High-Performance Virus Removal Filter Paper for Drinking Water Purification

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Access to drinking water is one of the greatest global challenges today. In this study, the virus removal properties of mille-feuille nanocellulose-based filter papers of varying thicknesses from simulated waste water (SWW) matrix are evaluated for drinking water purification applications. Filtrations of standard SWW dispersions at various total suspended solid (TSS) content are performed, including spiking tests with 30 nm surrogate latex particles and 28 nm ΦX174 bacteriophages. Filter papers of thicknesses 9 and 29 µm are used, and the filtrations are performed at two different operational pressures, i.e., 1 and 3 bar. The presented data using SWW matrix show, for the first time, that a filter paper made from 100% nanocellulose has the capacity to efficiently remove even the smallest viruses, i.e., up to 99.9980–99.9995% efficiency, at industrially relevant flow rates, i.e., 60–500 L m⁻² h⁻¹, and low fouling, i.e., $V_{\text{max}} > 10^3–10^4$ L m⁻². The filter paper presented in this work shows great promise for the development of robust, affordable, and sustainable water purification systems.

1. Introduction

Urbanization and population growth lead to increased and inhomogeneous consumption of water resources. The problem of drinking water is very complex and multifaceted, and, therefore, it is relevant for both high- and low-income countries. This is why access to safe drinking water and sanitation is among the UN’s Sustainable Development Goals, i.e., Agenda 2030.[1] In low-income countries, when a centralized water supply is not available or the quantity is limited, poor households have to rely on water obtained from rivers, rain, or shallow wells.[2] The lack of safe water is a part of a vicious cycle, where poverty and poor healthcare feed each other. Water safety is also an important issue in high-income countries with centralized water supply since water-borne outbreaks continue to occur[3–10] and many more incidents are expected, especially among susceptible population groups, including increasing elderly population above the age of 65, children below the age of 5, chronically ill patients, recipients of immunosuppressive therapies, and pregnant women.[11] Globalization of commerce, expansion of food supply sources from regions with unimproved irrigation, people migration from areas with poor hygiene, and natural sporadic mutations contribute to the spread of water-borne diseases.[11]

There are about 140 known water-borne pathogens, including viruses, bacteria, and protozoa. Viruses are the smallest and probably the most difficult to deal with from them all. They have the greatest infectivity among all known water-borne pathogens and high resistance to disinfection.[11] They are further excreted with feces in relatively large quantities ($>10^{11}$ g⁻¹) and generally exhibit longest survival potential in the environment. It is estimated that known viral pathogens account for nearly 8% of water-borne infection outbreaks. It should however be noted that water-borne infections of undetermined etiology, which account for almost half of all documented cases, are probably due to viruses for which currently there are no suitable laboratory host cultures.[11] According to the World Health Organization (WHO), moderately to severely infectious water-borne viral pathogens are numerous and include adenovirus, astrovirus, echovirus, hepatitis viruses, rotavirus, caliciviruses (including norovirus), enteroviruses (including coxsackieviruses and polioviruses), polyomaviruses, cytomegalovirus, papillomaviruses, influenza viruses, and coronaviruses.[12–14] Massive virus outbreaks due to poor water quality can occur in any country, including countries with centralized water supply and sustained vigilance.[15–10] Only in the USA, there are estimated 4 million to 33 million cases of acute gastrointestinal illnesses each year due to drinking water.[15,16] Virus clearance is more difficult than that of bacteria, and, therefore, bacteriologically safe water may not be necessarily virally safe.

Traditionally, the main source for water-borne pathogens has been considered to be surface water, since all surface waters contain water-borne pathogens, with a substantial number of zoonotic infections. Animal farms and sewage discharges in close proximity are well-known sources of contamination for surface water.[11] Recently, it is recognized that another potential source of virus contamination, previously considered safe and therefore unmonitored, is groundwater, which is utilized...
directly by small communities and private homeowners. Several surveys in the USA documented evidence of viruses in ground water, which is often associated with proximity of septic systems to water sources due to landfill leaks, improper waste-water disposal, and septic tank contamination. It is estimated that about half of the water-borne disease outbreaks in the USA is due to untreated groundwater. Also in the European Union (EU), where there are ≈85 000 decentralized small supplies providing drinking water to ≈65 million citizens, almost 40% of them are not properly monitored or do not deliver drinking water complying with microbiological quality standards.

