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Determination of Boron Content Using a Simple and Rapid Miniaturized Curcumin Assay

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Abstract

To determine boron quantity in soil, water and biological samples, several protocols are available. Colorimetric assays are the simplest and cheapest methods which can be used to determine boron concentration. However, published protocols do not give straightforward guidance for beginners to adopt these protocols for routine use in the laboratory. Based on a previously published available procedure, we present a detailed and modified version of a curcumin based colorimetric protocol to determine boron concentration extracted from any sample. Our modified protocol is able to determine up to 0.2 nmole of Boron in a sample volume of 300 μ l.

Keywords

Boron; Curcumin method; Protoplasts; *Arabidopsis*; Yeast

Background

Boron (B) can be quantified using spectrometric and colorimetric methods. Inductively coupled plasma mass spectrometry (ICP-MS) is the most sensitive method currently available having a detection limit of 0.01 mg/L (Kmiecik *et al.*, 2016) but requires a sample volume of 5 ml. However, this technique requires sophisticated and expensive equipment which is not affordable for smaller laboratories. Alternatively, colorimetric based assays using curcumin or Azomethine-H dyes can be used for the routine analysis of boron in all laboratories with access to a spectrophotometer that can measure absorbance at 550 nm. Bingham (1982) reported that the curcumin assay is more efficient than the Azomethine-H based assay. Therefore, we present a modified version of simple and rapid curcumin assay to quantify B based on a protocol originally published by Wimmer and Goldbach (1999). We have used this protocol to determine the intracellular boron in yeast cells and *Arabidopsis* protoplasts. Additionally, this protocol can be used to study the uptake, root to shoot translocation-mechanisms of Boron in plants.

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Authors declare no conflict of interest.

Materials and Reagents

Note: As boron leaches from lab glassware, all chemicals should be prepared in plastic bottles.

1. Falcon tubes (15 ml and 50 ml) (SARSTEDT, catalog number: 62.547.254)
2. 1.5 ml tubes
3. Pipette tips
4. 96-Well UV Microplate (Thermo Fisher Scientific, Thermo Scientific™, catalog number: 8404)
5. Boric acid (H_3BO_3) (Sigma-Aldrich, catalog number: 15663)
6. 0.1 N hydrochloric acid (HCl)
7. 2-Ethyl-1,3-hexanediol (Acros Organics, catalog number: 118512500)
8. Chloroform (Fisher Scientific, catalog number: 10122190)
9. Concentrated sulfuric acid (H_2SO_4) (Fisher Scientific, catalog number: 10294300)
10. Concentrated acetic acid (Fisher Scientific, catalog number: 10304980)
11. Curcumin (Alfa Aesar, catalog number: B21573)
12. Methyl-Isobutyl-Ketone (MIBK) (Avantor Performance Materials, MACRON, catalog number: 6247-04)
13. Extraction solution (see Recipes)
14. Acid mixture (see Recipes)
15. Curcumin solution (see Recipes)

Equipment

1. Centrifuge
2. Chemical fume hood
3. Pipette
4. Microplate reader (BMG LABTECH, model: CLARIOstar)

Procedure

1. Harvest the yeast/protoplasts samples in a 50 ml Falcon tube by centrifugation at $4,000 \times g$ (for yeast)/ $100 \times g$ (for protoplast) for 5 min. Wash the pellet with 25 ml ice cold water to remove boric acid from the pellet. Resuspend the pellet in 400 μl of Milli-Q water in a 1.5 ml tube. Heat the samples at 90°C for 30 min and vortex to release intracellular B into solution. Centrifuge at maximum speed

for 2 min at room temperature and collect 300 μ l of supernatant for B quantification.

For detailed boron uptake assay in yeast, users may refer to Takano *et al.*, 2002 ; and to process the protoplasts from plant tissue please refer to Yoo *et al.*, 2007.

Note: We recommend that users test the assay with their own buffers/solvents if they do not use water to solubilise their samples.

