

Household water purification system comprising cartridge filtration, UVC disinfection and chlorination to treat turbid raw water

Paulo Marcos Faria Maciel^a, Natália de Melo Nasser Fava^a, Antônio Wagner Lamon^a, Pilar Fernandez-Ibañez^b, John Anthony Byrne^b, Lyda Patricia Sabogal-Paz^{a*}

^aDepartment of Hydraulics and Sanitation, São Carlos School of Engineering, University of São Paulo, Avenida Trabalhador São-Carlense 400, São Carlos, São Paulo, 13566-590, Brazil.

^bNanotechnology and Integrated Bioengineering Centre, School of Engineering, Ulster University, Jordanstown, BT37 0QB, Northern Ireland, United Kingdom.

* Corresponding author: lysaboga@sc.usp.br

Abstract

A system composed of cartridge membrane filters (25 µm, 10 µm and 1 µm opening size), UVC-lamp and chlorinator was constructed and tested with high turbidity water (up to 236 NTU). The proposed system treated around 180 L day⁻¹ of river water and the following parameters were analysed: pressure drop in filters, turbidity, apparent colour, true colour, *Escherichia coli*, total coliforms (TC), UV 254_{nm} transmittance, UV 254_{nm} absorbance and total organic carbon in both raw and treated water. The study was conducted in three phases. In Phase 1, the raw water was rested for 24 hours in a settling tank before filtration in a 10 µm cartridge filter followed by one cartridge of 1

27 μm , a UVC-lamp (17 W) and manual chlorination. In Phase 2, a 25 μm filter was added
28 before the 10 μm filter. In Phase 3, a blanket filtration step was added before the raw
29 water entered the rest tank. The filtration trials lasted 7, 21 and 69 days in Phases 1, 2
30 and 3, respectively. The turbidity reduction of the system ranged from 30% to 93%.
31 Disinfection by UVC-lamp was able to inactivate *E. coli* up to 5.00log, however the TC
32 reduction was lower (up to 4.24log). The final manual chlorination with a dose of 3 mg
33 L^{-1} of sodium hypochlorite increased the reduction of TC (up to 5.94log), regardless of
34 water turbidity. The system was effective in improving water quality aimed at
35 implantation in rural communities for domestic use at household level.

36

37 **Keywords:** cartridge membrane filters, decentralized, drinking water, low-cost
38 technologies, turbid raw water, rural communities.

39

40 **Highlights**

41

- 42 • Pre-filtration in non-woven synthetic fabric prolonged cartridge filters' lifespan.
- 43 • Surface on cartridge filter treating natural water formed microbiological community.
- 44 • UVC disinfection after cartridge filtration removed *E. coli* to non-detection level.
- 45 • Chlorination assured total coliform reduction to virtual absence.

46

47 **1. Introduction**

48

49 Worldwide, 785 million people still have no access to basic water services, consuming
50 water from unprotected sources or depending on travel times to collect potable water [1]. It is
51 estimated that globally 1.8 billion people use drinking water sources that have some type of

52 faecal contamination [2,3]. Therefore, inadequate access to safe water contributes to nearly
53 1.7 billion episodes of diarrhoea per year [4]. As well as drinking and food preparation, it is
54 important to have the availability of water for hygiene purposes such as handwashing, which
55 is crucial against diarrheal episodes [5] and it is imperative to protect human health during all
56 infectious disease outbreaks, including the COVID-19 outbreak [6]. In 2017, 18% of the
57 global population still had no handwashing facilities at home [1].

58 While water on premises is not a reality for many populations, water treatment at a
59 household level should be considered important as an interim and immediate solution [7].
60 Some interventions have provided point-of-use water treatment solutions based on ceramic
61 pot filtration, household slow sand filtration, solar disinfection, among others [8–10].
62 However, many of these technologies are designed to produce limited amounts of water per
63 day. Other solutions are developed to attend a small group of households [11] and are not
64 suitable to serve an isolated single-family location.

65 There is a lack of information in the scientific literature on water treatment systems for
66 serving family groups of up to 5 members based on materials that are easily available on the
67 market. Technologies such as cartridge filtration and UV disinfection are solid, and their
68 applications are widespread around the world. Notwithstanding, there is a lack of information
69 on evaluating the efficiency of commercial products based on these technologies.

70 A previous study investigated commercial cartridge filters treating a daily volume of
71 250 L, considering turbidity removal and a pressure drop of the filter [12]. The evaluated
72 system required less installation costs and shorter operation times when compared to other
73 filtration systems, such as ceramic filtration or rapid sand filtration. However, there is still a
74 gap in the literature regarding cartridge filtration technology considering turbid natural water
75 treatment with bacterial contamination. It is generally mentioned that the filters clog quickly

76 if the source water is cloudy [13], but a combination of filters with different porous and
77 options of pre-treatment are still to be investigated.

78 Cartridge filtration systems are the most common worldwide point-of-use water
79 treatment devices that use various types of filters in designated housing to produce potable
80 water [14]. They are often needed for prefiltering before other filters or disinfection since they
81 can reduce turbidity, are inexpensive, require small spaces and have a total reduced weight
82 when compared to sand or other media filtration [13,15].

83 The ease of operation and maintenance of cartridge filters makes them very attractive
84 for small water systems [16]. Particles can be retained on the membrane surface and form a
85 cake layer that progressively grows as the filtration cycle continues and it acts as another
86 physical barrier or sieve to pathogenic organisms [17,18].

87 UV-lamps are easy to install, they need little operator attention, no on-site storage or
88 use of potentially harmful chemicals, they are cost-effective and they have a high efficiency
89 for inactivation of various microorganisms [19,20]. There has been renewed interest in UV
90 irradiation with lamps in recent years because of its well-documented ability to extensively
91 inactivate two waterborne, chlorine-resistant protozoans, *Cryptosporidium parvum* oocysts
92 and *Giardia lamblia* cysts, at relatively low irradiation doses [21].

93 However, surface water can be cloudy or contain sediment that will limit UV light
94 penetration, and for that reason, the UV disinfection system is typically combined with other
95 treatment devices such as cartridge filters [22]. Although UV disinfection is effective and
96 promising in places with small distribution networks, it does not provide residual protection;
97 therefore, the water stored after its treatment may be subject to new contamination [20,23].
98 Moreover, some microorganisms can be reactivated after inactivation by UV [24]. In order to
99 achieve extended protection, chlorination can be performed to assure the residual protection
100 of the water post UV-treatment.

101 Chlorine is an attractive solution because of its availability, ease of use and
102 maintenance of residual in storage treated water. However, its effectiveness depends on the
103 water pH, concentration and contact time, temperature and chlorine demand of the water.
104 Moreover, natural water containing an elevated concentration of dissolved organic carbon and
105 bromide may represent a potential of carcinogenic disinfection byproduct (DBP) formation
106 [25]. Nevertheless, the correct chlorine dose and proper water pre-treatment reduce the
107 occurrence of DBP precursors [26].

108 Considering this context, a water treatment prototype (WTPt) was constructed
109 associating solid technologies that are applied in the context of isolated communities. The
110 WTPt was designed to treat 180 L of raw river water per batch, considering one batch per day,
111 with stressing conditions caused by elevated water turbidity. Sedimentation during 24 h
112 followed by cartridge filtration was included for sediment removal, UVC-lamp for
113 disinfection and chlorine for residual protection and safe storage. The purpose was to analyse
114 the efficiency of the WTPt in terms of bacteria reduction (*Escherichia coli* and total coliforms
115 – TC) and physical and chemical parameters (turbidity, apparent and true colour, absorbance
116 at 254_{nm} and total organic carbon). In addition, microscopic analyses were performed to
117 observe which microorganisms were retained in the cartridge filters. This study also aimed to
118 improve the sediment removal step including an additional cartridge filter and introducing a
119 fabric filtration with a non-woven blanket.

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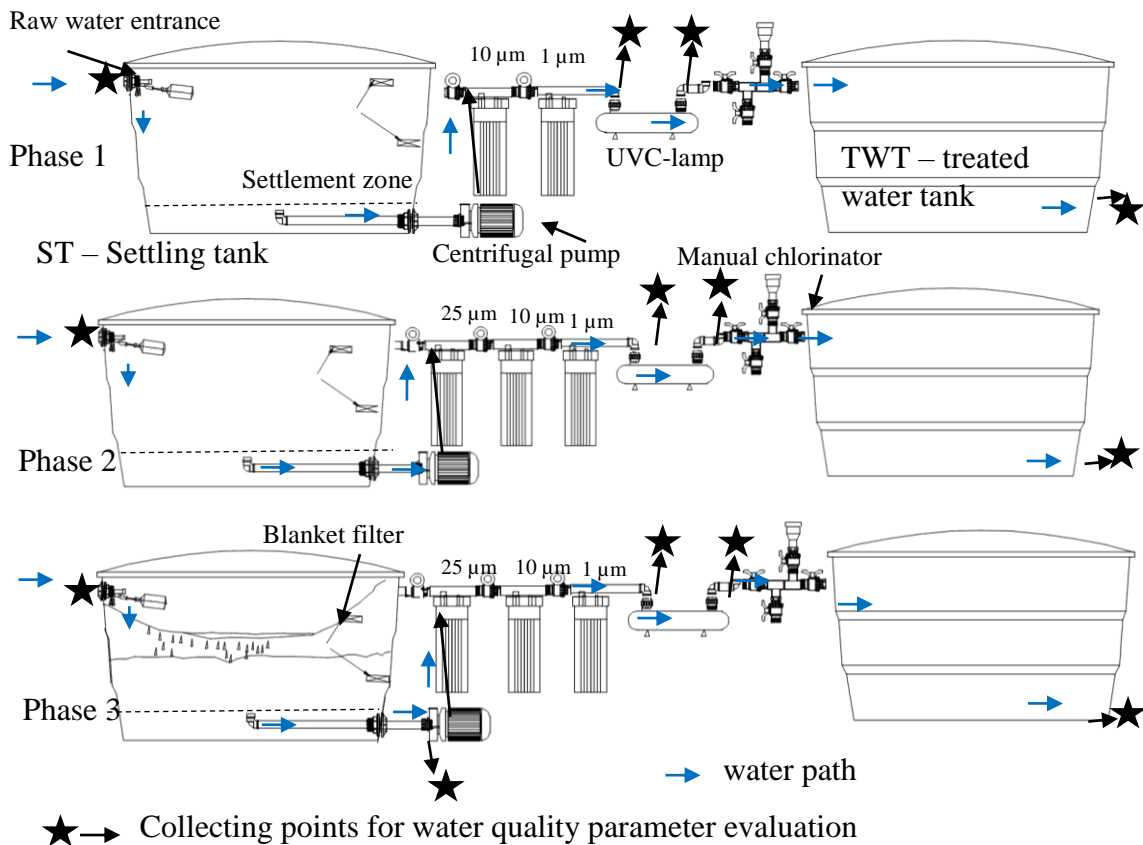
121 **2. Material and Methods**

122

123 **2.1. Prototype characteristics and operation**

124 The constructed prototype (Figure 1 and Figure S1 – Supplemental Material)
125 comprised two 310 L water tanks; one centrifugal pump (1/2 hp 40 L min⁻¹, 40 m maximum

126 height, *Amanco*); pressure gauges; a sequence of filter housings containing cartridge
 127 polypropylene pleated filters of 25 μm , 10 μm and 1 μm opening size; a 17 W UVC-lamp
 128 (*Polaris™ UV-4C, Polaris Scientific, USA*); and a manual chlorinator. The filter housing
 129 measured 30 cm (height) x 12 cm (diameter). The UV dose commercial information of the
 130 UVC-lamp is presented in Figure S2 (Supplemental material).



