

Order-no: 843150

RIVM - (Corona Advies)

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| Description | : 1695-7 | | |
|-------------------|---------------|-------------------------|-----------------------------|
| Client project-no | 5.1.2e | | |
| Sample received | : 06-07-2020 | Sample date | : |
| Report date | : 10-07-2020 | Start date microbiology | : 07-07-2020 |
| Sampling | : | Sample delivery | : Mail/Courier |
| Packing | : Plastic bag | Sample temperature | : Room temperature |
| Sealed | : No | Sample condition | : Sample and packing intact |

| Test | | | Result | | |
|-----------|---|---|------------|-----------------|--|
| Bacterial | Filtration Efficiency (BFE) | | | | |
| I 35000 | Testing BFE (n=5) (equiv. NEN-EN 14683+C1) | | | | |
| I 35050 | Test conditions | | | | |
| I 35051 | Dimensions of test specimens (width x height) | | Whole mask | | |
| I 35052 | Size of the area tested (width x height) | | 48.0 | cm ² | |
| I 35053 | Side facing the aerosol | | Face side | | |
| I 35054 | Flow rate during testing | | 28.3 | L/min | |
| I 35060 | Mean of the total plate counts of the two positive controls | а | 2,170 | cfu | |
| I 35070 | Mean plate count of the negative controls | а | 0 | cfu | |
| I 35100 | Bacterial Filtration Efficiency (BFE, equiv. NEN-EN 14683+C1) | | | | |
| I 35101 | BFE specimen 1 | а | 99 | % | |
| I 35102 | BFE specimen 2 | а | 99 | % | |
| I 35103 | BFE specimen 3 | а | 99 | % | |
| I 35104 | BFE specimen 4 | а | 99 | % | |
| I 35105 | BFE specimen 5 | а | 99 | % | |
| I 35199 | Average BFE | а | 99 | % | |

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Free State S



843150 Crder-no :

a) This result is possibly not representative for the microbiological composition of the sample at the moment of sampling. (The elapsed time between sampling and analysis is longer than allowed or not known.)

*) Performance requirements for medical face masks (acc. European Standard no. EN 14683:2019+AC):

Bacterial filtration efficiency (BFE) (%): Type I a >= 95, Type II >= 98, Type IIR >= 98 Differential pressure (Pa/cm2): Type I a < 40, Type II < 40, Type IIR < 60 Microbial cleanliness (cfu/g): Type I a <= 30, Type II <= 30, Type IIR <= 30

a) Type I medical face masks should only be used for patients and other persons to reduce the risk of spread of infections particularly in epidemic or pandemic situations. Type I masks are not intended for use by healthcare professionals in an operating room or in other medical settings with similar requirements.

This certificate of analysis is a test report. The tested samples are part of the mentioned batch/lot number. Batch validation is not the scope of this report.

Start date analysis: 07-07-2020, end date: 09-07-2020.

Findings are based on the sample as submitted. For more detailed information on applied methods please contact the operational manager.

Any interpretation of analytical results mentioned on this certificate lies outside the scope of accreditation.

With the unit % is meant: w/w%, unless otherwise stated. Nutrilab is not responsible for the information provided by the client.

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| Nutrilab bv Postbus 7, 4284 ZG Rijswijk Burgstraat 12, 4283 GG Giessen t. (0183) 44 63 05 f. (0183) 44 25 97 | info@nutrilab.nl www.nutrilab.nl KvK 18114291 BTW NL002007654B01 | Page 2/2 Deplanation of symbols: Q RM accreditated test (SO /IEC 17025) 1 Test performed by Nutritab BV E Test performed by sub-contractor |
|--|---|--|