

Micropropagation and its Potential in Vegetable Crops

**Ipsita Sahu¹,
Yogendra Singh²,
Reena Nair³,
Chandrabhan Ahirwar⁴
and Swapnil Mahana¹**

¹M.Sc Research Scholar,
Biotechnology Centre, JNKVV,
Jabalpur (M.P)

²Assistant Professor (Senior Scale)-
Biotechnology, Department of
Genetics and Plant Breeding,
JNKVV, Jabalpur (M.P)

³Assistant Professor , Department
of Horticulture, JNKVV, Jabalpur
(M.P)

⁴Ph.D Research Scholar,
Department of Genetics and Plant
Breeding, JNKVV, Jabalpur (M.P)



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*Corresponding Author

Yogendra Singh*

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INTRODUCTION

The concept of micropropagation was first introduced to the scientific community in 1960 by Georges Morel, who successfully used the technique to produce virus-free plants of the orchid *Cymbidium*. Micropropagation is an advanced and well-established technique used for the rapid multiplication of plants. Because of its high propagation rate, ability to produce disease-free plants, and generation of high-quality planting material, it has significant commercial potential. It is considered both an art and a science of plant propagation carried out under **in vitro** conditions. The micropropagation process involves several stages, including maintenance of stock plants, selection and sterilization of explants, manipulation of culture media for shoot proliferation, rooting of plantlets, acclimatization, and finally transferring the plants to field conditions. Since the procedure requires a sterile working environment, most of the steps are performed manually, which makes the process labor-intensive, repetitive, and relatively expensive.

Objectives of Micropropagation:

The primary objective of micropropagation is to produce a large number of **high-quality, genetically uniform, and disease-free plants** under controlled laboratory conditions. This technique helps in the rapid and efficient multiplication of plants and supports modern agricultural and horticultural practices.

The major objectives are as following:

- **Rapid Multiplication:**

To produce a large number of identical plant clones within a short period of time from a small piece of plant tissue.

- **Production of Pathogen-Free Plants:**

To obtain disease-free, especially virus-free plants, by using techniques such as meristem-tip culture.

- **Clonal Propagation of Elite Varieties:**
To maintain true-to-type characteristics of superior, high-yielding, or disease-resistant plant varieties.
- **Year-Round Plant Production:**
To enable continuous plant production throughout the year, independent of seasonal or environmental limitations.
- **Germplasm Conservation:**
To preserve rare, endangered, or valuable plant species and maintain plant genetic resources for future use.
- **Overcoming Propagation Limitations:**
To propagate plants that do not produce viable seeds or are difficult to multiply through conventional methods, such as through **in vitro tuberization in potato**

