Occurrence of *Microthrix parvicella* in sequencing batch reactors

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**ABSTRACT**

Sequencing batch reactors (SBRs) are known for high process stability and usually have a good sludge volume index (SVI). Nevertheless, in many SBRs in Germany for municipal wastewater treatment, scum and foam problems can occur, and SVI can be larger than 200 mL/g. The microscopic investigations of the activated sludge from plants with nitrogen and phosphorus removal have shown that *Microthrix parvicella* is dominant in the activated sludge in most of them. Studies showed that the optimum growth of *M. parvicella* is performed at a high sludge age (>20 d) and low sludge load in the range of 0.05–0.2 kg of biochemical oxygen demand per kg of total suspended solids per day (kg BOD₅/(TSS·d)). The investigations in 13 SBRs with simultaneous aerobic sludge stabilization (most of them are operated with a system called differential internal cycle strategy sequential batch reactor (DIC-SBR)) show that *M. parvicella* is able to grow in sludge loads less than 0.05 kg BOD₅/(kg TSS·d) as well. To optimize the operation of those SBRs, long cycle times (8–12 h) and dosing of iron salts to eliminate long-chain fatty acids are both recommended. This leads to better SVI and keeps *M. parvicella* at a low frequency.

**Key words** | DIC-SBR, *Microthrix parvicella*, SBR, simultaneous aerobic sludge stabilization

**INTRODUCTION**

The biological wastewater treatment is based on two main processes, the biological conversion of the pollutants caused by the activity of the bacteria in the biological reactor, and the separation of purified wastewater from the sludge. This separation is usually performed in the clarifier or can be combined with the biological process in the same unit such as in sequencing batch reactors (SBRs), which have been mainly used in the recent years as treatment plants for areas with more than 5,000 population equivalents (PE) (Lemmer & Lind 2000; Jenkins et al. 2004).

Most wastewater treatment plants (WWTP) require low sludge loads and the establishment of aerobic, anoxic and anaerobic zones to enable biological nitrogen and phosphorus removal. This has led to new biological communities in activated sludge. The most common filamentous bacteria which can live under these conditions have high flexibility and can exist in all the zones, for example *Microthrix parvicella*, nocardioforms, type 0092, type 1851 and *Nostocoida limicola* (Ekama et al. 1996; Lemmer & Lind 2000).

Large amounts of filamentous microorganisms may build bridges between or around the flocs, which has a negative effect on the settling characteristics of the activated sludge (high sludge volume index (SVI) >150 mL/g). This phenomenon is called filamentous bulking. As a result, many WWTP suffer under bulking-sludge or scum problems, which result in serious operational problems and might cause contamination of the rivers because of overflow of sludge from the WWTP (Casey et al. 1995; ATV-AG 2.6.1 1998; Lebek 2005; Jenkins et al. 2004).

The most common filamentous microorganism, which is responsible for bulking sludge, scum and foam problems in 93% of low load plants with nutrient removal in the north of Europe, is *M. parvicella* (Knoop 1997). The appearance of *M. parvicella* is associated with the following conditions (Slijkhuys 1983; Casey et al. 1995; Eikelboom et al. 1998; Kunst et al. 2000; Kunst 2002; Nielsen et al. 2002; Paris 2004; Rossetti et al. 2004):

- low sludge loading (0.05–0.2 kg of biochemical oxygen demand per kg of total suspended solids per day (kg BOD₅/(TSS·d))
- low temperatures (5–15 °C)
- large quantities of oil and grease
- the change between aerobic, anoxic and anaerobic phases

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• low oxygen concentration (less than 1 mg/L O2)
• recirculation of process water from anaerobic sludge treatment
• plants with primary clarifier.

There are many different studies about the methods to control the growth of *M. parvicella* in WWTP, but not all were successful. All studied methods can be summarized by the following points (Jenkins et al. 1993, 2004; Ekama et al. 1996; Knoop 1997; Kunst et al. 2000; Lemmer & Lind 2000; Kunst 2002; Lebek & Rosenwinkel 2002; Lebek 2003; Paris 2004):

• increasing the sludge load of 0.2 kg BOD5/(kg TSS · d) could have inhibiting effect on the growth of *M. parvicella*
• dosage of iron chloride (low impact on *M. parvicella*)
• dosage of acidic aluminum components (the most successful way)
• application of aerobic, anoxic and anaerobic selector (has no influence on the growth of *M. parvicella*).

