

Declaration of Conformity

Tobii Dynavox, a business unit and subsidiary of Tobii AB, herby declares that the product identified herein complies with the following directives:

FU Directives

- 93/42/EED Medical Device Directive (MDD), as amended 2007/47/EC, Annex IX, Rule 12
- 2011/65/EU Restrictions of Hazardous Substances Directive (RoHS 2)
- 2004/108/EC Electromagnetic Compatibility Directive (EMC)
- **REACH and WEEE Directives**

Product Information & Use

Type of product: Eye Tracker

Product name/Model: Tobii Dynavox EyeMobile Plus Product description: Remote eye tracking device

Intended use: The Tobii Dynavox EyeMobile Plus is a bracket with built-in eye tracker that allows the user to control a Windows based, tablet PC (e.g. Surface Pro by Microsoft) using eyegaze. The EyeMobile Plus is intended to be used in an indoor environment.

Other Directives & Standards

- Federal Communications Commission (FCC), CFR Title 47, Part15 Subpart B: 2015, CLASS B
- Industry Canada Equipment Standard (ICES-003), Issue 6: 2016 Class B
- Radiocommunication Standard 2008 (ACMA), AS/NZS CISPR 32:2013
- Korean Radio Waves Act, Clause 3, Article 58-2

The following standards have been used:

- ISO 14971:2007+2012 (Medical Device Risk Mgmt.)
- ANSI C63.4: 2104
- EN 55032: 2012+AC:2013 Class B
- EN 55024: 2010+A1:2015
- EN 61000-3-2:2014
- EN 61000-3-3:2013

- IEC 60950-1:2005
- IEC 60950-1:2005/AMD1:2009
- IEC 60950-1:2005/AMD2:2013

I hereby declare that the equipment named above has been designed to comply with the relevant sections of the above referenced specifications. The unit complies with all essential requirements.

Robert Cunningham, VP of Product Development

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Date

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