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Hydrophobicity Protocol

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¹In-house protocol

1 Works for me

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ABSTRACT

P-Xylene Hydrophobicity Assay

Note that I have written this before doing it. Changes most likely will be made.

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ABSTRACT

P-Xylene Hydrophobicity Assay

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Steps

1 Grow 10mls OG1RF in BHI using short-term or long-term supplementation

2	At 0.3 OD for long-term or 30 minutes after supplementation for short-term, spin cells and wash twice using PUM buffer
3	Aliquot 1 ml 6 times into glass tubes and mix with a range of P-xylene volumes • 0μl, 12.5μl, 25μl, 50μl, 100μl, and 200μl p-xylene in 1ml PUM buffer
4	Vortex vigorously for 30 seconds or place in a shaker for 2 minutes (for many samples)
5	Incubate static at 37°C for 15 minutes
6	Carefully extract 200µl of the aqueous phase and add to 800µl PUM buffer in a cuvette
7	Record OD ₆₀₀ for each P-xylene xoncentration
8	Present each sample s a percentage of cells in the aqueous phase relative to PUM buffer with no p-xylene
9	OD_{600nm} XµI p-xylene/ OD_{600nm} PUM only = % cells in aqueous phase