

Special Issue:

Special Issue on COVID-19 Aerosol
Drivers, Impacts and Mitigation (XI)

OPEN ACCESS 

Received: July 21, 2020

Revised: November 3, 2020

Accepted: November 24, 2020

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Publisher:

Taiwan Association for Aerosol
Research

ISSN: 1680-8584 print

ISSN: 2071-1409 online

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Efficacy Assessment of Newly-designed Filtering Facemasks during the SARS-CoV-2 Pandemic

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ABSTRACT

The SARS-CoV-2 pandemic resulted in shortages of production and test capacity of FFP2-respirators. Such facemasks are required to be worn by healthcare professionals when performing aerosol-generating procedures on COVID-19 patients. In response to the high demand and short supply, we designed three models of facemasks that are suitable for local production. As these facemasks should meet the requirements of an FFP2-certified facemask, the newly-designed facemasks were tested on the filtration efficiency of the filter material, inward leakage, and breathing resistance with custom-made experimental setups. In these tests, the facemasks were benchmarked against a commercial FFP2 facemask. The filtration efficiency of the facemask's filter material was also tested with coronavirus-loaded aerosols under physiologically relevant conditions. This multidisciplinary effort resulted in the design and production of facemasks that meet the FFP2 requirements, and which can be produced at local production facilities.

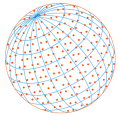
Keywords: Coronavirus filtration efficiency, Facemask, FFP2, NaCl particle filtration, Respirator

1 INTRODUCTION

The severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) is a novel coronavirus that was first identified in December 2019 in Wuhan, China, in patients suffering from acute respiratory syndrome (2019 coronavirus disease or COVID-19) (Zhu *et al.*, 2020). SARS-CoV-2 was declared pandemic by the WHO on March 11, 2020.

Healthcare professionals who are involved in aerosol-generating procedures on COVID-19 patients are required to use FFP2-classified filter facepiece respirators for respiratory protection against SARS-CoV-2 infection. FFP2-respirators (from now on called 'facemask') filter at least 94% of the submicron-sized test aerosols with NaCl, according to the NEN-EN 149:2001+A1:2009 standards (Table 1). These standards are used by the accredited test laboratories (the so-called notified bodies) within the European Union member states to test and certify the facemasks upon approval. According to these standards, FFP2-classified facemasks are also required to meet the criteria for inward leakage, maximal CO₂-content of inhaled air, and breathing resistance, which are summarized in Table 1.

As the global demand for FFP2-facemasks largely exceeded the production, distribution and test capacities of the conventional suppliers and notified bodies during the COVID-19 pandemic, we formed a Dutch collaborative initiative, consisting of the Reinier de Graaf hospital, Royal DSM,

**Table 1.** Requirements for FFP2-classified FFRs according to the NEN-EN149 standards.

Filter penetration test	FIT-test	CO ₂ content test	Breathing resistance test	
Max. penetration of 120 mg NaCl test aerosol at 95 L min ⁻¹	Max. inward leakage	Max. CO ₂ content of inhaled air	Inhalation	Exhalation
6%	8 ^a or 11 ^b %	< 1% on average	0.7 mbar at 30 L min ⁻¹ 2.4 mbar at 95 L min ⁻¹	3.0 mbar at 160 L min ⁻¹

^a for at least 8/10 individual wearer arithmetic means.^b for at least 46/50 individual exercise results (e.g., 10 subjects × 5 exercises).

Delft University of Technology, Leiden University Medical Center and Erasmus University Medical Center, with the aim to design facemasks that meet the FFP2-specifications. These facemasks were initially tested with custom-made test equipment on their filtration capacities of NaCl particles, fit, and breathing resistance under conditions that approximate the NEN-EN149 standards before being tested by a certified test laboratory. Those setups have been described in detail in Blad *et al.* (2020). The facemasks were also tested on virus filtration efficiency (VFE) with the mouse hepatitis virus (MHV), a beta coronavirus that causes lethal hepatitis in mice (Gledhill *et al.*, 1955), but that is non-pathogenic to humans. These studies identified mask designs that filter 99% of coronavirus-loaded aerosols, just as a commercial FFP2-facemask.

