

## EU Declaration of Conformity

Changzhou Shuangma Medical Devices Co. declares that this medical device complies with the following regulations:

<b>MANUFACTURER:</b>	<b>Changzhou Shuangma Medical Devices Co., Ltd.</b> San He Kou Development Zone, Zhenglu, Tianning, Changzhou, 213115, Jiangsu, China
<b>MEDICAL DEVICE:</b>	<b>MEDICAL DISPOSABLE NON STERILE FACE MASK FOR KID</b>
<b>MEDICAL DEVICE BRAND NAME:</b>	<b>BALTIC MASKS</b>
<b>MODEL:</b>	<b>BM-810</b>
<b>CLASSIFICATION:</b>	<b>CLASS I</b>
<b>PRODUCT DESCRIPTION:</b>	Medical device covering the mouth and nose providing a barrier to minimize the direct transmission of infective agents between staff and patient. This intended purpose is normally to limit the transmission of infective agents from staff to patients during surgical procedures and other medical settings with similar requirements. A medical face mask with an appropriate microbial barrier should also be effective in reducing the emission of infective agents from the nose and mouth of an asymptomatic carrier or a patient with clinical symptoms.
<b>QUALITY ASSURANCE AND CONFORMITY:</b>	
Directive:	<b>Medical Device Directive 93/42/EEC</b>
Standard:	<b>EN 14683:2019+AC:2019</b>
Type:	<b>II</b>
Test report:	<b>SL52045301107001TX (SGS October 27, 2020)</b>

This is hereby declared that following designated medical device complied with the essential requirements of above Council Directive(s) and standards.



VILNUS, LITHUANIA, DATE 2020 11 10  
InSpe (BALTICMASKS Brand holder) CEO,  
Guy Shapira



Changzhou, 2020-11-10

*Changzhou 2020-11-10*

Xu Aiping, General Manager

*Xu Aiping*

Document valid until 2021 11 10

This declaration is the responsibility of the Manufacturer