

EU Declaration of Conformity according to the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL <i>Class I Medical Device(non-sterile)</i> And the REGULATION (EU) 2016/425 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL <i>Category III Personal Protective Equipment</i>		
Manufacturer:	GUANGDONG KINGFA MEDICAL SCI.&TECH. CO., LTD.	
Address:	No.36 Delong Avenue, Shijiao Town, Qingcheng District, Qingyuan City, Guangdong Province, China	
Single Registration Number (SRN) of the Manufacturer:	CN-MF-000055146	
European Representative (ER):	Share Info GmbH	
Address:	Am Schulzentrum 12, 41564 Kaarst, Germany	
Single Registration Number (SRN) of ER:	DE-AR-000005132	
We, the manufacturer, declare under our sole responsibility that		
Product Name:	Nitrile examination gloves	
Type/model , identification of product allowing traceability (Where applicable):	KG-1101, KG-1107	
Intended Purpose:	The nitrile examination gloves are intended used for the health care personnel to prevent contamination during close contact with the patient. The products are single- use, powder-free and non-sterile.	
the medical device(s)	Classification: (Annex VIII of the MDR)	Class I Medical Device
	Basic UDI-DI:	KG-1101: 697988801KG-1101QK KG-1107: 697988801KG-1107QX
	Conformity assessment route:	EU Declaration of Conformity + Technical Documentation (Annex II) + Technical Documentation on Post-Market Surveillance (Annex III)

is/are in conformity with the relevant provisions and requirements of the Council and the parliament regulation (EU) 2017/745 for medical device and all applicable harmonized standards and Common Specification. All supporting documents are retained under the premises of the manufacturer.		
Applied harmonized standards and Common Specification	Regulation (EU) 2017/745	EN 455-1:2020+A2:2024
	EN ISO 14971:2019	EN 455-2:2024
	EN ISO 13485:2016	EN 455-3:2023
	EN ISO 20417:2021	EN 455-4:2009
	EN ISO 10993-1:2025	EN 455-5:2025
	MEDDEV 2.7.1: 2016	EN 62366-1:2015+A1:2020
	EN ISO 15223-1:2021+A1:2025	
the personal protective equipment(s)	Classification: (ANNEX I of the REGULATION (EU) 2016/425)	Category III Personal Protective Equipment
	Conformity assessment procedures:	EU type-examination (module B) set out in Annex V, and conformity to type based on quality assurance of the production process (module D) set out in Annex VIII.
is/are in conformity with the relevant provisions and requirements of the Council and the parliament regulation (EU) 2016/425 on personal protective equipment and all applicable harmonized standards and Common Specification. All supporting documents are retained under the premises of the manufacturer.		
Applied harmonized standards and Common Specification	Regulation (EU) 2016/425	EN ISO 21420: 2020 + A1:2024
	EN ISO 374-1:2016+A7: 2018	EN ISO 374-2: 2019
	EN16523-1: 2015+A1: 2018	EN ISO 374-4: 2019
	EN ISO 374-5: 2016	
Notified Body:	SATRA Technology Europe Limited	
Address:	Bracetown Business Park, Clonee, D15 YN2P Ireland	
Identification Number:	Notified body:2777	

EC Certificate(s):	EU type-examination (Module B) Certificate Number: 2777/15940-06/E00-00 (Expiry date: 09/03/2031) and is subject to the type based on quality assurance of the production process (module D) under surveillance of the notified body: SATRA Technology Europe Ltd, Bracetown Business Park, Clonee, D15 YN2PIreland; Notified Body Number: 2777
Signed on:  Place: Qingyuan, China 2026-05-25 Signature (on behalf of the manufacturer) : GUANGDONG KINGFA MEDICAL SCI.&TEC CO., LTD. Name of authorized signatory: Wang Zhonglin Position held in the company: General Manager 	