The Appendix is an integral part of
Certificate of Accreditation 269/2019 of 06/06/2019

Accredited entity according to ČSN EN ISO/IEC 17025:2018:

Univerzita Palackého v Olomouci
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Tests:

Ordinal number [1]	Test procedure/method name	Test procedure/method identification [2]	Tested object
1.	Test for *in vitro* cytotoxicity – **Test of direct contact**	SOP 1 (ČSN EN ISO 10993-5, cl. 1-3, 4.1, 4.3, 4.4, 5-7, 8, 8.3, 9,10; ČSN EN ISO 10993-12; ČSN EN ISO 7405, cl. 6.2.7, 6.2.8)	Insoluble solid samples with at least one flat surface[3]
2.	Test for *in vitro* cytotoxicity – **Test of extract**	SOP 2 (ČSN EN ISO 10993-5, cl. 1-3, 4.1, 4.2, 4.4, 5-7, 8, 8.2.1-8.2.8, 8.5, 9, 10; ČSN EN ISO 10993-12)	Insoluble solid samples[4]
3.	Test for *in vitro* cytotoxicity – **Test of soluble samples**	SOP 3 (ČSN EN ISO 10993-5, cl. 1-3, 4.3.3, 4.4, 5-7, 8, 8.2.1-8.2.8, 8.5, 9, 10; ČSN EN ISO 10993-12)	Soluble solid samples, liquid samples[5]

[1] Asterisk at the ordinal number identifies the tests, which the Laboratory is qualified to carry out outside the permanent laboratory premises.

[2] If the document identifying the test procedure is dated, only these specific procedures are used. If the document identifying the test procedure is not dated, the latest edition of the specified procedure is used (including any changes).

Explanation:

[3]Samples are medical devices defined in ČSN EN ISO 10993-1 cl. 3.1, insoluble in water.

[4]Samples are medical devices defined in ČSN EN ISO 10993-1 cl. 3.1, insoluble in water.

[5]Samples are medical devices defined in ČSN EN ISO 10993-1 cl. 3.1, soluble in water or liquid samples.