Although many scientists such as Knoop (1997), Lemmer & Lind (2000), Kunst et al. (2000), Kunst (2002), Lebek (2003) and Jenkins et al. (2004) mentioned that low sludge loads 0.05–0.2 kg BOD5/(kg TSS · d) favor the growth of *M. parvicella*, studies about the influence of the very low sludge load <0.05 kg BOD5/(kg TSS · d) on the growth of *M. parvicella* are still limited. These low loads correspond to plants with simultaneous aerobic sludge stabilization (Wanner et al. 1998; Lebek 2003).

In order to define the effect of low sludge loads (<0.05 kg BOD5/(kg TSS · d)) on the growth of *M. parvicella*, 13 SBRs in Germany with simultaneous aerobic sludge stabilization were studied.

According to the batch process in the SBR, the substrates can be stored by the flocc-forming and most of the filamentous bacteria in the form of glycogen and acids. There are two groups of these bacteria: high F/M bacteria (high food/microorganism ratio) and low F/M bacteria (Lemmer & Lind 2000). The high F/M bacteria such as *Sphaerotilus natans* and type 021 N can store the substrates as reserve materials only in limited amount. This causes a reduction in frequency of these high F/M bacteria in low sludge load plants. They are replaced by low F/M bacteria such as *M. parvicella*.

Most of the investigated SBRs are working with the differential internal cycle strategy sequential batch reactor method (DIC-SBR) (Figure 1). This method was developed by the company LimnoTec in Lübbecke, Germany, in order to optimize the wastewater treatment. Only 20 SBRs in Germany are operating with this method. The difference between this method and a conventional SBR is characterized by two fillings during one cycle with water having different C/N ratios. The different C/N ratios are adjusted in a multi-functional buffer reservoir before the SBR. The water in the buffer is separated from the solid particles by sedimentation. This supernatant water has a low C/N ratio (which can be called thin water) and a low amount of solids. During the first filling the thin water is applied to the reactor to an amount of 60–90% of the total filling amount during the cycle. During this filling anoxic and anaerobic zones exist, which encourage phosphorus removal and denitrification, if nitrate exists. Before the second filling the water in the buffer is mixed. The rest in the buffer (about 20% of the total amount) is characterized by a high C/N ratio. During the second filling the rest of the nitrate is denitrified (Rönner-Holm & Holm 2009; SH + E GROUP 2012).

It was proved that this method has a lot of advantages such as improvement in the elimination of nitrate and phosphorus, high flexibility and the opportunity of longer charging times.

**MATERIALS AND METHODS**

Thirteen SBRs in Germany (1,000–25,000 PE) were investigated during cold and dry weather conditions. From each plant just two samples from two locations were taken. One sample was taken from the plant’s inflow to measure...
BOD$_5$. The second sample was taken during the second aeration phase to measure SVI and to be investigated under a microscope. Unfortunately, the duration of the project was too short for taking another sample. But the results were compared with the results that were measured in the plant, in order to validate them.

The main task was the evaluation of the operation diaries of these plants and the identification of *M. parvicella* to give a statement about the best operation of SBR and the influence of the very low sludge loads on this bacterium.

**Microscopic investigations**

For microscopic investigations a Zeiss microscope type Axiolab was used. By using one drop of crystal violet solution on air-dried smear preparation the total filament abundance (filament category) was determined according to the seven total filament categories (0, 1, 2, 3, 4, 5, 6) (*Jenkins et al.* 2004). The microscopic investigation at 100× magnification using darkfield microscopy shows violet flocs and orange filaments (*Knoop & Kunst* 1998).

Some filamentous organisms can clearly be identified according to their morphology, mobility, branching, thread form, attached growth, length and width, appearance of the cells, etc. (*Schmid-Schmieder* 2006; *Remde* 2010). Some organisms are difficult to be identified by their morphology; in these cases Gram and Neisser staining have to be used.

By using Gram und Neisser staining the frequency of *M. parvicella* was determined according to Table 1. *M. parvicella* has a strong Gram-positive staining reaction, Neisser-positive granules, coiled growth, no branching and no movement. Its filaments are between 50 and 500 μm in length and 0.8 μm in width. They can be found inside the floc, surrounding the floc or dispersed (see Figure 2) (*Eikelboom 1975; Knoop 1997; Lemmer & Lind 2000; Paris 2004; Jenkins et al. 2004*).

