

SUPPLEMENTARY MATERIALS

Development and evaluation of culture media based on extracts of the cyanobacterium *Arthrospira platensis*

Kheirabadi, Elaheh.^{1,2}, Macia, Javier.^{1,*}

¹Department of Medicine and Life Sciences, Universitat Pompeu Fabra, Barcelona Biomedical Research Park. Avda. Dr. Aiguader 88, 08003 Barcelona, Spain.

²BioInspired Materials Company, c\Valencia 359, 08009 Barcelona, Spain.

CE COMPOSITION

Sugars		Amino acids	
Fructose	<0.10 g/100 g	Alanine	0.423 g/100 g
Galactose	0.12 g/100 g	Arginine	0.373 g/100 g
Glucose	0.47 g/100 g	Aspartic acid	0.570 g/100 g
Lactose	0.14 g/100 g	Cystein +Cystine	0.0430 g/100 g
Maltose	<0.10 g/100 g	Glutamic acid	0.769 g/100 g
Sucrose	<0.10 g/100 g	Glycine	0.280 g/100 g
		Histidine	0.0823 g/100 g
		Hydroxyproline	<0.2 g/100 g
Carbohydrates	4.6 g/100g	Isoleucine	0.258 g/100 g
		Leucine	0.425 g/100 g
Metals		Lysine	0.279 g/100 g
Calcium (Ca)	2.3 mg/100 g	Methionine	0.0870 g/100 g
Manganese (Mn)	<0.050 mg/100 g	Ornithine	<0.05 g/100 g
Magnesium (Mg)	22 mg/100 g	Phenylalanine	0.217 g/100 g
Zinc (Zn)	<0.050 mg/100 g	Proline	0.238 g/100 g
Potassium (K)	132 mg/100 g	Serine	0.269 g/100 g
		Threonine	0.281 g/100 g
		Thyptophane	<0.1 g/100g

	Tyrosine	0.219 g/100 g
	Valine	0.312 g/100 g

Table S1. Compositional analysis of CE.

LAB PREPARED MEDIA

Tryptic Soy Medium with 5% Defibrinated Sheep Blood

Tryptic Soy Broth (BD 211825) 30.0 g
Sheep Blood (defibrinated)..... 50.0 mL
DI Water..... 950.0 mL

Autoclave medium at 121°C. Cool to ~47°C.

Aseptically add 50 mL of room temperature defibrinated sheep blood.

YGC medium for bacterial cellulose production

Glucose.....50.0 g
Yeast extract.....5.0 g
CaCO₃12.5 g
Agar.....15.0 g
Distilled water.....1.0 L
Autoclave at 121°C for 15 minutes.

Mannitol Agar/Broth

Yeast Extract..... 5.0 g
Peptone..... 3.0 g
Mannitol..... 25.0 g
Agar (if required)..... 15.0 g
DI Water..... 1000.0 mL
Autoclave medium at 121°C.

GENETIC CONSTRUCT FOR GFP EXPRESSION

The plasmid pSB1C3 contains the gene encoding Green Fluorescent Protein (GFP) located downstream of a constitutive promoter. The genetic construct was created from genetic parts from the Registry of Standard Biological Parts using the Biobrick assembly method (<http://parts.igem.org/>). Specifically, the genetic construct was composed of the following genetic parts:

T14 + J23100 + RBS34 + GFP

where:

CODE	Description	Parts Registry ID
T14	Double terminator sequence	B0014
J23100	Strong constitutive promoter from Anderson's collection	J23100
RBS34	Strong Ribosomal Binding Site sequence	B0034
GFP	Green Fluorescence Protein	E0040

Table S2. Genetic components for GFP expression