

1 **Supplementary**

2 *Tables*

Type of substrates	Substrates	Type of solutions used for the experiment	Type of reactors*	Added <i>M. Aeruginosa</i> (g _{wet} /L)	Number of replicates	Time of experiment
vegetation	lichen	leachate	ER	1	2	16
			CR	none	1	
	moss	leachate	ER	1	2	14
			CR	none	1	
peat	active layer peat	leachate	ER	1	2	29
			CR	none	1	
	permafrost peat	leachate	ER	1	2	16
			CR	none	1	
lake water			ER	1	2	13
			CR	none	1	

3 **Tab. S 1:** General information on the experiment. All those reactors were submitted to the same
4 incubation's conditions, i.e. continuous light exposure, aeration through a porous stopper, a
5 constant temperature of 25 °C and continuous stirring by a magnetic bar. * ER = Experimental
6 Reactor; CR = Control Reactor.

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Substrate		moss	lichen	active layer peat	permafrost peat	lake water
<i>p</i> _value	pH	0.1097	0.2658	0.0006981**	0.07038*	0.01883**
(from	SUVA ₂₅₄	0.01565**	0.3924	0.03103**	0.002041**	0.9499
student test)	DOC	0.4009	0.2352	0.03318**	0.003663**	0.07593*

8 **Table S 2:** *p*_value of student test run between values of pH, SUVA₂₅₄ and DOC of experimental
9 and control reactors for each organic substrate. *The difference is significant with a confidence
10 interval of 90%. **The difference is significant with a confidence interval of 95%.

Z8 broth solution fabrication:

Step1: Stock solution Z8I		Step2: Stock solution Z8I	
Stock solution Z8I	Mass (g) for 1L*	Stock solution Z8I	Mass (g) for 1L*
NaNO ₃	46.70	K ₂ HPO ₄	3.10
Ca(NO ₃) ₂ ·4H ₂ O	5.90	Or K ₂ HPO ₄ ·3H ₂ O	4.10
Or Ca(NO ₃) ₂	4.10	Na ₂ CO ₃	2.10
MgSO ₄ ·3H ₂ O	2.50		

*add autoclaved MilliQ water (20min at 121 °C) to reach 1L

Step3: Stock solution Z8III		Solution B	
Solution A	Mass (g) to dissolve in 100 ml of HCl (0.1 M)	Solution B	Mass (g) to dissolve in 100 ml of HCl (0.1 M)
<u>Iron solution:</u> FeCl ₃ ·6H ₂ O	2.80	<u>EDTA solution:</u> EDTANa ₂ ·2H ₂ O	3.70

For 1L of stock solution Z8III mix 10 mL of solution A (iron solution), 10 mL of solution B (EDTA solution) and 980 mL of MilliQ water (pre-autoclaved at 121 °C for 20min)

Step4: Trace solution		
N°	Stock solution for trace solution	Mass (g) for 100 ml
1	Na ₂ WO ₄ ·2H ₂ O	0.33
2	(NH ₄) ₆ MO ₇ O ₂₄ ·4H ₂ O	0.88
3	KBr	1.20
4	KI	0.83
5	ZnSO ₄ ·7H ₂ O	2.87
6	Cd(NO ₃) ₂ ·4H ₂ O	1.55
7	CO(NO ₃) ₂ ·6H ₂ O	1.46
8	CuSO ₄ ·5H ₂ O	1.25
9	(NH ₄)Ni(SO ₄) ₂ ·6H ₂ O	1.98
10	Cr(NO ₃) ₃ ·9H ₂ O	0.41
11	KAl(SO ₄) ₁₂ H ₂ O	4.74
12	VO ₂ SO ₄	0.080
13	H ₃ BO ₃	3.1
	MnSO ₄ ·H ₂ O	1.6

For 1L of Trace solution, mix stock solutions as follow:

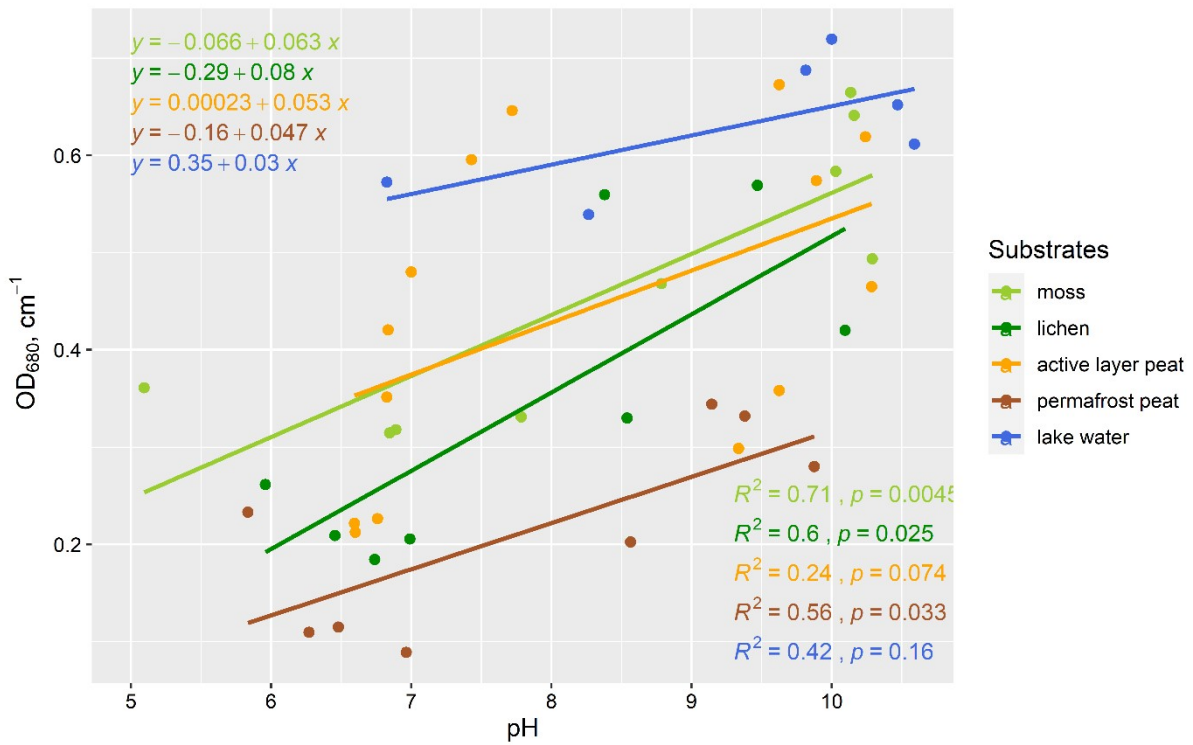
N° stock solution	Volume (ml) for 1L*
1	1
2	1
3	1
4	1
5	1
6	1
7	1
8	1
9	1
10	1
11	1
12	1
13	10

