

**MEXPO INTERNATIONAL, INC.**

2828 Faber Street  
Union City, CA 94587-1204, USA  
www.blossom-disposables.com

<b>Declaration of Conformity Latex Examination Gloves</b>		
<b>PRODUCT DESCRIPTION</b>		
1. Product Name: Latex Examination Gloves		
2. Product Classification: Class I under Medical Device Regulation (EU) 2017/745 Annex VIII Rule 1 & 5		
<b>ADDRESS:</b>		
<b>Mexpo International Inc.</b> 2828 Faber Street Union City, CA 94587-1204, USA Tel : +1 (510) 489-6800 Fax: +1 (510) 489-3111 E-mail : <a href="mailto:blossomglo@aol.com">blossomglo@aol.com</a> E-mail: <a href="mailto:tim@mexpo-glove.com">tim@mexpo-glove.com</a>	<b>EC Representative:</b> <b>Blossom Europe, S. L</b> Paseo de Recoletos, 37-41 28004 Madrid, Spain	<b>Brand Owner</b> white-med GmbH Marburger Straße 251 35396 Gießen, Germany
Single Registration Number (SRN) : US-MF-000032548		Single Registration Number (SRN) : ES-AR-000019689

We, **Mexpo International, Inc.** as the legal manufacturer declare under our sole responsibility that the medical devices listed below conform to the requirement of the Medical Device Regulation (EU) 2017/745.

**Latex Exames Gloves:**  
**Peppler Sensitive Aloe Vera (Examination Gloves, Powder Free) – Art.Nr. 412**

Size	UDI #
XS	4046144020814
S	4046144020821
M	4046144020838
L	4046144020845
XL	4046144020852

**Basic UDI-DI: 40461444717274**

It is declared that above devices meet the requirement of the Medical Device Regulation (EU) 2017/745.

The undersigned hereby declare that the disposable device(s) specified above are following the EU Type Examination and conformity with the provisions of the new PPE Regulation (EU) 2016/425- Cat III and, where such is the case, with the national standard transposing harmonized standard no. EN ISO 374-1:2016, EN 420:2003+A1:2009, EN 374-2:2014, EN 374-4:2013, EN ISO 374-5:2016.

The fulfilment of the applicable essential health and safety requirements set out in Annex II has been demonstrated under the supervision of the following notified body:

1. CENTEXBEL Textile Competence Centre, Technologiepark 70, BE 9052 Gent, Belgium is identical to the PPE EU Type Examination Certificate Nr. 049/2019/1262. Notified body No.: 0493,

In accordance with Annex VIII, Medical Device Regulation (EU) 2017/745, the device listed above are non-invasive transient devices and are Class I devices under Rule 1 & 5 as Rules 2, 3, and 4 do not apply.

The following person is responsible for signing this document:  
**Name and Address:** Mexpo International Inc., 2828 Faber Street, Union City, California 94587-1204, USA

**Authorized Signature:** 

**Date:** January 10, 2023

**Name of responsible Person:** Tim Thai

**Position:** President