

Micropropagation and Effect of Phloroglucinol on Rooting of *Diospyros crassiflora* Hiern

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Abstract. The jet-black to streaked ebony wood produced by the African ebony (*Diospyros crassiflora* Hiern, Ebenaceae) is exploited in Central and West Africa. A conservation effort is currently underway in Cameroon to propagate the tree through seedlings and cuttings. However, the intermittent availability of its seeds and the long time required for rooting formation of cuttings are limiting its propagation. This study aims to develop a successful protocol for ebony micropropagation. In vitro culture of nodal segments from seedlings was performed in half-strength Murashige and Skoog (MS) medium supplemented with either zeatin (0.0, 2.3, 4.6, 9.1, 13.7, 18.2, 22.8, and 27.4 μM) or 6-benzylaminopurine (BAP) (0.0, 2.2, 4.4, 8.8, 13.3, 17.8, 22.2, and 26.6 μM). After 12 weeks, all media allowed shoot budbreak. Shoots displaying the greatest budbreak were observed with 22.8 μM zeatin and 22.2 μM BAP. The shoots were then transferred to a medium supplemented with indole-3-butyric acid (IBA) and phloroglucinol (PG) at different concentrations for root induction. Root induction was observed on the shoots initially induced in the medium with BAP, but not in those grown in the medium with zeatin. In half-strength MS supplemented with 396.5 μM PG plus 14.2 μM IBA, the formation of a single tap root was observed on 79% of shoots, with an average root length of 2.8 ± 0.18 cm. Rooted plants were successfully acclimatized to the greenhouse, with a 50% survival rate. This is the first report on *Diospyros crassiflora* micropropagation, which paves the way for rapid ebony multiplication to respond to needed conservation efforts.

Diospyros crassiflora H. (Ebenaceae), commercially known as the African ebony, is a small forest tree found at elevations less than 1000 m in South Nigeria, Cameroon, Central African Republic, Equatorial Guinea, Gabon, Republic of the Congo, and Democratic Republic of the Congo (Lemmens et al., 2012). It is a shade-tolerant species characterized by a relatively low growth rate (Bongjoh and Nsangou, 2001). The hard,

dense black heartwood of the tree is exploited in Cameroon and other countries for the construction of musical instruments such as the violin, guitar, cello, contrabass, and piano keys (Lemmens et al., 2012; Taylor, 2012). The species is also used locally for medicinal purposes and pharmacological research, and has proven antifungal and antibacterial properties in stem bark extracts (Dzoyem et al., 2007; Tangmouo et al., 2006).

Diospyros crassiflora was listed as vulnerable on the International Union for Conservation of Nature Red List because it is threatened by the destruction of forests for agriculture and grazing, and the commercial logging of other species (Schatz et al., 2019). For now, the species has not been included in extended silvicultural or agroforestry systems. As part of the Congo Basin Institute's ebony conservation project, an integrated agroforestry model is currently being developed to compensate for the decrease of tree populations. The development of a cost-effective, reliable approach to large-scale production of the plant is becoming essential for long-term sustainability of the species.

Until now, propagation by seeds has been the main form of propagation of *D. crassiflora*. However, seed production is seasonal and year dependent. Moreover, reproduction through seeds does not allow genetic fidelity because of its heterozygosity. Vegetative propagation by leafy stem cutting is feasible, but the quantity of plant material available from a selected donor plant is limited and the cuttings maintain the plagiotropic growth of the lateral stem of the cutting (Tsobeng et al., 2011). Our study aimed at developing a protocol of ebony micropropagation that might circumvent the shortcomings of both seedling and cutting propagation.

Micropropagation of the close relatives *D. virginiana* (Palla et al., 2013) and *D. kaki* (Mitrofanova and Mitrofanova, 2004; Pijut, 2012) has shown that the development of axillary buds could be induced by cytokinin. PG is a phenolic compound known for its properties as a promoter of plant growth (Teixeira da Silva et al., 2013). The objective of this study was to investigate the effect of cytokinins on the development of axillary buds, and PG on root initiation of *D. crassiflora*.