**Calculation of sludge load**

To investigate the influence the low sludge load (<0.05 kg BOD$_5$/kg TSS · d) has on *M. parvicella*, the sludge load was calculated in every investigated SBR over some years using the following formula (DWA-M 210 2009)

\[
B_{TS,BOD} = \frac{B_{LBOD,in} \cdot (n \cdot V_R \cdot TSS_R)}{C_1 \cdot TSS_R} \cdot \frac{t_Z}{t_R}
\]

\[ (kg \text{ BOD}_5/(kg \text{ TSS} \cdot \text{d})) \]

$B_{LBOD,in}$: daily BOD$_5$ load in the input of SBR (kg/d), $n$: number of SBR basins (-), $V_R$: maximum volume of SBR (m$^3$), TSS$_R$: dry suspended solids content in relation to the maximum volume of the SBR (g/L), $t_Z$: cycle duration (h), $t_R$: duration of reaction phase (h).

**Table 1 | Determination of the frequency of one filament type** (*Jenkins et al.* 2004)

<table>
<thead>
<tr>
<th>Frequency (-)</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>No filaments were observed</td>
</tr>
<tr>
<td>1</td>
<td>Few filaments from the same type were observed</td>
</tr>
<tr>
<td>2</td>
<td>Some filaments from the same type were observed</td>
</tr>
<tr>
<td>3</td>
<td>Common filaments from the same type were observed</td>
</tr>
<tr>
<td>4</td>
<td>Very common filaments from the same type were observed</td>
</tr>
<tr>
<td>5</td>
<td>Plenty of filaments from the same type were observed</td>
</tr>
<tr>
<td>6</td>
<td>Excessive filaments from the same type were observed</td>
</tr>
</tbody>
</table>

**Figure 2 | M. parvicella in the plant Z:** (a) Gram-positive; (b) Neisser-positive granules; ×1000 enlargement.
According to the development of the sludge load over years and the frequency of *M. parvicella*, it is possible to determine the impact of the low sludge load (<0.05 kg BOD₅/(kg TSS · d)) on the growth of *M. parvicella*.

**RESULTS AND DISCUSSION**

SBRs are known to have SVI <150 mL/g with very compact flocs (Schreff & Hilliges 2013). The development of the SVI due to temperature in some SBRs such as in plant A (see Figure 3) shows that SVI can reach higher values in winter (>150 mL/g) and floating sludge may occur. The investigation under the microscope of activated sludge in plant A shows that *M. parvicella* is dominant.

In order to optimize the operation of these plants, 13 SBRs with aerobic sludge stabilization were investigated and analyzed and the frequency of *M. parvicella* was determined. All examined plants are treating municipal wastewater. Because of the operation with simultaneous aerobic sludge stabilization, these SBRs do not have a primary clarifier. The investigations of Lebek (2003) indicated that 10% of the long-chain fatty acids in urban wastewater are not dissolved. The pre-treatment in the primary clarifier is used to hydrolyze the fats, which improves the growth of *M. parvicella*. Due to the absence of primary clarifiers, the frequency of this bacterium in the systems with aerobic sludge stabilization could be lower. However, the investigations in some SBRs showed that this was not always the case (Helmreich et al. 2000).

Unfortunately, SBRs in Germany do not have any microscopic investigations in the plants. For this reason, as mentioned in ‘Materials and methods’, samples were taken during the second aeration phase in each investigated plant to measure SVI and to determine the existence of *M. parvicella* under the microscope. The self-measured SVI values, which are shown in Table 2, were compared with the plants’ measurements (operation diaries) and the results were very similar. In order to determine the impact of the sludge load on the growth of *M. parvicella*, the sludge load was calculated according to Equation (1). For this calculation the data for a minimum of 4 years were taken from the operation diaries. The results are shown in Table 2.

Table 2 shows that *M. parvicella* exists in all investigated SBRs, but in different frequencies and it can grow in plants with sludge loads more than 0.05 kg BOD₅/(kg TSS · d) and less than 0.05 kg BOD₅/(kg TSS · d) as well.

**Comparison of SBRs according to operational process**

Table 2 shows that *M. parvicella* can grow in most SBRs with different operations. All these SBRs, except F and U, are operated as DIC-SBR.

