Date: 05/04/2023

To: The Port and Airport Health Office (USMAF)
30123 Venezia, VE

Subject: Letter to acknowledge undertaking the procedure to update Maxtec's correct EU REP in the Italian medical devices Database for the product EyeMax2 - part numbers R300P01, R300P02 and R300P03.

Scope: This declaration is to acknowledge that the correct EU REP of Maxtec, LLC is-

EMERGO EUROPE
Westervoortsedijk 60,
6827 AT Arnhem
The Netherlands

We have reached out to Emergo and are in the process of getting this update completed in the Italian Ministry of Health Database and Repertorio before the goods are released for consumption.

We request you to kindly release the shipment as the update is in process.

Anvitha AnandRao

Digitally signed by Anvitha AnandRao
Date: 2023.05.04 12:32:36 -04'00'

Issued by: Anvitha Anand Rao
RA/Q Product Engineering Specialist