# Declaration of Conformity

We, ImmoGnost GmbH, herewith declare on our own responsibility, that the in vitro diagnostic product

**RACORDIAX COVID-19 Antibody Rapid Test**

is in conformity with the requirements of
**Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices.**

The conformity was assessed according to **Directive 98/79/EC Annex III.**
The RACORDIAX COVID-19 Antibody Rapid Test is CE marked.

CE

Göttingen, 2020-12-30

PD Dr. Frank Gessler
Quality Management Representative

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