CE Declaration of Conformity (DoC)

Sync-Think Inc.
Address - 2172 Staunton Court, Palo Alto, CA 94306, United States
Phone - +1 (650) 727-1819
Website - www.syncthink.com

Sync-Think Inc. hereby declares under the sole authority of the manufacturer, that the product
Type of Product - Eye Tracking Equipment
Product Name - EYE-SYNC
Designation - EYE-SYNC-X03
Description - Eye tracking system

is in accordance with the following European Directives

2017/745/EU	European Union Medical Devices and Clinical Investigations
2014/30/EU	European Union EMC Directive

and complies with the following European standards and requirements

ISO 13485:2016	Medical Devices
ISO 14791:2019	Risk Management
ISO 10993:2009	Biological Evaluation of Medical Devices
ISO 15223:2016	Medical Device Labeling
IEC 60601-1:2005 (Third Edition) + CORR. 1:2006 + CORR. 2:2007 + A1:2012 (or IEC 60601-1: 2012 reprint)	Medical Electrical Equipment
IEC 60601- 2-57 (First Edition): 2011, EN 60601-2-57: 2011, CAN/CSA 22.2 No. 60601-2-57:11	Medical Electrical Equipment

as a Class 1m Medical Device.

Signed for and on behalf of Sync-Think Inc. by:

Place of Issue	Palo Alto, CA 94306, United States
Date of Issue	June 24, 2022
Authorized Signature	
Name	Jan Henri van Merkensteijn IV
Position	Quality Manager

CE Declaration of Conformity (DoC)

Sync-Think Inc.
Address: 2172 Staunton Court, Palo Alto, CA 94306, United States
Phone: +1 (650) 727-1819 **Website:** www.syncthink.com

Sync-Think Inc., utilizing testing data and guidance obtained in conjunction with the accredited third party of TUV Rheinland Inc., hereby declares under the rightful and sole authority of the manufacturer, that the product

Type of Product - Eye Tracking Equipment
Product Name - EYE-SYNC
Designation - EYE-SYNC-X02
Description - Eye tracking system

is in accordance with the following European Directives

2017/745/EU	European Union Medical Devices and Clinical Investigations
2014/30/EU	European Union EMC Directive

and complies with the following European standards and requirements

ISO 13485:2016	Medical Devices
ISO 14791:2019	Risk Management
ISO 10993:2009	Biological Evaluation of Medical Devices
ISO 15223:2016	Medical Device Labeling
IEC 60601-1:2005 (Third Edition) + CORR. 1:2006 + CORR. 2:2007 + A1:2012 (or IEC 60601-1: 2012 reprint)	Medical Electrical Equipment
IEC 60601- 2-57 (First Edition): 2011, EN 60601-2-57: 2011, CAN/CSA 22.2 No. 60601-2-57:11	Medical Electrical Equipment

as a Class 1 Medical Device.

Signed for and on behalf of Sync-Think Inc. by:

Place of Issue	Palo Alto, CA 94306, United States
Date of Issue	July 8, 2022
Authorized Signature	
Name	Jan Henri van Merkensteijn IV
Position	Quality Manager