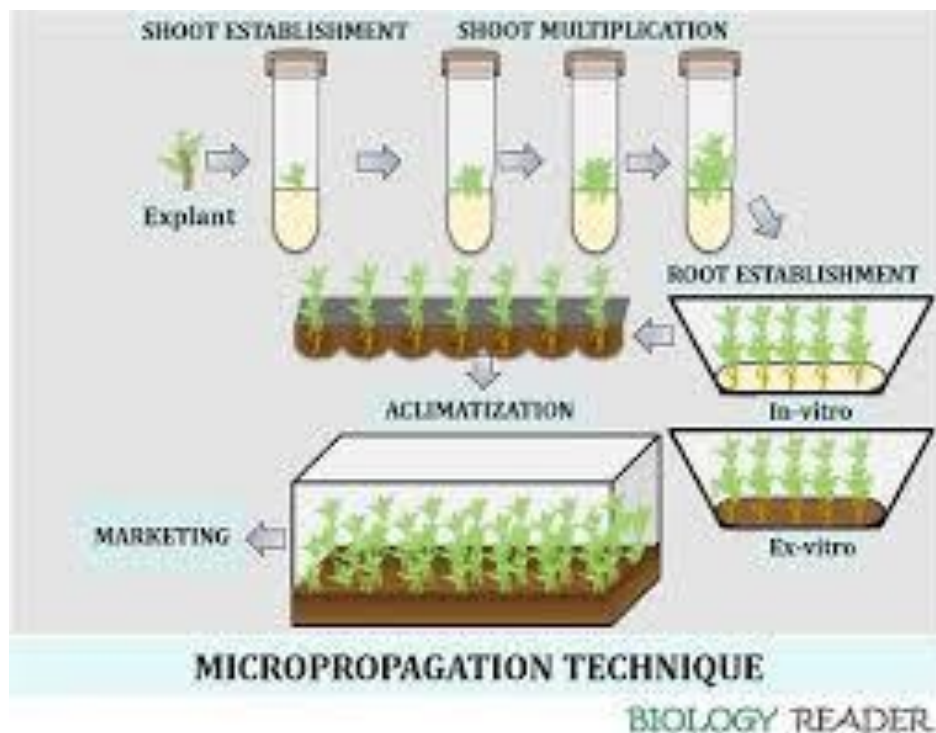


Fig: 01: Micropropagation Technique (Source: Biology Reader)

Applications of Micropropagation:

Micro propagation in gourds (*Trichosanthes dioica Roxb.*) through cuttings

Pointed gourds (*Trichosanthes dioica Roxb.*) from family *Cucurbitaceae* are dioecious perennial herbaceous vegetables. Fruits are the edible portion and are a good source of vitamin C and minerals. This crop is propagated conventionally by shoot cuttings using 60-90 cm long segments from basal portion of the vines. Seed propagation in gourds is not desirable due to its cross pollinated nature, poor germination, slow seedlings growth and segregation of male and female plants

Somatic embryogenesis in Carrots (*Daucus carota* subsp. *Sativus*)

Somatic embryogenesis is potentially one of the most efficient methods for plant micro propagation and in case of dicotyledonous culture, embryos at various developmental stages i.e. globular-, heart-, torpedo-, and cotyledonary-stage are frequently mixed in the suspension. Additionally, the rate of formation of embryos in each developmental stage alters correspondingly during the culture period in carrots. Somatic embryos harvested at the torpedo-stage, have better performance of plant conversions. Therefore, several attempts have been made to develop certain methods that results in obtaining a large number of torpedo-stage embryos of carrots with satisfactory degree of homogeneity.

***In vitro* tuber yields and multiplication rates of potato**

Potatoes are traditionally multiplied by tubers, with a multiplication rate of 6 to 10-fold annually. A major impediment to the introduction of new cultivars and to the production of high-quality tubers is the low production rate for tuber planting. Micro propagation can be a successful tool for enhancement of multiplication rates considerably. Under aseptic and controlled *i.e.* laboratory conditions, the rate of multiplication goes from 10 to 25-folds per 8 weeks, 8 to 84-fold per 40 days, and 4 to 7-fold per 3-5 weeks have been reported by researchers.

***In vitro* development of Cauliflower synthetic seeds**

The production of quality seeds is one of the main problems in cauliflower cultivation. Cauliflower, an open pollinated plant faces technical challenges to make in-bred lines with reliable self-incompatibility. Cabbage and cauliflower are two of numerous vegetables in the species *Brassica oleracea* of family *Brassicaceae*. These are grown, in Pakistan, for their edible value and are main cash crops of Pakistan. These are typically grown as a winter crop and sometimes also as summer vegetables. Vegetable production carries a valuable position in edible crops of Pakistan. Although the cabbage and cauliflower are among minor crops and are mostly grown in small farms even then Cauliflower is one of the most cultivated vegetable in the Punjab Province.

An innovative *in vitro* regeneration protocol in tomato

Tomato is among the most important vegetables and widely used as raw or cooked. Beside a source of antioxidants, minerals, fibers and vitamins it is also an excellent sculpt system for genetic studies, fruit development and ripening process. Insects, pests and different pathogens can reduce tomato yield in the field. Microbial pathogens

include lepidopteron, *Helicoverpa armigera* and common fruit borers which largely attack on fruit while *Spodoptera litura* destroy leaves. Incorporation of Bt genes in tomato has showed considerable resistance against lepidoterons. Tomato has served as an excellent model for plump fruit development and ripening. Well characterized ripening mutants, high density genetic maps, small genome size, short life cycle, efficient and stable transformation made tomato an excellent sculpt for studying viruses, development and fruit ripening process through genetic modification.

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