2 METHODS

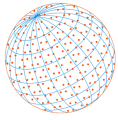
2.1 Facemask Designs

Three different types of in-house designed facemasks were tested in triplicate in various custom-made experimental setups. The “Reinier 0.1” facemask was constructed with a dental facemask, that consisted of two layers polypropylene nonwoven fabric (40 and 20 gr m⁻²) and a single layer of 20 g m⁻² melt-blown fabric, to which two extra layers of spun-bond polypropylene filters (100 and 20 g m⁻²) was added. The “DSM 1.0” facemask consisted of five polypropylene nonwoven filter layers that consisted of 55 g m⁻² spunmelt, 20 g m⁻² melt-blown, 30 g m⁻² melt-blown, 20 g m⁻² melt-blown, and 47 g m⁻² spunmelt polypropylene filters. The name of this mask refers to the collaboration with the Dutch nutrition and health company Royal DSM during the development of this facemask. The “Reinier 1.0” facemask consisted of three layers of filter material, of which two layers of spun-bond polypropylene (100 and 20 gr m⁻²) and a single 20 gr m⁻² melt-blown polypropylene layer. These facemasks were benchmarked against the FFP2-certified 3M Aura 1862(+) facemask. All layers of filter material were used throughout the masks and were therefore considered as homogeneously distributed. The electrostatic charge of the aerosols and material was not measured.

2.2 The Dry Particle Penetration Test

The dry environmental particle penetration test was performed to provide an initial indication of the filtering capacity of the in-house designed facemasks. The particles were counted by a Solair 3200 particle counter (Lighthouse), which is normally used for quality control measurements of the operating rooms. The particle counter was connected with a particle chamber, on which a facemask was fixed in an airtight manner. The particle counter generated a flow of 56.6 L min⁻¹, which created an air velocity of 0.25 m s⁻¹ through the facemask’s filter material. Particles in the range of 0.3–0.5 μm, 0.5–1.0 μm, 1.0–3.0 μm, 3.0–5.0 μm, and 5.0–10.0 μm in size were counted. First, a reference count of the number of environmental particles (# particles_{Ref}) was performed in absence of a facemask and subsequently, the number of environmental particles was counted after fixing a facemask on the particle chamber (# particles_{mask}) and averaged over three one-minute measurements. For each of the five particle size ranges, the filter capacity was calculated according to the following formula:

$$\text{Filter capacity (\%)} = ((\# \text{ particles}_{\text{Ref}} - \# \text{ particles}_{\text{mask}}) / \# \text{ particles}_{\text{Ref}}) \times 100 \quad (1)$$



2.3 NaCl Particle Penetration Test

For the NaCl particle penetration test a PVC-tube system was constructed with a 90° bend, going from a vertical to a horizontal direction (the design and instructions to build are online available at <https://projectmask.nl/testing/filter-material-penetration/build/>). The vertical part of the tube system was connected with an Atomizer Aerosol Generator ATM 226, and the distal end of the horizontal part contained a PMMA tube with a sample holder in which a facemask was fixed in an airtight manner (Figs. S1 and S2). The aerosol generator produced NaCl particles from a 2% NaCl solution, resulting in a particle concentration of $2.5\text{--}3.5 \times 10^4$ particles cm^{-3} in the tube system. The number of NaCl particles that passed through the facemask's filter material was counted by a TSI PortaCount Pro 8030 particle counter, which detects particles in a 0.02 to $> 1 \mu\text{m}$ size range and generated an air velocity of 0.1 m s^{-1} through the filter material of the facemasks. The number of particles that passed through the filter material was analysed with TSI FitPro⁺ software.

2.4 The Fittest

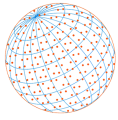
The fittest was performed to determine the leakage of particles around the edges of the facemasks. First, a probe was placed in the facemask by which it was connected to the PortaCount particle counter. To obtain representative results, each facemask was tested by three different individuals, who performed eight different exercises: normal breathing, deep breathing, moving the head side to side, moving the head up and down, talking, grimace, bending over and normal breathing, as described in the NEN-149 standards. The particle leakage was determined by comparing the number of particles in front and behind the facemask's filter material and was recorded with a PortaCount particle counter. The data were analysed with the TSI FitPro⁺ software.

2.5 The Breathing Resistance Test

The breathing resistance of the facemasks was determined with a custom-made experimental setup (Fig. S3; the design and instructions to build are online available at <https://projectmask.nl/testing/breathing-resistance/build/>). The facemask was placed on a manikin head (described in Zhuang and Bradtmiller, 2005) that was connected to a tube system, in which the pressure drop after the filter material was recorded. Before testing the facemasks, the pressure sensors were calibrated without any specimen at inlet or exit (open flow), blockage of the exit (blocked flow), or with a known flow resistance. When adding the masks, the increasing airspeed led to an increased airflow through the facemask and a pressure drop in the system, which was related to the resistance.