Materials and Methods

Plant material and source of explant. Mature seeds were harvested on 35-year-old *D. crassiflora* trees in Ekombite, Central Region, Cameroon (lat. 3°16'N, long. 11°19'E). Voucher specimens of one mother plant (V. Deblauwe no. 376) were deposited at the herbaria of the Higher Teacher Training College, University of Yaounde I, Yaounde, Cameroon; Université Libre de Bruxelles, Belgium; and the Missouri Botanical Garden, St. Louis, MO. The seeds were germinated and maintained in a shade house at the International Institute of Tropical Agriculture campus in Yaounde, Cameroon. Germination was conducted on moist Whatman paper n° 3 (GE Healthcare Bio-Sciences, Pittsburgh, PA) in a 500-mL plastic culture vessel for 2 weeks. Each vessel, containing 10 seeds, was incubated in a growth chamber at 25 ± 1 °C with an irradiance of 40 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ provided by cool white fluorescent tubes (40 W, Philips) with a 16-h photoperiod. After germination, seedlings were subcultured in plastic bags (100 mm \times 150 mm \times 50 μm) containing a mixture of sterilized black soil and river sand

Table 1. Effect of zeatin and 6-benzylaminopurine (BAP) on shoot formation from nodal explants of *D. crassiflora* after 12 weeks of culture in half-strength MS medium.

Plant growth regulators (μM)	Budbreaking (%)	Mean no. of leaves/shoot	Mean no. of shoots/explant	Explant with more than one shoot (%)	Shoot length (cm)	Callus formation (%)
Control	0	0.0 \pm 0.0	0.0 \pm 0.0	0	0.0 \pm 0.0	0
Zeatin						
2.3	87	4.7 \pm 0.4 b	0.9 \pm 0.1 b	0	2.9 \pm 0.2 ab	17
4.6	85	4.7 \pm 0.4 b	0.9 \pm 0.1 b	2	2.0 \pm 0.1 c	10
9.1	92	5.3 \pm 0.4 ab	0.9 \pm 0.1 b	0	2.1 \pm 0.1 c	14
13.7	91	4.8 \pm 0.5 b	1.0 \pm 0.0 b	6	2.0 \pm 0.1 c	18
18.2	91	6.1 \pm 0.5 ab	1.2 \pm 0.1 ab	15	2.4 \pm 0.1 bc	18
22.8	95	6.8 \pm 0.4 a	1.4 \pm 0.1 a	24	2.9 \pm 0.1 a	67
27.4	90	5.5 \pm 0.4 ab	1.1 \pm 0.1 ab	17	2.0 \pm 0.1 c	78
BAP						
2.2	78	5.2 \pm 1.0 b	1.7 \pm 0.3 ab	24	1.9 \pm 0.1 a	0
4.4	86	5.0 \pm 0.8 b	1.9 \pm 0.3 c	33	1.8 \pm 0.1 a	0
8.8	85	5.7 \pm 0.6 ab	2.2 \pm 0.3 c	53	1.7 \pm 0.1 a	0
13.3	87	6.7 \pm 0.7 b	2.4 \pm 0.3 c	72	2.0 \pm 0.1 a	0
17.8	85	6.2 \pm 0.7 ab	2.7 \pm 0.4 bc	64	1.9 \pm 0.1 a	0
22.2	92	7.0 \pm 0.8 a	3.1 \pm 0.3 a	78	1.9 \pm 0.1 a	0
26.6	91	6.1 \pm 0.6 ab	2.5 \pm 0.2 c	74	2.0 \pm 0.1 a	0

Values represent means \pm SE. Lower case letters indicate significant differences within each treatment according to Tukey's honestly significant difference test ($P \leq 0.05$).

