



EU DECLARATION OF CONFORMITY

We hereby declare that the magnifying spectacles produced on our behalf as per below information is in conformity with the relevant requirements: **(EU) 2017/745 regulation**

Class 1 low risk medical device

Description of the products	GMDN Code	Basic UDI-DI
Magnifying Spectacles	42832	5036079ReadyReaders02FJ

Product Names:

Readezee 125 Readezee 150 Readezee 200 Readezee 250 Readezee 300 Readezee 350
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The products meet the requirements of the relevant Union harmonisation legislation:

(EU) 2017/745 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No. 178/2002 and Regulation (EC) No. 1223/2009 and repealing the directives 90/385/EEC and 93/42/EEC of the Council.

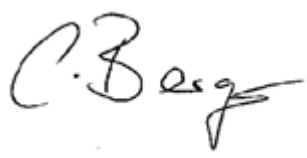
The harmonised standards used are the following:

EN ISO12870:2024, EN14971:2020-07, EN16128:2016-02

Name and Address of the Manufacturer:

The Optoplast Actman Eyewear Co Limited, 83 Sefton Lane, Maghull, Liverpool, L31 8BU, UK

Authorised by Clare Berger for and on behalf of Victoria Collection (EU authorized Representative)



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