2.6 Virus Filtration Efficiency Test

As no standardized test procedures exist to determine the virus filtration efficiency of facemasks, an in-house designed experimental setup was developed that consisted of a curved tube with a 0.45 m vertical and a 0.9 m horizontal part with at the distal end a sample holder for airtight placement of a facemask (Fig. S4). This tube was connected to a mixing chamber, in which the virus-loaded aerosols were mixed with mist droplets of bidest, that was generated by an ultrasonic mist maker, for more efficient virus collection. The mixing chamber was connected to three SKC BioSampler impingers, in which the collected virus was impinged into 45 mL virus transport medium (VTM) (HMEM (Lonza), 12% v/v glycerol, 0.5% w/v Lactalbumin enzymatic hydrolysate, 0.02 mg mL^{-1} Polymyxin B sulfate, 0.01 mg mL^{-1} Nystatin, Penicillin/Streptomycin mixture 240/240 U mL^{-1} and 0.3 mg mL^{-1}). The maximal airflow in each SKC BioSampler was 12.5 L min^{-1} , generating a total maximal flow of 37.5 L min^{-1} . The facemasks were challenged with mouse hepatitis virus (MHV), a beta coronavirus that causes hepatitis in mice, to be able to perform these tests under laboratory biosafety level (BSL) 2 conditions. 10^8 plaque-forming units (PFUs) of MHV were aerosolized with an Aerogen Solo nebulizer, which produced small inhalable particles (MMAD $2.1 \mu\text{m}$) in the vertical part of the tube. The particles were then directed through the 90° bend, and subsequently the horizontal part of the tube system with the airtight fixed facemask. The facemasks were placed in the sample holder in an airtight manner between two sanitary rings with a diameter of 40 mm, which created an air velocity of 0.42 m s^{-1} at a continuous airflow of 31.5 L min^{-1} ($3 \times 12.5 \text{ L min}^{-1}$ minus 6 L min^{-1} for the mist maker). The self-made facemasks were tested in triplicate and benchmarked against the FFP2-certified 3M Aura



1862(+) facemask. The aerosolized virus was also collected in the absence of a facemask as a reference. Between each test, the tube system was flushed with HEPA-filtered air for 15 min at a flow of 37.5 L min⁻¹.

2.7 Virus Quantification

Viral RNA copy number was quantified by a quantitative reverse-transcription polymerase chain reaction (RT-qPCR) analyses and the number of infectious virus particles by plaque assays.

For RT-qPCR, viral RNA was isolated from 135 µl VTM with the QIAamp Viral RNA Mini Kit (Qiagen) according to the manufacturer's protocol. Equine arteritis virus (EAV) was added to the lysis buffer as an internal control for the RNA isolation and RT-qPCR efficiency as described in Scheltinga *et al.* (2005). The isolated RNA was converted to copy DNA (cDNA) and quantified in a TaqMan Fast Virus 1-step master mix (Applied Biosystems) in the presence of 450 nM primers (MHV-FPr1: ACGCCGCCTTATTAAGATG, MHV-RPr1: GGCATAGCAGGATCACATTT) and 200 nM probe (TexRed-TCCTGTACTCATGGGTT GGGACTATCC-BHQ2) that targets the viral gene *nsp12*, coding for RNA dependent RNA polymerase (RdRp). The primers and probe for the EAV internal control were also added and have been described in Loens *et al.* (2012). The reactions were run in a CFX384 Q-PCR Thermocycler (BioRad) with a two-step protocol: Cycle 1 (1×): 50°C, 5 min. and 95°C for 20 sec., and Cycle 2 (45×): 95°C for 5 sec. and 60°C for 30 sec. An MHV standard was generated by *in vitro* transcription with the T7 mMessage mMachine kit (ThermoFisher) according to the manufacturer's protocol and was also analysed as described above to determine the copy number of viral RNA in the samples.

The number of infectious virus particles was determined by a plaque assay. 8×10^5 17CL1 cells, derived from mouse (*Mus musculus*) BALB/c fibroblasts, were seeded in six-wells plates and incubated overnight at 37°C. The cells were inoculated with an undiluted or one of the diluted MHV samples from a ten-fold serial dilution for 1 hr at 37°C. After inoculation, the cells were washed twice with PBS, overlaid with an Avicell (Sigma) overlay, and incubated for an additional 24 hrs at 37°C. The cells were subsequently fixated with 3.4% formaldehyde in PBS for 1 hr at room temperature and stained with 0.75% crystal violet staining solution for 5 min at room temperature. After removal of the staining solution, the wells were washed with water, and the number of plaques was counted.

