RETROSUPERFUTURE

EU Declaration of conformity Prescription frames

Milano 01/09/2020

The company, Retrosuperfuture SRL, with operational headquarters at 7, Via Ferdinando Bocconi, Milan, 20136, with VAT No. IT06267680962, Milan Chamber of Commerce No. 1881995, as a manufacturer of class I medical devices pursuant to Article 13 of Legislative Decree No. 46 of 24 February 1997 and subsequent amendments, declares that the following medical devices:

Model: MONDO

Colors: DBI, V1Q, W3B, 6MJ, IBG, WJN, 61W, F9W, SN9

comply with the applicable provisions for Class I Medical Devices pursuant to Article 1, Part III of Annex IX of Directive 93/42/CEE and subsequent amendments, implemented in Italy by Legislative Decree No. 46 of 24 February 1997, Legislative Decree No. 95 of 25 February 1998 and Legislative Decree No. 37 of 25 January 2010 and to the national standard that transposes the harmonised standard ISO 12870:2012.

This declaration of conformity is issued under the sole responsibility of the manufacturer.

Milano 01 / 09 / 2020

Stefano Pomogranato (CEO)

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T +39 02 869 969 71 F +39 02 869 932 26 E rsf@retrosuperfuture.com

TAX ID IT06267680962

RETROSUPERFUTURE

EU Declaration of conformity Prescription frames Milano 01/09/2020

The company, Retrosuperfuture SRL, with operational headquarters at 7, Via Ferdinando Bocconi, Milan, 20136, with VAT No. IT06267680962, Milan Chamber of Commerce No. 1881995, as a manufacturer of class I medical devices pursuant to Article 13 of Legislative Decree No. 46 of 24 February 1997 and subsequent amendments, declares that the following medical devices:

Model: NUMERO 49

Colors: 7PY, 87E, OJU, 22L, RMR, YP2, 3NG, IF4, OCH

comply with the applicable provisions for Class I Medical Devices pursuant to Article 1, Part III of Annex IX of Directive 93/42/CEE and subsequent amendments, implemented in Italy by Legislative Decree No. 46 of 24 February 1997, Legislative Decree No. 95 of 25 February 1998 and Legislative Decree No. 37 of 25 January 2010 and to the national standard that transposes the harmonised standard ISO 12870:2012.

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Model: NUMERO 53

Colors: 6R4, PQ1, WPD, 8P3, D3F, ZZW, 6O2, MEB, OOP

comply with the applicable provisions for Class I Medical Devices pursuant to Article 1, Part III of Annex IX of Directive 93/42/CEE and subsequent amendments, implemented in Italy by Legislative Decree No. 46 of 24 February 1997, Legislative Decree No. 95 of 25 February 1998 and Legislative Decree No. 37 of 25 January 2010 and to the national standard that transposes the harmonised standard ISO 12870:2012.

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