In water treatment, virus inactivation, aimed at reducing the infectivity of viruses, can be achieved both by physical (e.g., UV-C and y-irradiation and boiling) and chemical means (e.g., oxidants and coagulants). Chemical disinfection, e.g., with chlorine, is the most common way of disinfection. However, higher doses are normally required to destroy virus as compared to bacteria, and chlorination may also lead to toxic by-products (e.g., trihalomethanes and haloacetic acids) as well as residual infectivity depending on the water quality. An increased reliance on UV light for water disinfection has raised the concerns for viral resistance, especially related to adenovirus infection, which is especially problematic for immunocompromised population, e.g., in cancer and bone marrow transplant patients. Chemical coagulation using metal ions, e.g., alum (Al2(SO4)3), salts of iron, lime (Ca(OH)2), and various polyelectrolytes is the second most common method of inactivating viruses from water after disinfection. However, research suggests that flocculation does not remove all viruses exhaustively, showing varying results between log10 virus titer reduction of 1 and 2.86.

Size-exclusion filtration is a very robust and highly efficient method of clearing viruses, which does not require using chemicals, heat, or radiation. Most common filters, including ceramic- and biosand-based filters, used in water treatment, while effective against bacteria (typically >6log10 reduction) feature low virus removal capacity (typically, <1log10 reduction). Therefore, to be able to completely remove viruses from water, only advanced filters can be used featuring pore sizes of around 10–20 nm. In this respect, affordable and efficient ultralifers, which are preferably produced from sustainable raw materials, i.e., nonplastic, are highly demanded.

In the context of above discussion, cellulose-based filters for water treatment would be useful due to the abundance, sustainability, and cost efficiency of this raw material. Traditional general-purpose filter paper has too large pores to remove bacteria or viruses from water. Various adsorptive-type virus retentive cellulose-based depth filters have been developed. However, because viruses possess highly variable isoelectric points, ranging between 1.9 and 8.4, adsorptive-type virus retentive depth filters show limited robustness and their function can be interfered by other competing charged species. On the other hand, the size-exclusion virus removal filters based on cuprammonium-regenerated cellulose, e.g., Planova filters by Asahi Kasei, used for purification of protein-based drugs and which are manufactured by phase inversion processing, remain unaffordable for water treatment applications due to their high price.

Virus removal filters should combine high flow rates, low fouling, and high virus removal capacity, which explains the high price of such filters. Controlling the pore-size distribution of paper filter in the region suitable for virus removal is not

Table 1. CP-DSC analysis of nanocellulose filter papers.

| Filter paper thickness [μm] | T_{pk}^{b)} [°C] | ΔT_{on-pk}^{b)} [°C] | r_{p} [nm] |
|---------------------------|-----------------|------------------|----------|
| 9                         | −1.06 ± 0.08    | −1.67 ± 0.08     | 13.5 ± 0.7 |
| 29                        | −1.03 ± 0.08    | −1.64 ± 0.08     | 13.7 ± 0.6 |

\( a) T_{pk} \): Peak temperature for melting of water confined in pores; \( b) \Delta T_{on-pk} \): Temperature depression.

Figure 1. BJH N2 gas desorption pore-size distribution of nanocellulose filter papers.

Figure 2. Particle-size distribution from dynamic light scattering analysis of SWW. The distribution has a peak value of \( d_{\text{particle}} = 0.87 \) μm and no particles were found at \( d_{\text{particle}} \leq 0.59 \) μm and \( d_{\text{particle}} \geq 31.1 \) μm.
easy. This is because during drying nanocellulose irreversibly agglomerates into a compact, impermeable film due to a process known as “cellulose hornification.”[38] Alternatively, solvent exchange, critical point drying, or freeze-drying can be used to produce cellulose aerogels but these are expensive processes, too, for industrial-scale production.[38]