3 RESULTS AND DISCUSSION

3.1 NaCl Particles were more Efficiently Filtered than the Environmental Particles of between 0.3–0.5 µm

The filtering capacities of the new facemasks were initially determined by measuring the filtration of dry environmental particles in a broad particle size range (0.3–10 µm) with a Solair 3200 particle counter by airtight placement of one of the facemasks on the particle chamber. A technical limitation of using the Solair 3200 particle counter is that only particles larger than 0.3 µm were counted. The broad particle size range was chosen to obtain a general impression on the filtering capacities of the new facemasks. Furthermore, it is currently unknown what particle sizes are relevant for the airborne transmission of the SARS-CoV-2. The size of the virus particle is estimated to be 0.1 µm, but to which extend transmission takes places via aerosols (< 5 µm droplets) or larger respiratory droplets (> 5 µm) remains to be determined. When the FFP2-certified facemask was placed on the particle chamber, 99.4–99.9% of the environmental particles of between 0.3–10.0 µm was filtered (Fig. 1(a)). Two models of the self-designed masks showed significantly lower filtration efficiencies for particles of between 0.3–0.5 µm than the certified mask, in which the Reinier 0.1 filtered $74.6 \pm 0.4\%$ and the Reinier 1.0 $88.2 \pm 0.8\%$. The DSM 1.0 performed better than the other two models in this size range as this mask filtered $97.9 \pm 0.3\%$ of these particles. At least 96.9% of the particles larger than 0.5 µm were filtered by each of the three models. These observations indicate that the various types of masks are able to filter a significant amount of environmental particles. However, as the most penetrating particle size (MPPS) for fibrous filters has been shown in literature to vary between 0.03–0.5 µm (Lee and Liu, 1980; Podgorski *et al.*, 2006; Rengasamy *et al.*, 2009), further testing with submicron-sized NaCl particles under standardized conditions is required.

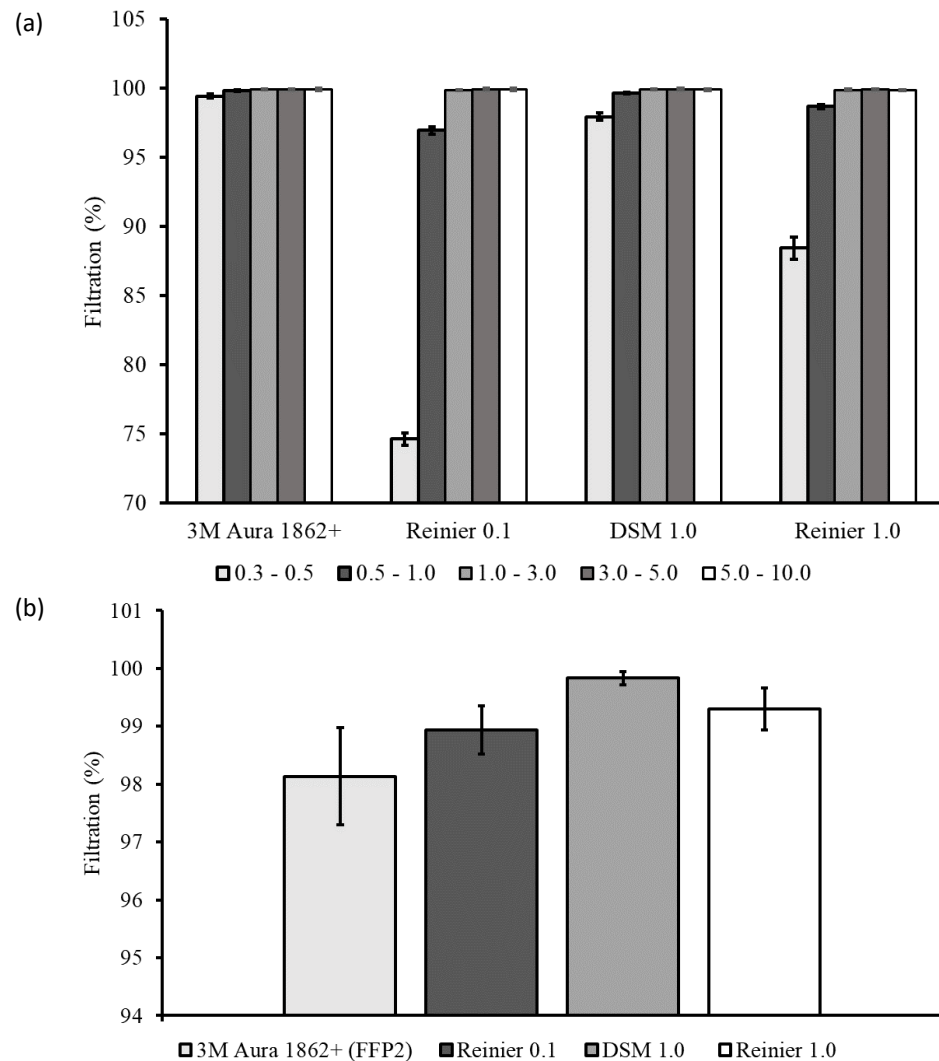
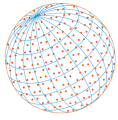
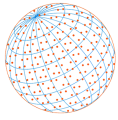


Fig. 1. (a) Filtration efficiency (mean \pm sd.) of environmental dry particles and (b) non-neutralized polydisperse NaCl particles by the manually-produced facemasks and benchmarked against an FFP2-certified facemask.

