



## In vitro Propagation of pomegranate (*Punica granatum L.*)

Hadeer Hassan Ali<sup>1</sup>  
[Hasanhadeer521@gmail.com](mailto:Hasanhadeer521@gmail.com)

Jassim Mohammed Khalaf<sup>2</sup>  
[Jasim\\_agob@uokirkuk.edu.iq](mailto:Jasim_agob@uokirkuk.edu.iq)

Ali Mohammed Noori<sup>3</sup>  
[aloky1515@uokirkuk.edu.iq](mailto:aloky1515@uokirkuk.edu.iq)

<sup>1</sup> Kirkuk Agriculture Directorate, Ministry of Agriculture, Kirkuk, IRAQ.

<sup>2,3</sup> Department of Horticulture and land scape, College of Agriculture, University of Kirkuk, Kirkuk, IRAQ.

• Date of research received 15/2/2024 and accepted 31/3/2024.

### Abstract:

This study was conducted in the Plant Cell and Tissue Culture Laboratory of the Department of Horticulture and Landscape Engineering/College of Agriculture/University of Kirkuk, to demonstrate the study of the effect of adding benzyl adenine (0.0, 0.5, 1.0, 1.5, and 2.0 mg L<sup>-1</sup>) and kinetin (0.0, 2.0, 4.0, 6.0, and 8.0 mg L<sup>-1</sup>) on the growth of pomegranate plants in Woody Plant Medium (WPM) According to the results, the medium containing 1.0 mg L<sup>-1</sup> (BA) had the highest average number of branches at 3.3 branches. part<sup>-1</sup> in contrast, the medium containing 1.5 mg L<sup>-1</sup> had the longest branches, on average, measuring 1.5 cm, while the medium containing 2.0 mg L<sup>-1</sup> (BA) produced a highest average of 20 leaves. The results of adding kin to agricultural media showed that the highest average number of branches was 2.5 branches. Part-1 when treated with two concentrations (8.0 and 4.0) mg L<sup>-1</sup>, the length of the longest branch is 2.13 cm, and the highest average number of leaves is 17.6 leaves. Plant part when grown in medium supplemented with 2.0 mg L<sup>-1</sup>.

**Keywords:** *Punica granatum L.*, Micropropagation, WPM, Kin, BA

**Citation:** hasan, H., khalaf, J., & Noori, A. (2024). In vitro Propagation of pomegranate (*Punica granatum L.*). *Kirkuk University Journal for Agricultural Sciences*, 15(1), 325-330. doi: 10.58928/ku24.15129

**Correspondence Author:** Hadeer Hassan Ali<sup>1</sup>-[Hasanhadeer521@gmail.com](mailto:Hasanhadeer521@gmail.com)

**Copyright:** This is an open access article distributed under the terms of the creative common's attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original author and source are created.

## Introduction:

*Punica granatum* L., the pomegranate, is in the puniceae family, which means "apples with many seeds." Pomegranates are one of the oldest fruits that can be eaten. Both Jews and Christians have written about them in their holy books. The pomegranate was talked about three times in the Quran [1, 2]. There are carbs in pomegranate seeds, mostly sugars, as the percentage of sugars in their juice reaches 0.16%. The juice also contains 0.5% protein and 1.1% citric acid, in addition to 0.3% fatty substances, in addition to mineral elements such as calcium, phosphorus, iron and potassium. The fruit's peel also contains 28% tannin, which is an astringent. Therefore, it is used to treat diarrhea and combat leukemia by reducing the effectiveness of leukemia cells and their spread [3]. The Sulaimi variety is one of the most valuable pomegranate types in Iraq. It is also the most common, grown and harvested in farms in the center and northern regions. [4] [5] [6]. This variety is characterized by the fact that its fruits are large, round, with a thick peel, and the color of the peel is green with a red tint. Then a dark red color covers all the fruits at the end of the season (when fully ripe). The red seeds are juicy and have a bitter taste. As the fruits mature, their sweetness increases and their acidity decreases [6],[7]. Different sciences are progressing and thriving, and this depends on the technologies available to them that can be adopted to expand the scope of experiments, including plant tissue culture technology. The concept of plant tissue culture is expressed by culturing various plant cells or tissues in glass or plastic vessels containing artificial nutrient environments consisting of food materials under controlled conditions and are completely sterilized [8]. At present, with the development of science, development in the cultivation of plant cells and tissues has reached such an extent that it has become possible to grow different plant parts on nutrient media and obtain complete plants from them, as well as use them in research

and application. Including its fields of plant breeding and improvement, the production of medicines and medicinal drugs, and the production of seedlings. From medical injuries [9].

