

Apr 21, 2016

Buffers for Chloroplast Isolation from Diatoms

 In 1 collection

DOI

[dx.doi.org/10.17504/protocols.io.ea2bage](https://doi.org/10.17504/protocols.io.ea2bage)

P. Dreux Chappell¹, Bethany D. Jenkins¹

¹Old Dominion University, University of Rhode Island



P. Dreux Chappell

Old Dominion University, University of South Florida

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Protocol Citation: P. Dreux Chappell, Bethany D. Jenkins 2016. Buffers for Chloroplast Isolation from Diatoms. **protocols.io**
<https://dx.doi.org/10.17504/protocols.io.ea2bage>

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Protocol status: Working

Created: December 15, 2015

Last Modified: March 10, 2018

Protocol Integer ID: 2106

Abstract

Guide for reagent/buffer preparation for protocol to separate chloroplasts from diatom cells using ammonium fluoride to permeate the silica frustule and a percoll gradient to separate the plastid from other cellular components.

Guidelines

Buffers and reagents can be mixed up ahead of time and stored cold (refrigerated). Percoll gradients (layering of 40% and 80% percoll solutions) should be done the same day of the extraction shortly before use.

Isolation Buffer (40 mL)

1 Isolation Buffer (40 mL)

1) Mix:

- 20 ml 1 M sorbitol
- 400 µl 0.6 M Na₂-EDTA
- 200 µl 1 M MgCl₂
- 400 µl 1 M KCl
- 40 µl 1 M MnCl₂
- 2 ml 1 M HEPES-KOH pH = 8.0
- 16.96 ml Sterile H₂O.

2) Add bovine serum albumen (BSA) 1% (w/v) just before use to subset of isolation buffer needed at the beginning of the procedure

PEG-6000 Solution

2 PEG-6000 Solution

Mix:

- 6 g PEG
- 10 ml water

Percoll Solution (20 mL Stock)

3 Percoll Solution (20 mL Stock)

Mix:

- 19 ml Percoll
- 1 ml PEG – 6000 solution
- 0.2 g Ficoll
- 0.2 g BSA

Gradient Mixture (10 mL Stock)

4 Gradient Mixture (10 mL Stock)

Mix:

- 250 µl 1 M HEPES-KOH pH = 8.0
- 1 ml 0.1 M EDTA
- 6.26 ml 1 M sorbitol
- 2.49 ml water

3X Bottom Layer (80% Percoll)

5 3X Bottom Layer (80% Percoll)

Mix:

- 5.7 ml Percoll Solution
- 1.01 ml Gradient Mixture

3X Top Layer (40% Percoll)

6 3X Top Layer (40% Percoll)

Mix:

- 5.04 ml Percoll Solution
- 6.96 ml Gradient Mixture

10M Ammonium Fluoride (NH₄F)

7 10M Ammonium Fluoride (NH₄F)

Mix:

- 3.7 g NH₄F
- 10 ml H₂O