The advances in nanocellulose science allow today for cost-efficient production of this material.[39] Recently, a highly cost-efficient virus removal filter paper, aka mille-feuille filter paper, was described for applications in biotechnology.[40,41] It is produced by traditional paper-making processing, which contrasts starkly from the membrane technology relying on phase inversion. This nonwoven filter paper, made from 100% cellulose nanofibers, features a stratified internal architecture, consisting of numerous cellulose nanofiber sheets—hence, its name is “mille-feuille” (or thousand leaves) filter paper. The separation efficiency of the mille-feuille filter was verified for surrogate nanoparticles, e.g., gold nanoparticles or fluorophore-labeled latex nanobeads,[41–43] and real viruses, e.g., influenza virus, i.e., swine influenza A virus (100 nm);[44] retrovirus, i.e., murine leukemia virus (100 nm);[44] and parvovirus, i.e., minute virus of mice (20 nm).[40] In this article, the properties of the mille-feuille filter paper are evaluated for water treatment applications for the first time. Furthermore, this work discusses the effect of process parameters such as filtration pressure and virus load on clearance efficiency.

2. Results and Discussion

2.1. Pore-Size Characterization of Nanocellulose Filter Papers

The pore-size distribution of the nanocellulose filter papers was evaluated both in the dry state, using nitrogen gas sorption, and in the wet state, using differential scanning calorimetry (DSC) cryoporometry (CP-DSC). The resulting pore-size distributions from nitrogen sorption measurements of the nanocellulose filter papers of thicknesses 9 and 29 µm are presented in Figure 1. A shift in the pore-size distribution toward pores of a greater width was observed for the thinner filter papers, consistent with previous observations.[40,42]

The results from CP-DSC of the nanocellulose filter papers of thicknesses 9 and 29 µm are summed up in Table 1, where peak temperature for melting of water confined in pores, temperature depression, and calculated peak pore mode radius for the nanocellulose filter papers is presented. No statistically significant difference in the peak pore mode radius, \( r_p \), between the two filter thicknesses was found from the CP-DSC measurements. Representative heat flow curves from CP-DSC measurements of the nanocellulose filter papers can be found in Figure S1 in the Supporting Information.

2.2. Simulated Waste Water Filtration

The nanocellulose filter papers were challenged with simulated waste water (SWW) dispersions containing two different amounts of total suspended solids (TSS), i.e., 0.251 and 2.51 mg L\(^{-1}\). The source of the TSS in the SWW was diatomite and formazin was the turbidity contributing additive. Dynamic light scattering (DLS) analysis of the particle diameter, \( d_{\text{particle}} \), in the SWW revealed a particle size distribution with a peak value of 0.87 µm for the suspended solids, as presented in Figure 2. No particles at cutoff values \( d_{\text{particle}} \leq 0.59 \) µm and \( d_{\text{particle}} \geq 31.1 \) µm were found.

Compared to the feed dispersion, a visible decrease in the turbidity of the permeate was noticed in the filtrations of SWW for both TSS contents. In Figure 3 representative pictures of feed and permeate are presented.

![Figure 3](image-url) Feed (left) and permeate (right) in filtrations of SWW through nanocellulose filter papers. The total solid content of the feed dispersion was 0.251 mg L\(^{-1}\).

![Figure 4](image-url) a) Postfiltration image of a nanocellulose filter paper challenged with SWW with a TSS content of 2.51 mg L\(^{-1}\). b) SEM image of a nanocellulose filter paper postfiltration of SWW. Deposited solids from the SWW are visible on top of the filter structure. The image was taken at 21 × 10\(^3\) times magnification.
As seen in Figure 4a, a clear accumulation of solids on the filter surface was observed after filtration of SWW. From the scanning electron microscopy (SEM) image presented in Figure 4b, the solid deposit on the filter surface does not appear to be clogging the porous structure of the filter, and exposed areas of the filter are visible through the deposited solids. The measured transmittance at 500 nm of feed and collected permeates is presented in Tables 2 and 3, where $T_{\text{permeate}}$ and $T_{\text{feed}}$ are the measured transmittances for permeate and feed, corrected for water background.

