## EU DECLARATION OF CONFORMITY



manufacturer: ZARYS International Group sp. z o.o. sp.k. address: ul. Pod Borem 18, 41-808 Zabrze, Poland tel. +48 32 271 69 91, fax +48 32 274 72 84, contact:

e-mail: zarys@zarys.pl, website: www.zarys.com

SRN: PL-MF-000000410

We declare under our sole responsibility that a medical device/personal protective equipment:

## easyCARE latex PF - latex examination gloves, powder-free, non-sterile

sizes\*: from XS to XL

(\*detailed list of products covered by this declaration is available in document TD-49-I.1.1.b-1.2 – Identification – Annex 1, batch code - certificate of analysis of production batch DZDO-01- Annex 2)

## classification:

medical device: class I, rule 5 (in accordance with Annex VIII of Regulation (EU) 2017/745)

personal protective equipment: category III

Basic UDI-DI: 59079968T010201GM

intended purpose: The device is intended for patient diagnostics, providing a barrier to the transmission of microorganisms between the operator and the patient, minimizing the risk of cross-contamination.

## is in conformity with:

- Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices;
- Regulation (EU) 2016/425 of the European Parliament and of the Council of 9 March 2016 on personal protective equipment;
- Commission Regulation (EU) No. 10/2011 of 14 January 2011 on plastic materials and articles intended to come into contact with
- Regulation (EC) No. 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food.

The medical device described above meets all applicable provisions of the Annex I of Regulation (EU) 2017/745. Conformity assessment procedure has been performed in accordance with Article 52 (7).

For the personal protective equipment the notified body SATRA Technology Europe Limited (2777) performed the EU typeexamination (Module B) and issued the EU type-examination certificate:

number: 2777

notified body: SATRA Technology Europe Limited

Bracetown Business Park, Clonee,

D15 YN2P, Ireland

certificate no.: 2777/10467-05/E21-01 expiry date: 2027-01-10

The PPE is subject to the conformity assessment procedure conformity to type based on quality assurance of the production process (Module D) under surveillance of the notified body SATRA Technology Europe Limited (2777).

The medical device covered by this declaration complies with standards EN 455-1:2020+A1:2022, EN 455-2:2024, EN 455-3:2015, EN 455-4:2009, EN ISO 13485:2016+EN ISO 13485:2016/A11:2021, EN ISO 14971:2019+EN ISO 14971:2019/A11:2021, EN 62366-1:2015+EN 6256-1-EN 6256-1-EN 625-1-EN 1:2015/A1:2020, EN ISO 15223-1:2021, EN ISO 20417:2021, EN ISO 10993-1:2020, EN ISO 10993-5:2009, EN ISO 10993-10:2023, EN ISO 10993-23:2021, EN ISO 11737-1:2018+EN ISO 11737-1:2018/A1:2021, the personal protective equipment complies with standards EN ISO 21420:2020+A1:2024, EN ISO 374-1:2016+A1:2018, EN ISO 374-2:2019, EN 16523-1:2015+A1:2018, EN ISO 374-4:2019, EN ISO 374-5:2016, ASTM F1671, ASTM D3578.

The device intended for food contact meets the requirements specified in Annex 3.

place and date of issue: Zabrze, 20.01.2025 Aleksandra Stec name:

Quality and Regulatory Compliance position:

Specialist

SPECJALISTA DS. JAKOŚCI I ZGODNOŚCI REGULACYJNEJ ZARYS International Group sp. z o.o. sp.k.

Aleksandra Stec .....

signature

(on behalf of the President of the General Partner's

Management Board)