(1:1, v/v). Plantlets were allowed to grow in a nonmist propagator [developed by Leakey et al. (1990)] installed in a shade house with a shade cloth (88% shade protection, ultraviolet stable polypropylene, water permeable for rainfall and irrigation).

Cuttings from seedlings were used to establish a protocol that will be later extended to cuttings derived from mature trees with good commercial characteristics (i.e., size of the black wood). Cuttings with a length of 0.5 to 1 cm with a single internode were harvested from 6-month-old seedlings and used as explants. The explants were washed under running tap water for 30 min, followed by treatment with 0.3% (w/v) of Banko[®] Plus (Arysta LifeScience, Paris, France) fungicide; a suspension of 550 g·L⁻¹ chlorothalonil and 100 g·L⁻¹ carbendazim for 20 min. Explants were rinsed three times with sterile distilled water in a laminar flow hood and were disinfected successively with 0.1% (w/v) mercuric chloride solution for 10 min and 50% commercial bleach (2.4% sodium hypochlorite) containing 0.1% Tween 80 for 20 min. Last, they were rinsed three times with sterile distilled water. Shoot segments (length, 0.5–1.0 cm) containing single nodes were excised from the surface-sterilized materials and used as explants. The explants were cultured in glass culture tubes (20 ×

150 mm) containing 10 mL medium for shoot outgrowth.

Culture media and culture conditions. The basal medium consisted of half-strength MS (Murashige and Skoog, 1962) salt solution containing Morel and Wetmore (1951) vitamins and 3.0% sucrose. For shoot growth, media were solidified with 0.3% (w/v) gelrite (Duchefa Biochemie B.V., Haarlem, The Netherlands), whereas for root induction 0.7% Bacto agar (Liofilchem s.r.l., Roseto degli Abruzzi, Italy) was used. In each successive subculture, the basal medium was supplemented with different concentrations of plant growth regulators (see next section). The pH of the media was adjusted to 5.8 with 1 N sodium hydroxide or hydrogen chloride. The media were distributed in test tubes (10 mL in each) and autoclaved at 121 °C at 1.2 kg·cm⁻². The cultures were incubated at 25 \pm 2 °C in a growth chamber, with an irradiance of 40 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ provided by cool white fluorescent tubes (40 W), for a 16-h photoperiod.

Shoot growth. Explants were cultured on the medium described earlier with eight concentrations of cytokinins: zeatin (0.0, 2.3, 4.6, 9.1, 13.7, 18.2, 22.8, and 27.4 μM) or BAP (0.0, 2.2, 4.4, 8.8, 13.3, 17.8, 22.2, and 26.6 μM). Sixty-five, 45, and 50 explants were used for each concentration of zeatin, BAP, and control, respectively. Explants were transferred to fresh media every 3 weeks. After four subcultures (i.e., 12 weeks), the percentage of plants producing shoots, the number of leaves per explant, the number of shoots per explant, the shoot length, and the rate of callus formation were recorded.

Root induction. Shoots of at least 1.5 cm in length were sampled and transferred to half-strength MS basal medium designed for root induction and differentiation. The basal medium was supplemented with six concentrations of IBA (0.0, 2.5, 4.9, 9.8, 14.7, and 19.6 μM), and 0.0 or 396.5 μM PG. Twenty-four explants were used for each treatment. The percentage of shoot-forming roots, mean

number of roots per shoot, and root length were recorded after 12 weeks of culture in the root induction medium.

Acclimatization. Plantlets were removed from the culture flasks and washed gently to remove all media attached to the roots. They were then transplanted in a sterilized mixture of locally collected black soil and river sand (1:1, v/v). The mixture was set in recycled half-liter water plastic bottles cut into two parts. The upper and lower parts served as the lid and pot, respectively. The cultures were incubated at 25 \pm 2 °C in a growth chamber, with an irradiance of 40 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ provided by cool white fluorescent tubes (40 W), for a 16-h photoperiod for 6 weeks. Relative humidity was slowly reduced by gradually removing the lids. Plantlets were transferred to a mini propagator, where they were stored for 2 months. From the mini propagator, they were transferred to black plastic bags containing a sterilized mixture of locally collected black soil and river sand (1:1, v/v) and were moved to the shade house. To keep a high humidity level, plants were watered every 3 d. The percentage of surviving plants was recorded after 14 weeks of growth in the propagator.

