Electronic Supplementary Material (ESI) for Lab on a Chip. This journal is © The Royal Society of Chemistry 2021

Supplementary Information

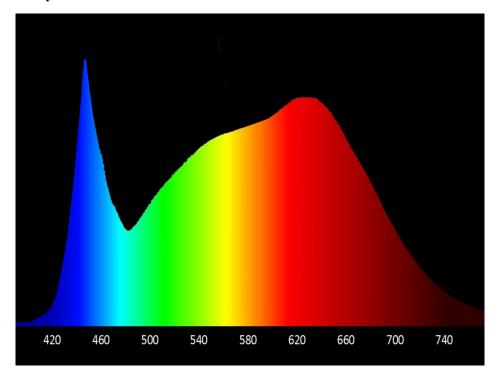


Figure S1. Spectrum of the 45W LED-light lamps used for the growth experiment of the control PBR batch cultures.

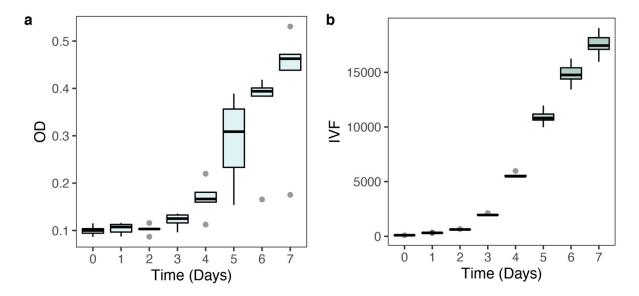


Figure S2. Box-plot representation of the within-well variability over a 7-days' time period. Results presented here are measurements from one randomly chosen well (B6) used as an example. Both optical density (A) and in vivo chl a fluorescence (B) were measured by using the multiple reads per well option where 5 different measurements points were performed for each well. The five measurements are represented by the filled circles.

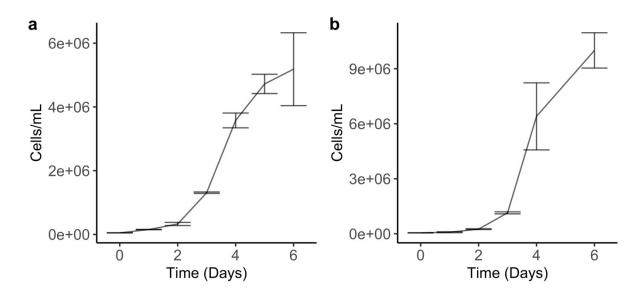


Figure S3. Growth curves of control cultures of P. tricornutum acclimated to low light (~ 20 μ mol photons $m^{-2}s^{-1}$) (a) and high light (~ 200 μ mol photons $m^{-2}s^{-1}$) (b) at 16 °C \pm 1. Growth measurements were done by cell counting by using the BD Accuri C6 Flow Cytometer (BD Bioscience) every 24h for 7 consecutive days (including day 0). Results are presented as a mean of three biological replicates \pm SD.

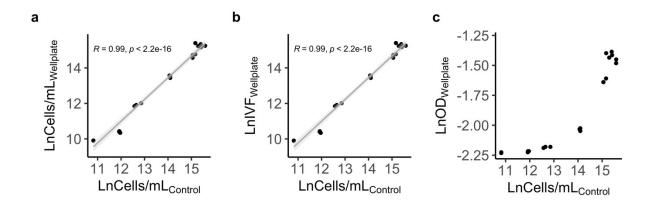


Figure S4. Scatter plot and best-fitting regression line illustrating the relationships between growth measurements performed in the well plate setting versus the control (Batch culture: 125 mL). (a) Cell count/mL_{Control} vs. Cell count/mL_{Wellplate}; (b) Cell count/mL_{Control} vs. IVF_{Wellplate}; (c) Cell count/mL_{Control} vs. OD_{Wellplate}, Regression line is only drawn for (a) and (b). Grey surrounding area: 95% confidence belt for the fitted line.

Table S1. Table summarizing the specific growth rate (μ, d^{-1}) estimates based on the measurement performed. Results are presented for the quantification methods performed in the 96 well plate versus: control (a): represented by a standard batch PBR illuminated with the Nanocosm LED-light board and control (b): batch PBR illuminated with another LED light source (spectrum Figure S1). Values presented are from calculation performed between day 2 and 3 when growth is known to be highest.

Light source	Nanocosm			Nanocosm	algae growth room
Culturing method	96 well plate			Batch PBR	Batch PBR
Biomass quantification method	Cell count	IVF	OD	cell count	Cell count
Specific growth rate (μ, d-1)	1.56 d ⁻¹	1.45 d ⁻¹	0.14 d ⁻¹	0.97 d ⁻¹	1.52 d ⁻¹
$ln(x_t/x_{t-1})$	- 133 %				-1.5 - 4.