The masks were therefore also challenged with non-neutralized polydisperse NaCl-particles (0.02–2.0 μm ; MMD: 0.6 μm) under conditions that closely approximate the NEN-149 standards. According to these standards, FFP2-facemasks are required to filter at least 94% of the NaCl-particles. The FFP2-certified facemask filtered $98.13 \pm 0.84\%$ of the NaCl particles (Fig. 1(b)). The Reinier-0.1 and -1.0 facemasks filtered respectively $98.9 \pm 0.42\%$ and $99.3 \pm 0.36\%$ of the NaCl particles. This is significantly more than observed for the dry environmental particles of between 0.3–0.5 μm . These two models contained a single layer of melt-blown polypropylene and the above-described observations indicate that the two layers nonwoven fabric of the three-layer dental facemask of the Reinier-0.1 model can be removed without affecting the facemask's filtering performance. The DSM 1.0-facemasks is the only type with three layers of melt-blown polypropylene filters and showed the highest NaCl-particle filtration efficiency of $99.83 \pm 0.12\%$ (Fig. 1(b)). This model also performed best in the environmental particle filtration test (Fig. 1(a)).

The submicron-sized environmental particles were filtered less efficiently by the newly-designed facemasks than the NaCl particles, which might be explained by the higher air velocity during the environmental particle filtration test. It has been well described in literature that a higher air velocity results in a lower filtration efficiency (Rengasamy *et al.*, 2010, 2011, Miguel, 2013, Gao *et al.*, 2016, Mukhametzanov *et al.*, 2016). Another explanation is that the MPPS of the Reinier-0.1 and -1.0 mask coincidentally fell between 0.3–0.5 μm during the environmental



particle filtration test, while under the standard conditions with NaCl particles the MPPS might be an order of magnitude smaller as previously observed by Rengasamy *et al.* (2009), since the MPPS is dependent on aerosol type (He *et al.*, 2013) and air velocity (Lee and Liu, 1980). Furthermore, it has been shown by others that particles of other sources might have higher penetration rates than the NaCl-particles (Penconek *et al.*, 2013; Grima-Olmedo *et al.*, 2014, Serfozo *et al.*, 2017). Although the NaCl particle filtration test might overestimate a mask's filtering performance, this test is currently the golden standard when assessing the facemask's filter capacity. The NaCl-particle filtration efficiency of the Reinier 0.1- and DSM 1.0-facemasks was also verified by notified bodies and showed similar filtration efficiencies as in this study (Figs. S5 and S6), confirming that the custom-made setup indeed results in similar observations as under the conditions of the NEN149 standards. Hence, a good indication has been obtained that the filter capacity of the three types of facemasks meets the FFP2 requirement under standardized conditions.

3.2 Reinier-0.1 and -1.0 Facemasks have Acceptable Inward Leakage and Breathing Resistance

A proper fit and acceptable breathing-resistance are also crucial for the protective quality of a facemask. FFP2-certified facemasks are allowed to have an average inward leakage of maximal 8 or 11%, depending on the number of tests and exercises, according to the NEN-149 standards. (Table 1). As our facemasks were tested by only three different persons, not the average, but maximal observed inwards leakage was taken as a measure for the fit of a facemask (Table 2), thus defining a worst-case scenario. The FFP2-certified facemasks showed a maximal inward leakage of 0.5% (Table 2). Of the self-designed masks, the Reinier-0.1 and -1.0 models have acceptable fits of below 8%, whereas the DSM 1.0 shown significant leakage of 14.6%. For the breathing resistance tests, the facemasks were placed on a manikin head and the pressure drop was recorded while the airflow was increased. The maximal pressure drop at an inhalation flow of 30 or 95 L min⁻¹ according to the NEN-149 standards is 0.7 and 2.4 mbar respectively. The pressure drops varied between 0.14 and 0.16 mbar at a continuous flow of 30 L min⁻¹, and at a flow of 95 L min⁻¹, the pressure drop varied between 0.74 and 0.85 mbar for all three facemask models (Table 3). Exhalation at a continuous flow of 160 L min⁻¹ resulted in a pressure drop of between 1.11 and 1.44 mbar for the three different designs, which is below the 3.0 mbar that is allowed according to the NEN-149 standards. These observations indicate that all three designs meet the NEN149 standards regarding breathing resistance. However, for the DSM-facemask this result might be due to the significant amount of inward leakage.

