Digitally signed by Ermicev Alexandr
Date: 2024.05.30 09:16:48 EEST
Reason: MoldSign Signature
Location: Moldova

Manufacturer's Declaration

in relation to Regulation (EU) 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices, in particular with respect to

- the validity of certificates issued under Council Directive 90/385/EEC on Active Implantable Medical Devices (AIMDD) or Council Directive 93/42/EEC on Medical Devices (MDD) (Directive Certificates) *and/or*[1]
- the compliance of the devices and us as their manufacturer with the conditions for the continued placing on the market and putting into service

Manufacturer name	CHIRANA T. Injecta, s.r.o.
Manufacturer address and contact details	Komoranska 2148, 143 00 Prague 4, Czech Republic
Single Registration Number (SRN) (if available)	CZ-MF-000034353

Authorised Representative name (if applicable)	N/A
Authorised Representative address and contact details	N/A
Single Registration Number (SRN) (if available)	N/A

Notified body name (if applicable)	☒See attached schedule
Notified body number (if applicable)	☒See attached schedule

[1] The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body.

Directive Certificate number(s) to which this confirmation is made (if applicable)	☒See attached schedule
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	☒See attached schedule
End date of extended validity/transition period	☒See attached schedule

We, as the manufacturer declare under our sole responsibility:

- for the above listed **Directive Certificate** (or see attached schedule, if multiple certificates) the conditions for the legal extension of validity as required in Article 120.2 of the MDR are met *and/or*[2]
- the listed **device(s)** in the attached schedule and we as their manufacturer are in compliance with the conditions listed in Article 120.3c of the MDR for continued placing on the market and putting into service,

namely by fulfilling the following conditions:

➢ **Directive Certificate(s)** as listed above or in the attached schedule

- Directive Certificate(s) covering the listed device(s) was/were issued after 25 May 2017, was/were valid on 26 May 2021 and have not been withdrawn afterwards.

  Choose applicable statements:

  ☐ Expired *before* 20 March 2023:

  ☐ Before the original date of expiry as indicated on the Directive Certificate(s), we and the notified body have signed written agreement(s) in accordance with Section 4.3, second subparagraph of Annex VII to this Regulation for the conformity assessment(s) in respect of the device(s) covered by the expired certificate(s) or in respect of a device(s) intended to substitute that/those device(s), or

  ☐ A Competent Authority has granted a derogation from the applicable conformity assessment procedure in accordance with Article 59(1) MDR (may be provided upon request), or

[2] The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body

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☐ A Competent Authority has required the manufacturer, in accordance with Article 97(1) MDR, to carry out the applicable conformity assessment procedure (may be provided upon request)

Choose one of the following statements only if a derogation per Article 59(1) or a requirement per Article 97(1) has been granted by a Competent Authority:

☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

☒Expired/expires *after* 20 March 2023:

Choose one applicable statement:

☒ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

➢ **Upclassified devices**

In case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body:

Choose one applicable statement:

☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitutes and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

➢ **Quality Management System (QMS)**

Choose one applicable statement:

☒ A QMS in accordance with Article 10(9) MDR will be put in place by no later than 26 May 2024.

☐ A QMS in accordance with Article 10(9) MDR is in place.

☐ A notified body has issued the attached certificate for the MDR-compliant QMS.

➢ **Device(s) as listed in the attached schedule**

- The device(s) continue to comply with the AIMDD or MDD.
- There are no significant changes in the design and intended purpose.
- The device(s) do not present an unacceptable risk to health or safety of patients, users or other persons, or to other aspects of the protection of public health.

Signed for and on behalf of the manufacturer:

CHIRANA T. Injecta, s.r.o.

Komoranska 2148, 143 00 Prague 4, Czech Republic

4.3. 2024

CHIRANA T. Injecta, s.r.o.
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RNDr. Anar Mamytbekova

Person responsible for regulatory compliance

E-mail: anar.mamytbekova@t-injecta.cz

Phone: + 420 241 097 902, + 420 731 540 982

Schedule of Devices

The above Manufacturer's Declaration is valid for the following devices:

Identification of the device(s)[3] (e.g., device name, family/group name device model or catalogue number)	**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number that issued the Directive Certificate** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device (s)** (if applicable)
Surgical suture Trade name: Chirasorb braided UDI-DI: 8596165ChirasorbV5	**MED 190018 MED 190021**	**26.05.2024**	**Electrotechnical Testing Institute, s.p., 1014**	**3EC International a. s., 2265**	**31.12.2027**	**N/A**
Surgical suture Trade name: Chirasorb rapid braided UDI-DI: 8596165ChirasorbrapidNS	**MED 190018 MED 190022**	**26.05.2024**	**Electrotechnical Testing Institute, s.p., 1014**	**3EC International a. s., 2265**	**31.12.2027**	**N/A**
Surgical suture Trade name: Chirasorb Plus braided UDI-DI: 8596165ChirasorbPlusBQ	**MED 190018 MED 190027**	**26.05.2024**	**Electrotechnical Testing Institute, s.p., 1014**	**3EC International a. s., 2265**	**31.12.2027**	**N/A**

[3] for devices with AIMDD/MDD certificate(s) the identification should be as in the certificate, and only if the certificate has a generic scope it should be as defined above)

Surgical suture Trade name: Chirlac braided Alternative trade name: C-TEC Alfatec braided UDI-DI: 8596165ChirlacA8	MED 190018 MED 190019	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A
Surgical suture Trade name: Chirlac rapid braided UDI-DI: 8596165Chirlacrapid5R	MED 190018 MED 190020	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A
Surgical suture Trade name: Monolac monofilament Alternative trade name: C-TEC Caprotec monofilament UDI-DI: 8596165MonolacKF	MED 190018 MED 190024	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A
Surgical suture Trade name: Polydox monofilament Alternative trade name: C-TEC Cynadox monofilament UDI-DI: 8596165PolydoxPW	MED 190018 MED 190023	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A
Surgical suture Trade name: Chiralen monofilament UDI-DI: 8596165ChiralenXB	MED 190018 MED 190025	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A

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Surgical mesh – partially absorbable Trade name: Capromesh UDI-DI: 8596165CapromeshX4	MED 190018 MED 190026	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A
Non-absorbable surgical mesh Trade name: Chiralen mesh UDI-DI: 8596165ChiralenMeshYL	MED 190017	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2027	N/A
Surgical suture Trade name: Silon braided UDI-DI: 8596165SilonbraidedT9	MED 190017	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2028	N/A
Surgical suture Trade name: Silon monofilament Alternative trade name: C-TEC Celon monofilament UDI-DI: 8596165Silonmonofil6X	MED 190017	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2028	N/A
Surgical suture Trade name: Tervalon braided UDI-DI: 8596165TervalonDV	MED 190017	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2028	N/A
Surgical suture Trade name: Silk braided UDI-DI: 8596165SilkN4	MED 190017	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2028	N/A
Surgical suture Trade name: Chiraflon monofilament UDI-DI: 8596165ChiraflonSA	MED 190017	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2028	N/A
Eyed needles – non-sterile Trade name: Eye-needles UDI-DI: 8596165EyeNeedlesSU	MED 200068	26.05.2024	Electrotechnical Testing Institute, s.p., 1014	3EC International a. s., 2265	31.12.2028	N/A

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