Statistical analysis. Normality and homoskedasticity of the replicated measurements were verified with the Shapiro-Wilk test and Bartlett and Levene test, respectively. An analysis of variance (ANOVA) was performed with the Agricolae package in R version 1.09j (Mangiafico, 2015). When the ANOVA indicated statistical significance ($P < 0.05$), Tukey's test for multiple comparisons was used to distinguish the differences among treatments.

Results and Discussion

Shoot proliferation. The first budbreak was observed on explants after 3 weeks grown in half-strength MS medium supplemented with plant growth regulators (data not shown). After 12 weeks, no shoot induction

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was observed for explants in the control media. Shoot growth, number of shoots and leaves on cuttings, and callus formation on explants were affected by the type and the concentration of cytokinins (Table 1, Fig. 1A).

The greatest percentage of shoot induction (95%) was observed in the medium

supplemented with 22.8 μM zeatin. The mean number of leaves and the average shoot length observed in that treatment were 6.8 ± 0.4 and 2.9 ± 0.1 , cm respectively (Table 1, Fig. 1B). Callus induction was observed at the sectioned base of shoots in all zeatin treatments. These calli were compact and black (Fig. 1C).

BAP stimulated budbreak and multiple shoot regeneration (Fig. 1D). The best shoot proliferation and growth were observed on medium containing 22.2 μM BAP. In this treatment, 92% of the explants displayed budbreak and 78% of cuttings produced more than one shoot. On average, an explant produced 3.1 ± 0.3 shoots with an average length of 1.9 ± 0.1 cm (Table 1). Our results support previous findings of the positive effect of zeatin on shoot induction in *Diospyros* species (Cooper and Cohen, 1985; Fukui et al., 1989; Palla et al., 2013; Tao and Sugiura, 1992). Cooper and Cohen (1985) reported that MS medium with 4.56 mM or 13.7 mM zeatin produced the best shoot growth from nodal explants of *D. kaki* compared with the medium with 6-benzyladenine, kinetin, isopentenyladenine, or benzyl-tetrahydropyranlyl adenine. Fukui et al. (1989) established shoot cultures of 95 Japanese persimmon cultivars on half-strength nitrogen MS medium plus 100 mM zeatin. Tao and Sugiura (1992), using dormant buds of *D. kaki*, noted that if cultures were initiated in BAP, those cultures needed to be transferred to 25 mM 6-(γ,γ -dimethylallylamino) purine or 3 mM zeatin to ensure shoot elongation. Shoot cultures were established with nodal stem explants from *D. virginiana* by Palla et al. (2013) in MS medium containing 10 μM zeatin.

Contrary to results reported for *D. kaki* in which BAP had little or no effect (Fukui et al., 1989), our results show that, for *D. crassiflora*, BAP medium produced a greater number of shoots per explant than zeatin. BAP promoted multiple shoot formation, whereas cell elongation was limited. BAP has been used to induce budbreak and multiple shoot formation in several woody plant species, such as *Balanites aegyptiaca* (Anis et al., 2010; Khamis et al., 2016), *Melia azedarach* (Husain and Anis, 2009), *Salix tetrasperma* (Khan et al., 2011), and *Shorea tumbergaia* (Shukla and Sharma, 2017).