The transmittance measurements suggest that all solids were removed from the SWW when filtered through the nanocellulose filter paper, in accordance with what was observed in Figure 4a and in SEM postfiltration imaging in Figure 4b. Considering the particle distribution in the SWW, where no particles were found at $d_{\text{particle}} \leq 0.59 \, \mu m$, no particles should be small enough to pass the nanometer-sized pores of the nanocellulose filter paper.

The observed fluxes during filtrations of SWW through the nanocellulose filter papers are presented in Figures 5 and 6. The manufacturer recommends the TSS content of 0.251 mg L$^{-1}$ for validation studies. No or little decline in flux was observed during filtration at the reported pressures and amounts of TSS. The flux seemingly remained unaffected by the amount of TSS in the feed dispersion, i.e., 0.251 and 2.51 mg L$^{-1}$. The average flux values for 9 µm filter papers at 1 and 3 bar were 200 and 500 L m$^{-2}$ h$^{-1}$, respectively. The average flux values for 29 µm filter papers at 1 and 3 bar were 65 and 150 L m$^{-2}$ h$^{-1}$, respectively.

The fouling behavior during filtration of the SWW was further analyzed and quantified through the so-called $V_{\text{max}}$ analysis.$^{[45]}$ $V_{\text{max}}$ was calculated from the slope of the linear fit associated with each flux data curve and is available in Figures S2 and S3 as well as Tables S1 and S2 in the Supporting Information.

As seen from both the flux data presented in Figures 5 and 6 as well as the result of the $V_{\text{max}}$ analysis, the flux remained stable during filtration of SWW through the nanocellulose filter paper. This would indicate that no significant clogging of the filter occurs when challenged with the SWW. When a value for $V_{\text{max}}$ was possible to achieve, i.e., a positive value for $1/V_{\text{max}}$, the $V_{\text{max}}$ analysis suggests that volumes in the order of $10^4$ L m$^{-2}$ can be filtered before complete clogging of the nanocellulose filter paper occurs.

### 2.3. Filtration of Latex Nanoparticles in SWW

The nanocellulose filter papers were further challenged with SWW dispersions containing latex particles with a diameter of 30 nm. The TSS content in the dispersions was 0.251 mg L$^{-1}$ and the concentration of 30 nm latex particles was in the order of $10^{13}$ mL$^{-1}$. The resulting removal efficiency of the 30 nm latex particles is presented in Figure 7 for pressures 1 and 3 bar, and for filter thicknesses 9 and 29 µm. The limit of detection (LOD) in fluorescence spectroscopy of the 30 nm latex particles corresponds to a logarithmic reduction value (LRV) of 2.0, and no latex particles where detected in permeates at the reported pressures for neither of the two filter thicknesses.

The observed fluxes during filtrations of 30 nm latex particles in SWW are presented in Figure 8. In contrast to the filtrations performed with only SWW, a decline in the flux can

### Table 2. Measured transmittance for SWW filtrations through nanocellulose filter papers. TSS content was 0.251 mg L$^{-1}$. $T = 100\%$ corresponds to the transmittance of pure water.

| Filter paper thickness [µm] | Overhead pressure [bar] | $T_{\text{feed}}$ 500 nm [%] | $T_{\text{permeate}}$ 100 nm [%] |
|----------------------------|-------------------------|-------------------------------|-------------------------------|
| 9                          | 1                       | 94.7 ± 0.4                    | 100 ± 0                        |
| 9                          | 3                       | 94.9 ± 0.3                    | 100 ± 0                        |
| 29                         | 1                       | 94.2 ± 0.6                    | 100 ± 0                        |
| 29                         | 3                       | 94.5 ± 0.1                    | 100 ± 0                        |

### Table 3. Measured transmittance for SWW filtrations through nanocellulose filter papers. TSS content was 2.51 mg L$^{-1}$.

