

# EU Representative Agreement

Document Number: JH-ERA-20682V00

This agreement will be valid for 1 years from 2020.04.21 to 2021.04.1. Part A could choose to renew the agreement by then, otherwise this agreement will be terminated automatically. 此合同有效期为1年，自2020年04月21日至2021年04月1日，到期前由甲方选择续约或本合同自动失效。

<b>Part A (甲方)</b>	
Name(名称):	Jinan Shangran Tongda Composite Material Co., Ltd
Add(地址):	No.28, Kailuan Road, Shanghe Economic Development Zone, Jinan City, Shandong Province, China.
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<b>Party B (乙方)</b>	
Name(名称):	Lutus Lebenswelt GmbH
Add(地址):	Kochstr. 1, 47877, Willich, Germany
DIN/DI Code	DE.0000047791
Tax Number	DE306829099
Contact Person:	Lin Sun
Tel/Fax:	0049-1715605732
E-mail:	info.mg@lulusw.de
<b>Competent authority (主审机构信息)</b>	
Name	Bezirksregierung Düsseldorf, Dorothea 24
Federal state	Nordrhein-Westfalen
City	Düsseldorf
Postal code	40474
Street, house no.	Coelienallee 2
Phone/Fax	+49-211-4750 / +49-211-4750971
E-mail	dez24.mpg@brd.nrw.de

Party A hereby appoints Party B as the authorized European Representative for their Medical Device with CE mark, Party B accepts the appointment to be the authorized European Representative for Party A in the market of European Union (E.U), EEA and Switzerland, Turkey. Both parties enter this agreement as follows, the appointed product categories set out in below form.

甲方任命乙方为CE认证产品欧盟授权代表，乙方接受甲方任命，为甲方在欧洲、EEA、瑞士、土耳其市场的CE认证产品授权代表。双方签署下列协议，委托的产品类别如下表。

No.	Product Name(产品名称)	Model/型号	Classification/分类
1	Disposable Medical Mask		I



# EC Declaration of Conformity

*Manufacturer:*

Jinan Shangrun Tongda Composite Material Co., Ltd.  
Shanghe Economic Development Zone, Jinan City,  
Shandong Province, China  
Scarlett Liu  
Tel: 86 531 56773122  
E-mail: 2597675051@qq.com

*whose single Authorized EU-Representative:*

Luxus Lebenawelt GmbH  
Kochstr. 1, 47877, Willich, Germany  
DINDEI DE/0000047791  
Lin Sun  
Tel: 0049- 1715606732  
E-mail: info.m@luxuslw.de

We, the manufacturer, herewith declare that the products

Name: Disposable medical mask  
Type: NON-STERILE  
Model: 17.5cm x 9.5cm

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class I according to Annex IX of the Directive 93/42/EEC. It bears the mark



following the procedure relating to the EC Declaration of Conformity set out in Annex VII of Directive 93/42/EEC.

This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration.

The above mentioned declaration of conformity is exclusively under the responsibility of  
Jinan Shangrun Tongda Composite Material Co., Ltd.  
Shanghe Economic Development Zone, Jinan City, Shandong Province, China

*For and on behalf of*  
Jinan Shangrun Tongda Composite Material Co., Ltd.  
济南尚润通达复合材料有限公司

Place, date Apr 16th 2020

Legally binding signature, Function

*Authorized Representative*