

# Declaration of Conformity

**We, the manufacturer, under compliance to Article 19 of EU MDR 2017/745,  
declare under our sole responsibility that the medical device:**

Mission® Lancing Device (C121-3051)  
Insight® Lancing Device (C121-3055)  
On Call® Lancing Device (G124-11A)  
On Call® GenTouch Lancing Device (G124-17A)

**of class I according to Rule 13 of Annex VIII of regulation (EU) 2017/745,  
is in conformity with EU MDR 2017/745.**

**This declaration is based on:**

**Manufacturer's Name:** ACON Laboratories, Inc.

**Manufacturer's Address:** 5850 Oberlin Drive, #340 San Diego, CA 92121

**Manufacturer's SRN:** US-MF-000023913

**Authorized Representative Name:** MDSS GmbH

**Authorized Representative Address:** Schiffgraben 41, 30175 Hannover, Germany

**Authorized Representative SRN:** DE-AR-000005430

**Basic UDI-DI:** 8260799999900013V

**Intended Purpose of device:** The device is intended for injuring the fingertip in combination with a disposable lancet for obtaining a small amount of blood sample.

Signed this 15 day of June 2023  
in San Diego, CA USA



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Senior Staff, Regulatory Affairs & Clinical Affairs  
ACON Laboratories, Inc.