| Filter paper thickness [µm] | Overhead pressure [bar] | $T_{\text{feed}}$ 500 nm [%] | $T_{\text{permeate}}$ 500 nm [%] |
|----------------------------|-------------------------|-------------------------------|-------------------------------|
| 9                          | 1                       | 46.9 ± 0.6                    | 100 ± 0.0                     |
| 9                          | 3                       | 46.9 ± 0.6                    | 100 ± 0.0                     |
| 29                         | 1                       | 45.7 ± 1.7                    | 100 ± 0.1                     |
| 29                         | 3                       | 45.7 ± 1.7                    | 100 ± 0.1                     |

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**Figure 5.** Observed fluxes at overhead pressures 1 and 3 bar for filtrations of SWW through nanocellulose filter papers of thicknesses a) 9 µm and b) 29 µm. TSS content in SWW was 0.251 mg L$^{-1}$. Gray area indicates standard deviation of the measured flux for three separate filtrations.
Figure 6. Observed fluxes at overhead pressures 1 and 3 bar for filtrations of SWW through nanocellulose filter papers of thicknesses a) 9 µm and b) 29 µm. TSS content in SWW was 2.51 mg L$^{-1}$. Gray area indicates standard deviation of the measured flux for three separate filtrations.

Figure 7. Logarithm reduction values (LRV) for filtration of 30 nm latex particles in SWW through nanocellulose filter papers of thicknesses a) 9 µm and b) 29 µm. Arrows indicate that the obtained values are below the limit of detection (LOD) in fluorescence spectroscopy of the 30 nm latex particles. The results are the average of two measurements.

Figure 8. Observed fluxes at overhead pressures 1 and 3 bar for filtrations of 30 nm latex particles in SWW through nanocellulose filter papers of thicknesses a) 9 µm and b) 29 µm. Gray area indicates standard deviation of the measured flux for two separate filtrations.
Another interesting result from the be seen for the thinner filters, as compared to the thicker filters. A tendency of fouling behavior occurring at a faster rate can be observed for the thinner filter papers, and the concentration of ΦX174 bacteriophages was in the order of 10^6 mL^-1. The resulting LRVs for the filtrations are presented in Figure 9, and the virus removal efficiency is summed up in Table 5.

As seen from the results presented in Figure 9 and Table 5, thinner filter papers exhibit lower virus removal efficiency compared to the thicker filters. In general, better removal efficiency for both filter thicknesses is achieved at the higher overhead pressure, i.e., 3 bar. It is further seen from Figure 9 that the efficiency of removal is progressively decreasing with increased load, especially for 9 μm filter papers. Nonetheless, even for the thinner filter at 1 bar, the removal efficiency never falls below 99.2% for the largest load volumes.

Under the experimental conditions presented in this work, two cases should be scrutinized in more detail, i.e., 9 μm filter paper at 1 bar and 29 μm filter paper at 3 bar. Both filters show comparable pore-size modes and flow rates at respective overhead pressure, i.e., 200 L m^-2 h^-1 for 9 μm filter paper at 1 bar and 150 L m^-2 h^-1 for 29 μm filter paper at 3 bar. However, the performance of these two filter papers with respect to virus removal efficiency and fouling is very different. In particular, the 29 μm filter paper at 3 bar exhibits steady LRV ≥ 5 and V_{max} = (1.8 ± 0.1) × 10^3 L m^-2, measured with spiked 30 nm latex nanoparticles in SWW, whereas for 9 μm filter paper at 1 bar the observed virus LRV gradually decreases from 3.5 to 2.2 and V_{max} is around (9.7 ± 4.6) × 10^2 L m^-2, under similar experimental conditions. The latter difference could potentially be due to compaction of the 29 μm filter paper at 3 bar as discussed earlier. It should be noted that the 9 μm filter exhibits higher variability of V_{max} values, which is likely to be due to larger spread of initial flux values and extrapolation.

