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# Tangential flow filtration (TFF) concentration of phytoplankton Version 3

#### **Daniel Vaulot**

## **Abstract**

Concentrate phytoplankton samples about 100-fold typically from 5L down to 20 mL. Takes about 1 hour per sample. Samples can be used for flow cytometry sorting or for cultures. Enrichment by TFF usually keep growing for a longer time than unconcentrate samples.

#### Reference

Marie, D., Shi, X.L., Rigaut-Jalabert, F. & Vaulot, D. (2010). Use of flow cytometric sorting to better assess the diversity of small photosynthetic eukaryotes in the English Channel. *FEMS Microbiology Ecology*. 72. p.pp. 165–178.

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## **Guidelines**

In order to estimate actual concentration efficiency measure Synechoccus, pico, and nanoeularyote concentration before and after TFF.

## **Before start**



# **Necessary equipement**

- Vivaflow Cartridge 100,000 MWCO (Regenerated Cellulose RC) VF20C4 for viruses
- Vivaflow Cartridge 0.2 μm (PES) VF20P7 for plankton
- Masterflex Pump 6-600 rpm (ref Bioblock F39671) It is critical to have a 600 rpm pump, lower speed will not work. The rate of the pump must be adjustable.
- Rotor 3 "galets" (ref F39110)

- 1 Head high throughput (ref F40103) (Can be replaced with quick load head)
- Replace tube provided by stronger tube with two connectors (see picture)
- Bottle 6 L
- Bottle 1 L
- Conical tube 50 mL (Falcon tube)
- Masterflex Tygon tubing size 16
- Plastique pipettes (1 mL) this is used to plunge in the sample
- Clamps with screw (to control retentate speed)

#### **Solutions**

MilliQ water : 1LNaOH 0.1 N : 500 mLEtOJ 10% : 500 mL

## **Materials**

- MilliQ water by Contributed by users
- Filtered Seawater (0.2 μm) by Contributed by users
- ✓ 0.1 M NaOH by Contributed by users
- ✓ Ethanol 10% by Contributed by users

## **Protocol**

## Rinsing cartridge

#### Step 1.

Get Vivaflow catridge out of storage

# Rinsing cartridge

## Step 2.

Mount Vivaflow catridge as Fig. 1 (image de C. Brussaard) in open circuit.

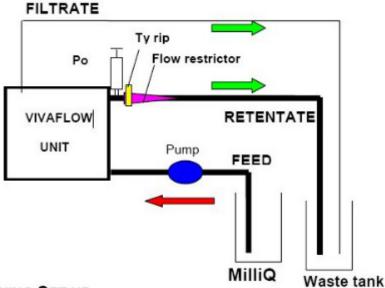


FIG 1. FLUSHING SET UP

## Rinsing cartridge

## Step 3.

Remove the clamps

# Rinsing cartridge

## Step 4.

Set the pump to maximum speed. Manometer should be at about 2.5 bars (with a new cassette sometimes the manometer get stuck, if the value is too low there is a leak in the system)

#### Rinsing cartridge

#### Step 5.

Rinse the cartridge with about 250 mL of MilliQ water (longer is cartridge has been stored in ethanol)



250 ml Additional info:



**REAGENTS** 

✓ MilliQ water by Contributed by users

**O DURATION** 

00:10:00

## Rinse cartridge

## Step 6.

Replace MilliQ water by sample in 6 L bottle

## Rinse cartridge

## Step 7.

Take sample for flow cytometry to compute concentration factor.

# Rinse cartridge

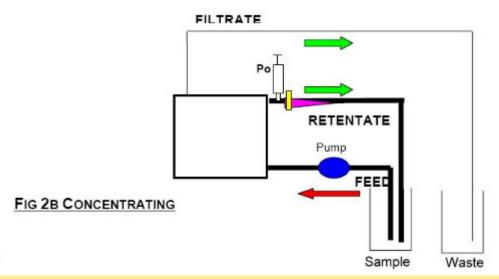
## Step 8.

Rinse cartridge with about 250 mL of sample

#### Concentrate

#### Step 9.

Put the retentate line into the sample bottle (Fig. 2B - Recirculation mode). Record sample volume and start pump increasing to macimum speed.



#### Concentrate

#### Step 10.

Clamp the retentate line to increase filtrate flow so that Manometer gets up to 2.5 bars.

#### Concentrate

#### **Step 11.**

Concentrate sample until about 250 mL remains (6 L takes about one hour)

© DURATION

01:00:00

#### Concentrate

#### **Step 12.**

Transfer sample to a smaller bottle (250 mL) then finally to a 50 mL tube with conical bottom

O DURATION

00:10:00

## Concentrate

#### **Step 13.**

Continue to concentrate very carefully, lowering the pump speed until the sample volume is reduced to 15-20 mL

#### NOTES

#### Daniel Vaulot 08 Dec 2016

It is very important to lower the pump speed inonder to avoid loosing the sample.

#### Recirculation

## Step 14.

When final volume is about 10 mL, clamp filtrate tube and recirculate slowly (no change of volume should take place)

## Recirculation

## **Step 15.**

Leaving the filtrate tube clamped, get the feed line out of the sample in order to get back the total volume of concentrated sample

#### Recirculation

#### **Step 16.**

Take sample for flow cytometry count (to compare with original sample concnetration and estimate concentration factor)

#### Recirculation

#### **Step 17.**

Store concentrated sample for later use (e.g. flow cytometry sorting, culture etc...)

## Rinsing

## **Step 18.**

Go back to Fig. 1 configuration (Open circuit)

#### Rinsing

## Step 19.

Rinse 1 min with filtered sea water



**REAGENTS** 

Filtered Seawater (0.2 μm) by Contributed by users

**O DURATION** 

00:01:00

## Rinsing

#### Step 20.

Rinse 1 min with distilled water



**REAGENTS** 

Distilled Water by Contributed by users

**O DURATION** 

00:01:00

#### Rinsing

#### Step 21.

Rinse with 50 mL NaOH 0.1 M



50 ml Additional info:



**REAGENTS** 

✓ 0.1 M NaOH by Contributed by users

## Rinsing

#### Step 22.

Put all three tubes (feed, retentate, filtrate) in bottle containing NaOH 0.1 M



REAGENTS

✓ 0.1 M NaOH by Contributed by users

# Rinsing

#### Step 23.

Recirculate for 20 min (to get rid of everything on the cartridge filter)

**O DURATION** 

00:20:00

## Rinsing

#### Step 24.

Rinse with 250 mL of MilliQ water (Fig. 1)



250 μl Additional info:



✓ MilliQ water by Contributed by users

#### Storage

# Step 25.

Stop the pump and clamp all three tubes

# Storage

# Step 26.

Store at 4°C. For a storage beyond 1 day, store with Ethanol 10%



✓ Ethanol 10% by Contributed by users