131
 132 Figure 1 – Configuration of the Water Treatment Prototype in the three phases of the study
 133 and water collecting points indicated

134 The components of the prototype were assembled in a metal structure. The pump
 135 pumped water from the Settling Tank - ST to the elevated Treated Water Tank - TWT,
 136 passing through the filtration units and the disinfection units. The system inlet piping was
 137 positioned 15 cm above the base of the ST to provide a sedimentation zone for 24 hours
 138 between the raw water entrance and the pump start (Figure 1). One semi-automatic tap float
 139 was installed in each water tank to control the minimum and maximum level of water and the

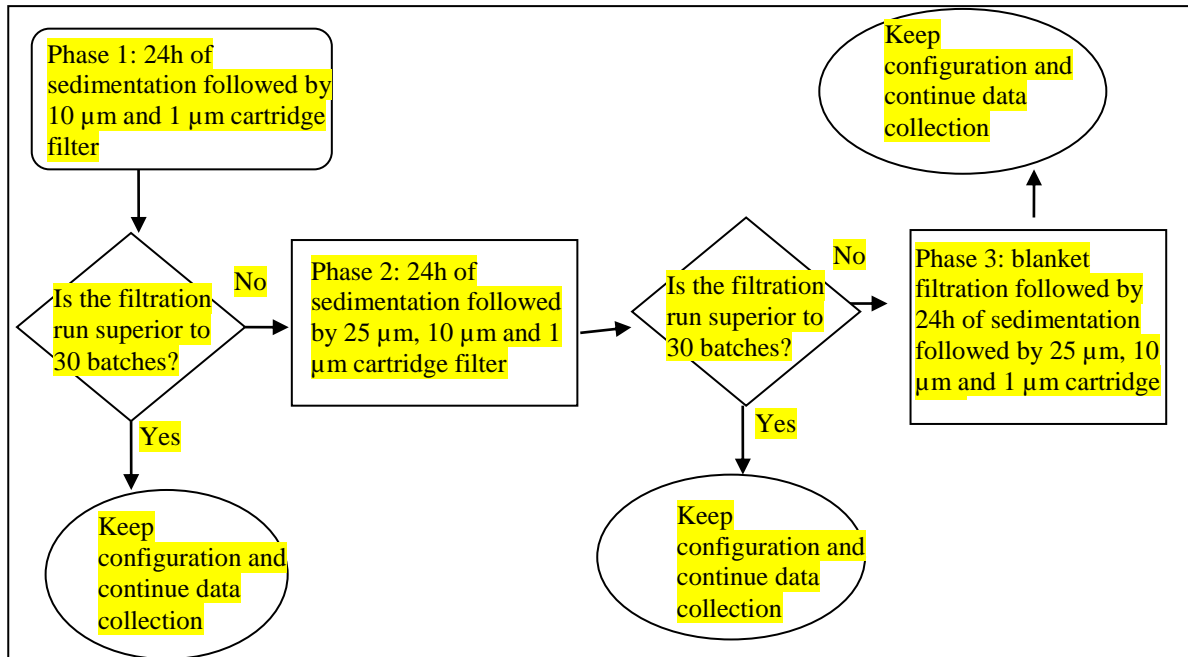
140 pump operation. The pump operation was related to one batch, which corresponded to the
 141 treatment of 180 L of raw water and the filtration rate during the batches ranged from 825 to
 142 180 L m⁻² h⁻¹, according to the filter clogging. One batch was performed per day of operation.
 143 The pump operation pressure was kept between 0.6 bar, at the first operation batch and 3.0
 144 bar.

145 The study was conducted in three phases, with differences in the sediment removal
 146 step. Table 1 summarizes the phases of the study and Figure 2 shows a decision flowchart
 147 regarding the experimental procedure. In Phase 1, a sequence of two pleated filters was
 148 adopted (10 µm and 1 µm opening porous size) after the sedimentation time. In Phase 2, a 25
 149 µm opening porous size pleated filter was added before the 10 µm filter. In Phase 3, two
 150 layers of a non-woven blanket were installed (specific gravity: ±0.2 g cm³, composition:
 151 100% polyester and thickness = 2.8 mm with 25 µm fibres) on the top of the ST to filter the
 152 raw water before the sedimentation time and a sequence of filters was used in Phase 2. The
 153 configuration of each phase is illustrated in Figure 1. The disinfection step performed by the
 154 UVC-lamp and manual chlorinator was used in the three phases.

155 Table 1– Experimental phases of the study

Phase	sediment removal step	disinfection step	water collection points
1	Sedimentation (24h) + cartridge filtration 10 µ and 1 µm	UV disinfection and chlorination	raw water entrance, after cartridge filters, after UVC-lamp, treated water tank
2	Sedimentation (24h) + cartridge filtration 25 µm , 10 µm and 1 µm	UV disinfection and chlorination	raw water entrance, after cartridge filters, after UVC-lamp, treated water tank
3	Blanket filtration + Sedimentation (24h) + cartridge filtration 25 µm, 10 µm and 1 µm	UV disinfection and chlorination	raw water entrance, before cartridge filters , after cartridge filters, after UVC-lamp, treated water tank

156 **Bold:** Modification from the previous phase.



157

158 Figure 2 – Decision Flowchart of the experimental procedure

159 A semi-automatic control system was designed for the prototype operation. A block
 160 diagram of the design of the electrical control of the system is shown in Figure S3 and a
 161 model of the operation protocol is represented in Figure S4, both in the supplemental material
 162 of this manuscript.

163

164 **2.2. Source water**

165 The raw water was daily collected from the Monjolinho River (21.9869S; 47.8760W)
 166 in the city of São Carlos, state of São Paulo, Brazil, from August 2019 to February 2020. The
 167 water was moved directly to the ST, without storage time interval. This water is used as a
 168 source for the water treatment plant in the city of São Carlos. Characteristics of the raw water
 169 are presented in Table 2.

170 Table 2: Raw water characteristics

Characteristic	Mean ± standard deviation (median) [range]
Turbidity (NTU)	28.83 ± 29.83 (17.30) [7.34 – 236]

Apparent colour (Hu)	86.72 ± 61.74 (57.90) [25.9 – 316]
True colour (Hu)	38.60 ± 20.07 (30.70) [18.9 – 98.2]
UV ₂₅₄ Absorbance	0.131 ± 0.057 (0.109) [0.057 – 0.2812]
UV ₂₅₄ Transmittance (%)	55.23 ± 16.59 (59.80) [29.30 – 74.6]
Average Particle Size (nm)	414.93 ± 198.83 (347.25) [194 - 1068]
Total organic carbon (mg L ⁻¹)	3.55 ± 1.20 (3.23) [1.10 – 7.81]
<i>Escherichia coli</i> (CFU 100 mL ⁻¹)	14,911 ± 51,793 (1,400) [750 – 300,000]
Total coliforms (CFU 100 mL ⁻¹)	69,535 ± 205,691 (16,200) [4,900 – 1,000,000]

171

172 **2.3. Sample collection and analyses**

173 Performance analyses of the following parameters were considered for the evaluation
174 of the WTPt: turbidity (2100N Turbidimeter – *Hach Company*, USA); apparent colour and
175 true colour (DM-COR Colorimeter – *Digimed*, Brazil); absorbance at $\lambda=254$ nm and
176 transmittance at $\lambda=254$ nm (Nanocolor® UV VIS II, *Macherey-Nagel*, Germany); average
177 particle size (Zeta Sizer Nano Z90, *Malvern Company*, UK); total organic carbon – TOC
178 (TOC-L, *Shimazu*, Japan); *Escherichia coli* and TC (Chromocult® coliform, *Merck*,
179 Germany). Standard Methods (APHA et al., 2012) were followed to evaluate the above-
180 mentioned parameters.

181 In Phases 1 and 2, samples were taken from the raw water and from treated water
182 (after filters, after UVC disinfection and after chlorination in the TWT). In Phase 3, one extra
183 collecting point was added before the membrane filtration system to evaluate the blanket
184 filtration step. These water collecting points in the three phases are shown in Figure 1.

185 The three pressure gauges positioned before each cartridge filtration unit were read in
186 each batch operation to evaluate the pressure drop. A ball valve after the pump regulated the
187 flow rate of the system, which was measured daily. The initial flow rate was kept under 4.5 L

188 min^{-1} . The ball valve was completely open at the end of the filter run. In the last 17 batches of
189 Phase 3, the flow rate was deliberately reduced to near 3 L min^{-1} to investigate the effect of
190 increasing the UVC irradiation in the treatment.

191 The filter run was defined here as the number of batches, in which one batch is equal
192 to one day of operation, between starting the operation and stopping it to clean or replace the
193 first membrane filtration unit. The path taken by the water after the centrifugal pump defined
194 the order of the filters.

195 **2.4. Non-woven blanket cleaning**

196 The blanket used in Phase 3 was cleaned when clogging was observed. Maintenance
197 consisted of removing the blanket from the tank, placing it on the floor and scraping the solids
198 using a broom for this procedure only. While the blanket was scraped, it was washed with the
199 water produced by the WTPt. Pictures of the dirty blanket and its cleaning are presented in
200 Figure S5 of the Supplemental Material of this manuscript.

201

202 **2.5. Microscopic analysis**

203 At the end of Phase 3, microorganisms housed on the surfaces of the non-woven
204 blanket and the three pleated filters ($25 \mu\text{m}$, $10 \mu\text{m}$ and $1 \mu\text{m}$) were microscopically
205 identified. To visualise and identify organisms, one drop of the pellet of centrifuged material
206 of each sample was placed in a glass slide, covered with a coverslip and observed in an optic
207 microscopic (BX51, *Olympus*®, Japan), in the bright field under the 40x objective (400x
208 magnification).

209

210 **2.6. Statistical Analyses**

211 The effect of the WTPt on water quality parameters was evaluated by statistical tests
212 performed using the PAST software – PAlaeontological STatistics, published by Hammer et

213 al. [27]. Shapiro-Wilk's test was used to assess data normality ($p > 0.05$). The two-tailed
214 Mann-Whitney U test was used to test the medians of the influent water and the treated water
215 ($p < 0.05$ means different medians). The Kruskal-Wallis test compared the medians of three or
216 more groups (i.e., influent water, water after blanket filtration and filtered water).

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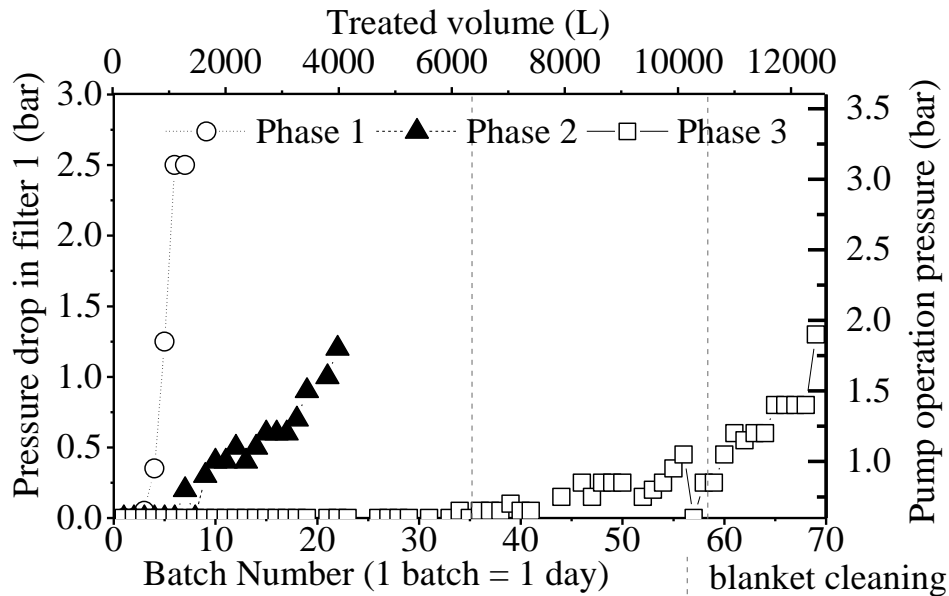
218 **3. Results and Discussion**

219 **3.1. Source Water**

220 The collected river water represented a scenario where the surface water is the only
221 available alternative to a household. In Brazil, surface water is the main source of supply for
222 56% of the cities [28]. This sort of source is susceptible to surface runoff and contamination.
223 The studied raw water was intended for the stressed challenge phase of testing of the WTPt,
224 as well as being one real source of supply.

225 **3.2. Pressure drop in cartridge filters**

226 The pressure drop in filter 1 in the three phases is presented in Figure 3, as well as the
227 pump operation pressure. In Phase 1, Filter 1 (10 μm) started to clog on the fourth day; after
228 this, the pressure drop raised abruptly for the next three days. The operation in this phase was
229 interrupted on the seventh day because the flow rate was too low even though the ball valve
230 after the pump was completely open. The pressure drop in the second filter is shown in Figure
231 S6 of the Supplemental Material of this manuscript.



