

DECLARATION OF CONFORMITY

released according to §13, item 2 of the Act No. 22/1997 Coll. about products technical requirements as amended and according to the Government Regulation No. 54/2015 Coll. that defines technical parameters for medical devices as amended, (Hereinafter referred to as Medical device) according to Law No. 268/2014 Coll. about medical devices and amending Law no. 634/2004 Coll., on administrative fees amendment of some related legislation, as amended.

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Declares that

Product **Bedside cabinet**

Type **NS-1-O, NS-2-O, NS-3-O, NS-20-O, NS-21-O, NS-40-O, NS-41-O, NS-50-O, NS-51-O, NS-60-L, NS-60-P, NS-61-L, NS-61-P, NS-99-P, HANDY**

is not classified as medical device defined by the Act. No. 268/2014 Coll. about medical devices and amending Law no. 634/2004 Coll., on administrative fees amendment of some related legislation, as amended. Therefore the above mentioned product is not subject to obligation to issue the CE declaration of conformity with the technical requirements determined in the Government Regulation No. 54/2015 Coll. that defines technical parameters for medical devices as amended (according to European council directive 93/42/EEC). Above mentioned product is not subject to obligatory registration as medical equipment at the Ministry of health of the Czech Republic.

Producers also confirms that

1. This product is safe when used for intended purpose and according to the User Manual and complies all hygienic requirements,
2. This products is intended for use in využívání zdravotně postiženými osobami ke zmírnění nebo kompenzaci zdravotního postižení nebo neschopnosti,
3. This product is suitable for use in interiéru,
4. That provisions were taken to provide the conformity of all products in market with the relevant technical documentation,
5. The technical documentation is stored in the company headquarters.

In Česká Skalice, 13.8.2019

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Company stamp

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Zdeněk Jakubský
CEO



We are entitled to use the above shown protected marks