eyelevel

CE EC Declaration of Conformity MDD eyelevel® diagnostik ADHS/OPP

Manufacturer	eyelevel GmbH
Address of Manufacturer	Oberanger 44 D-80331 München
Product and version	eyelevel diagnostik ADHS/OPP 1.x
Council (EU) Directive	Medical Device Directive MDD 93/42/EWG
Classification	Class I
Rules applied	Rule 12: All other active devices are in class I
Conformity Assessment Procedure according to Article 11 of the MDD	Declaration of conformity issued under the sole responsibility of the manufacturer according to Annex VII of the directive

We, eyelevel GmbH, declare under our sole responsibility conformity to the Medical Device Directive MDD 93/42/EWG specified above.

The product specified above is a medical device according to Article 1 of the Medical Device Directives MDD 93/42/EWG.

The product complies with the essential requirements as of Annex I of the Medical Device Directives MDD 93/42/EWG.

The procedure to assess conformity specified above has been followed, all provisions of the Medical Device Directives MDD 93/42/EWG and the harmonized standards (DIN EN ISO 14971, IEC 62366, IEC 62304) are met.

The declaration is valid through the date of signature below.

Munich, 21.05.2021

Benjamin Luther