

## UKCA DECLARATION OF CONFORMITY

We,

MEDICOM SAS  
Boulevard de la Chanterie – 49 124 Saint Barthélemy d’Anjou – France

**Declare that the declaration of conformity is issued under our sole responsibility and relates to the following product:**

**SafeTouch<sup>®</sup> Advanced<sup>™</sup> Black Lite**  
**Category III Personal Protective Equipment – Protective gloves**  
**Class I Medical Devices – Medical gloves**

Reference	Brand	Powder	Color	Packaging
MBKN3	MEDICOM	Powder-free	Black	10 boxes of 100 units

**MD Intended purpose:** Single-use, non-sterile, powder-free, nitrile medical glove, intended to cover the hands and the wrists of the healthcare professional during medical cares or examinations, in order to prevent the risk of cross-contamination.

**PPE intended purpose:** Single-use, non-sterile, powder-free, nitrile protective glove, intended to protect the bearer against tested chemicals (type C) and microorganisms (bacteria, moulds and viruses).

**The objects of the declaration described above comply with the following Union harmonisation legislations:**

- Regulation 2016/425 on personal protective equipment, as amended to apply in GB
- UK Medical Device Regulation 2002

**The following harmonised standards and technical specifications have been applied:**

Medical Device	Personal Protective Equipment
EN 455-1	EN ISO 21420
EN 455-2	EN ISO 374-1
EN 455-3	EN ISO 374-2
EN 455-4	EN ISO 374-4
	EN 16523-1
	EN ISO 374-5
	ISO 16604

MEDICOM SAS

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SAS au capital de 1 214 000 € - RCS ANGERS 523 019 354 – Siret : 52301935400035 – Code APE 1395Z - TVA intra FR42523019354

**Conformity assessment procedure:**

- **Personal protective equipment:**

The notified body CTC (0075) performed the EU type-examination (Module B) and issued the EU-type examination certificate n°0075/1467/162/11/22/2311.

The product is subject to the conformity to type assessment procedure based on internal production control plus supervised product checks at random intervals (Module C2) under surveillance of the notified body CTC (0075).

As stated by the UK government on November 14th 2022, products being CE marked by a notified body before December 31st 2024, do not require UKCA type examination certification by an Approved Body (UK). CE conformity assessment can be used as a basis for UKCA marking and the product can be placed on the UK market until the validity of the CE certificate expires or for three years (up to December 31st 2027).

- **Medical device:**

The product is subject to the procedure set out in UK Medical Device Regulation 2002 and does not require a UKCA type examination certificate by an approved body.

**Signed for and on behalf of: Sandrine Engels, President of Medicom Europe**

**Name:** Yannick CHEVALIER

**Place of issue:** Saint Barthélemy d'Anjou

**Function:** European Quality, Regulatory and R&D Director

**Date of issue:** 06/02/2024

**Signature:**



**Expiry date:** 30/12/2027

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