2. Prepare standards containing 0, 0.02, 0.05, 0.10, 0.19, 0.39, 0.77 mg boric acid/L using Milli-Q water in a 15 ml Falcon tube (refer to Table 1).
3. Aliquot 300 μ l (which is equivalent to 0.1, 0.2, 0.5, 0.9, 1.9, 3.7 nmole B) of each standard and samples into 1.5 ml tubes.
4. Acidify the samples and standards by adding 100 μ l of 0.1 N HCl. Mix well by vortex and incubate for 5 min at room temperature.

Note: From Step 3 onwards perform all the reactions in chemical fume hood.

5. Add 70 μ l extraction solution (see Recipes), mix vigorously by vortexing for 30 sec. After 2 min, repeat the vortex for another 30 sec.
6. Centrifuge the samples at 15,000 \times g speed for 5 min to facilitate clear separation of organic phase.
7. Pipette 50 μ l of lower organic phase (Figure 1A) into a new 1.5 ml tube containing 200 μ l of acid mix (see Recipes) and mix well by vortexing.

Notes:

- a. *Be aware that the lower phase contains chloroform and that chloroform leaks out from plastic pipette tips. To avoid pipetting errors, make sure that the tip is tightly attached to pipette and transfer liquids quickly.*
 - b. *The mixture of sulphuric acid and acetic acid is a viscous solution. Therefore, care must be taken while pipetting to avoid pipetting errors and hazards to users from corrosive solutions. We recommend cutting the tip of the 1 ml tip and pipette slowly in a fume hood.*
8. Add 250 μ l of curcumin solution (see Recipes) and shake well until it forms homogenous dark purple colour (Figure 1B).
 9. Centrifuge the tubes at 15,000 \times g for 1 min and incubate the reaction mixture at room temperature for 1 h.
 10. After 1 h, stop the reaction by adding 500 μ l of Milli-Q water and mix well by inverting the tubes.
 11. Centrifuge the tubes at 15,000 \times g for 1 min to facilitate clear phase separation.
 12. Carefully pipette 200 μ l of upper phase (Figure 1C) into a 96-well UV Microplate and measure the absorbance at 550 nm using a microplate reader.

Note: Chloroform in the reaction mixture melts many types of plastic microplates and interferes in absorbance reading. Therefore, it is very important to use microplates resistant to organic solvents (for example, Thermo Scientific™ 96-Well UV Microplate or a quartz microplate).

13. Plot the absorbance against the quantity of B and prepare the calibration curve as shown in Figure 2. To obtain calibration curve, plot the absorbance values (A_{550} nm) against the quantity of boron (nmole B) using a spreadsheet (left panel of Figure 2).

Note: Jennings et al., 2007 have used both curcumin method and ICP-MS to quantify boron in yeast cells. They found that using the two methods the results were indistinguishable.

14. Use the obtained calibration curve to determine the quantity of B in unknown samples.

Recipes

Note: Every time make fresh solutions in a falcon tube.

1. Extraction solution
Dissolve 2-ethyl-1,3-hexanediol 10% (v/v) in chloroform
2. Acid mixture
Mix sulphuric acid (conc.) and acetic acid (conc.) in 1:1 (v/v) ratio in a Falcon tube
3. Curcumin solution
Dissolve curcumin (2 mg/ml) powder in MIBK (Methyl-Isobutyl-Ketone)

