

# Declaration of Conformity

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.

<b>Manufacturer</b>	Vivo Red Light Therapy B.V.
<b>Address</b>	Constructieweg 90-K, 3641 SP, Mijdrecht, The Netherlands
<b>Product</b>	Red Light Therapy Panel
<b>Trade Mark</b>	Vivo Red Light Therapy
<b>Model(s)</b>	BioPhoton 3.0, BioPhoton+, BioPhoton Go, BioPhoton Max Bi-Tone

The object of the declaration described above is in conformity with the relevant Union harmonisation legislation:

- EMC Directive 2014/30/EU
- Low Voltage Directive 2014/35/EU
- RoHS Directive 2011/65/EU and amendment Directive (EU) 2015/863

Conformity is demonstrated by compliance with the following standards and test reports:

Category	Test Standards	Test Report
<b>EMC standards</b>	EN IEC 55014-1:2021 EN IEC 55014-2:2021 EN IEC 61000-3-2:2019+A1:2021+A2:2024 EN 61000-3-3:2013+A1:2019+A2:2021+AC:2022	DACE260617003EC001
<b>LVD standards</b>	EN 60335-1:2012 + AC:2014 + A11:2014 + A13:2017 + A1:2019 + A14:2019 + A2:2019 + A15:2021 EN 60335-2-27:2013 + A1:2020	DACE260617005SA001
<b>RoHS standards</b>	IEC 62321-3-1:2013 IEC 62321-4:2013+AMD1:2017 IEC 62321-5:2013 IEC 62321-7-1:2015 IEC 62321-7-2:2017 IEC 62321-6:2015 IEC 62321-8:2017	DTI2026EE200625

**Signed on behalf of:** Vivo Red Light Therapy B.V.

<b>Name</b>	Michel Peters
<b>Position</b>	Director
<b>Place</b>	Mijdrecht, The Netherlands
<b>Date</b>	9-7-2026
<b>Signature</b>	