The presented data using SWW show that nanocellulose filter paper has the capacity to efficiently remove even the smallest viruses at industrially relevant flow rates and low fouling. Thus, it is expected that the filter can act as a total microorganism removal filter, removing not only viruses but also all other larger microbes. Compared to other available advanced water treatment membranes, the mille-feuille filter is

### Table 4. Results from V_{max} analysis for filtration of 30 nm latex particles in SWW through nanocellulose filter papers.

| Filter paper thickness [μm] | Filtration overhead pressure [bar] | Slope of linear fit (1/V_{max}) | V_{max} [L m^-2] |
|-----------------------------|----------------------------------|--------------------------------|-----------------|
| 9                           | 1                                | 1.3 (±0.6) × 10^{-4}            | 9.7 (±4.6) × 10^2 |
| 9                           | 3                                | 2.5 (±0.1) × 10^{-4}            | 4.0 (±0.2) × 10^2 |
| 29                          | 1                                | 1.4 × 10^{-4}                  | 7.4 × 10^1      |
| 29                          | 3                                | 5.7 (±0.3) × 10^{-4}            | 1.8 (±0.1) × 10^1 |

*a) Values for V_{max} were obtained at a particle concentration of 10^{11} mL^-1. b) The second filtration exhibited a negative slope in the V_{max} analysis.

2.4. Filtration of ΦX174 Bacteriophage in SWW

The nanocellulose filter papers were also challenged with SWW dispersions spiked with the 28 nm large ΦX174 bacteriophage, used as a surrogate model for worst-case small-sized mammalian viruses. TSS content of the dispersions was 0.251 mg L^-1.

![Figure 9](https://example.com/figure9.png)  
**Figure 9.** Logarithm reduction values (LRV) for filtration of ΦX174 bacteriophages in SWW through nanocellulose filter papers of thicknesses a) 9 μm and b) 29 μm at overhead pressures 1 and 3 bar. Arrows indicate that obtained values are below the limit of detection (LOD) in the plaque-forming unit (PFU) assay, and no phage particles were detected in large volume plating. The results are the average of three measurements.
attractive for point-of-use applications as an affordable, single-use filter paper, similar to coffee filters used in household for brewing. Because traditional water treatment membranes are used over an extended period of time, i.e., 5–15 years after installation, they need to be regularly backflushed and treated with chemical disinfectants (hypochlorite) in order to avoid biofilm formation on the surface of the membrane. The biofilm formation is a major issue for the membranes, as it is deteriorating the key processing parameters over time, including flux and permeate quality. In contrast, an affordable single-use filter paper with excessive microorganism removal capacity and high flux eliminates the risk of biofilm formation and thereby the need for regular cleaning and disinfection. Therefore, we suggest that the mille-feuille filter paper is an interesting point-of-use, sustainable alternative to address the global challenges related to drinking water purification.

### 3. Conclusion

In this work, the filtration of SWW matrix spiked with surrogate latex nanobeads and ΦX174 bacteriophages as model for worst-case small-sized mammalian virus is shown using the nanocellulose-based filter paper for drinking water purification applications. The presented data using SWW matrix show for the first time that a filter paper made from 100% nanocellulose is capable of efficiently removing even the smallest viruses, i.e., up to 99.9980–99.9995% efficiency, at industrially relevant flow rates, i.e., 60–500 L m⁻² h⁻¹, and low fouling, i.e., $V_{\text{max}} > 10^{4}–10^{6}$ L m⁻². The filter paper presented here shows great promise for development of robust, affordable, and sustainable water purification systems.

### 4. Experimental Section

**Materials:** Cladophora algae cellulose was provided by FMC BioPolymer (batch G3828-112). SWW matrix was obtained from Sigma (MMW001-250ML, lot number LRAB5412). Fluorescent latex beads (30 nm; carboxylate-modified polystyrene; L5155), calcium chloride dihydrate (C5080), magnesium chloride hexahydrate (M2670), and sodium chloride (S5886) were purchased from Sigma-Aldrich.

**Biological Materials:** Escherichia coli (Migula) Castellani and Chalmers (E. coli, 13 706) bacteria strain and ΦX174 bacteriophage (13 706-B1) were purchased from the American Type Culture Collection (ATCC). Agar (214 530) and yeast extract (212 750) were purchased from BD. Tryptone broth powder (T-broth) (J870) was purchased from Amresco.

