



Effect of hormones on callus induction in Maize (*Zea mays L.*)

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Abstract: Callus induction from explants is a critical process in regeneration, micropropagation and transformation of maize (*Zea mays L.*) plants. Formation of callus from plant tissues on culture is affected by several factors. This study revealed to establish the effect of genotype, source of explants and auxin concentration on callus induction from five genotypes UMI 757 (G₁), UMI 615 (G₂), UMI 112 (G₃), UMI 285 (G₄) and CO 1 (G₅) and one hybrid CO H (M) 5 (G₆). Callus induction of the six maize varieties was investigated using immature embryos (E₁), leaf bits (E₂), root tips (E₃), hypocotyls (E₄) and seeds (E₅) as explants with different concentrations of hormones. In this study, immature embryo was taken from 10 to 12 days after pollination (DAP) to get maximum response. The highest percentage of callus induction was observed (99.10) in immature embryo culture and seed culture gave the highest percentage of rhizogenic callus formation when compare to immature embryo. Among the genotypes tested, CO H (M) 5 recorded the highest callus induction percentage on (2D2K2) medium composition.

Keywords: Callus induction, Hormones, Immature embryo, Maize, Seed.

INTRODUCTION

Maize (*Zea mays L.*) with $2n = 20$, is the third most widely distributed crop of the world after Rice and Wheat (Devi, *et al.*, 2016) being grown in diverse seasons and ecologies with highest production and productivity among food cereals. It is an important source of carbohydrate, protein, iron, vitamin B and minerals (P and K). In Sudan, maize is produced using traditional or mechanical methods and is mainly used for food, forage and is a potential source of foreign exchange through export (Omer *et al.*, 2008). Maize production is affected by biotic and abiotic factors. These constraints can be overcome through development of varieties that can tolerate (or) resist the stress. This can be done through complementing conventional breeding and genetic transformation. Success in plant transformation is dependent on the ability to regenerate a whole plant from transformed tissue (Ahmadabadi *et al.*, 2007) plant regeneration through tissue culture of maize was first reported by Green and Philips (1975). *Agrobacterium* mediated transformation of maize needs efficient regeneration systems. In this study, six maize genotypes were evaluated for their response to tissue culture at different hormone concentrations using five different explants sources.

MATERIALS AND METHODS

Plant Materials: Seeds of maize (*Zea mays L.*) genotypes, UMI 757, UMI 615, UMI 112, UMI 285 and

CO 1 and one hybrid CO H (M) 5 were used in this study. These seeds were surface sterilized with 70% ethanol for 2 minutes followed by 0.1% HgCl₂ for 5 minutes and then washed with three to four times with sterilized distilled H₂O under aseptic conditions. The sterile seeds were used as source of explants for callus induction. To establish plants to obtain leaf bits, root tip, Hypocotyl explants, sterile seeds were planted in sterile Jam jars containing MS basal salts (Murashige and Skoog, 1962). The Jar were kept in a growth room and maintained at a temperature of 28°C and sterile seeds. Immature embryo also used as explants. Immature embryos of 1.0- 2.0 mm size were aseptically excised from surface sterilised kernels. The Immature embryos were placed on the semisolid MS medium with the rounded scutellar side exposed and the plant Plumule radical axis side in contact with the medium.

Preparation of callus induction: Callus induction were performed on MS medium comprising of MS salts and vitamins supplemented with Macronutrients, Micronutrients, Micronutrient and Vitamins, 3% (w/v) sucrose. The pH of the medium was adjusted to 5.6 to 5.8 with IM NaOH or 0.1 M HCl and 0.8% (w/v) agar added before autoclaving to sterilize. The sterilized medium was allowed to cool before adding 2,4-D. The medium was dispensed in sterile Petridishes in volumes of 30 ml and allowed to solidify. Explants were cultured on the medium and plates sealed with parafilm. Sixteen levels of treatment combinations of 2,4-D and kinetin were tested to establish their efficacy in