Root induction. Shoots derived from explant growing in zeatin did not produce roots in all the treatments (data not shown). Shoot explants grown in medium supplemented with BAP failed to produce roots in the control medium. IBA alone (2.5–19.6 μM) did not induce roots (data not shown). Root induction was observed, however, when IBA was supplemented with 396.5 μM PG. The greatest frequency of root formation (79%) and root length (2.8 ± 0.18 cm) was obtained in half-strength MS medium containing 396.5 μM PG and 14.2 μM IBA (Table 2). When rhizogenesis was observed, the number of roots induced per explant was always limited to one tap root without secondary root hairs. This root displayed a characteristic black color similar to the taproot produced from seed-grown plants (Fig. 2). A positive rooting response using PG and IBA has been reported for several tree genera, including *Diospyros*, *Malus* (Dobránszki and Teixeira da Silva, 2010), *Prunus* (Hammatt and Grant, 1997), *Jatropha* (Daud et al., 2013), and *Juglans* (Licea-Moreno et al., 2015).

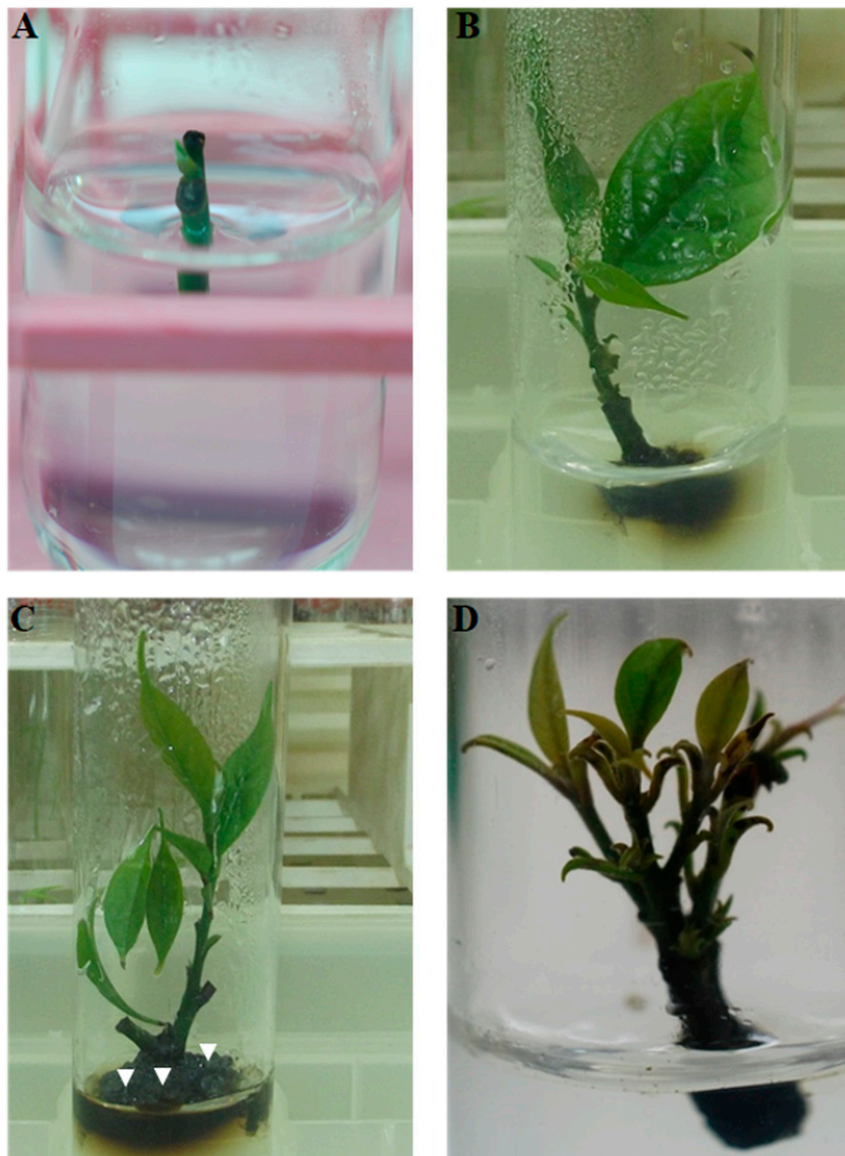


Fig. 1. Shoot induction from explants growing in half-strength MS medium. (A) Budbreaking of explant cultured in 22.8 μM zeatin. (B) Shoot elongation of explant cultured in 22.8 μM zeatin. (C) Callus formation at the cut extremity of explant cultured in 22.8 μM zeatin (arrows). (D) Shoot proliferation in 22.2 μM 6-benzylaminopurine.

