EC-Declaration of Conformity

We, the company

retec® Kunststofftechnik GmbH
Industriestraße 2
D-61191 Rosbach v.d.Höhe (Rodheim)

hereby declare in full responsibility as designer, manufacturer and first supplier, that the quality and safety relevant properties of our product variant from the medical device

"Acrylics for the manufacturing of dentures"

SHERAPRESS
SHERAPRESS FLEX
SHERADON
(including all color variations)

classification according to 93/42/EEC annex IX, section III, rule 5 (duration of use > 30 days):

Class IIa

fulfill the following fundamental requirements:

Directive 93/42/CEE concerning medical devices - Annex II

Notified Body: **DQS Medizinprodukte GmbH**, August-Schanz-Straße 21, 60433 Frankfurt am Main, CE 0297

Certificate Registration No.: 059034 MR2

This EC-Declaration of Conformity is valid until: **December 31th 2028**

Rosbach, 16.05.2024

Dr. Manfred Steinbach
PRRC (Person responsible for regulatory compliance)
retec® Kunststofftechnik GmbH

Productlist „Acrylics for the manufacturing of dentures“

Artikel-No.	Product name
SHE-502030	SHERAPRESS, klar, 1kg
SHE-502050	SHERAPRESS, rosa transparent veined, 1 kg
SHE-502010	SHERAPRESS, rosa, 1 kg
SHE-502040	SHERAPRESS, rosa veined, 1 kg
SHE-502020	SHERAPRESS, rosa opak, 1kg
SHE-508020	SHERAPRESS, liquid, 500 ml, klar
SHE-508021	SHERAPRESS, liquid, 1000 ml, klar
SHE-502060	SHERAPRESS, FLEX rosa, 1 kg
SHE-508030	SHERAPRESS, Flex liquid, 500 ml, klar
SHE-501030	SHERADON, rosa transparent veined, 1 kg
SHE-501010	SHERADON, rosa, 1 kg
SHE-501020	SHERADON, rosa veined, 1 kg
SHE-508010	SHERADON liquid, 500 ml