## Harvest of Lemna minor:

## **Principle**

This protocol describes the usual steps taken after an experiment to harvest plant and solution samples for subsequent analysis.

# **Equipment and Reagents**

Machine/Product	Reference (Company, Type,)
Disposable Straining Filters	
5 mm Petri dishes	
Long tweezers	
Disposable plastic weighing tray	
Analytical scale	0.00001 g precision
Pb(NO <sub>3</sub> ) <sub>2</sub> solution	
1 mM (= 331,21 mg in 1 L)	Sigma 467790
Drying oven	Memmert

### **Protocol**

Sampling the growth medium:

Carefully take a 5 ml sample from the pots before moving them.

This is to prevent re-suspension of possible precipitate.

### Pictures:

Take pictures of the pots for frond number and area analysis.

## Harvest:

Remove the Lemna plants using long tweezers

Carefully pat them dry on absorbent paper

Weigh the plants on a balans (total Fresh Weight)

Optional: Subsampling for Pigment analysis

Weigh 2-3 plants (Fresh weight between 10 and 20 mg), record the weight and transfer the plants in Eppendorf microtubes containing 500 µL DMF.

Keep in the dark at 4°C for 24 hours (3 days max) before measuring pigments.

# Washing for elemental analysis:

The plants are transferred to a disposable strainer filter.

Put the filter 50 mm Petri dish containing 25 ml Pb(NO3)2 (1 mM)

Wait for 10 minutes, and occasionally wash the plants using a pipette

Pat dry the filter and transfer it to a 50 mm Petri dish containing 25 ml demin water

Wait for 10 minutes, and occasionally wash the plants using a pipette

Repeat the last rinsing step.

Remove the plants from the filter and pat them dry.

Transfer the plants to a small tarred weighing tray

Immediately weigh on an analytical scale (0.00001 g)

Dry the plants at 60° for a week

The plants are now ready for mineralization (refer to the specific protocol)

pH of the solution Measure the pH of the growth medium using a pH electrode. Optionally, the conductivity of the solution can be recorded too.