Table 2. Effect of phloroglucinol and indole-3-butyric acid (IBA) concentrations on root formation of shoot of *D. crassiflora* after 12 weeks.

Phloroglucinol (μM)	IBA (μM)	Rooting (%)	Root length (cm)
396.5	2.5	0	0.0 ± 0.00
	4.9	17	1.6 ± 0.14 a
	9.8	50	1.2 ± 0.15 a
	14.7	79	2.8 ± 0.18 a
	19.6	71	1.1 ± 0.2 b

Values represent means \pm SE. Different lower case letters indicate significant differences between treatments according to Tukey's test ($P \leq 0.05$).

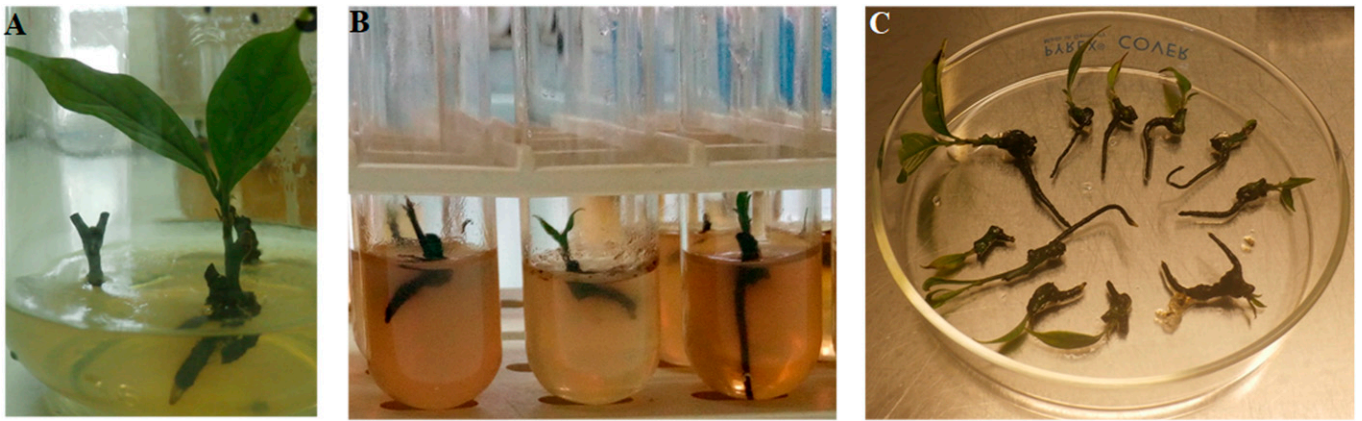


Fig. 2. Root induction in *D. crassiflora* plants cultured in various media. (A) Shoot in half-strength MS medium with 14.7 μM indole-3-butyric acid (IBA) + 396.5 μM phloroglucinol (PG). (B) Shoot in half-strength MS medium with 19.6 μM IBA + 396.5 μM PG. (C) Rooted shoots in half-strength MS medium with 14.7 μM IBA + 396.5 μM PG.