3.3 The Prototype Facemasks Filter at Least 98% of Virus-loaded Aerosols

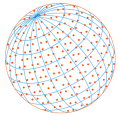
The facemask were also challenged with virus-loaded aerosols as an extra testcase for the filtration capacities of the new facemasks in addition to the conventional test with NaCl particles.

Table 2. Maximal and minimal observed inward leakage.

Facemask	Max. observed inward leakage (%)	Min. observed inward leakage (%)
3M Aura 1862+ (FFP2)	0.5	0.5
Reinier 0.1	4.8	4.2
DSM 1.0	14.6	6.7
Reinier 1.0	0.8	0.5

Table 3. Pressure drop over the self-designed facemasks at continuous inhalation flows of 30 or 95 L min⁻¹ and an exhalation flow of 160 L min⁻¹.

Facemask	Inhalation 30 L min ⁻¹ (0.7 mbar max.)	Inhalation 95 L min ⁻¹ (2.4 mbar max.)	Exhalation 160 L min ⁻¹ (3.0 mbar max.)
Reinier 0.1	0.14 ± 0.003	0.74 ± 0.009	1.44 ± 0.05
DSM 1.0	0.14 ± 0.003	0.84 ± 0.008	1.31 ± 0.06
Reinier 1.0	0.16 ± 0.002	0.85 ± 0.01	1.11 ± 0.06



Furthermore, this test was strongly demanded by the healthcare professionals. 10^8 PFU of MHV, a beta coronavirus that infects mice, was aerosolized in our custom-designed tube system and passed through the filter material of each airtight fixated mask. The high number of PFU was required to observe at least 2 logs decrease in the number of PFUs, equivalent to 99% filtration, as it was expected that most virus particles might be lost after aerosolization into the tube system. This number of virus particles is orders of magnitude larger than the number of infectious SARS-CoV-2 particles that was sampled in a patient room with two COVID-19 patients (Lednický *et al.*, 2020). The virus was aerosolized with an Aeroneb Solo nebulizer, which creates small inhalable particles (MMAD 2.1 μm). When those particles were passed through a tube over 0.9 m distance, 99% of the particles were $< 5.0 \mu\text{m}$ in size (Fig. 2). Particles smaller than $5.0 \mu\text{m}$ are generally defined as aerosols that can be easily inhaled (Heyder *et al.*, 1980). The three types of facemask were tested in two separate sessions, which resulted in two independent datasets (Fig. 3) In the no mask-control, MHV was sampled in absence of a facemask in the sample holder to determine the maximum amount of virus that could be recovered from the air. This resulted in virus recovery of $4.09 \pm 0.21 \log \text{PFU mL}^{-1}$ in the first, and $5.89 \pm 0.21 \log \text{PFU mL}^{-1}$ in the second session (Figs. 3(a) and 3(c)), indicating that the virus collection efficiency in the latter session was higher than in the first session. Nevertheless, the collection efficiency of infectious virus particles was high enough to observe at least a 2-logs decrease in virus collection after placing a facemask in the sample holder. When an FFP2-certified facemask was placed in the sample holder, the amount of infectious virus and viral RNA copies that were recovered behind the facemask were respectively on average 2.3- and 2.4-logs lower than in the no-mask control in the first session. This corresponds to filtration efficiencies of 99.4 and 99.6% (Fig. 3(b)). When this type of facemask was used in the second session, the infectious virus and viral RNA recovery decreased by respectively 3.1- and 3.5-logs on average, which corresponds to 99.92 and 99.96% filtration efficiency (Fig. 3(d)). The Reinier-0.1 facemasks reduced the recovery rates of infectious virus and RNA copies on average by 2.1 and 2.8-logs respectively compared to the no mask-control (Fig. 3(a)). This corresponds with the filtration efficiencies of 99.2 and 99.9% (Fig. 3(b)). 2.3 and 2.6-log reductions were observed with the DSM 1.0-facemasks, which corresponds with 99.3 and 99.6% filtration efficiency. The Reinier-1.0 facemasks were tested in the second session, which reduced the infectious virus and total viral RNA copy numbers with 3.1- and 2.6 logs on average, corresponding to 99.92 and 99.71% filter efficiency (Figs. 3(c) and 3(d)). The RNA copy nr to PFU-ratio with and without facemask are similar within each experiment, indicating that the mask's filter material does not inactivate the virus upon passage.