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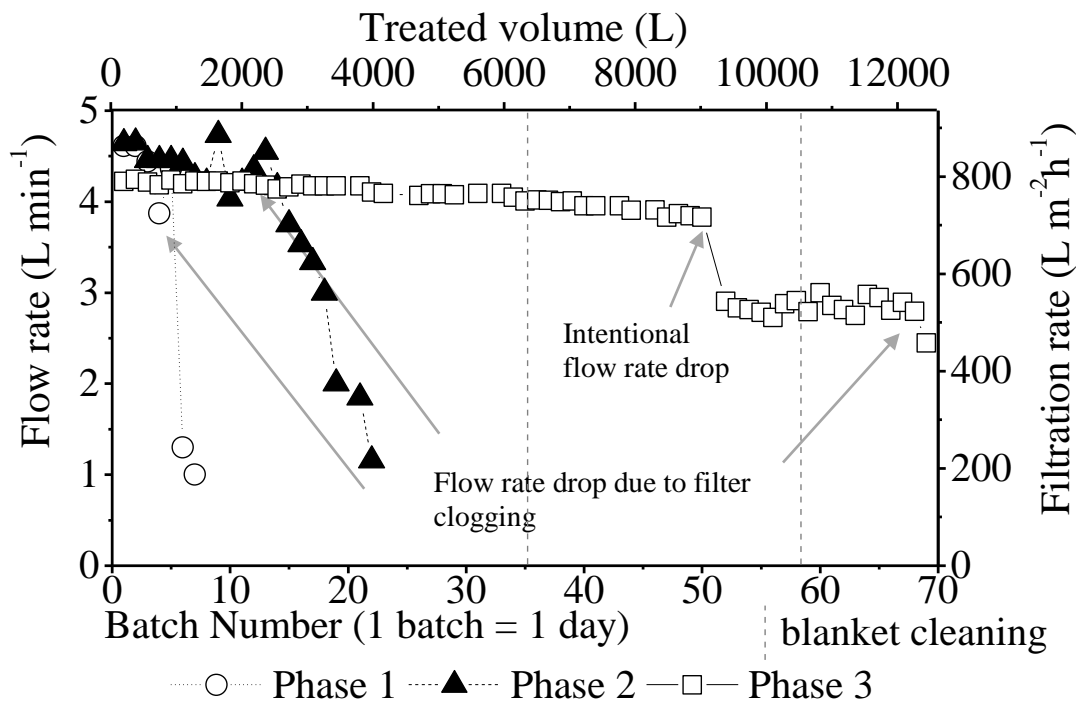
233 Figure 3. Pressure-drop in filter 1 in Phase 1 (2 filters – filter 1: 10 μm), Phase 2 (3 filters –
 234 filter 1: 25 μm) and Phase 3 (blanket + 3 filters – filter 1: 25 μm); and the pump operation
 235 pressure.

236 The cartridge filters in this study are generally used to protect a water purifier from
 237 damage and extend its life [29]. These filters remove suspended solids and should be chosen
 238 according to characteristics such as the particle size and flow rate requirements. It is expected
 239 that head losses increase significantly with prolonged use over time, due to the clogging
 240 particles retained on the outer surface of the cartridge, with a consequent reduction in the
 241 filtration capacity which indicates the need to replace the cartridge itself [30]. The 10 μm
 242 filter was inadequate to be the first filter of the system, despite succeeding the water
 243 decantation for 24 hours in the ST.

244 In Phase 2, the filter 1 (25 μm) started to clog on the eighth day and the filter run
 245 lasted for 22 days. In general, the largest rating size that will remove the intended
 246 contaminants will require the least maintenance [31]. Although more extended than Phase 1,
 247 the 22-day filter run was still insufficient for household drinking water treatment purposes.

248 The pressure drop on the second filter in Phase 2 is shown in Figure S7 of the Supplemental
 249 material of this manuscript.

250 Accordingly, in Phase 3 the pre-filtration was adopted in a non-woven blanket before
 251 the water decantation in the ST. The blanket filtration stage preserved the membrane filter's
 252 lifespan. Nonetheless, as the blanket accumulated suspended solids, it needed to be cleaned,
 253 as shown by the dashed line in Figure 4. The cleaning procedure took 20 min of one person's
 254 work. Phase 3 lasted 69 days and the first membrane filter started to clog around the 40th day.
 255 In this phase, the flow rate was maintained in the desired value for a long period and the need
 256 to increase the opening of the ball valve until the 50th day was not observed (Figure 4). The
 257 pressure drop in filter 2 in Phase 3 is presented in Figure S8 of the Supplemental material of
 258 this manuscript.



259
 260 Figure 4 – Flow rate and filtration rate behaviour of the WTPt in Phase 1 (2 filters), Phase 2
 261 (3 filters) and Phase 3 (blanket + 3 filters).

262 The pleated cartridge filters are surface filters, therefore the filter layer restrains
263 particles higher than the mean pore size. The filter relies on the mechanism of straining. This
264 mechanism is still dominant when the cake layer is formed above the surface [32]. Previous
265 research hypothesized that other deposition mechanisms, such as inertial impact, interception
266 and Brownian diffusion, also occurs on pleated filters as they observed a removal of particles
267 smaller than the filter pores [12]. As particles were retained, the interstitial water velocity
268 increased. This phenomenon can lead to particle breakthrough and Afkhami [12] observed
269 turbidity removal decline with an increase of pressure and consequent increase of interstitial
270 water velocity. Particle breakthrough was not investigated in the present study since there was
271 considerable variation in the turbidity of the natural study water.

272 **3.2. Water quality parameter evaluation**

273 The value of the water quality parameters after the cartridge filters and after UVC
274 disinfection in Phase 1 and Phase 2 are shown in Table 3, as well as the p-value for statistical
275 comparisons between the medians of the raw water parameters and treated.

276 In both Phase 1 and Phase 2, significant differences were observed for the reduction of
277 turbidity, apparent colour and average particle size. The treatment did not significantly
278 change the true colour, absorbance at $\lambda=254$ nm and TOC, in both phases. Indeed, TOC is
279 reported to not be removed at perceptible levels by cartridge filtration and UVC irradiation
280 [31,33].

281 The cartridge filtration was reported to remove nearly 54% of turbidity elsewhere [34].
282 In Phase 1, turbidity removal ranged from 36%, on the first day to 98% on the last day, when
283 the filter was almost completely clogged. In Phase 2, the turbidity removal ranged from 48%
284 to 90%. Data from routine measurements are presented in the figures of the Supplemental
285 Material (Figure S9 to S19).

286 The reductions and variations of water quality parameters from Phase 1 and Phase 2
287 were similar. Nevertheless, the p-values of Phase 2 indicate a statistically stronger difference
288 from this phase compared to the p-values of Phase 1 (Table 3). This was due to the greater
289 filtration run of Phase 2, which enabled greater data collection than the previous phase.

290 Water after cartridge filters presented elevated values of *E. coli* and TC. A significant
291 reduction of these two groups was observed before disinfection only in the last measurement
292 of Phase 1 (2.87 log for *E. coli* and 2.34 log for TC) when the filters were almost clogged and
293 the flow rate was too low (0.86 L min⁻¹). More details can be seen from Figure S14 to Figure
294 S18 (Supplemental Material).

295 The evaluated WTPt was challenged to treat 180 L day⁻¹. Achieving such a daily
296 volume in a household treatment unit can be a difficult task considering the use of compact
297 devices. In the present study, there was still an aggravating factor such as the treatment of
298 water with high peaks of turbidity.

299 The treatment of high turbidity water is usually carried out by decanting the water.
300 However, this operation depends on product dosing, which can be complicated, and mixing
301 units, that take up a lot of space. As the cartridge filters showed to be sensitive to turbidity
302 stress and ineffective, a non-woven blanket was proposed to pre-filter the raw water, which
303 increased the sustainability of the WTPt.

304 In Phase 3, the difference between the three groups of samples was evaluated,
305 therefore, in Table 4, the average value of physical and chemical parameters of the raw water,
306 the water after blanket filtration and settling, and the water after cartridge filtration are
307 presented. The microbiological parameters in these three groups of samples and after the
308 UVC disinfection stage are also presented. Similar to what was observed in Phase 1 and Phase
309 2, the treatment in Phase 3 presented a statistical effect on turbidity, apparent colour and
310 average particle size (Table 4).

311 Table 3 – Treated water characteristics in Phase 1 and Phase 2, after cartridge filtration and after UVC disinfection.

Parameter	Phase 1						Phase 2					
	After cartridge filtration			After UVC disinfection			After cartridge filtration			After UVC disinfection		
	Value (M± SD)	Reduction or variation	p-value	Value (M± SD)	Reduction or variation	p-value	Value (M± SD)	Reduction or variation	p-value	Value (M± SD)	Reduction or variation	p-value
Turbidity (NTU)	7.92 ± 6.59	68 ± 23%	0.0215	N.E.	N.E.	N.E.	4.96 ± 1.42	65 ± 11%	<0.0001	N.E.	N.E.	N.E.
Apparent colour (Hu)	30.60 ± 21.07	57 ± 27%	0.0215	N.E.	N.E.	N.E.	25.83 ± 6.22	47 ± 16%	<0.0001	N.E.	N.E.	N.E.
True colour (Hu)	21.53 ± 6.91	18 ± 15%	0.3837	N.E.	N.E.	N.E.	26.42 ± 4.69	34 ± 11%	0.0606	N.E.	N.E.	N.E.
UV ₂₅₄ Absorbance	0.074 ± 0.017	16 ± 13%	0.3827	N.E.	N.E.	N.E.	0.068 ± 0.012	27 ± 6%	0.1124	N.E.	N.E.	N.E.
Average Particle Size (nm)	306 ± 25	31 ± 14%	0.0369	N.E.	N.E.	N.E.	292 ± 74	27 ± 18%	0.0312	N.E.	N.E.	N.E.
Total organic carbon (mg L ⁻¹)	2.84 ± 0.12	1 ± 2%	1.0000	N.E.	N.E.	N.E.	2.78 ± 0.26	6 ± 25%	0.8852	N.E.	N.E.	N.E.
<i>Escherichia coli</i> (CFU 100 mL ⁻¹)	1,423 ± 1,253	0.55 log median (0.02 – 2.87 log)	Not applica- ble*	<1	3.61 log median (2.87 – 4.06 log)	Not applicable*	371 ± 321	0.58 median (0.20 – 1.26 log)	0.0515	<1	2.92 log median (2.41 -4.08 log)	<0.0001
Total coliforms (CFU 100 mL ⁻¹)	5,835 ± 8,152	1.22 log median (0.10 – 2.33 log)	Not applica- ble*	79 ± 46	2.50 log median (2.08 – 3.10 log)	Not applicable*	6,720 ± 4,276	0.40 log median (0.15 – 0.85 log)	0.03671	50 ± 24	2.47 log median (2.35 -2.94 log)	0.0051

312 *Insufficient data to perform statistical comparison of median. N.E.: parameters not evaluated after UVC-disinfection, as they were not expected to change. M ± SD: mean ±

313 standard deviation. The p-value < 0.05 indicates that there is a difference between the median of parameter of the treated water compared to the raw water.

314 Table 4 – Water characteristics in different stages in Phase 3

Parameter	Raw water (M ± SD)	After Blanket filtration and settling (M ± SD)	After Cartridge filtration (M ± SD)	p-value (equality of medians)	After UVC disinfection	Total reduction or variation
Turbidity (NTU)	37.28 ± 42.63	14.73 ± 16.79	11.55 ± 13.44	<0.0001	N.E.	67% ± 17%
Apparent colour (Hu)	103.66 ± 71.11	62.81 ± 54.44	30.60 ± 21,71	<0.0001	N.E.	51% ± 17%
True colour (Hu)	53.94 ± 21.24	47.48 ± 23.94	40.67 ± 20.36	0.2448	N.E.	26% ± 13%
UV ₂₅₄ Absorbance	0.183 ± 0.060	0.164 ± 0.065	0.143 ± 0.059	0.3306	N.E.	22% ± 15%
Average Particle Size (nm)	377 ± 183	270 ± 34	256 ± 25	<0.0001	N.E.	24% ± 20%
Total organic carbon (mg L ⁻¹)	4.29 ± 1.29	3.86 ± 0.98	3.87 ± 0.92	0.4397	N.E.	8% ± 16%
<i>Escherichia coli</i> (CFU 100 mL ⁻¹)	20,206 ± 62,201	8,958 ± 28,887	8,286 ± 29,110	<0.0001	<1	3.23 log median (2.47 – 5.17 log)
Total coliforms (CFU 100 mL ⁻¹)	95,662 ± 249,306	24,078 ± 57,869	20,076 ± 51,022	<0.0001	47 ± 38	2.91 log (median) (1.62 – 4.24 log)