**Preparation of Nanocellulose Filter Paper:** A dispersion of Cladophora cellulose (0.1 wt%) was prepared. The dispersion was then run twice in succession through 200 and 100 µm sized chambers at 1800 bar, using an LM20 Microfluidizer. Filter papers of two different thicknesses, i.e., 9 and 29 µm, were then prepared by adjusting the solids content of dispersion. The resulting nanocellulose dispersion was then drained over a nylon filter membrane (Durapore, 0.65 µm DVPP, Merck Millipore) fitted in a funnel using vacuum. The resulting wet cellulose mass was then dried at 80 °C using a hot press (Carver, USA).

**Nitrogen Gas Sorption:** Pore-size distribution evaluation of the nanocellulose filter papers with the Barret–Joyner–Halenda (BJH) method was performed based on the desorption branch of the nitrogen sorption isotherm. This was done using an ASAP 2020 (Micrometrics, USA) instrument. The filter sample was degassed at 90 °C in vacuum for 4 h and then analyzed by nitrogen sorption, carried out at 77 K. The performance of the instrument was validated using Micrometrics Silica-Alumina SSA 210 m² g⁻¹ (lot number: A-501-49) standard prior to analysis. The deviation between the pore-size mode of the calibration data from the nominal standard values was 0 nm.

**Cryoporometry DSC:** Pore-size distribution evaluation of the nanocellulose filters was performed using cryoporometry DSC, using a Mettler Toledo DSC 3 (Switzerland). The filter samples were soaked in water for 1 h prior to DSC analysis to ensure substantial wetting of the filter structure. DSC analysis was performed by cooling the samples to -25 °C at a rate of 15 K min⁻¹ followed by heating of the samples to 4 °C at a rate of 0.7 K min⁻¹. Five separate measurements were performed for each filter thickness. The peak value for the melting of bulk water was determined from repeated measurements on deionized water. Deionized water (1, 2, 4, 6, and 10 µL) was analyzed in five separate measurements per volume, and the mean peak value for the water was determined to be 0.61 °C.

Cryoporometry is a method for determining the pore size in porous materials from the liquid-solid transformation of a probing medium. With the use of DSC, the pore radius can be related to the temperature where freezing or melting of a probing liquid in the pores occur. If melting of the probing liquid is considered, transition from solid to liquid phase confined in pores will occur at a temperature $T_m$ below the melting temperature $T_m$ of the bulk liquid. This temperature depression, $\Delta T$, is related to the pore radius $r_p$ as shown in Equation (1).

### Table 5. Virus removal efficiencies for filtration of ΦX174 bacteriophages in SWW through nanocellulose filter papers.

| Filter paper thickness [µm] | Filtration overhead pressure [bar] | Load, =3 L m⁻² [%] | Load, =58 L m⁻² [%] | Load, =93 L m⁻² [%] | Load, =133 L m⁻² [%] | Load, =148 L m⁻² [%] |
|-----------------------------|-----------------------------------|---------------------|---------------------|---------------------|---------------------|---------------------|
| 9                           | 1                                 | 99.9700             | 99.5000             | 99.6800             | 99.3700             | 99.2100             |
|                             | 9                                 | 99.800              | 99.8400             | 99.7500             | 99.6000             | 99.5000             |
|                             | 9                                 | 99.900              | 99.5000             | 99.3700             | 99.3700             | 99.3700             |
| 9                           | 3                                 | 99.990              | 99.9000             | 99.8700             | 99.7500             | 99.6800             |
|                             | 9                                 | 99.990              | 99.9800             | 99.9500             | 99.9000             | 99.9000             |
|                             | 9                                 | 99.990              | 99.9400             | 99.9400             | 99.8700             | 99.8400             |
| 29                          | 1                                 | 99.9940             | 99.9900             | 99.9700             | 99.9800             | 99.9200             |
|                             | 29                                | 99.9980             | 99.9940             | 99.9800             | 99.9800             | 99.9700             |
|                             | 29                                | 99.9960             | 99.9800             | 99.9500             | 99.9700             | 99.9600             |
|                             | 29                                | 99.9995             | 99.9995             | 99.9995             | 99.9995             | 99.9995             |
|                             | 29                                | 99.9994             | 99.9994             | 99.9994             | 99.9994             | 99.9994             |
|                             | 29                                | 99.9992             | 99.9992             | 99.9992             | 99.9992             | 99.9880             |
\[ \Delta T = \frac{2T_0 \gamma \cos \theta}{\rho \Delta H_f} - T_p \]  

\[ r_p (\text{nm}) = -\frac{19.082}{\Delta T + 0.1207} + 1.12 \]

The peak pore mode radius was determined from the DSC, using Equation (2). The difference between the peak maximum for melting of pore confined solid phase and the peak value for melting of bulk water (0.38 °C), \( \Delta T_{pk} \), was used as the temperature depression.