In SBR F wastewater is fed continuously during the cycle time. In plant U there is no buffer and wastewater is fed parallel to both reactors. The cycle time in this plant is always 6 h and does not depend on the influent amount. The reaction phase is 3 hours; the sedimentation and decanting phase is 3 hours too. During the sedimentation phase in the first tank the water is charged to the second one.

**Comparison of SBRs according to the sludge load**

The frequency of *M. parvicella* in SBRs which permanently have low sludge load <0.05 kg BOD₅/(kg TSS · d) is 1–4,
except for plant U. It is less than in plants with higher loads of >0.05 kg BOD$_5$/kg TSS · d (frequency = 3–6). Although the plant U works with sludge load less than 0.05 kg BOD$_5$/kg TSS · d, it still has a high frequency of *M. parvicella* and nocardioforms as well. Plant U shows very high SVI: >200 mL/g.

**Comparison of SBRs according to the precipitant**

However, the growth of *M. parvicella* can also be influenced by the used precipitant (see Table 2). The aluminum salts have different effects on *M. parvicella*. In SBR N the dosing of FeCl$_3$ over the complete year reduces the amount of long-chain fatty acids, and aluminum salts are also dosed in winter when *M. parvicella* begins to occur. This variation shows the best effect on *M. parvicella* and the other filaments. This fact corresponds to the investigations of Lebek (2003).

At the same time the sodium aluminate has obviously no influence on this bacterium (plants P, R and S). This trend matches with the results of Misera (2002) and Lebek (2003). On the other hand, in low load plants iron salts show a good influence on *M. parvicella* but these salts cannot completely remove this bacterium from the plant (plants A, B, H and O) (Bidder 1999; Lebek 2003).

The dosing amount of the precipitant is variable. It depends on the phosphorus load in the input of the SBR. In order to give a better statement about the recommended precipitant which should be used to optimize the SBR, the dosing amount of the previous named precipitants was determined depending on the 85% value of TSS and the active substance content (see Tables 3 and 4).

Table 3 indicates that the dosage of 0.28 g Fe/(kg TSS · d) during both aeration times has the best effect against the growth of *M. parvicella* (plant H). A higher dosage amount (1.0 g Fe/(kg TSS · d)) during the second aeration can have good results too (plant O). On the other hand, Table 4 shows that high dosage amount of sodium aluminate (during the second aeration) can affect the growth of this bacterium.

### Table 2 | Sludge load, SVI, used precipitant, filament category and frequency of *M. parvicella* in the investigated SBRs

<table>
<thead>
<tr>
<th>Plant</th>
<th>Sludge load (kg BOD$_5$/kg TSS · d)</th>
<th>SVI (mL/g)</th>
<th>Precipitant</th>
<th>Filament category</th>
<th>Frequency of <em>M. parvicella</em></th>
<th>PE</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>&lt;0.05</td>
<td>96</td>
<td>FeCl$_3$</td>
<td>2–3</td>
<td>2–3</td>
<td>5,000</td>
</tr>
<tr>
<td>H</td>
<td>&lt;0.05</td>
<td>72</td>
<td>FeCl$_3$</td>
<td>2–3</td>
<td>1–2</td>
<td>10,000</td>
</tr>
<tr>
<td>C</td>
<td>&lt;0.05</td>
<td>79</td>
<td>–</td>
<td>2</td>
<td>2</td>
<td>4,500</td>
</tr>
<tr>
<td>D</td>
<td>&lt;0.05</td>
<td>74</td>
<td>NaAl(OH)$_4$</td>
<td>1–2</td>
<td>2–3</td>
<td>5,000–10,000</td>
</tr>
<tr>
<td>P</td>
<td>&lt;0.05</td>
<td>90</td>
<td>NaAl(OH)$_4$</td>
<td>3</td>
<td>3–4</td>
<td>16,000</td>
</tr>
<tr>
<td>B</td>
<td>0.03–0.072</td>
<td>74</td>
<td>FeCl$_3$</td>
<td>2–3</td>
<td>3</td>
<td>9,200</td>
</tr>
<tr>
<td>O</td>
<td>0.03–0.1</td>
<td>86</td>
<td>FeCl$_2$</td>
<td>2</td>
<td>1–2</td>
<td>16,000</td>
</tr>
<tr>
<td>S</td>
<td>0.05–0.12</td>
<td>122</td>
<td>NaAl(OH)$_4$</td>
<td>3–4</td>
<td>4</td>
<td>22,500</td>
</tr>
<tr>
<td>Z</td>
<td>0.006–0.09</td>
<td>142</td>
<td>–</td>
<td>4–5</td>
<td>5</td>
<td>5,200</td>
</tr>
<tr>
<td>N</td>
<td>–$^a$</td>
<td>86</td>
<td>aluminum salts + FeCl$_3$</td>
<td>2</td>
<td>1</td>
<td>7,850</td>
</tr>
<tr>
<td>F$^b$</td>
<td>–$^a$</td>
<td>129</td>
<td>(Al$_2$(SO$_4$)$_3$)</td>
<td>3</td>
<td>3</td>
<td>4,000</td>
</tr>
<tr>
<td>U$^b$</td>
<td>&lt;0.05</td>
<td>259</td>
<td>FeCl$_3$</td>
<td>5</td>
<td>4–5</td>
<td>8,200</td>
</tr>
</tbody>
</table>