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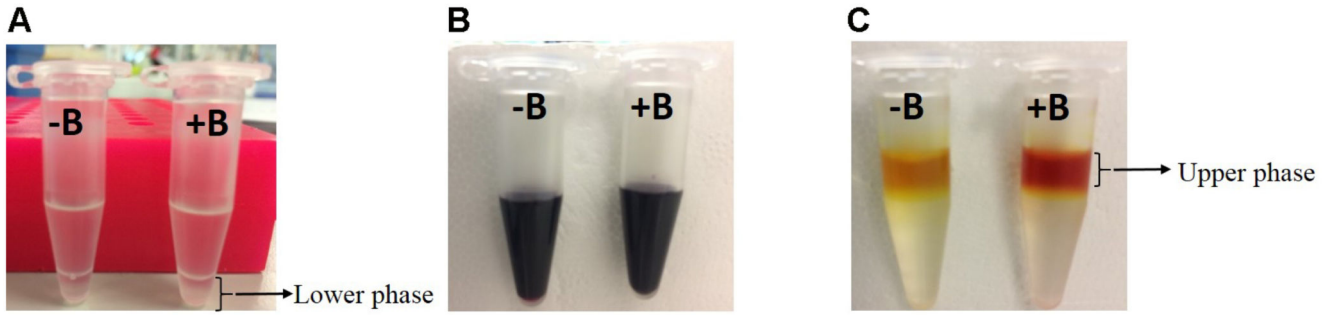


Figure 1. Steps showing phase separation and colour development in the absence (-B) or presence (+B) of boron

nmole B	A _{550nm}
0	0
0.1	0.02
0.2	0.05
0.5	0.10
0.9	0.20
1.9	0.37
3.7	0.76

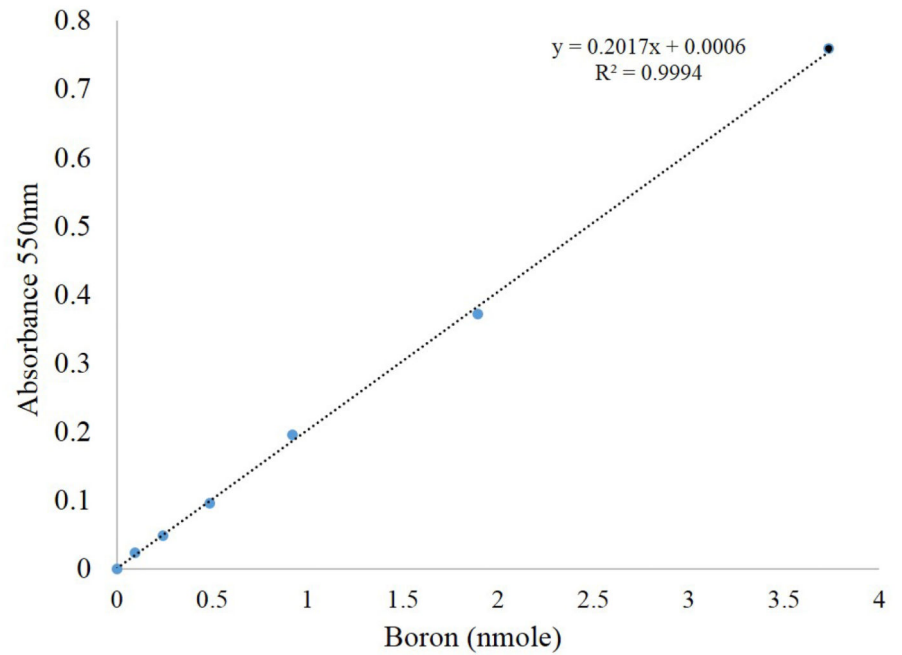


Figure 2. Calibration curve for the determination of boron concentration.
Absorbance measured at 550 nm (A_{550} nm).

Table1
Boric acid standards presented in different units

Boric acid μM	Boric acid mg/l	nmole B/300 μl
0.00	0.00	0.0
0.39	0.02	0.1
0.78	0.05	0.2
1.56	0.10	0.5
3.13	0.19	0.9
6.25	0.39	1.9
12.50	0.77	3.7

Representative calculation

Mol wt of Boric acid (H_3BO_3) = 61.83

61.83gm=1 mole

0.77 mg = 12.5 μmole

12.5 μmole (0.77 mg) of H_3BO_3 in 1000 ml = 12.5 μM

Therefore aliquot of 300 μl from 12.5 μM boric acid contain 3.7 nmole of Boron.

$$\frac{12.5\mu\text{mole}}{1000\text{ml}} \times 300\mu\text{l} = 3.7 \text{ nmole of Boron (B)}$$