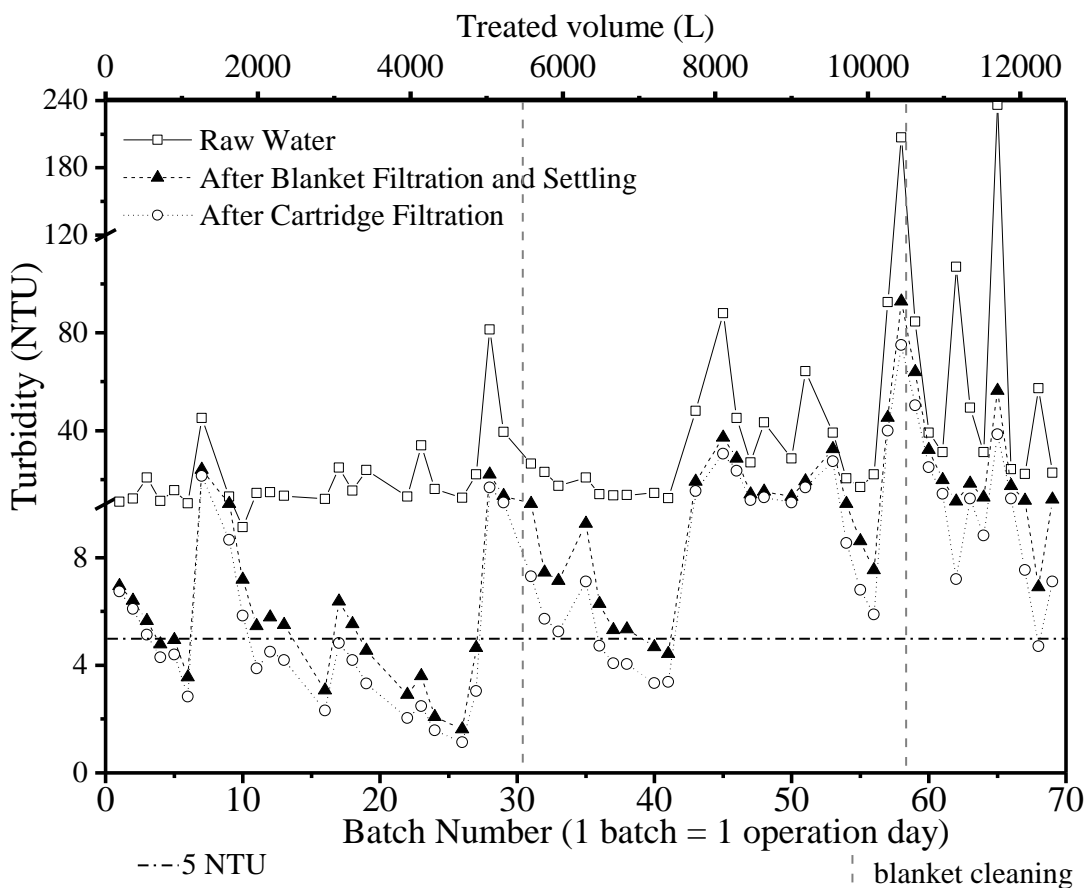
315 N.E.: parameters not evaluated after UV disinfection. The p-value < 0.05 indicates that at least one of the three compared groups has a different

316 median; it means a statistical effect of the treatment. M ± SD: mean ± standard deviation.

317

318 In Phase 3, when comparing the water samples after blanket filtration and settling, and
 319 after cartridge filtration, no statistically significant differences were observed for turbidity
 320 ($p=0.0955$), apparent colour ($p=0.0593$), true colour ($p=0.4306$), average particle size
 321 ($p=0.0634$), total organic carbon ($p=0.9215$), *E. coli* ($p=0.5633$) and TC ($p=0.6798$). Turbidity
 322 explains the effect of the blanket filtration in Figure 5, in which removal in this phase ranged
 323 from 30% to 93% (mean $67\% \pm 16\%$).

324 The cartridge filtration step showed little additional removal of turbidity after the
 325 previous stage (Figure 5). This result was expected since the average particle size influent to
 326 the cartridge filtration (270 ± 34 nm) was inferior to the minimum opening size porous ($1 \mu\text{m}$)
 327 of the filters used.



328
 329 Figure 5 – Turbidity in raw water, water after blanket filtration and settling and after cartridge
 330 filtration in Phase 3.

331 Occasionally, water may contain fine suspended material, which may be too small to
332 be removed by typical cartridge filtration [31]. Thus, the WTPt was not able to produce water
333 with turbidity below the acceptance level of 5.00 NTU [35] when the water after blanket
334 filtration and sedimentation was above 6.50 NTU.

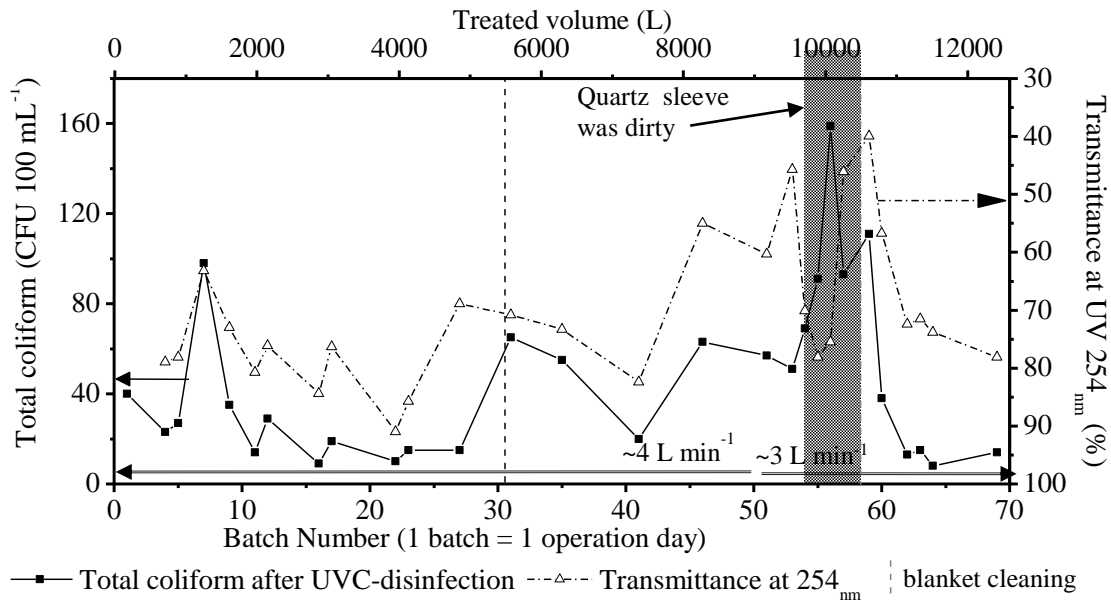
335 On the other hand, in Phase 3 the system proved to be efficient when the raw water
336 showed turbidity values near 10 NTU. On day 26, when the turbidity of the raw water was
337 12.5 NTU, the water after cartridge filtration presented 1.13 NTU of turbidity. Pontius [34]
338 showed effective turbidity removal with filters which had smaller size openings, such as 0.45
339 μm and 0.10 μm . Afkhami et al. [12] showed high removal of turbidity with pleated cartridge
340 filters, however they adopted a pressure drop in the system limited to 1 bar and water
341 treatment with artificial turbidity, caused by the addition of kaolinite.

342 The mean turbidity removal by blanket filtration combined with the settlement was 59
343 \pm 18%, varying from 17% to 90%. As the ST possessed a sedimentation zone (Figure 1), the
344 remaining water from the previous day was mixed with the water that was recently filtered by
345 the blanket. Therefore, the wide range of turbidity removal in the blanket filtration stage is
346 explained by the raw water quality variation (Figure 5).

347 This removal variation can also be explained by the ripening period that may have
348 improved the performance of the filtration by the blanket [36]. A substantial reduction in the
349 turbidity is expected to occur when the solids are retained on the surface of the fabric [37].

350 The filtration by fabric is shown to have a better performance in removing
351 contaminants as the number of layers increases. In our research, only two layers of non-
352 woven blanket were considered, mainly because of the large area of the upside of the 310 L
353 tank. Nonetheless, Siwila and Brink [38] reported improvements of more than 40 percentage
354 points from one to eight layers of non-woven geotextile, in a bench-scale experiment.
355 Therefore, the optimization of fabric filtration for producing a large volume has yet to be

356 evaluated. The turbidity of the treated water met the minimum acceptable value of 5 NTU in
 357 some moments of Phase 3, however the TC values were not below the detection limit (1 CFU
 358 100 mL⁻¹) after UVC disinfection (Figure 6).



359
 360 Figure 6 – Total coliform remaining after UVC disinfection and Transmittance at $\lambda=254$ nm
 361 in Phase 3 related to batch number

362 The efficiency of the UV radiation depends on the transmittance at $\lambda=254$ nm of the
 363 medium and the flow rate. The best results for TC inactivation with the flow rate of 4 L min⁻¹
 364 were attained when the transmittance was near 90%. This agrees with the recommended
 365 values of transmittance-254 for water to use UVC disinfection by the experts in the area,
 366 which is recommended to be > 75-80 % [39]. After the 50th day of the study, the flow rate was
 367 changed to 3 L min⁻¹, aiming to achieve better TC inactivation.

368 Between days 54 and 57, relatively high values of transmittance were noticed,
 369 however less inactivation of TC was obtained, which indicated that the quartz sleeve that
 370 involved the UVC bulb was dirty (Figure 6). Fouling on the quartz-water interface is a linear
 371 process caused by a broad spectrum of inorganic metals and anionic ligands [20,40]. After
 372 cleaning the quartz sleeve, the distance between the lines of "Transmittance UV at 254 nm"

373 and "Total coliform after UVC disinfection" in Figure 6 increased in the part of the
374 experiment of flow rate 3 L min⁻¹ compared to the one of 4 L min⁻¹. The difference in distance
375 between these lines indicates the effect of the flow rate on the total coliform removal.

376 The recommended UV dose for routine drinking water disinfection is set by most
377 regulatory bodies at 40 mJ cm⁻² for a 4-log inactivation, which is sufficient for all bacteria,
378 protozoans and viruses except adenovirus [41,42]. The flow rates applied here promoted
379 doses greater than 40 mJ cm⁻² of UV radiation (Figure 2). However, increasing levels of
380 turbidity, particulate matter and natural organic matter absorb more UV light and make it less
381 available for disinfection [22]. A turbidity value greater than 1 NTU is appointed to shield
382 microorganisms from UV radiation [29].

383 The *E. coli* count after UVC-disinfection was almost always below the detection limit
384 of 1 CFU 100 mL⁻¹. The exception to this was observed in batch numbers 53, 54 and 56,
385 when the quartz sleeve was dirty (Figure S16, Supplemental Material). This group of bacteria
386 is sensitive to UV irradiation, even when the water presented low transmittance at $\lambda = 254$ nm.
387 As an example, in the 57th batch, the water transmittance was 46% and the *E. coli* count was
388 below the limit detection after UVC disinfection. This disinfection step showed to be
389 protective for drinking water household interventions, i.e. above 2 *E. coli* log reduction [43],
390 in all the analyses performed. When the influent water presented *E. coli* contamination as 10⁵
391 CFU 100 mL⁻¹, the UVC-disinfection reached above the 4-log reduction, which is considered
392 highly protective [43].

393 The chlorine dose of 3 mg L⁻¹ was applied daily by the manual chlorinator in all tests,
394 regardless of the water turbidity. This simple device was used to insert the chlorine at a point
395 before the TWT (Figure 1). The flow through the pipe promoted the successful mixing of
396 chlorine. The treated water rested for 30 minutes in the TWT after the end of the filtration to

397 ensure sufficient contact time for the indicator bacteria to be inactivated by the chlorine
398 action.

399 After the chlorination stage, the TC count was below the detection limit of 1 CFU 100
400 mL⁻¹, as was expected. Even when the turbidity of 40 NTU was measured in the filtered water
401 (Phase 3, 57th batch), the inactivation of up to 6 log of TC was attained by chlorination. As an
402 effect of high turbidity of the filtered water, low residual chlorine values were observed, for
403 instance, the value of 0.4 mg L⁻¹ in the 57th batch of Phase 3. Hence, the final chlorination step
404 ensured that the treated water presented bacterial contamination indicators (*E. coli* and TC)
405 below the detection limit, even when the raw water presented total coliforms at concentrations
406 of 10⁶ CFU 100 mL⁻¹.

407 It was not necessary to increase the chlorine dose according to the increase in
408 turbidity, contrary to what was previously recommended by the literature [44]. As
409 householders cannot measure the water turbidity before chlorination [44], the present study
410 aimed to ensure that the dose of 3 mg L⁻¹ would be sufficient to inactivate bacteria, regardless
411 of turbidity. On the other hand, if the householder realizes that the chlorinated water is turbid,
412 it is recommended to be consumed within 8 h post-chlorination [45], because the residual
413 chlorine might not be maintained.

414 Although there is a concern regarding the formation of potentially carcinogenic by-
415 products when chlorine is dosed in turbid water, Lantagne et al. [46] did not observe a
416 concentration of trihalomethanes (THM) that exceeded the WHO guidelines when adding
417 chlorine to turbid waters with TOC concentrations ranging between 0 and 9.8 mg L⁻¹.
418 Besides, Abdullah et al. [47] did not find a correlation between water turbidity and THM
419 formation.

