



## Declaration of Conformity

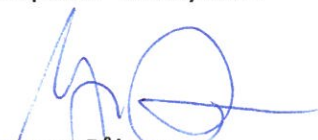
Manufacturer **Actim Oy**  
Klovinpellontie 3, FI-02180, Espoo  
Finland  
IVD product **Actim® CRP** (Product code 31031ETAC)

**We hereby declare that the above mentioned device complies with the requirements of Directive 98/79/EC on *in vitro* diagnostic medical devices (IVD Directive) and the corresponding Finnish National Act 629/2010 and to the following standards**

Standards **SFS-EN ISO 13485:2016, AC 2016, AC 2018**  
Medical devices – Quality Management Systems – Requirement for the regulatory purposes.  
**SFS-EN ISO 14971: 2019**  
Medical Devices. Application of risk management to medical devices.  
**SFS-EN 13612:2002 + AC**  
Performance evaluation of *in vitro* diagnostic medical devices.  
**SFS-EN ISO 23640:2015 (ISO 23640:2011)**  
In vitro diagnostic medical devices. Evaluation of stability of *in vitro* diagnostic reagents.  
**ISO 15223-1:2016**  
Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied Part 1: general requirements.  
**ISO 15223-2:2010**  
Medical devices -- Symbols to be used with medical device labels, labelling, and information to be supplied -- Part 2: Symbol development, selection and validation.  
**SFS-EN ISO 18113-1:2012**  
*In vitro* diagnostic medical devices – Information supplied by the manufacturer (labelling) – Part 1: Terms, definitions and general requirements.  
**SFS-EN ISO 18113-2:2012**  
*In vitro* diagnostic medical devices – Information supplied by the manufacturer (labelling) – Part 2: *In vitro* diagnostic reagents for professional use.  
**SFS-EN 13975:2003**  
Sampling procedures used for acceptance testing of in vitro diagnostic medical devices – Statistical aspects

IVDD classification **General class**

In Espoo 6<sup>th</sup> of May 2021

  
**Magnus Pålsson**  
Managing Director