

# EC DECLARATION OF CONFORMITY

General In vitro Diagnostic Medical Devices in  
accordance with Directive 98/79/EC

**Manufacturer** : Chil Tibbi Mal. San. Tic. Ltd. Şti.  
**Head Office** : 10028 Sok. No:11 AOSB 35620 Çiğli/İzmir-Turkey  
**Manufacturing Plant** : 10028 Sok. No:11 AOSB 35620 Çiğli/İzmir-Turkey  
**Product name** : COVID-19 Antigen Rapid Test  
**Brand** : CHIL®  
**Classification** : Self-test IVD  
**Route** : Annex III.6

We, here with declare that the above mentioned products meet the provisions of the council directive 98/79/EC for In-Vitro Diagnostic Medical Devices. All supporting documentation is retained under the premises of the manufacturer.

**Applied Standards & regulations** : Applied Standards/Regulations List is attached.

**Notified Body** : Institut Pro Testování A Certifikaci, a. s.  
T. Bati 299 Louky, 76302 ZLIN, Czech Republic  
Notified Body number : 1023

**Start of CE Marking**

**Issued Date/Place** : 14.03.2022 / Izmir-Turkey

**Revision Date-No** : 14.03.2022-0

**Signature: Name** : ŞENAY ÇİL

**Position** : MANAGEMENT RESPONSIBLE

CE 1023

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