420

421 **3.3. Microbiological community on the surface of filters and blanket**

422 According to the microscopic analysis of filter’s surfaces, algae was the most
 423 predominant group, both in the variety of genera and number, with some genera such as
 424 *Melosira*, *Navicula*, *Pleurosigma* and *Trachelomonas* occurring in all samples. However,
 425 cyanobacteria, helminths and protozoa were also identified. Protozoa was the second most
 426 prevalent group founded herein. Among them, we highlight *Corythion* spp. and *Giardia* spp.
 427 found in all samples. The visualized genera retained in each filter of the Phase 3 experiment
 428 are presented in Table 5.

429 The methodology used observed the pre and post content of the blanket, the 25 µm
 430 filter and the 10 µm, since the sediments retained on the surface of one barrier represented
 431 what passed through the previous one. Hence, it was not possible to observe the
 432 microorganisms post the 1 µm filter.

433 Table 5 – Microorganisms in the sediments from blanket and cartridge filters identified by
 434 bright field microscopy

Biological class	Microorganisms	Blanket	Pleated filters		
			25 µm	10 µm	1 µm
Algae	<i>Acanthosphaera</i> spp.		x		
	<i>Achnanthydium</i> spp.	x		x	x
	<i>Ankistrodesmus</i> spp.	x	x		
	<i>Aulacoseira</i> spp.	x	x		
	<i>Asterocystis</i> spp.				x
	<i>Chilomonas</i> spp.		x		
	<i>Chlorella</i> spp.	x	x	x	x
	<i>Chlamydomonas</i> spp.		x		
	<i>Closterium</i> spp.				x
	<i>Coelastrum</i> spp.	x	x		
	<i>Cyathomonas</i> spp.		x		
	<i>Cyclotella</i> spp.			x	
	<i>Desmodesmus</i> spp.		x		x
	<i>Diatoma</i> spp.				x
	<i>Euastrum</i> spp.		x		
	<i>Euglena</i> spp.	x	x		
	<i>Kirchneriella</i> spp.	x			
	<i>Melosira</i> spp.	x	x	x	x
	<i>Micrasterias</i> spp.		x		
	<i>Navicula</i> spp.	x	x	x	x
<i>Nitzschia</i> spp.	x	x			
<i>Oocystis</i> spp.	x				

	<i>Palmella</i> spp.	x			
	<i>Phacus</i> spp.		x		
	<i>Pleurosigma</i> spp.	x	x	x	x
	<i>Rhodomonas</i> spp.		x	x	
	<i>Scenedesmus</i> spp.	x	x	x	
	<i>Sphaerocystis</i> spp.	x	x		
	<i>Staurodesmus</i> spp.		x		
	<i>Synedra</i> spp.	x	x	x	x
	<i>Tetrastrum</i> spp.	x			
	<i>Trachelomonas</i> spp.	x	x	x	x
Helminths	<i>Tabellaria</i> spp.	x			
	<i>Hymenolepsi</i> spp. (egg)		x		
	Nematode (larvae)	x		x	
Protozoa	<i>Blastocystis</i>				
	<i>Coleps</i> spp.	x			
	<i>Corythion</i> spp.	x	x	x	x
	<i>Entamoeba</i> spp. (cyst)			x	x
	<i>Euplotes</i> spp.		x		
	<i>Giardia</i> spp. (cyst)	x	x	x	x
	<i>Paramecium</i> spp.	x			
	<i>Vorticella</i> spp.		x		
Others	<i>Anabaena</i> spp. (cyanobacteria)	x	x	x	x
	Rotifera (animalia)	x			

435 (x): surface above where the microorganisms were identified.

436 One study has demonstrated infective-stage larvae being able to traverse
437 polypropylene cartridge filters of 20 μm , 10 μm and 1 μm filtration ratings [48]. Even though
438 filtration rating as 1 μm is expected to completely retain larger organisms such as nematodes,
439 nominal pore size ratings are the average pore size rather than the largest; particles larger than
440 the nominal pore size may pass through the filter [16]. Many filter manufacturers attest that
441 filter micron sizing is based on nominal particulate ratings of >85% of a given size as
442 determined from single-pass particle counting results [48]. This characteristic partially
443 explains the presence of some organisms such as nematode larvae in the surface of 10 μm
444 filter and *Corythion* spp. in the surface of 1 μm filter, even though they are organisms larger
445 (\cong 1mm and \cong 45 μm , respectively) than the porosity of the filters used in the system. As
446 shown in Table 5, nematode was identified on the surface of the 10 μm filter, which was not
447 identified after the 10 μm filter.

448 While some organisms require inactivation doses within the spectrum offered by the
449 UVC system used, such as *Cryptosporidium* [42] and nematode larvae [49], some nematode

450 eggs are hardly susceptible to this type of water treatment [49]. Thus, it is essential to establish
451 periodic cleaning of the quartz sleeve to inactivate a wide variety of microorganisms, because
452 clogging reduces UV light transmittance [42] and it is essential to retain the egg form of
453 helminths by physical barriers.

454 Despite the evasion of some microorganisms through physical barriers (25 μm and 10
455 μm) the use of these blanket and three filters in sequence, among other things, increases their
456 lifespan since the accumulation of microorganisms is partitioned, according to the size, which
457 postpones the clogging of the filters, which last longer. The partitioned grouping of
458 microorganisms also results in more than one biofilm. The gelatinous aspect of biofilm
459 favours the retention of microorganisms which are removed from the water, helping to
460 improve its quality.

461 **3.4. Operation and cost evaluation**

462 The potential user of the WTPt can have autonomy to operate the system with specific
463 training and periodic follow-ups. Attention should be paid to the functioning of the UVC-
464 lamp, indicator lights monitor the lamp operation and the user can easily interpret if it is or
465 not working properly. The commercial UVC-lamp is made of a stainless-steel cylinder that
466 protects the user from any UV radiation. The mercury bulb is kept inside a quartz-sleeve.
467 Instructions should be provided to the household user on how to periodically clean the quartz
468 sleeve and how to change and dispose of the bulb after the end of its lifespan. In the case of
469 interventions, it would be better to be accompanied by specialized personnel for periodic
470 maintenance. Maintenance by households would be required to ensure efficient and correct
471 application of the system.

472 The WTPt cost US\$ 1,114 (February, 2021). The most expensive item of the system
473 was the UVC-lamp, acquired in Brazil for US\$ 415. Each cartridge filter cost US\$ 5.90 and
474 the replacement UV bulb was budgeted at US\$ 99.00. Considering the system working 1h

475 per day, it would expend monthly 11.20 KWh and 0.51 KWh of power consumption of the
 476 pump and UVC-lamp, respectively. The power consumption of this system could be reduced
 477 by 75% by replacing the centrifugal pump by one 80W diaphragm pump, which fits in the
 478 configuration of the WTPt.

479 The operation and maintenance costs of the system are presented in Table 6. The
 480 major expense would be the replacement of the three cartridge filters after 59 days of
 481 operation. By comparison, the flocculant-disinfectant sachet from Protec & Gumble®, which
 482 can treat turbid water, would cost US\$ 10 per 1,000 L of water treated [50].

483 The proposed system can treat daily larger volumes than presented here. Perhaps a
 484 household could not afford the initial price of the WTPt, nevertheless the cost per litre of
 485 treated water is compatible with the Brazilian minimum income. It could also be a solution in
 486 for small schools, farms, or any isolated facility.

487

488 Table 6 – Operation and maintenance costs of the water treatment prototype

Expense item	1,000 L (US\$)	1 month (US\$)
UVC bulb (9,000 h per bulb)	0.06	0.32
Cartridge filters (12,000 L)	1.42	7.67
Chlorine	0.07	0.38
UVC-lamp: expenses on power consumption*	0.01	0.05
Pump: expenses on power consumption*	0.23	1.23
Total	1.79	9.66

489 *considering US\$ 0.11 KWh⁻¹ (average price of electricity in rural areas in Brazil)

490

491 4. Conclusions

492 This work has experimentally assessed the capacity and effectiveness of a treatment
 493 prototype based on cartridge filtration, UVC disinfection and chlorination to provide the
 494 potable water daily needs for a household of 5 members sourcing from a local river.

495

496 It was observed that commercial cartridge filters were severely impacted by stressing
497 conditions of turbidity, as direct filtration through 10 and 1 μm cartridge filters resulted in
498 only 7 batches treating 180 L each. Nevertheless, pre-treatment with fabric filtration and 25
499 μm filter increased the sustainability of the evaluated system, resulting in 69 batches (days of
500 operation). From microscopic observation, an active biological layer was observed on the
501 surface of filters and the blanket, which could have contributed to both the filter clogging and
502 retaining of particles.

503 The proposed system can be an attractive solution considering source water with
504 turbidity below 10 NTU. When considering sources with higher values of turbidity, more
505 studies should be conducted to optimize the water clarification since the UVC disinfection
506 was not carried out properly in case of filtered water turbidity higher than 1 NTU and
507 transmittance at UV 254 nm higher than 75%. The chlorine disinfection step was one barrier
508 of safety in the case of the present study as the final water presented *E. coli* and the total
509 coliform count was below the detection limit (virtually absence) in all tests performed.

510

511 **5. Declaration of competing interest**

512

513 The authors declare that they have no known competing financial interests or personal
514 relationships that could have appeared to influence the work reported in this paper.

515

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517

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520

521 **7. Supplementary Material**

522

523 The scheme of the water treatment prototype evaluated the UVC-lamp dose
524 information, the semi-automatic operation protocol for the WTPt, the block diagram of the
525 WTPt electrical control, the steps for the blanket cleaning procedure, the pressure drop per
526 filter, turbidity and colour and coliform removals per operation phases are provided as
527 supplementary material.

528

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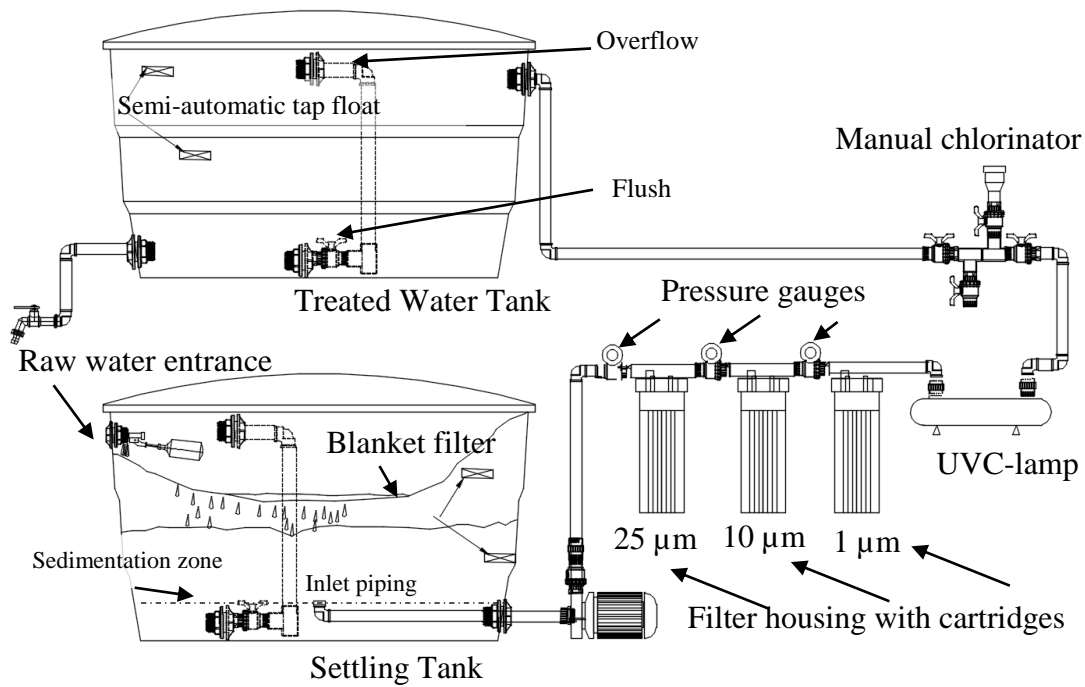
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Supplemental material

