

# EU Declaration of Conformity

## 1 Device identity

<b>Device name:</b>	Evira
<b>Device version:</b>	1.0

## 2 EU Declaration of Conformity

<b>Name of the manufacturer:</b>	Evira AB
<b>SRN:</b>	SE-MF-000024616
<b>Address of registered place of business:</b>	Triewaldsgränd 2, 111 29 Stockholm
<b>Basic UDI-DI:</b>	7300009080851QR
<b>Intended purpose:</b>	Evira is to be used as aid in clinical behavioral obesity treatment. An app is used regularly at home by end-users. A web app is used by supporting health care professionals.
<b>Risk class:</b>	MDR Class I
<b>Reference to any CS used and in relation to which conformity is declared:</b>	–
<b>Name and identification number of the notified body:</b>	N/A
<b>Conformity assessment procedure performed:</b>	2017/745, Annex I, II
<b>Identification of the certificate or certificates issued:</b>	N/A
<b>Additional information:</b>	–

Stockholm, 2024-03-05



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Andreas Drangel  
CEO  
Evira AB

This declaration of conformity is issued under the sole responsibility of the manufacturer.

The device that is covered by the present declaration is in conformity with Regulation (EU) 2017/745.