establishing callus from five different explants. The mature and healthy seeds of maize were taken for callus induction. The seeds were sterilized with 70% ethanol for 2 min followed by 0.1 percent Hg Cl₂ for 5 min. Sterilized seeds were inoculated under aseptic conditions in the callus induction medium. The culture tubes were incubated in darkness at 25 ± 2°C for callus induction. The Immature embryo has proven to be the best source for the establishment of embryogenic callus and plant regeneration in maize. The Immature Embryo from maize cob were collected 10-16 days after pollination. The kernels were surface sterilized under Laminar Air Flow Chamber, the Immature embryos were aseptically excised from the kernels of the ear by cutting off kernels using scalpel blade and removing the endosperm. The immature embryo was placed on semisolid medium and was incubated for callus induction. About one cm explants were cut from seven day old seedlings collected from aseptically germinated seedlings. Inoculation of leaf bits, hypocotyls and roots were incubated on to the different callus induction media, such as T₁ MS + 0.5 mg l⁻¹ 2,4 - D + 0.3 mg l⁻¹ kin + 1g l⁻¹ CH, T₂ MS + 0.5 mg l⁻¹ 2,4 - D + 0.2 mg l⁻¹ kin + 1g l⁻¹ CH, T₃ MS + 0.5 mg l⁻¹ 2,4 - D + 0.1 mg l⁻¹ kin + 1g l⁻¹ CH, T₄ MS + 1.0 mg l⁻¹ 2,4 - D + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH, T₅ MS + 1.0 mg l⁻¹ 2,4 - D + 0.2 mg l⁻¹ kin + 1 g l⁻¹ CH, T₆ MS + 1.0 mg l⁻¹ 2,4 - D + 0.1 mg l⁻¹ kin + 1 g l⁻¹ CH, T₇ MS + 1.5 mg l⁻¹ 2,4 - D + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH, T₈ MS + 1.5 mg l⁻¹ 2,4 - D + 0.2 mg l⁻¹ kin + 1 g l⁻¹ CH, T₉ MS + 1.5 mg l⁻¹ 2,4 - D + 0.1 mg l⁻¹ kin + 1 g l⁻¹ CH, T₁₀ MS + 2.0 mg l⁻¹ 2,4 - D + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH, T₁₁ MS + 2.0 mg l⁻¹ 2,4 - D + 0.2 mg l⁻¹ kin + 1g l⁻¹ CH, T₁₂ MS + 2.0 mg l⁻¹ 2,4 - D + 0.1 mg l⁻¹ kin + 1 g l⁻¹ CH, T₁₃ MS + 0.5 mg l⁻¹ IAA + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH, T₁₄ MS + 1.0 mg l⁻¹ IAA + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH, T₁₅ MS + 1.5 mg l⁻¹ IAA + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH, T₁₆ MS + 2.0 mg l⁻¹ IAA + 0.3 mg l⁻¹ kin + 1 g l⁻¹ CH.

Statistical analysis: The observations recorded were statistically analyzed by subjecting the data to Factorial Completely Randomized Design, designed by (Gomez and Gomez, 1984). Level of significance was determined by using standard analysis of variance. Differences among mean values were assessed by LSD. The data obtained with per cent values were subjected to arc sine transformation.

RESULTS AND DISCUSSION

Effect of hormones on callus induction: Auxins were used to induce cell division and root differentiation in tissue culture medium. Among them, 2,4-D is widely used for callus induction. In the present study, MS medium containing 1.0 g l⁻¹ casein hydrolysate was supplemented with various levels of 2,4-D *viz.*, 0.5, 1.0, 1.5 and 2.0 mg l⁻¹ and also IAA at 0.5, 1.0, 1.5 and 2.0 mg l⁻¹ combination with cytokinin (kinetin). A

differential influence of various concentrations of 2,4-D, IAA in callus behaviour was observed. Addition of adequate levels of synthetic auxins such as 2,4-D in to a basal medium resulted in prolific callus formation in maize tissue culture. Callus could not be induced in N₆ medium in the absence of 2,4-D (Binott *et al.*, 2008). The influence of different concentration of 2,4-D, IAA alone or in combination with 0.3 mg l⁻¹ kinetin were recorded maximum, callus induction percentage, number of days for callus induction and fresh weight of callus on 28th day. This is in accordance with the findings of Ansari (1997) and Shohaël *et al.* (2003) in maize. The complex action of kinetin was observed by Inoue and Maeda (1982) in rice. They explained the effect of kinetin on callus induction could be promotive or inhibitory depending upon the kind and concentration of auxin in the medium.

In the present study, MS medium containing 1.0 g l⁻¹ Casein Hydrolysate has been added with different levels of kinetin *viz.*, 0.1, 0.2 and 0.3 mg l⁻¹ in combination with auxin 2,4-D and IAA. The result showed that among different concentrations tried, 0.3 mg l⁻¹ kinetin recorded higher values (60.44%) of callus induction per cent when compared to other levels. Since kinetin is susceptible to interactions with 2,4-D, there was variation in response. But when 0.3 mg l⁻¹ of kinetin was combined with 2,4-D 1.5 mg l⁻¹ (T₇), IAA 1.5 mg l⁻¹ (T₁₅) recorded the higher values when compared to either 2,4-D or IAA alone. Similar results reported by Shohaël *et al.* (2003), embryogenic calli formation was high when N₆ medium supplemented with L-Proline 2.3 g l⁻¹, casein hydrolysate 200 mg l⁻¹, 2,4-D 1.0 mg l⁻¹ and Kinetin 0.1 mg l⁻¹. Abebe *et al.* (2008) also reported, when 3 mg l⁻¹ of 2,4-D was combined with 0.5 mg l⁻¹ kinetin recorded highest callus induction per cent. Among the genotypes the maximum percentage of callus induction was recorded by CO H (M) 5 (59.61%) followed by UMI 285 (56.96%) and UMI 757 (55.25%). The maximum percentage of callus induction response was recorded by seed followed by immature embryo the explants leaf bits recorded the poorest response for callus induction. Among the 16 treatments studied, T₇ showed the maximum of 60.44% followed by T₁₅ (56.79%) and T₁₆ (56.33%) (Table 1).