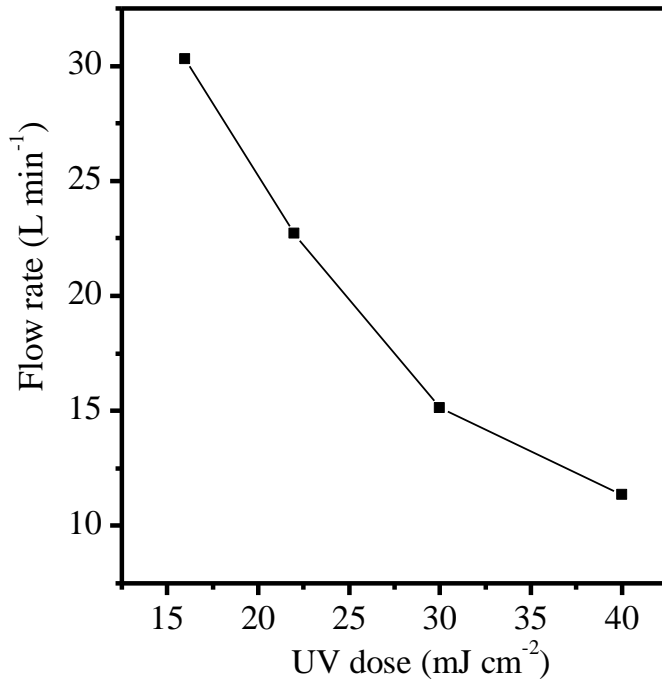


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Figure S1 – Scheme of the Water Treatment Prototype evaluated

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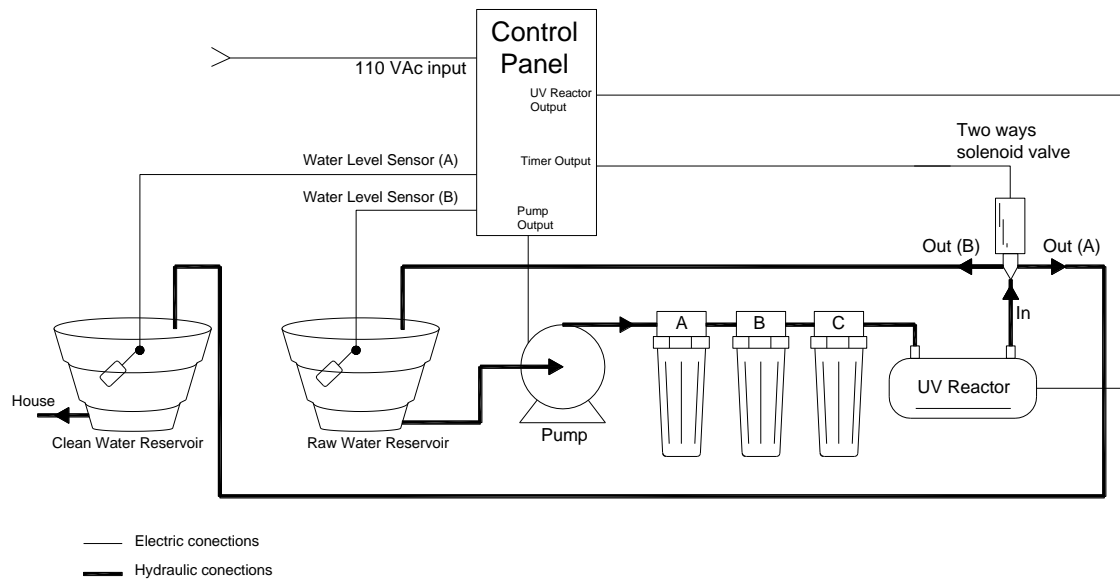
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Figure S2: UVC-lamp Polaris™ UV-4C information dose according to the flow rate, adapted

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from HYDRONIX (2018)

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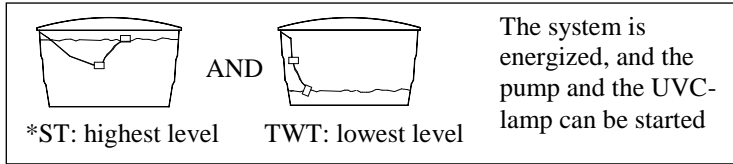


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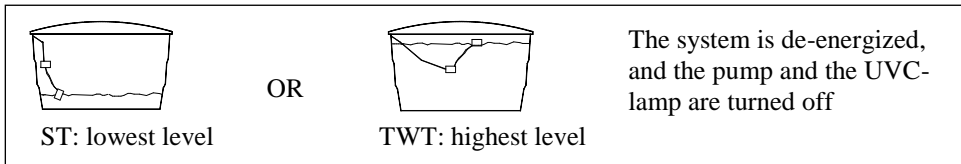
699 Figure S3 – Block diagram of the electrical control of the Water Treatment Prototype

700

- 8:00 am: The pump and the UVC-lamp are turned on (manual action).



- 8:00-8:05 am**: The water flows through the filters, the UVC-lamp, the two-way solenoid valve, and returns to the ST. The UVC-bulb warms up.
- The operator adds chlorine (sodium hypochlorite or calcium hypochlorite) to the manual chlorinator (manual action).
- 8:05 am: A timer sends a signal to the solenoid valve to close the returning way and open the way to the TWT. The water flows through the filters, the UVC-system, the manual chlorinator and to the TWT (automatic action).
- ~8:50 am: The ST goes to the lowest level, and the pump and the UVC-lamp are turn-off (automatic action).



- After 8:50 am: The operator opens the raw water entering gauge and the water decants in the ST until the next operation day.
- The treated water is collected from the TWT, which is then empty.

* ST: Settling Tank; TWT: Treated Water Tank
 **During the first 5 minutes of the pump operation, the water passed through the UVC-lamp and returned to the ST in the control of a two-way solenoid valve. This action was performed to remove any air bubble and debris and to allow the warming of the UVC-lamp, as a manufacturer UVC-lamp recommendation.

701
 702 Figure S4 – Model of the semi-automatic operation protocol for the Water Treatment
 703 Prototype.
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707 Figure S5- Steps for the blanket cleaning procedure

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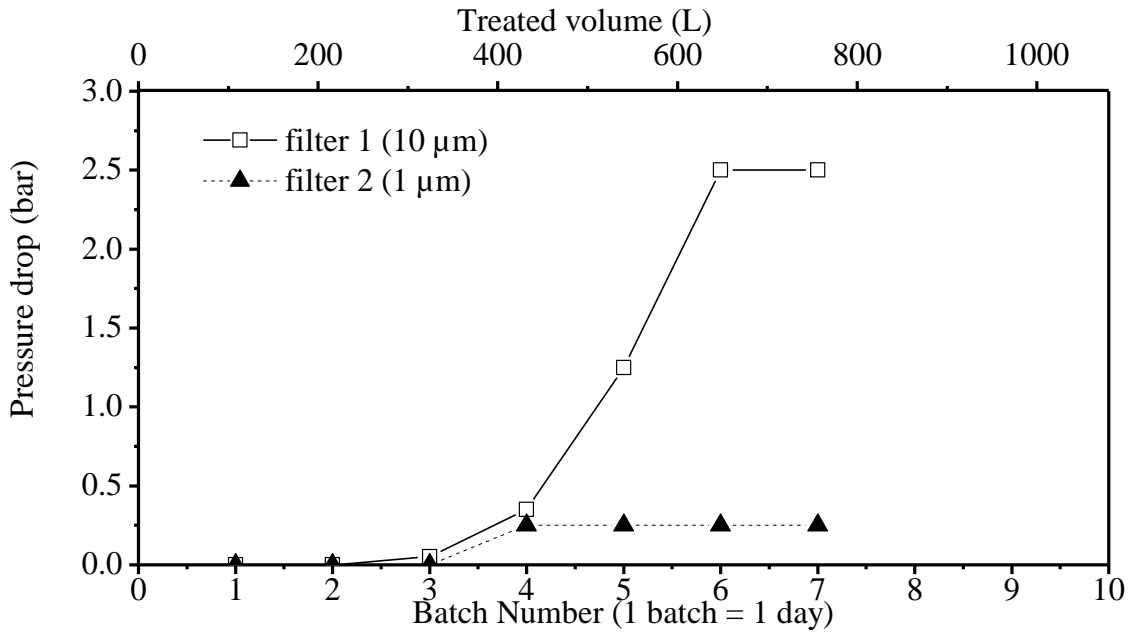
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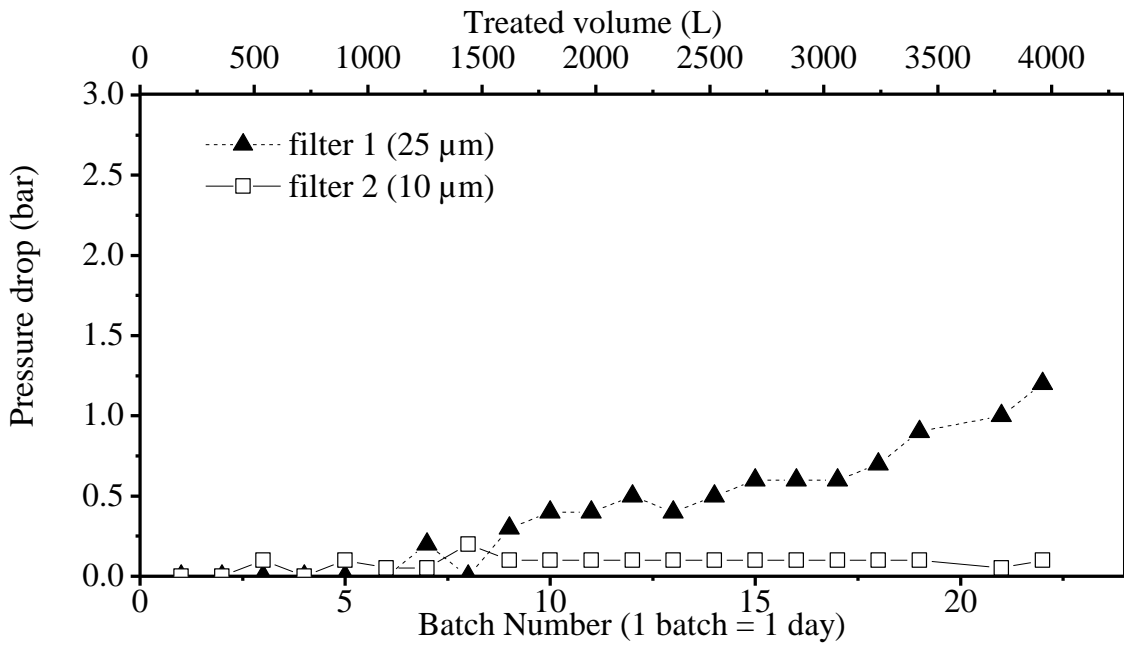
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715 Figure S6 – Pressure drop in filter 1 (10 μm) and filter 2 (10 μm) during Phase 1 operation

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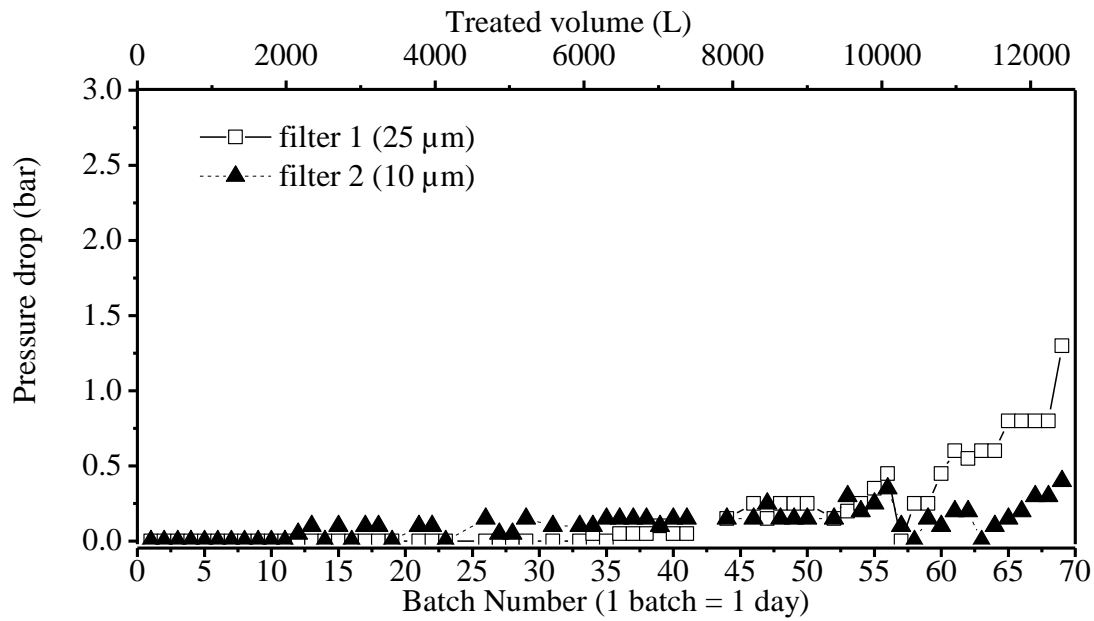


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718 Figure S7 – Pressure drop in filter 1 (25 μm) and filter 2 (10 μm) during Phase 2 operation