The virus filtration efficiency of the facemasks was determined at a continuous air velocity of 0.42 m s^{-1} , which is significantly higher than during the NaCl particle penetration test, that was performed under conditions according to the NEN-149 standard. Similar filtration efficiencies

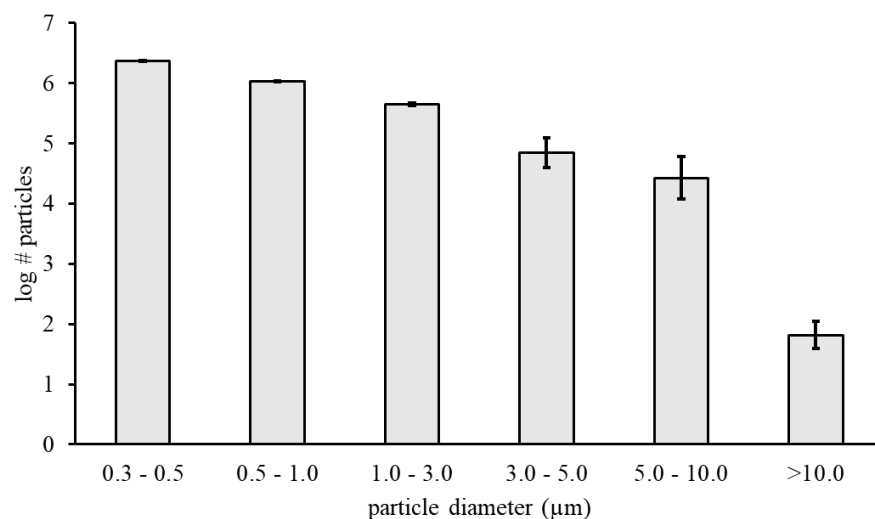


Fig. 2. Particle size range distribution (mean \pm sd.) after aerosolization of $100 \mu\text{L}$ virus diluent by the Aeroneb Solo nebulizer and passage through a 0.9 m tube.

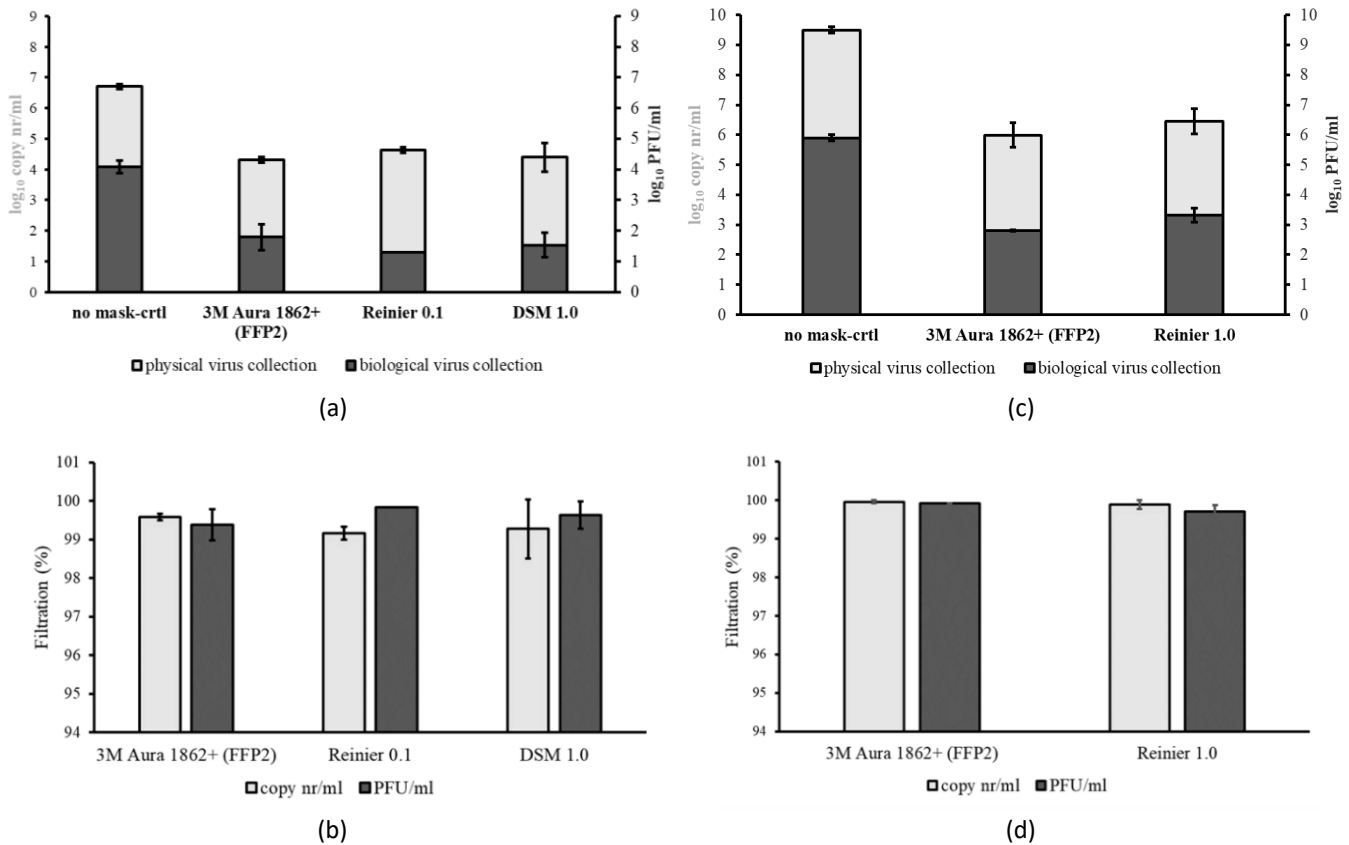
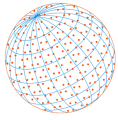
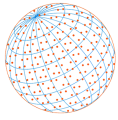