Among the G x E interactions, the maximum callus induction percentage was observed in G₄E₁ (99.50%) followed by G₅E₁ (99.20%) and G₁E₁ (98.46%) with the treatment of T₁₅ (MS + 1.5 mg / l IAA + 0.3 mg / l K + 1 g/l (H). The treatment T₁₂ showed poor performance in all the explants and in all the genotypes. Among the treatment combination G₄ E₁ T₁₅ recorded the maximum value of (99.50%) (9.20%) and (98.46%) for the genotypes COH (M) 5, UMI 285 and UMI 757 respectively. The genotypes UMI 615 and CO 1 had a slightly lower value compared to the other genotypes.

Effect of explant: In the present study, investigations were made using five explants seed (E1), immature embryo (E2), leaf bits (E3), root (E4) and hypocotyls (E5). These explants were cultured with various levels of hormones and organic supplements and variation in their capacity in callus response was studied.

The studies indicated that maximum percentage of callus induction response was observed in seed followed by immature embryo and leaf bit recorded the poorest response of callus induction. Similar results were already reported by Shohael *et al.* (2003) in maize. The high callus responding nature and culturability of mature seed was in accordance with the reports of Delporte *et al.* (2001) in wheat and Bijy (2002) in rice. Al-Abed *et al.* (2006) and Sikandar *et al.* (2007) reported that the mature seed scutellum is the best explant for high totipotent embryogenic callus initiation and they also reported that the plant regeneration from coleoptile and root segments was unsuccessful. The high culturability of immature embryo was in agreement with the reports of Oduor *et al.* (2006). Benson, (2000) and Huang and Wei (2004) reported the juvenile tissues are usually more responsive to tissue culture than mature ones.

In the present study, studies regarding explants performance immature embryo as well as seed was more (or) less similar and best. Hypocotyl, root and leaf bits were lagging behind and the poor response of leafbits was also reported by Vinothini (2004) in rice. Chand and Sahrawat (2000) reported high callus induction and plant regeneration using root explants of barley. In contrast, Khaleda and Al-Forkan (2006) reported that root explant showed poor response be due to the fact that the calli derived from those explants were not totipotent for plant regeneration in rice.

Number of days for callus induction (days): In callusing duration studies, for early callus induction inferred that the use of 1.5 mg l⁻¹ IAA in combination with 0.3 mg l⁻¹ kinetin (T₁₅) followed by 2.0 mg l⁻¹ IAA + 0.3 mg l⁻¹ kinetin (T₁₆) and 1.5 mg l⁻¹ 2,4-D + 0.3 mg l⁻¹ kinetin (T₇) was the best. Among the explants, seed recorded the earlier induction of callus followed by immature embryo. The leaf bit recorded maximum days for callus induction.

Among the genotypes the earlier induction of callus was observed in CO H (M) 5 (G₄) (17.68 days) followed by UMI 285 (G₅) 17.73 days. CO 1 (G₆) recorded the maximum days for callus induction (18.22). The influence of genotypes on callus induction in maize was reported already by Green and Philips, (1975) and Tomes and Smith, (1985). Bronsema *et al.* (1997) indicated that the genetic information needed for embryogenic callus formation of A 632 as female parent, was transferred through A 188 pollen. The results revealed that among the explants, seed (E₁) recorded the earlier induction of callus (16.22 day) followed by immature embryo (E₂). The leaf bit explant took maximum days

for callus induction (19.27 days). Among the G x E interactions, G₁E₁, G₄E₁, G₂E₁ recorded the earlier days with T₁₅ (MS + 1.5 mg / l IAA + 0.3 mg / l K + 30 g / l maltose) 15.08 days, 15.10 days and 15.56 days respectively. The interactions G₆E₅ with T₉ recorded longer days for callus induction (20.80 days). The results on number of days for callus induction for the hormonal combination revealed significant 1.39 differences among the treatments. MS + 1.5 mg / l IAA + 0.3 mg / l K + 30 g/l maltose (T₁₅) treatment recorded the minimum number of days for callus induction 17.20 days followed by T₋₁₂ (18.67 days) (Table 2).