$^a$No information available.

$^b$Not working according to DIC-SBR (see ’Comparison of SBRs according to operational process’).

### Table 3 | Comparison of DIC-SBRs which use ferrous precipitants

<table>
<thead>
<tr>
<th>Plant</th>
<th>Precipitant</th>
<th>Fe content (g Fe/kg)</th>
<th>Dosing amount (g Fe/(kg TSS · d))</th>
<th>Frequency of <em>M. parvicella</em></th>
<th>Notice</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>FeCl$_3$</td>
<td>135$^a$</td>
<td>0.13</td>
<td>2–3</td>
<td>Dosing in buffer</td>
</tr>
<tr>
<td>B</td>
<td>FeCl$_3$</td>
<td>135$^a$</td>
<td>0.26</td>
<td>3</td>
<td>Dosing after the second aeration</td>
</tr>
<tr>
<td>H</td>
<td>FeCl$_3$</td>
<td>123</td>
<td>0.28</td>
<td>1–2</td>
<td>Dosing during the first and second aeration</td>
</tr>
<tr>
<td>O</td>
<td>FeCl$_2$</td>
<td>85</td>
<td>1.0</td>
<td>1–2</td>
<td>Dosing after the second aeration</td>
</tr>
</tbody>
</table>

$^a$Fe content was calculated according to ATV-DVWK A 202 E (2004).
The microscopic investigations show that flocs are still compact when using iron salts; in contrast, the flocs in SBR with sodium aluminate are more disrupted. Figure 4 shows that the filament category of the activated sludge in the plant O is 2. The iron salt does not affect the floc morphology and it has a good influence against the other filamentous bacteria. That is not the case in plant S, which uses sodium aluminate.

According to the results in Table 2, it can be concluded that the operation of DIC-SBR is better than the other SBRs. The outflow quality of plant F may suffer because of the continuous filling. The mode of operation of the plant U encourages the growth of M. parvicella and nocardioforms and the dosage of FeCl₃ is not effective in this plant.

In addition to that, low sludge load of <0.05 kg BOD₅/(kg TSS · d) could have negative effect on the growth of M. parvicella. To prove this theory, one SBR with high load and high frequency of M. parvicella was chosen and the operation of this plant was converted (plant S).

### Characteristics of SBR S

Plant S operates as a DIC-SBR. The elimination rates of BOD₅, NH₄-N and Pₜₐₒₜ in plant S are usually >80% (see Table 5). The change in SVI in 2010 due to the temperature is shown in Figure 5. It shows that SVI increased in winter, while it decreased in summer. This development is a sign that M. parvicella is dominant in activated sludge (Knoop 1997).

Figure 6 shows the summation curve of sludge load in SBR S over 8 years (2005–2012). Although, SBR S is operated with simultaneous aerobic sludge stabilization, the sludge load in this plant is more than 0.05 kg BOD₅/(kg TSS · d) (>75%). The 85% percentile of sludge load is 0.1221 kg BOD₅/(kg TSS · d).

The aforementioned results were calculated according to the operation diaries of the plant S. The microscopic investigations of the sludge in SBR S show that the filament category is 3 and the frequency of M. parvicella is 4.
Reducing the sludge load in SBR S

To study the influence of low sludge load on the growth of *M. parvicella* in plant S, TSS was increased from 2.5 g/L to a value more than 4 g/L. This caused the reduction of the sludge load to a value less than 0.05 kg BOD$_5$/(kg TSS · d). This investigation was carried out in the period 01.10.2012–06.12.2012. Within the conversion period, two samples were taken once a week (see ‘Materials and methods’). TSS, SVI and sludge load were self-determined.