## Materials and methods:

Experiments were conducted in the Plant Tissue Culture Laboratory – Department of Horticulture and Landscape Design- University of Kirkuk-IRAQ. This study used plant parts (explants) taken from vegetative branches of pomegranate (*Punica granatum* L.) seedlings obtained from a nursery in Kirkuk- IRAQ. The plant parts were prepared by cutting the plant branches into parts of equal length and placing them inside a glass cup in preparation for washing them under running water for 5 minutes to remove any remaining dust and dirt. Add regular washing powder (Brite) and let sit for a few minutes. After that, place in a strainer and soak for 12 minutes under running water. The explant were passed on to a 250 ml glass container and an air chamber with laminar flow, then immersed in a commercial sodium hypochlorite (NaClO) solution at three concentrations (2, 4 and 6%) for (2, 4 and 6) minutes. Until the explant are submerged. Completely to ensure complete sterilization. After the sterilization ended, to eradicate any potential hazardous effects of the sterile chemical, the plant portions were washed three times consecutively about 5 minutes of distilled water. The plant parts were cut into lengths (1 cm) to be ready for transplantation in Woody Plant Medium (WPM) containing Benzyl Adenine (BA) at concentrations of (0.0, 0.5, 1.0, 1.5 and 2.0) mg L<sup>-1</sup> and Kinetin (Kin) at concentrations of (0.0, 2.0, 4.0, 6.0 and 8.0 mg L<sup>-1</sup>)., which was previously prepared by placing 400 ml of twice-distilled water in a 1000 ml glass beaker, then adding sucrose in an amount of 30 gL<sup>-1</sup>, then adding agar in an amount. 7.5 gL<sup>-1</sup> (agar), then (WPM) is added to it according to the recommended amount, and the volume is added to 1000 ml of sterile

distilled water, then heated on a device (Magnetic stirrer hot plant), then the pH is adjusted. Measurement. Accordingly, between (5.7) use (HCl and NaOH), and after the solution is completely homogeneous, it is poured into tubes designated for cultivation until used. The coefficients were compared using Duncan's multinomial test at a probability level of 5%. [10].

### Results and discussion:

The results in Table (1) indicate that there is a significant effect of sterilization concentrations and sterilization periods on

the contamination rate. The 2% concentration significantly outperformed the rest of the concentrations, as it is noted that the percentage of pollution reached the lowest possible level at this concentration, in which the plant parts were not exposed to any damage or pollution. The reason for this may be due to the percentage of the active ingredient in the sterilization (NaOCL). Which may be suitable for sterilizing the plant part without harming its components [11]. While the highest percentage of contamination of plant parts was at a concentration of 6% for 2 minutes

Table 1: The effect of sodium hypochlorite (NaClO) and the duration of sterilization on the percentage of contamination of pomegranate plant parts *Punica granatum L.*

Duration (min)	Concentration effect			
Concentration (mL)	2	4	6	(mL)
2	100 a	100 a	100 a	100 a
4	50 b	40 b	70 b	51.72 b
6	70 b	10 b	60 b	76.67 b
Duration effect	73 a	80 a	76 a	

Similar letters do not differ in terms of statistical significance at 5%.

The results of Table (2) note that (BA) showed a significant effect on the opening of buds of the parts that were grown on media supplement with different concentrations of WPM media, as the concentration of 1.0 mgL<sup>-1</sup> was superior to the rest of the concentrations, and the comparison treatment gave an average number of branches of 3.3 branch. While the concentration of 1.5 mgL<sup>-1</sup> Ba led to an increase in the length of the branches to reach 1.57 cm, the highest average number of leaves reached 20.4 leaves at a concentration of 2.0 mgL<sup>-1</sup> (BA).

The effects of (Kin) treatment on plant parts growing in (WPM) medium were observed in Table (3). The maximum average number of branches reached 2.5 branches when treated with two concentrations (4.0 and 8.0) mg L<sup>-1</sup>. (Kin), as there were no significant differences between the two concentrations, and they outperformed the comparison treatment and the rest of the treatments. The highest average number of longest branches reached 2.13 cm when treated with a concentration of 2.0 mg L<sup>-1</sup> (Kin), and the highest number of leaves reached 17.6 leaves when treated with a concentration of (2.0) mg L<sup>-1</sup> (Kin).