*add autoclaved MilliQ water (20min at 121 °C) to reach 1L

Step5: Final Z8 solution				
N° stock solution	Z8I solution	Z8II solution	Z8III solution	Trace solution
Volume (ml) for 1L*	10	10	10	1

*add autoclaved MilliQ water (20min at 121 °C) to reach 1L

optimal pH~7.5-7.6, can be adjusted before sterilization (autoclaved at 121 °C for 20 min)

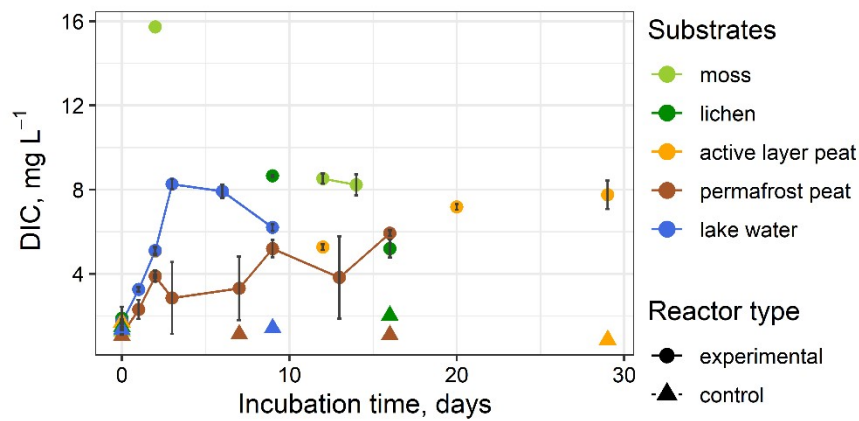


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15 **Fig. S 2:** Correlation between pH and optical density at 680nm (biomass' proxy) for each
 16 substrate.

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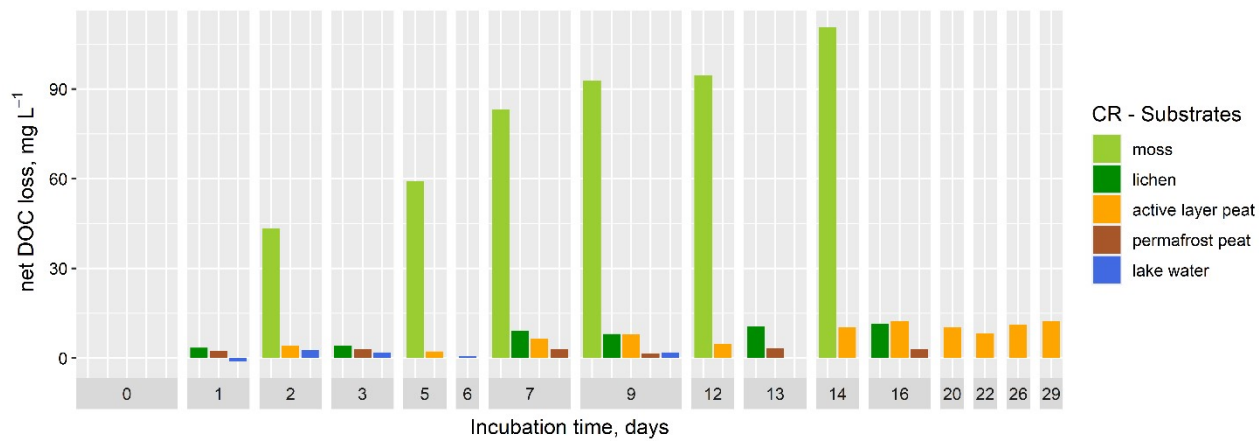


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20 **Fig. S 3:** Dissolved inorganic (DIC) carbon concentrations over the experiments. The absence
 21 of data indicates a non-detectable amount of DIC (i.e. < 1mg.L-1).

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25 **Fig. S 4:** Net DOC loss for all CR during the incubation relative to DOC at day 0 of incubation.