### rpc00005

# Phytolacca americana PAR callus culture

#### Components

• A 9-cm plastic Petri dish, containing cells placed on semi-solid medium

### Notice

- Subculture the cells to fresh medium immediately after arrival [Notes I].
- Do not store the cell culture in a refrigerator and a freezer.
- Maintain aseptic conditions of the cell culture, and work in a laminar flow cabinet.

## Method

- Culture medium: MS medium, 1 mg/L 2,4-D, 1.2% (w/v) agar, pH 5.8 (medium no. 4) [Materials III]
- Culture conditions: 27°C, dark [Methods II]
- Subculture: 28-day intervals [Methods I]

### Citation of cell line

When results obtained by using this cell line are published in a scientific journal, it should be cited in the following manner: "*Phytolacca americana* PAR cell line (rpc00005) was provided by the RIKEN BRC through the National BioResource Project of the MEXT, Japan."

Experimental Plant Division RIKEN BioResource Research Center (BRC) Koyadai 3-1-1, Tsukuba, Ibaraki 305-0074 Japan FAX: +81 29 836 9053 E-mail: plant.brc@riken.jp http://epd.brc.riken.jp/en/

# Introduction

Pokeweed PAR cell line was established from *Phytolacca americana* L. (Sakuta *et al.* 1986). The PAR callus cells are red and accumulate high amount of betacyanin. The PAR cells are grown on a Murashige and Skoog (MS) medium supplemented with 1 mg/L 2,4-dichlorophenoxyacetic acid (2,4-D), and solidified with 1.2% (w/v) agar, pH 5.8. Our PAR cell culture has been maintained in the dark at 27°C and subcultured at 28-day intervals.

# **Materials**

## Chemicals and stock solutions

- (All stock solutions are stored at  $4^{\circ}$ C)
  - A) MS salt mix

Murashige and Skoog Plant Salt Mixture, FUJIFILM Wako Pure Chemical Corporation (#392-00591)

- B) Sucrose
- C) MS\_VT

Nicotinic acid
Pyridoxine·HCl
<b>Thiamine</b> ·HCl
Glycine

0.5 mg/mL 0.5 mg/mL 0.1 mg/mL 2 mg/mL

D) MS\_inositol

*myo*-Inositol

40 mg/mL

E) 2,4-D (0.2 mg/mL)

2,4-D sodium monohydrate 0.236 mg/mL (2,4-Dichlorophenoxy)acetic acid sodium salt monohydrate, Sigma-Aldrich (D6679)

- F) Agar, powder
- G) KOH (1 N)

## **Glassware and equipment**

- A) Erlenmeyer flask (300 mL), capped with two layers of aluminum foil
- B) Forceps, sterilized before use

#### Preparation of MS medium (medium no. 4)

1. Dissolve the following chemicals in approximately 800 mL of distilled water.

MS salt mix	1 bag (1 L)
Sucrose	30 g

2. Add following stock solutions, and fill up to approximately 950 mL with distilled water.

MS_VT	1 mL
MS_inositol	2.5 mL
2,4-D (0.2 mg/mL)	5 mL

- 3. Adjust the pH of the solution to 5.8 with KOH (1 N), and fill up to 1 L with distilled water.
- 4. Pour 100 mL of the medium into a 300-mL flask containing 1.2 g of agar.
- 5. Autoclave the flask at 121°C for 20 min.

# Methods

- 1. Pick up an appropriate amount of callus cells from a 28-day-old culture with a forceps and place the cells onto fresh MS medium.
- 2. Incubate cell cultures under the dark condition at 27°C.

# Notes

- We send PAR cells on semi-solid MS medium in a 9-cm disposable Petri dish. The cells should be subcultured to fresh MS medium immediately after arrival.
- In order to maintain PAR callus culture stably, it is essential to observe the growth of cells carefully. Because proliferation of PAR cells is affected by culture conditions, such as a room temperature, aeration conditions of the culture and so on, an amount of cells transferred to fresh medium and the subculture intervals may vary from one lab to another. We usually inoculate three pieces of PAR callus (about 10-mm in diameter) on 100 mL of MS medium in a 300-mL flask, and culture them for 28 days.

## References

Sakuta M, Takagi T, Komamine A (1986) Growth related accumulation of betacyanin in suspension cultures of *Phytolacca americana* L. Journal of Plant Physiology 125: 337–343. DOI: 10.1016/S0176-1617(86)80155-9

Chemical	Concentration (mg/L)
KNO <sub>3</sub>	1900
NH <sub>4</sub> NO <sub>3</sub>	1650
$CaCl_2 \cdot 2H_2O$	440
$MgSO_4 \cdot 7H_2O$	370
KH <sub>2</sub> PO <sub>4</sub>	170
H <sub>3</sub> BO <sub>3</sub>	6.2
$MnSO_4 \cdot 4H_2O$	22.3
$ZnSO_4 \cdot 7H_2O$	8.6
KI	0.83
$Na_2MoO_4 \cdot 2H_2O$	0.25
$CuSO_4 \cdot 5H_2O$	0.025
$CoCl_2 \cdot 6H_2O$	0.025
$FeSO_4 \cdot 7H_2O$	27.8
Na <sub>2</sub> -EDTA	37.3
Nicotinic acid	0.5
Pyridoxine·HCl	0.5
Thiamine·HCl	0.1
Glycine	2
myo-Inositol	100
Sucrose	30000
2,4-D sodium monohydrate	1.18
Agar	12000

Table A.1. Murashige and Skoog medium (medium no. 4)

# **Appendix A: Formulation of culture medium**