**Legal**
- Legal framework: N/A
- Authorizing legislation: N/A
- Guidelines: N/A

**National Regulatory Authority**
- National Regulatory Authority present: N/A
- Name: N/A
- Responsibilities of the NRA: N/A

**Medical device definition**
- Medical device defined: N/A
  - Text: N/A
- In vitro diagnostic medical device (IVD) defined: N/A
  - Text: N/A

**Medical device classification**
- Classification: N/A
- Categories: N/A
- Classification rules: N/A
- Classification rules details: N/A

**Essential principles**
- Essential principles: N/A
- Details: N/A

**Conformity assessment**
- Conformity assessment bodies: N/A
- Details: N/A
- Pre-marketing / procedure: N/A

**Reliance**
- Reliance: N/A
- Details: N/A
- Jurisdictions: N/A

**Clinical investigation**
- Clinical investigation controls: N/A
- Details: N/A

**Registration and listing**
- Registration of establishment: N/A
- Details: N/A
- Listing of medical devices: N/A
- Details: N/A

**Import controls**
- Import controls: N/A
- Details: N/A
Post market controls

Post Market Surveillance: N/A
Details: N/A

Inspection (QMS): N/A
Details: N/A

Enforcement: N/A
Details: N/A

Adverse event reporting: N/A
Details: N/A

Field safety corrective action monitoring: N/A
Details: N/A

Advertising: N/A
Details: N/A

Labelling: N/A
Details: N/A