Under various levels of organic supplements, significantly superior callus induction per cent on immature embryo was noticed in CO H (M) 5, UMI 285 and UMI 757 when compared to other types. Study in seed emphasized the capacity of CO H (M) 5 and UMI 285 with higher per cent values and indicated its superiority over other genotypes. Similarly CO H (M) 5 and UMI 285 genotypes showed significant values in the case of root and hypocotyl, where as other genotypes stood inferior. When study was conducted on duration of callusing, minimum number of days was observed for immature embryo in CO H (M) 5, UMI 285 and UMI 757 when compared to other genotypes. Seed culture confirmed again minimum duration induction for COH (M) 5 followed by UMI 285 and UMI 112 and they stood superior to other genotypes. In the case of root and hypocotyl, significantly higher values were noticed in COH (M) 5 followed by UMI 112 as compared to others.

In all the above studies, COH (M) 5 was performing extremely well under all treatments. This can be attributed to its already high responding nature and it is a tissue culture friendly type. UMI 285, UMI 757, UMI 112, UMI 615 and CO 1 also high response, where as CO 1 lagged behind all treatments. Such type of varietal variation to culture response was reported by other researchers Hodges *et al.* (1986) in maize, Bronsema *et al.* (1997) in maize Agarwal *et al.* (2006) in rice and Binott *et al.* (2008) in maize.

Fresh weight of callus on 28th day (mg): Callus fresh weight on 28th day, significantly superior at T₁₅ (1.5 mg l⁻¹ IAA) followed by T₁₆ (2.0 mg l⁻¹ IAA) and T₇ (1.5 mg l⁻¹ 2,4-D) (Table 20). Among the five explants, seed recorded maximum fresh weight of callus followed by immature embryo. This was in agreement with the report of Ansari, (1997), he cultured on 2,4-D ranged from 1.5 mg l⁻¹ to 2.0 mg l⁻¹ with different media *viz.*, (MS) Murashige and Skoog, (N₆) Chu, (B₅) Gamborg B5 Medium, (YP) Yeast Media and (LS) Linsmaier and Skoog Medium. Callus induction was high in (MS) Murashige and Skoog and (N₆) Chu medium when compared to other media, which was due to the differences on major nutrients present in these basal media.

Among the genotypes, the maximum callus weight was

observed in COH (M) 5 (G₄) 1753.13 mg followed by UMI 285 (G₅) (1750 mg) and UMI 757 (G₁) (1745.2 mg). The genotype CO 1 (G₆) recorded the minimum callus weight of 1444.62 mg. The results revealed that among the explants, the seed explant (E₁) recorded the maximum weight of 3400.73 mg followed by immature embryo (E₂) 3264.51 mg. The leaf bit explant (E₃) had minimum fresh weight 392.83 mg. The results of fresh weight on 28th day of the hormonal combination revealed significant differences among the treatments. The treatments T₁₅ (MS + 1.5 mg /l IAA + 0.3 mg / l K 30 g/l maltose), T₁₆ (MS + 2.0 mg /l IAA + 0.3 mg / l K + 30 g/l maltose) and T₇ (MS + 1.5 mg / l 2,4 D + 0.3 mg / l K + 30 g/l maltose) recorded the maximum weight of 1813.9 mg, 1777.1 mg and 1744.2 mg respectively. The treatment T₃ recorded the minimum weight of callus (1690.90 mg). Among the G x E interactions G₄ E₁ recorded the maximum weight of 3443.13 mg, followed by G₅E₁ (3441.06 mg) and G₅E₂ (3440.75 mg). The treatment combination G₆ E₅, G₂E₅, G₃E₅ recorded the minimum weight of 375.25 mg, 387.50 mg and 3930.06 mg respectively (Table 3). Among the G x E x T interactions G₄E₁ T₁₅ recorded the maximum weight of callus (3538 mg) followed by G₅E₁T₁₅ (3536 mg). The treatments G₆E₅T₉, G₆E₅T₃ and G₂E₅T₃ recorded the minimum of callus of 280 mg, 310 mg and 315 mg respectively. Among the G x E x T interactions G₁E₁T₁₅ recorded the minimum value of 15.02 days followed by G₄E₁T₁₅ (15.10 days), while the treatment combination G₆E₅T₉ recorded the maximum number of days for callus induction (20.80 days).

Conclusion

Transformation of maize plants was made successfully by standardization of tissue culture experiments. The study concluded that 0.3 mg/l of kinetin combined with 1.5 mg/l 2,4-D and 1.5 mg/l IAA has recorded highest callus induction percentage. The genotypes COH(M) 5 with Immature embryo as a explant exhibited earlier callus induction and maximum callus weight. These optimized tissue culture protocol, will be used for transformation of resistance to biotic and abiotic stress in maize in future.

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