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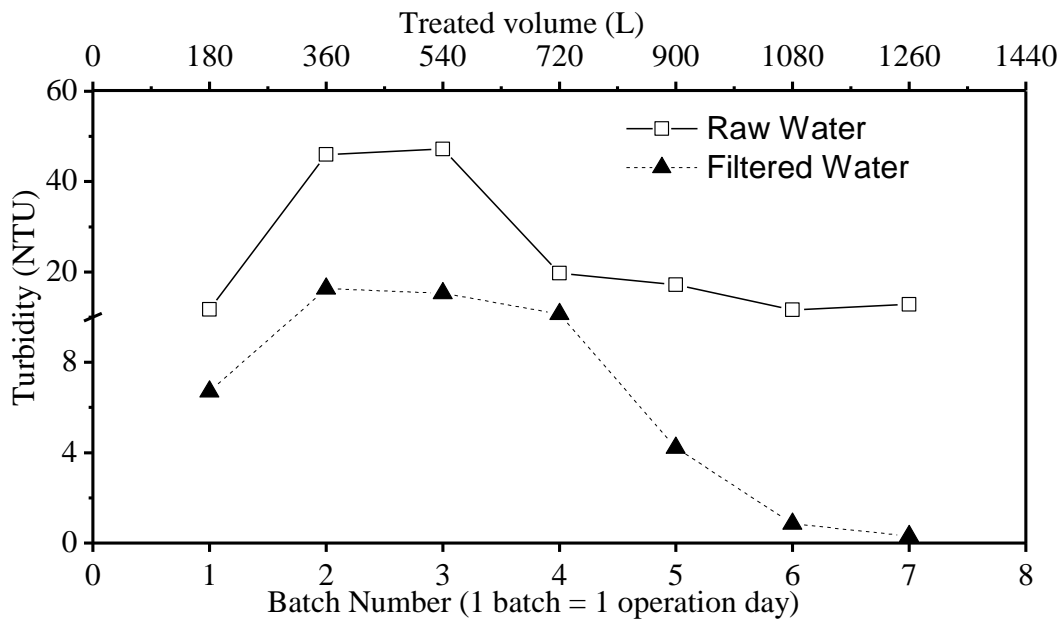
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722 Figure S8 – Pressure drop in filter 1 (25 μm) and filter 2 (10 μm) during Phase 3 operation

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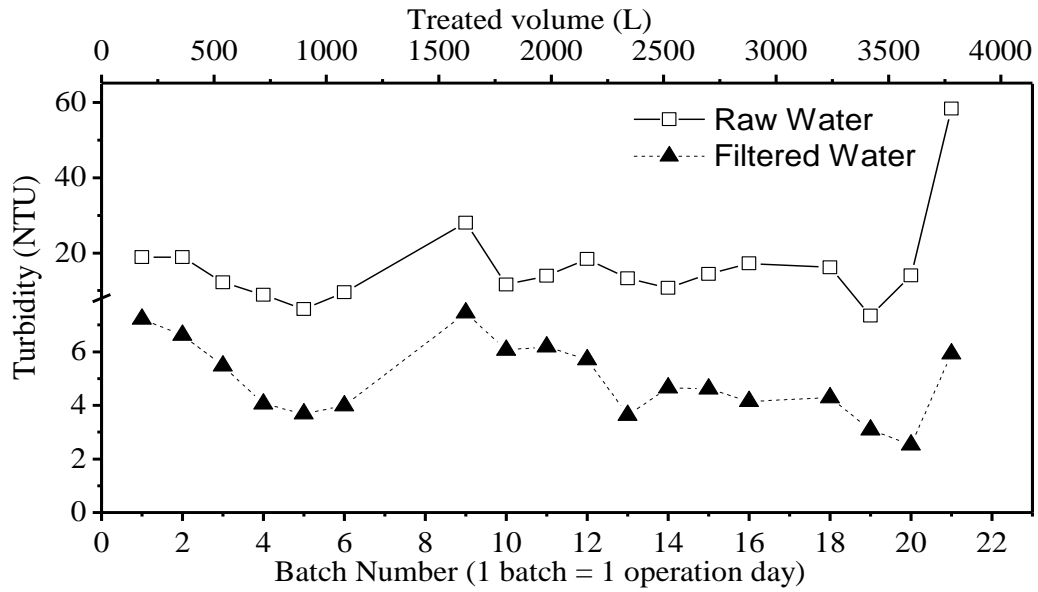
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725 Figure S9 – Turbidity of raw and filtered water during Phase 1 operation

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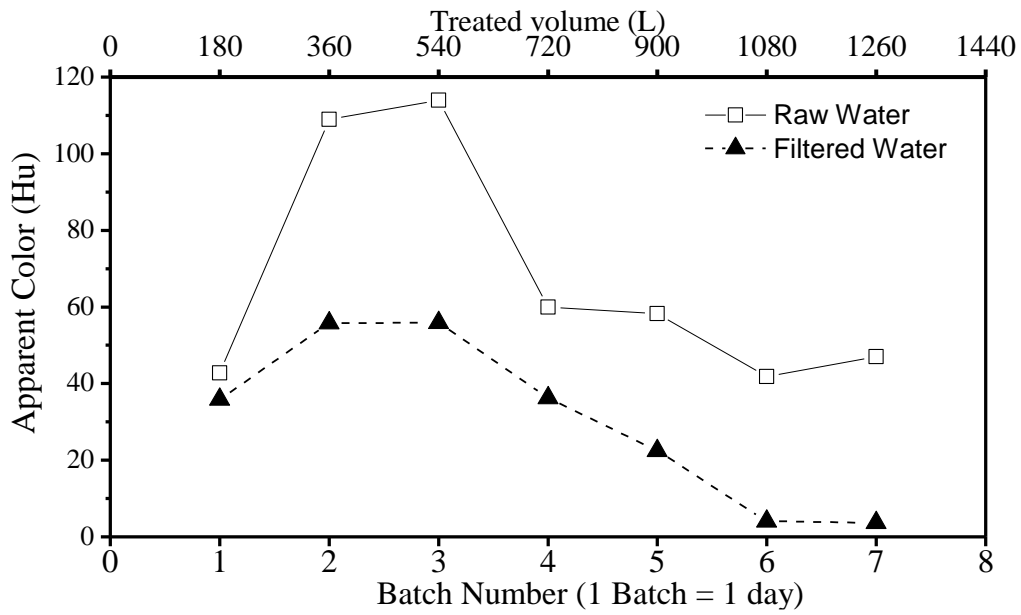
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730 Figure S10 – Turbidity of raw and filtered water during Phase 2 operation

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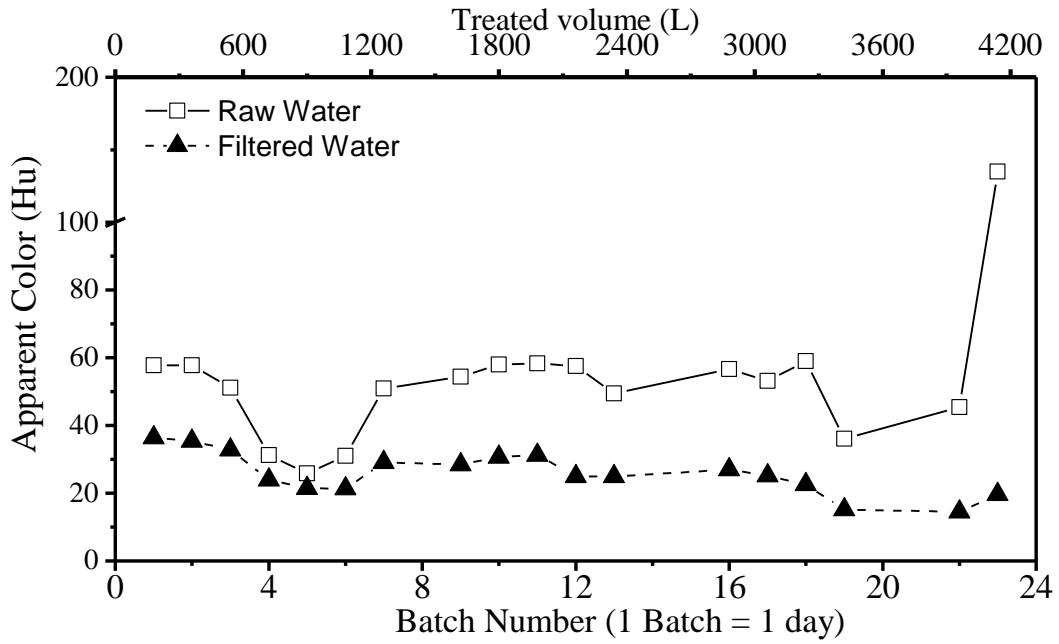
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734 Figure S11 – Apparent colour of raw and filtered Water during Phase 1 operation

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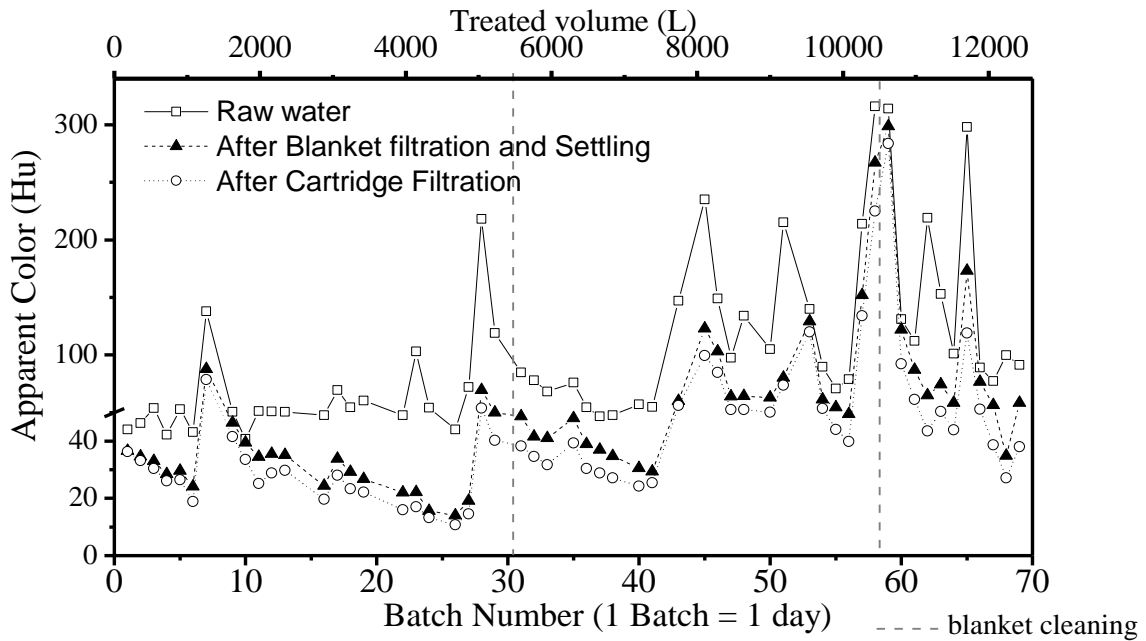
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739 Figure S12 – Apparent colour of raw and treated water during Phase 2 operation

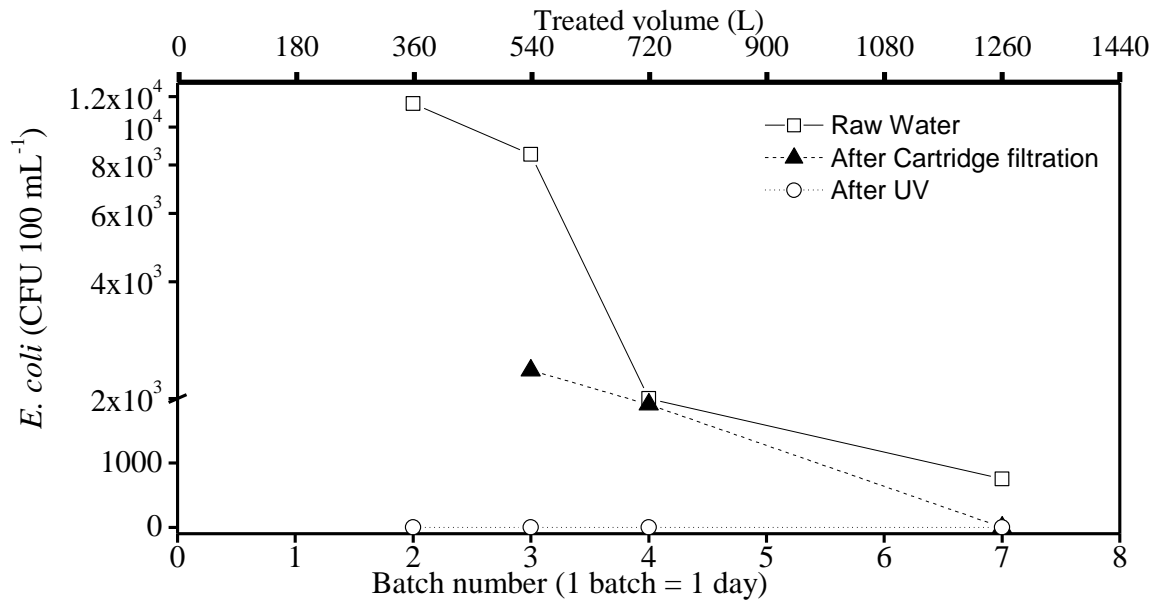
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741