**Dynamic Light Scattering:** Particle-size distribution of the suspended solids in the SWW was analyzed through DLS using a Malvern Mastersizer 3000. Refractive index of the dispersant was set to that of solids in the SWW-simulated matrix with deionized water to a final volume (200 mL). Final TSS contents were achieved (0.251 and 2.51 mg L\(^{-1}\)) of SWW-simulated matrix with deionized water to a final volume of 200 mL. Filtrations were carried out at overhead pressures (1 and 3 bar). The permeate solution was collected in fractions (40 mL). Concentrates were inoculated into Luria–Bertani medium (1% tryptone broth, 0.5% yeast extract, 1% NaCl, 1 \times 10^{-1} M CaCl\(_2\)-2H\(_2\)O, and 1 \times 10^{-1} M MgCl\(_2\)-6H\(_2\)O in deionized water) for 5 h at 35 °C with an agitation (120 rpm). Bacteriophage was harvested by centrifugation at 5000 \( \times \) g for 10 min, and the suspension was collected and stored at 4 °C prior to use.

The X174 bacteriophage titer was determined by plaque-forming units (PFU) assay and expressed as PFU mL\(^{-1}\). Briefly, to perform an assay, tenfold dilutions of a bacteriophage stock were prepared and inoculated into E. coli host on the agar plates. Each dilution was plated in duplicate to enhance the accuracy, and PFU mL\(^{-1}\) was calculated using Equation (4).

\[ \log_{10} \left( \frac{\text{PFU}}{\text{ml}} \right) = \log_{10} \left( \frac{\text{number of plaques}}{10^{\text{dilution factor}}} \right) \]

where 0.1 is the volume (mL) of the added virus.

To supplant the air from the inner pores of the nanocellulose-based filter, prefiltration with deionized water was performed. The filter was fixed in an Advantec KST-47 filter holder, 30 mL of deionized water was added, and the filter was driven by the overhead air pressure of 1 bar. Prefiltration was stopped after 20 mL of the water was filtered to avoid the ingress of the air into the filter.

SWW was spiked with X174 bacteriophage stock dispersion to obtain the titer of \( \approx 10^6 \) PFU mL\(^{-1}\). Feed solution (200 mL) was filtered through the nanocellulose-based filter at the overhead air pressure of 3 bar in a dead-end setup. Permeate samples were collected (40 mL fractions) and stored at 4 °C before PFU assay. Bacteriophage removal capacity was expressed by LRV

\[ \text{LRV} = \log_{10} \left( \frac{\text{PFU}}{\text{ml}} \right)_{\text{feed}} - \log_{10} \left( \frac{\text{PFU}}{\text{ml}} \right)_{\text{permeate}} \]

The limit of detection (≤0.7 PFU mL\(^{-1}\)) of PFU assay current experimental design refers to ≤55 bacteriophages mL\(^{-1}\), corresponding to a single detectable plaque in one of the plates for nondiluted duplicate samples. To detect any possibly omitted bacteriophage in the permeate fractions, the filtrates were concentrated by centrifuging at 4500 \( \times \) g in 100 kDa cutoff protein concentrator tubes (Thermo, 88 533) to the final dead stop volume (≤50 \( \mu \)L). Concentrates were inoculated into E. coli host culture at exponential growth phase in Luria–Bertani medium for 5 h at 35 °C with an agitation (120 rpm). Bacteriophage presence was determined by decreased OD\(_{600}\) values compared to control samples.

**Supporting Information**

Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest

The corresponding author is the inventor behind IP related to the virus removal filter paper.

Keywords

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