Influence of sludge load <0.05 kg BOD$_5$/(kg TSS · d) on SVI

During the conversion process (reducing the sludge load), it was noticed that SVI increased (see Table 6 and Figure 7). Actually, the temperature was decreasing in this period too (from 14 to 10 °C) (see Table 6). This could provide a more suitable environment for the growth of *M. parvicella*.

But at the beginning of year 2012 the temperature was very low (<10 °C) and in this period SVI was better than the values during the conversion (see Figure 8 and Table 6). This indicates that both the temperature and low sludge load affected the sludge negatively and led to high SVI.

The elimination rates of BOD$_3$, NH$_4$-N and total phosphorus (P$_{total}$) were not affected by the conversion (see Figure 9). The main problem during this conversion was the appearance of floating sludge.

A few weeks after the raising of biomass concentration started, it was noticed that floating sludge appeared (see Figure 10). The risk of sludge overflow in the case of...
strong rain was high, which could lead to deterioration of the outflow quality. For this reason and because of the risk that a further reduction of the sludge load might cause a further increase of SVI, the conversion process was stopped after 10 weeks.

The aim of the operational conversion was to determine the influence of the very low load sludge (<0.05 kg BOD₅/(kg TSS·d)) on the growth of *M. parvicella*. Figures 11 and 12 show that the frequency of *M. parvicella* increased by decreasing the sludge load to a value <0.05 kg BOD₅/(kg TSS·d), and the filament category as well increased from 3 to 4 which led to higher SVI. The effluent quality of plant S was deteriorated and also floating sludge problems appeared.

Reducing the sludge load in SBR S showed that *M. parvicella* can grow in low loads of <0.05 kg BOD₅/(kg TSS·d). Sodium aluminate could not affect this bacterium. Table 7 shows a comparison between SBR S (22,500 PE) and the other DIC-SBR plants which have sludge load <0.05 kg BOD₅/(kg TSS·d) and phosphorus is eliminated biologically or by using sodium aluminate. Small plants (C, D and P) (1,000–10,000 PE) have fewer problems than the bigger ones (R and S). This may be due to the existence of higher concentration of long-chain fatty acids in bigger plants because of the longer flow times to the more centrally located WWTP. In this case it is recommended to increase the sludge load to over 0.05 kg BOD₅/(kg TSS·d) or to use suitable precipitants such as iron salts or acidic aluminate precipitants (PAC or AlCl₃). However, these aluminum precipitants must be used continuously (Paris 2004), which results in very high costs. For this reason, using iron salts is recommended in these plants during the regular operation. This causes improvement of SVI and reducing the other filamentous bacteria. In the case of using sodium aluminate and the appearance of filamentous bulking the dosing amount should be raised.

**Subdominant filamentous bacteria in SBR**

The microscopic investigations show that in addition to the growth of *M. parvicella* in SBR, many other filamentous bacteria, which can cause scum and bulking problems, are present and appear subdominant. The most common ones are type 0041/0675, type 0581, type 0914, *Haliscomenobacter hydrossis*, and in some plants also the sulfate bacteria (type 021N and *Thiothrix* sp.). The filling with low chemical oxygen demand content encourage the growth of filamentous bacteria (Chudoba et al. 1973). Figure 13 shows some of these bacteria in the investigated SBRs. The favorable conditions which encourage the growth of these filaments are different,
but they still have the following connected factors (Kunst et al. 2000; Lemmer & Lind 2000):

- low oxygen concentration
- low sludge load \(<0.2 \text{ kg BOD}_5/(\text{kg TSS} \cdot \text{d})\)
- unsettled C:N:P ratio.

DIC-SBR provided oxygen until the concentration in the water is 2 mg/L O_2. Then the aeration stops until the concentration becomes \(<1 \text{ mg/L O}_2\). This offers a suitable environment for many of these bacteria to grow in these plants. Kunst et al. (2000), Lemmer & Lind (2000), Jenkins et al. (2004) and Remde (2010) mentioned that the optimization of the aeration in the plants can prevent the growth of these bacteria.

**Influence of the long cycle time**

In the majority of the investigated DIC-SBRs the cycle time is 8 h. In SBR O it is 12 h. Table 8 shows a comparison between the occurrence of *M. parvicella* in plant O and the other plants which use precipitants with iron content. Plant O shows low frequency of *M. parvicella* and a lower filament category.