The increase in the percentage of live, uncontaminated plant parts resulting from immersion in sodium hypochlorate solution may be due to the effectiveness of this substance when using it to sterilize plant components. The results can be explained by considering chlorine as a sterilising chemical. When applied to plant parts at the right concentration and for the right amount of time, it kills microorganisms like fungi and bacteria. This substance is characterized by its ease of removal from the explant by repeated washing, as this substance decomposes into a less toxic substance and thus is easier to remove from the plant part. [12] The results of tables (2 and 3) can be interpreted to indicate the major role that cytokinins play in stimulating cell divisions and the formation of adventitious branches on plant organs, as

well as encouraging the growth of buds. The benefits after eliminating the phenomenon of apical dominance [13]. It was shown from calculations that BA had better results than Kin. This is attributed to the composition of BA, which is linked to its chain by a number of evidence, three double bonds, which makes it superior in its activities to other organizations. The bonds of BA are personally effective and compound active, which makes them more influential in terms of cell divisions and their expansion in size and differentiation, that is, more influential in the development of growth and development. This makes it one of the most prominent cytokinins used in the propagation of many plant species [12] [14]., and these results are in line with [15], [16], [17], [18] and [19].

Table 2: The effect of )BA( on Initiation Stag of pomegranate *Punica granatum* L.

(BA) mg L <sup>-1</sup>	Number of branches	Length of longest branch (cm)	number of leaves
0.0	0.7 b	0.58 b	3.7 c
0.5	2.9 a	1.25 a	14.5 b
1.0	3.3 a	1.45 a	18.7 ab
1.5	2.6 a	1.57 a	16.6 ab
2.0	2.7 a	1.50 a	20.4 a

Similar letters do not differ in terms of statistical significance at 5%

Table 3: Effect of (Kin) on Initiation Stage of pomegranate *Punica granatum* L.

(Kin) mg L <sup>-1</sup>	Number of branches	Length of longest branch (cm)	number of leaves
0.0	0.7 b	0.58 c	3.8 b
2.0	2.4 a	2.13 a	17.6 a
4.0	2.5 a	1.8 ab	16.4 a
6.0	1.8 a	1.55 b	15.2 a
8.0	2.5 a	0.92 c	15.4 a

Similar letters do not differ in terms of statistical significance at 5%

## Conclusions:

It was found that WPM medium was more effective than M S, as it led to an increase in the number of branches, their length, and the number of leaves of the plant parts growing in it.

## References:

- [1] Mohammed, M. H. M (2006). Fruits and Herbs Mentioned in the Holy Quran and the Sunnah of the Prophet. Dar Al Nada for Cultural Production and Distribution. Alexandria. Egypt. In Arabic.
- [2] Alwan, J. M (2017). Technology of deciduous fruit. Its propagation, cultivation, care, and production (Part One). Dar Al-Wadah Publishing. Amman. Jordan. In Arabic.
- [3] Khurshid, C. and Ahmed, O. (2016). Functional Variation of Soluble Polyphenols in Oak Apple Gall and Pomegranate Peels and their Inhibition Activity in Leukemia K562 Cells. *J Exp Food Chem*, (101), 2472-0542.
- [4] Al-ezzi, M. A. J (1990). The pomegranate fruit worm, its life, its harms, and its control - guidance bulletin - Ministry of Agriculture. Iraq. In Arabic.
- [5] Youssef, H. Y and Al Nuaimi, J. H (1980). Deciduous fruit production. Basra. Ministry of Higher Education and Scientific Research. Iraq. In Arabic.
- [6] Nasar, T. A (1991). Fruits Are Evergreen and Deciduous. Production. Most Important Types in the Arab World. Dar Al Maaref. Egypt 2<sup>ed</sup>; 324.
- [7] Al-dowry, A. H and Al-rawy, A. K (2000). Fruit production for specialized departments in horticulture. Dar Al-Kutub for Printing and Publishing, University of Mosul, Iraq. In Arabic.
- [8] Shukri, W. M and Al-Muaeyqal, R. M (2013). Plant Cell and Tissue Culture. Al-Mutanabbi Library. Dammam - Kingdom Saudi Arabia.
- [9] Hartman, H. T., Kester, D. E and Davies, F. T (2001). Plant Propagation: Principles and Practices. 7<sup>th</sup> Edition. Prentice Hall Publishers. New Jersey.
- [10] Al-Rawy, K. M and Khalaf Allah, A. M (1980). Design and analysis of agricultural experiment. Dar Al Kutub printing press. Mosul University, Iraq.
- [11] Al-Kinani, F. R (1987). Tissue culture and plant cells. University of Mosul. Iraq. In Arabic.
- [12] Salman, Muhammad Abbas (1988). Fundamentals of plant cell and tissue culture. Ministry of Higher Education and Scientific Research Dar Al-Kutub for printing and publishing. University of Baghdad\_Iraq.
- [13] Dodds, J. H and Roberts, L. W (1985). Experiment in Plant Tissue Culture. Washington D.C.
- [14] Geroge, E. F (1993). Plant propagation by Tissue Culture. Part1: The Technology. Exegetics limited.
- [15] Anurdaha, S., Vijayluxmi, S. K and Bhat, S (2016). Effect of Growth Regulators on in Vitro Root Formation in Strawberry, *Res. Environ. Life Sci* 9(13); 16-18.
- [16] Budiono, R. T., Setiawati, G. G., Pitaloka, L., Anggreini, M and Mutaqin, A. Z (2016). Micropropagation Strober (*Fragaria x Ananases* Var. Earlibrite) Degnan Penambahan BA (Benzyl Adenine) Dan IBA (Indole Butyric Acid) Pada Media MS (Murashing and Skoog). *Presiding Seminar Nasional II Tohun*. 1 (11): 26-38.
- [17] Buntsevich, L. L., Bamatov, I. M and Vinter, M. A (2018). Improvement of the Efficiency of Sanitation and Primary Propagation Technology of Gaden Strawberry in Vitro Culture. *Journal of Pharmaceutical Sciences and Research* .1(10): 79-84.
- [18] Khatik, P. V and Kaushik, R. A (2019). In Vitro Propagation of Strawberry *International Journal of Current Advanced Research*, no. October.
- [19] Nor, N. Y and Junkasiraporn, S. (2019). Effect of Cytokinins and Auxin on in Vitro Propagation of *Fragaria x Ananases* Duch. Cv. Pharaohatan 80. *Burapha Science Journal* 3(24): 1190-1204



