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# Start-up of a drinking water biofilter physical, chemical and bacteriological changes



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## INTRODUCTION

When producing drinking water from groundwater, some waterworks use biofilters as the heart of the treatment process. In biofilters, microorganisms are allowed to populate granular filter media and to carry out the work of purifying the water (de Vet, 2009). This process is gaining attention because of a number of attractive features including 1) low price, 2) no addition of chemicals and 3) increase in the microbiological stability of the finished water.

One drawback of biofilters is the long start-up period when new filter medium is commissioned. During the start-up period, an inorganic coating and a biofilm are established on the filter medium, after which the treated water complies with drinking water criteria. This period typically lasts two or more months (Cai, 2015; Stembal, 2004; Zeng, 2010). Disadvantages of a long start-up period include: 1) the need to discharge water to the environment since the finished water does not comply with drinking water standards, 2) the use of energy and the waste of a precious resource, and 3) the need for an alternative drinking water source for the consumers for the start-up period. If the start-up process is to be optimized, a thorough knowledge of the development of fullyfunctioning biofilters is required.

This poster elucidates the start-up process through a holistic monitoring approach at a newly-constructed full-scale waterworks in Denmark. This poster documents a natural start-up, using only inherent inoculation from microorganisms that are present in the raw water and the water used for backwash (no pro-active inoculation with old filter media or backwash water sludge was utilized).

#### **METHODS Raw water quality Filtration** Average Std. Dev. Parameter Filter 2 0.22 mgL<sup>-1</sup> Oxygen 1.40 mgL<sup>-1</sup> Iron 0.45 mgL<sup>-1</sup> Manganese Ammonium 0.21 mgL<sup>-1</sup> 0<sub>2</sub> Magnesium 7.6 mgL<sup>-1</sup> 7.3 bН 58 mS/m Conductivity 3375 mm 8.9°C Temperature Support material 2 Quartz sand Support material 1 Tap - Sampling

One of the production lines at Truelsbjerg waterworks, Denmark (Søborg et al., 2015).

Shading indicates concentrations not in compliance with Danish drinking water criteria.

0.12

0.19

0.07

0.04

0.3

0.06

0.8

0.1

### Sampling

Water samples (unfiltered) were collected from stainless steel taps at 16 locations: 13 different depths on Filter 1 as well as raw water, water between filters and finished water.

## **RESULTS and DISCUSSION**

### Chemical

Separate phases for removal of individual compounds during start-up were observed in this study.

concentrations achieved compliance Iron immediately, complete ammonium almost removal required about 6 weeks while complete manganese removal required about 10 weeks.

### Fe, NH4<sup>+</sup> & Mn (mg·L<sup>-1</sup>) 1.0





(Modified from Frischherz et al., 1985)

- Clear stratification with depth was observed in water samples collected from Filter 1:
- Iron was removed at the top of the filter (top 30 cm).

- Filter media samples were collected from 4 different depths in Filter 1 using a hollow stainless steel probe.
- Backwash water samples were collected at one minute intervals during selected backwash events.

### Analyses

Physical	Chemical	Microbial
Continuous measurements <ul> <li>Flow</li> <li>Temperature</li> <li>Pressure</li> <li>Turbidity</li> </ul>	Continuous measurements <ul> <li>Dissolved oxygen</li> <li>In-line pH</li> <li>Conductivity</li> <li>At-line ammonium</li> </ul>	Continuous measurements <ul> <li>Bacterial counts (Grundfos</li> <li>BACMON)</li> </ul>
<ul> <li>Grab samples</li> <li>Grain size, surface area and particle shape using Camsizer®64, Retch Technology GmbH</li> </ul>	<ul> <li>Grab samples</li> <li>Iron, manganese and ammonium using Hach DR3900 spectrophotometer</li> <li>Nitrite</li> </ul>	<ul> <li>Grab samples</li> <li>Heterotrophic plate counts</li> <li>DNA isolation (PowerBiofilm, MoBio Laboratories Inc.)</li> <li>qPCR (Eubacteria and relevant bacterial groups)</li> </ul>



### **Biological**

- Ammonium removed was immediately below the iron strata (30-70 cm).
- Manganese was mainly removed close to the bottom of the filter (130-210 cm).

The central portion of Filter 1 appeared to be less active.

The bacterial community on samples of filter medium was investigated at various times during the start-up period using qPCR methods. Total bacteria (Eubacteria) as well as ammonium oxidizing bacteria (AOB), Nitrospira and Leptotrix were quantified.



Numbers of specific bacterial groups increased over time, while EUB remained nearly constant. The highest number of EUB was found in the top layer of the filter. At the end of the start-up period, EUBs outnumbered the sum of the other bacterial groups by about 10:1.



event. Note that the goal of backwashing during start-up is to avoid pressure build-up, not to ensure clean finished water.

Colour changes in filter media. Over time, the colour of the filter media changed in the four depths that were sampled.



Parameter	calcium carbonate	Filter 2		Both filters	
Description		manganese oxide	quartz sand	support material	
Particle density (kgL <sup>-1</sup> )	2.41	3.32	2.51	-	
Porosity (%)	40	44	45	-	
Grain size (mm, 10-90% fractile)	2.3-4.1	1.6-3.2	0.5-0.8	3-5/1.6-2.5	
Layer thickness (mm)	2290	200	2090	310	

Filter media properties. Precipitation of iron oxides (red color) and manganese oxides (black color) were seen on filter media samples collected over time from four depths of each filter (Breda et al., 2016).

## **CONCLUSIONS**

- Using inherent inoculation, full-scale start-up was complete after a period of approximately 10 weeks.
- The change from virgin filter media to fully functioning mature filter media is a complex mix of physical, chemical and microbiological processes. Holistic monitoring of these processes using water, filter media and backwash water samples provided a more clear understanding of the start-up period.
- Total bacteria (Eubacteria) were most abundant in the top 40 cm of Filter 1. Selected bacterial groups (AOB, Nitrospira, Leptotrix) represented only a small percentage of the total bacteria.
- Results from this work have important implications for optimizing the start-up process such as when and where to inoculate and what to inoculate with.

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