances

No

Presence of

a substance Not applicable

which, if

used

separately,

may be

considered

to be a

medicinal

product:

Presence of

a substance Not applicable

which, if

used

separately,

may be

considered

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

Clinical Investigation(s)

Clinical Investigation

Clinical No Investigation, if applicable:

Certificate information