Fig. 3. Physical (copy nr mL^{-1}) and biological (PFU mL^{-1}) collection of (a and c) aerosolized mouse hepatitis virus and (b and d) filtration efficiencies for the self-designed facemasks. The FFP2-certified mask was used as a benchmark.

were nevertheless observed as in the NaCl particle filtration test, which might be explained by larger sizes of the virus-loaded aerosols compared to the NaCl particles. The penetration rates of specifically the submicron-sized virus-loaded aerosols could unfortunately not be determined in this experiment. The biological origin of the virus-loaded aerosols is not relevant for the filtration efficiency by a facemask (Rengasamy *et al.*, 2017). The air velocity used in present study was also higher than described in various other studies where the virus filtration efficiencies of facemasks were tested (Borkow *et al.*, 2010; Harnish *et al.*, 2013, 2016; Rengasamy *et al.*, 2017; Zhou *et al.*, 2018). It was chosen here to test at 0.42 m s^{-1} as this is a more physiological relevant air velocity during inhalation as the maximal air velocity during an inhalation cycle reaches up to 1 m s^{-1} (Tang *et al.*, 2013). Despite the higher air velocity and the usage of a different virus, at least 98% of the virus-loaded aerosols were filtered by the FFP2-certified facemask, similarly as observed in the prementioned studies. However, significant lower virus filtration efficiencies (79% on average) were observed in a recently published airborne simulation experiment at an air velocity of 2 m s^{-1} (simulating coughs) when testing airtight fixated N95 facemasks on a recipient mannikin head (Ueki *et al.*, 2020). The virus-loaded aerosols can apparently be forced through the filter material at this air velocity, which severely affected the filtering efficiency a facemask. Furthermore, when the N95-mask was not airtightly fixed to the mannikin-head, which would be the case in practice, the virus filtration efficiency was found comparable with that of a surgical mask. So although our self-designed masks were shown here to filter submicron-sized NaCl particles and virus-loaded aerosols, and showed acceptable breathing resistance and fits, more research under realistic conditions is needed to assess the protective capacity of the new facemasks.

4 CONCLUSION

It is concluded that under standardized conditions the Reinier-0.1 and -1.0 facemasks meet



the requirements of the NEN-149 standards in terms of filtration capacity, breathing resistance and fit. The self-designed masks were also shown to filter 99% of virus-loaded aerosols after airtight fixation, but more research is needed to assess the protective capacity of these masks.

ACKNOWLEDGMENTS

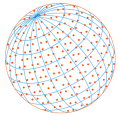
This work was supported by the NIH/NIAID (contract number HHSN272201400008C) and European Union's Horizon 2020 research and innovation program VetBioNet (grant agreement No 731014). Part of this research was supported by the Leiden University Fund (LUF), the Bontius Foundation, and donations from the crowdfunding initiative "wake up to corona". S.H. was funded in part by an NWO VIDI grant (contract number 91715372). Part of this research was supported by ZonMw initiative "creative solutions against CoVID-19" (5001-0013). The authors also wish to thank Monique Elsing-van Olphen for supporting this research and for supplying information on the facemask designs.

SUPPLEMENTARY MATERIAL

Supplementary data associated with this article can be found in the online version at <https://doi.org/10.4209/aaqr.2020.07.0424>

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