

## EU DECLARATION OF CONFORMITY

(no. EC 27-2024)

We, **CHIRANA T. Injecta a.s.**,  
Nám. Dr. A. Schweitzera 194, 916 01 Stará Turá, Slovakia

Single registration number (SRN): SK-MF-000003702

manufacturer of: SINGLE USE MEDICAL DEVICES

declare in our own responsibility, that the product:

### **BLOOD COLLECTION NEEDLE HOLDER FOR SINGLE USE (NON-STERILE)**

brand name: **CHIRAVAC™**

versions: various types

intended purpose: blood collection in connection with blood collection needle  
and vacuum tube

classified as Class: I according to classification set out in Annex VIII, Rule 1  
of the EU Medical Device Regulation (MDR) 2017/745

is in compliance:

- with provisions of Annex I of the Regulation (EU) 2017/745 of the European parliament and of the council of 5<sup>th</sup> April 2017 on medical devices which apply to them
- with provisions of Regulation (EU) 2017/745, Article 52 (7) relating to the Technical Documentation set out in Annex II and Annex III
- with provisions of specification PS 310-0718-02, EN ISO 14971:2019, EN ISO 13485:2016/AC:2018 standards selected from the group EN ISO 14644 and EN ISO 10993 and other relevant ISO/EN standards mentioned in specification and list of standards for product

Product models are marked with CE.

EU declaration of conformity is issued under the sole responsibility of the manufacturer.  
Signed for and on behalf of manufacturer.

in Stará Turá, date 02.02.2024



  
Rastislav Broska, PhD.  
Quality Manager

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*Product:*

**BLOOD COLLECTION NEEDLE HOLDER – CHIRAVAC™  
(NON-STERILE)**

### **Product variants - CHIRAVAC™:**

<b>Product REF code</b>	<b>Product model</b>	<b>Basic UDI-DI (GMN)</b>
<i>CHBCNH01</i>	<i>Standard</i>	<i>858601504HOLDER272L</i>
<i>CHBCNHS01</i>	<i>Safety</i>	
<i>CHBCNHQ01</i>	<i>Quick release</i>	