Gabb et al. (1996) showed that a long aeration phase reduces the frequency of *M. parvicella*. The total sludge
age in the investigated SBRs is 25 d. In DIC-SBR with 8 h cycle time the reactor should be aerated in the first filling for about 2.5 h and in the second filling for 1 h. This means it is aerated for about 3.5 hours per cycle according to the ammonium concentration. In this case the aerated sludge age is about 11 d (44% of the total age). In DIC-SBR with 12 h cycle time the aeration time is more (aerated sludge age is about 50% of the total age).

In the majority of the investigated SBRs in Germany the reactor is aerated until the dissolved oxygen

![Figure 12](https://iwaponline.com/wst/article-pdf/69/10/1984/471410/1984.pdf)

**Figure 12** Development of the frequency of M. parvicella during the conversion in plant S.

<table>
<thead>
<tr>
<th>Plant</th>
<th>Sludge load (kg BOD₅/(kg TSS · d))</th>
<th>Precipitant</th>
<th>Frequency of M. parvicella</th>
<th>PE</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>&lt;0.05</td>
<td>–</td>
<td>2–3</td>
<td>3,000</td>
</tr>
<tr>
<td>D</td>
<td>&lt;0.05</td>
<td>–</td>
<td>2</td>
<td>4,500</td>
</tr>
<tr>
<td>P</td>
<td>&lt;0.05 NaAl(OH)₄</td>
<td>2–3</td>
<td>5,000–10,000</td>
<td></td>
</tr>
<tr>
<td>R</td>
<td>&lt;0.05 NaAl(OH)₄</td>
<td>3–4</td>
<td>16,000</td>
<td></td>
</tr>
<tr>
<td>S</td>
<td>&lt;0.05 NaAl(OH)₄</td>
<td>4–5</td>
<td>22,500</td>
<td></td>
</tr>
</tbody>
</table>

*aDuring the conversion.*

![Figure 13](https://iwaponline.com/wst/article-pdf/69/10/1984/471410/1984.pdf)

**Figure 13** Filamentous bacteria in different SBRs: (a) Haliscomenobacter hydrossis; (b) type 0041/0675 (original magnification, ×100); (c) type 0581; (d) type 0914 (Gram staining, ×1000).
concentration (DO) is 2 mg/L O₂ and then it is stopped. In order to optimize the operation of these plants longer cycle times and longer aeration phases can be used (such as in plant O). It is recommended that the reactor can be aerated until DO reaches 1 mg/L O₂ and for a longer time (aerated sludge age >44%) until the nitrification is completed. This means more energy costs for the aerator, but at the same time the frequency of bulking filaments can be reduced (D. Schreff, Office for Water, Wastewater and Energy, Germany, personal communication).

**CONCLUSION**

From the aforementioned discussion, it can be concluded that *M. parvicella* can grow in plants with nitrogen and phosphorus removal in all zones (aerobic, anaerobic and anoxic) and low sludge loads. Many studies showed that the optimal sludge load for *M. parvicella* is 0.05–0.1 kg BOD₅/(kg TSS · d). To study the influence of very low loads of <0.05 kg BOD₅/(kg TSS · d) on *M. parvicella* 13 SBRs with simultaneous aerobic sludge stabilization in Germany were investigated. *M. parvicella* was found with different frequencies in most of the plants in addition to other filaments. These filamentous bacteria can cause scum and bulking sludge under some conditions.

The conversion of plant S showed that *M. parvicella* can grow in low loads of <0.05 kg BOD₅/(kg TSS · d). Its frequency is higher in plants with 10,000–25,000 PE because of the existence of higher concentration of long-chain fatty acids. Due to the high costs of PAC or AlCl₃ the use of iron salts in these plants is recommended. The dosing of the precipitant should be carried out during the aeration, the dosing amount can vary between 0.28 and 1 g Fe/(kg TSS · d). This can lead to better SVI, low filament category and low frequency of *M. parvicella*.

Additionally, even though DIC-SBR shows in comparison to the other types of SBR better nitrogen elimination, biological elimination of phosphorus and good effluent quality, the filling with different C/N ratios can improve the development of other filamentous bacteria such as type 0041/0675, type 021 N and type 0581. This can be considered as a drawback for DIC-SBR. To prevent the growth of these filaments, longer cycles of 8–12 h and longer aeration times are recommended.

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