742 Figure S13 – Apparent colour in raw water, water after blanket filtration and settling, and
 743 water after cartridge filtration, during Phase 3 operation

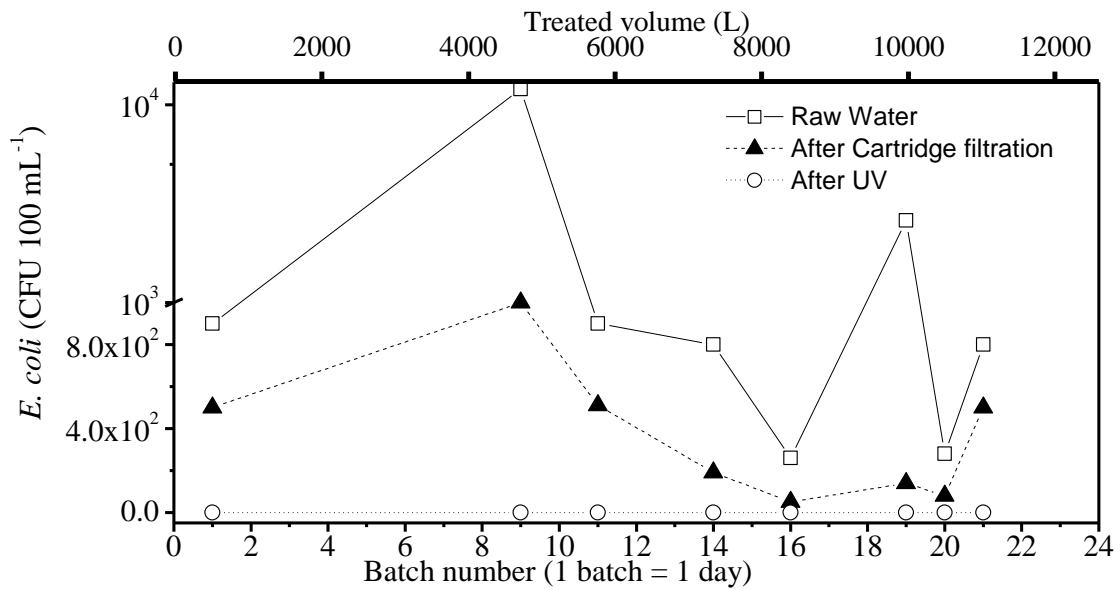
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746 Figure S14 – *Escherichia coli* in raw water, after cartridge filtration, and after UV irradiation
 747 during Phase 1 operation

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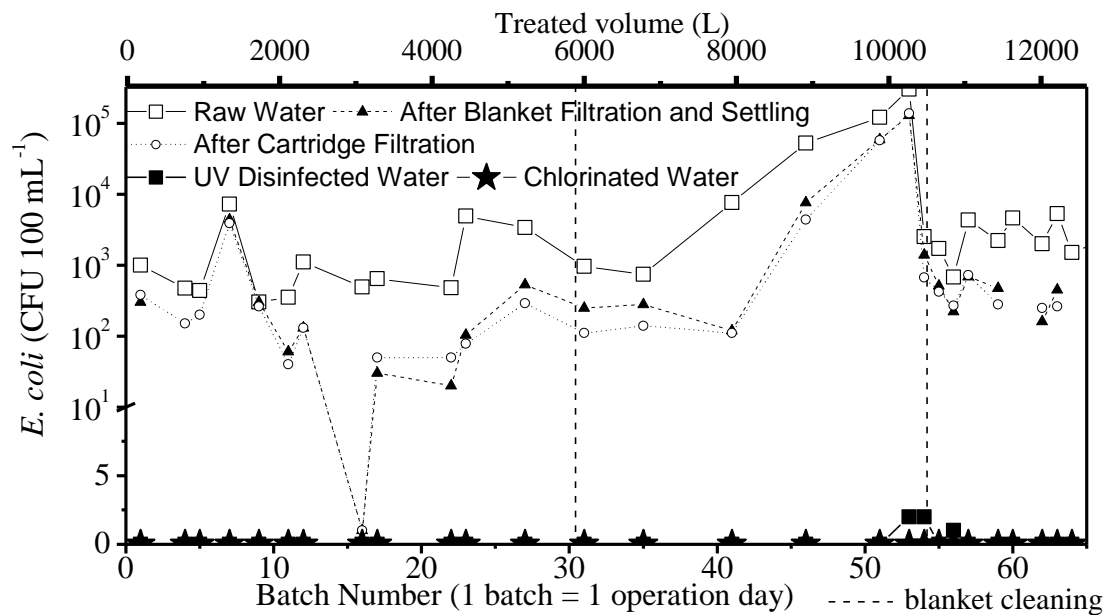
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750 Figure S15 - *Escherichia coli* in raw water, after cartridge filtration, and after UV irradiation
 751 during Phase 2 operation

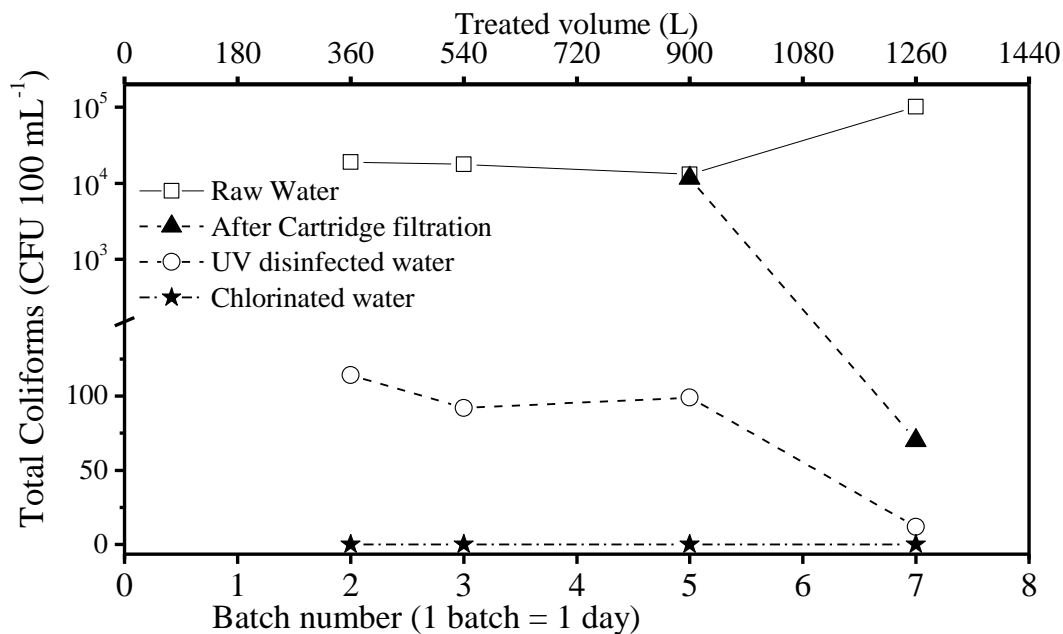
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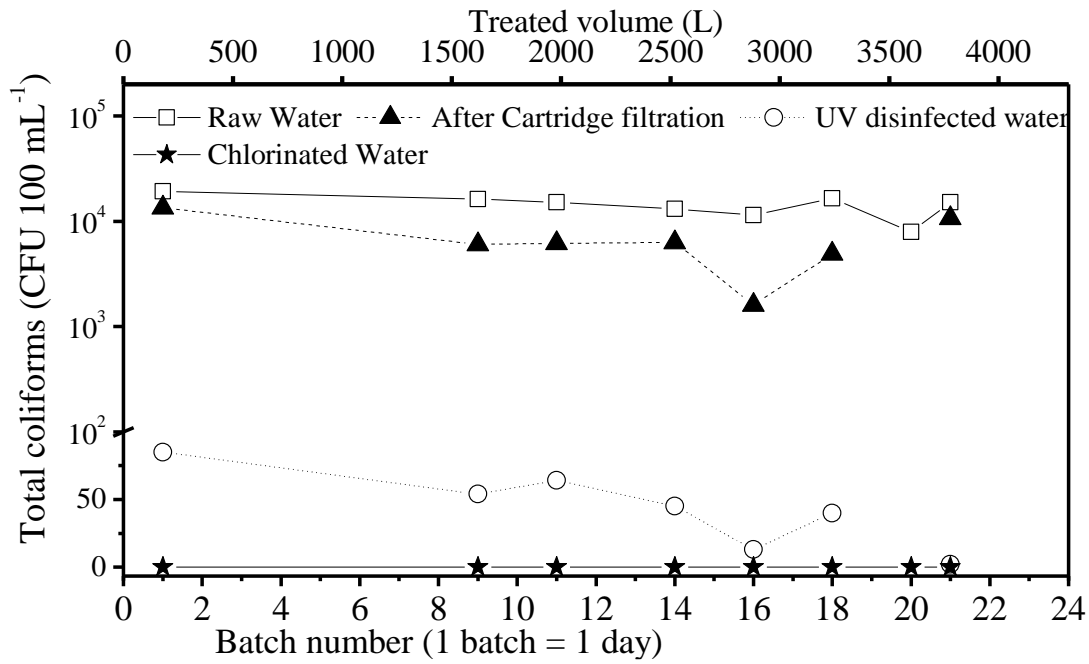
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755
 756 Figure S16- *Escherichia coli* in raw water, after blanket filtration and settling, after cartridge
 757 filtration, and after UV irradiation during Phase 3 operation
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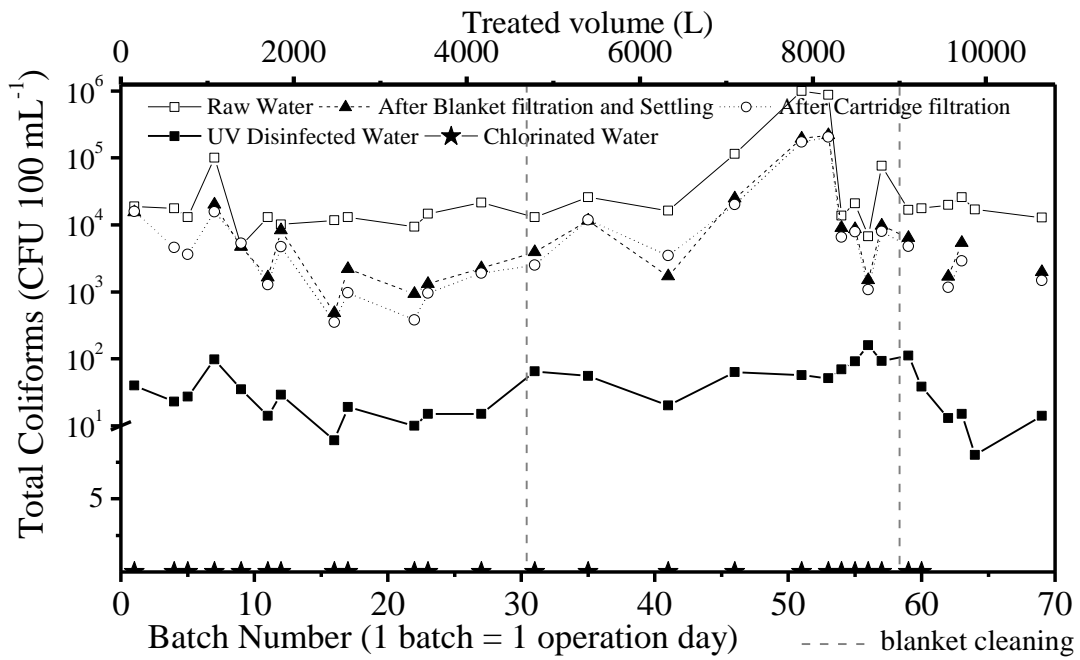


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 760 Figure S17 – Total coliforms in raw water, after cartridge filtration, after UV irradiation, and
 761 after chlorination, during Phase 1 operation



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763 Figure S18 - Total coliforms in raw water, after cartridge filtration, after UV irradiation, and
 764 after chlorination during Phase 2 operation



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766 Figure S19 - Total coliforms in raw water, after blanket filtration and settling, after cartridge
 767 filtration, after UV irradiation, and after chlorination during Phase 3 operation

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