

## **CERTIFICATE OF CONFORMANCE**

Acting as Regulatory Authorized Representative in Europe for **OrCam Ltd.** from **Israel**, we hereby declare that:

The following products listed hereunder or in the "Products Schedule" has been registered with the European Competent authority in the UK (MHRA) on **12 October 2015** Ref. no. **CA015099** as Class I – **Low vision Aids.** 

By submitting the information to the MHRA, the legal requirements for registration have been met. **OrCam Ltd**. complies with the requirements of the Medical Device Directive 93/42 EEC amended by 2007/47 and the above products can be CE marked according to the above directive.

## **Product Schedule**

Sr. No.	Product Name / Family	
1	OrCam MyEye 1	
2	OrCam MyEye 2.0	

For Medes Ltd.

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Benny Arazy General Manager Medes Ltd.

Date: 26.03.18