## إكثار الرمان *Punica granatum* L. صنف سليمي خارج الجسم الحي

<sup>3</sup>علي محمد نوري

<sup>2</sup>جاسم محمد خلف

<sup>1</sup>هدير حسن علي

[aloky1515@uokirkuk.edu.iq](mailto:aloky1515@uokirkuk.edu.iq)

[Jasim\\_agob@uokirkuk.edu.iq](mailto:Jasim_agob@uokirkuk.edu.iq)

[Hasanhadeer521@gmail.com](mailto:Hasanhadeer521@gmail.com)

<sup>1</sup>مديرية زراعة كركوك، وزارة الزراعة، كركوك، العراق.

<sup>2,3</sup>قسم البستنة وتنسيق الحدائق، كلية الزراعة، جامعة كركوك، كركوك، العراق.

• البحث مستل من رسالة الماجستير للباحث الاول.

• تاريخ استلام البحث 2024/2/15 وتاريخ قبوله 2024/3/31.

### الملخص:

أجريت هذه الدراسة في مختبر زراعة الخلايا والأنسجة النباتية التابع لقسم البستنة وهندسة الحدائق، كلية الزراعة جامعة كركوك، لبيان تأثير إضافة (0.0, 0.5, 1.0, 1.5, 2.0) ملغم/لتر  $BA^{-1}$  و (0.0, 2.0, 4.0, 6.0, 8.0) ملغم/لتر  $Kin^{-1}$  على وسط (WPM) لنبات الرمان *Punica granatum* L. أظهرت النتائج تسجيل أعلى معدل لعدد الأفرع 3.3 فرع/جزء نباتي<sup>1</sup> في الوسط المزود ب1.0 ملغم/لتر  $BA^{-1}$  في حين كان أعلى معدل لطول أطول فرع 1.5 سم في الوسط المزود ب1.5 ملغم/لتر وأعلى معدل لعدد الأوراق 20 ورقة/جزء نباتي<sup>1</sup> عند الزراعة بالوسط المزود ب2.0 ملغم/لتر  $BA^{-1}$  وبينت نتائج إضافة  $kin$  إلى الأوساط الزراعية تسجيل أعلى معدل لعدد الأفرع 2.5 فرع/جزء نباتي<sup>1</sup> عند المعاملة بتركيزي (4.0 و 8.0) ملغم/لتر وطول أطول فرع 2.13 سم وأعلى معدل لعدد الأوراق 17.6 ورقة/جزء نباتي<sup>1</sup> عند الزراعة بالوسط المزود ب 2.0 ملغم/لتر  $BA^{-1}$ .

الكلمات المفتاحية: الرمان،  $Kin$ ،  $BA$ ، أوساط زراعة، الإكثار الدقيق.