03-2-09

## Regeneration of Bottle Gourd Inbred lines and Induction of Putative Embryogenic Callus

Hyo Soon Kim, Jung Suk Lee, Ji Soo Shin, Sung Jegal, Sang Lyul Han, Yoon Sup Shin, Tae Ik Lee, Seung Gyun Yang and Chee Hark Harn

Biotechnology Center, Nong Woo Bio Co., Ltd., Jeongdan, Ganam, Yeoju, Kyonggi, Korea;

## **Objectives**

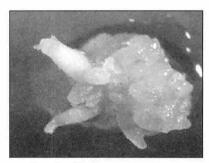
The objective is to develop a regeneration system of gourd inbred lines and to obtain an induction system for embryogenic callus. The ultimate objective for this research is to establish a transformation system of gourd crops.

## **Materials and Methods**

- 1. Cotyledons from 10 inbred lines of gourd (*Lagenaria* siceraria Standl.) were used for explants
- 2. To obtain the regeneration condition, cytokinin (BA 0, 0.5, 1.0, 2.0, 4.0 mg/l) and auxin (IAA 0, 0.1, 0.5, 1.0 mg/l) concentrations were combined in MS medium.
- 3. For embryogenic callus induction, 2,4-D (0, 1.0, 2.0, 4.0, 6.0 mg/l) was supplied in MS medium.

## **Results and Discussion**

- 1. A total of 10 inbred lines were tested for the regeneration rate and the rate was dependent on each line. Generally, cytokinin over 2 mg/l and IAA 0.5 mg/l was required for the shoot formation.
- 2. For the first time, embryogenic calli, although not confirmative yet, were obtained from the inbred line #8 and #10 by inducing callus at 2,4-D 2mg/l.
- 3. Putative embryogenic cell aggregates in suspension culture were able to undergo normal embryo mass propagation on 2,4-D 2mg/l over and zeatin at 0.2 mg/l. In solid culture (on the plate), the putative embryogenic callus was propagated well.



Putative Embryogenic Callus

<sup>\*</sup> Corresponding author: Chee Hark Harn; 031-883-7055; chharn@nongwoobio.co.kr