Certificate of Conformity

Huizhou Lexuslance Technology Co.Ltd

Phase I Plant,EilingXiechang Company,Wuyi Village,Chenjiang Sub-district Office,Zhongkai High-tech Zone,Huizhou City,Guangdong Province,China.

The following products have been tested by us with the listed standards and found in compliance with the European Community Directive (EU) 2016/425
Assessment of compliance of the product with the requirements relating to was based on the following standards:

EN 149:2001 +A1:2009

Product: **KN95 Anti Bacterial Respirator**

Model No.: **LK-003**

Parameters: **FFP2**

The statement is based on a single evaluation of one sample of above mentioned products. It does not imply an assessment of the whole production and does not permit the use of the test lab. Logo.
The manufacture should ensure that all product in series production are in conformity with the product sample detailed in this report. The applicant should hold the whole technical report at disposal of the competent all the right.

CE

After preparation of the necessary technical documentation as well as the conformity declaration the required CE marking can be affixed on the product.
Other relevant directives have to be observed.

Marks Licence No.: ACT20031213
Ref. Test Report: 68.5.13.10.2800.2695
Issued Date: 2020-03-12

ACT TESTING TECHNOLOGY AUTHORIZED

Steve Li
Chief Director

Approved by: ACT Testing Technology Co., Ltd.
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Huizhou Lexuslance Technology Co.Ltd

Phase I Plant,EilingXiechang Company,Wuyi Village,Chenjiang Sub-district Office,Zhongkai High-tech Zone,Huizhou City,Guangdong Province,China.

The following products have been tested by us with the for compliance with the Food and Drug Administration Regulations for Polyester resin articles intended for repeated use. It is only valid in connection with the test.

Test Standards: **Food and Drug Administration Regulations**

Product: **self-absorbed particulate respirator (MASK)**

Model No.: **LK-001**

Parameters: **FFP2 NR**

The statement is based on a single evaluation of one sample of above mentioned products. It does not imply an assessment of the whole production and does not permit the use of the test lab. Logo.

The manufacture should ensure that all product in series production are in conformity with the product sample detailed in this report. The applicant should hold the whole technical report at disposal of the competent all the right.

After preparation of the necessary technical documentation as well as the conformity declaration the required FDA marking can be affixed on the product.

Other relevant directives have to be observed.

Marks Licence No.: ACT20031212

Ref. Test Report: 68.5.13.10.2800.2694

Issued Date: 2020-03-12

TESTING TECHNOLOGY ACT AUTHORIZED

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The following products have been tested by us with the listed standards and found in compliance with the European Community Directive (EU) 2016/425
Assessment of compliance of the product with the requirements relating to was based on the following standards:

EN 149:2001 +A1:2009

Product: **Disposable Protective Mask**

Model No.: **LK-001**

Parameters: **FFP2 NR**

The statement is based on a single evaluation of one sample of above mentioned products. It does not imply an assessment of the whole production and does not permit the use of the test lab. Logo.
The manufacture should ensure that all product in series production are in conformity with the product sample detailed in this report. The applicant should hold the whole technical report at disposal of the competent all the right.

CE

After preparation of the necessary technical documentation as well as the conformity declaration the required CE marking can be affixed on the product.
Other relevant directives have to be observed.

Marks Licence No.: ACT20031211
Ref. Test Report: 68.5.13.10.2800.2693
Issued Date: 2020-03-12

ACT TESTING TECHNOLOGY * AUTHORIZED *

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The following products have been tested by us with the for compliance with the Food and Drug Administration Regulations for Polyester resin articles intended for repeated use. It is only valid in connection with the test.

Test Standards: **Food and Drug Administration Regulations**

Product: **KN95 Anti Bacterial Respirator**

Model No.: **LK-003**

Parameters: **FFP2**

The statement is based on a single evaluation of one sample of above mentioned products. It does not imply an assessment of the whole production and does not permit the use of the test lab. Logo.

The manufacture should ensure that all product in series production are in conformity with the product sample detailed in this report. The applicant should hold the whole technical report at disposal of the competent all the right.

FDA

After preparation of the necessary technical documentation as well as the conformity declaration the required FDA marking can be affixed on the product.

Other relevant directives have to be observed.

Marks Licence No.: ACT20031214

Ref. Test Report: 68.5.13.10.2800.2696

Issued Date: 2020-03-12

TESTING TECHNOLOGY ACT AUTHORIZED

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