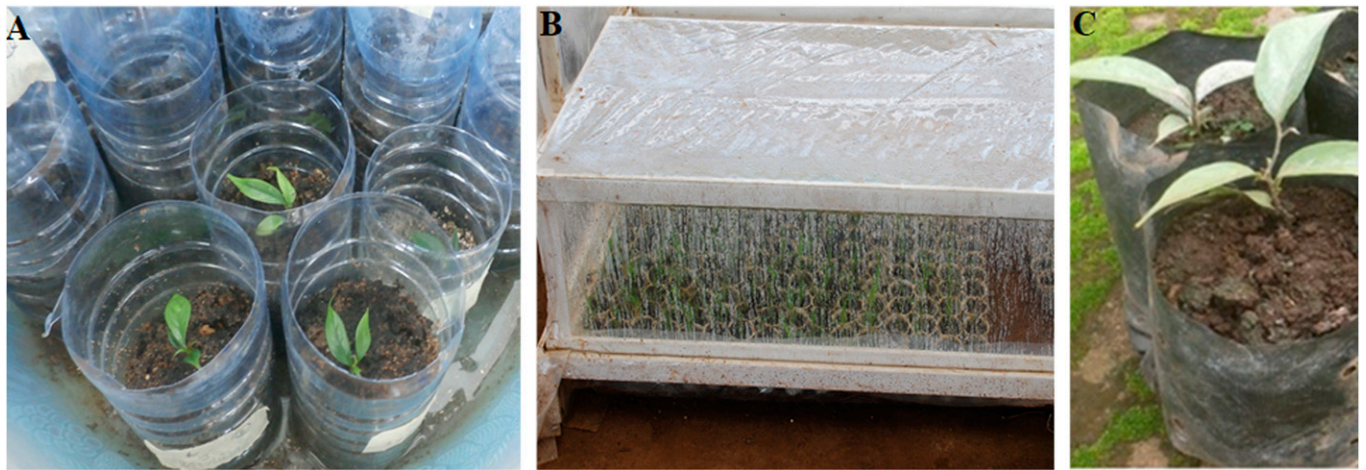


Fig. 3. Potted plantlets of *D. crassiflora* after 12 weeks on rooting medium of 14.7 μM indole-3-butyric acid + 396.5 μM phloroglucinol followed by 3 months of acclimatization in the greenhouse. Plantlets in culture room (A), mini green house (B), and acclimatized plants (C).

It has been reported that PG acts synergistically with auxin during the most sensitive phase of root initiation (Dobrąnski and Teixeira da Silva, 2010; Hammatt, 1994). The application of IBA alone did not result in root formation. Our results show that, in *D. crassiflora*, PG seems to be essential to induce root formation in the presence of an auxin (IBA, in our case). This confirms the view that PG is an auxin promoter (Daud et al., 2013; Pérez et al., 2016; Teixeira da Silva et al., 2013). A greater percentage of rooting of *D. virginiana* hybrid Rosseyanka (91%) was also achieved in MS medium containing 5 μM IBA and 0.1 $\text{g}\cdot\text{L}^{-1}$ PG (Palla et al., 2013).

Acclimatization. A major limitation in the application of micropropagation technology is high mortality experienced by in vitro-raised plants during laboratory-to-land transfer. When transferred ex vitro, micropropagated plants are exposed to conditions (altered temperature, light intensity, and water stress) that necessitate acclimatization for successful plantlet establishment and survival (Chandra et al., 2010; Kumar and Rao, 2012). The success of *D. crassiflora* acclima-

tization was one of the fastidious steps during our micropropagation process. All the in vitro plantlets displayed one black tap root without hair roots, which could explain their slow growth. Fifty rooted plants of *D. crassiflora* derived from different IBA and PG treatments were combined for acclimatization. They resumed their growth after 12 weeks of culture in an ambient growth condition (Fig. 3). The survival rate after 14 weeks was 50%.

Conclusion

This is the first report describing in vitro propagation of *D. crassiflora* through internode explants. Half-strength MS supplemented by zeatin was effective for shoot elongation and callus formation whereas BAP supplementation was effective for multiple shoot stimulation and development. PG had a positive effect on rooting and acclimatization of rooted shoots. The results obtained confirm that the micropropagation of African ebony is possible, opening up opportunities for domestication of the species and rapid multiplication to meet the demands of conservation efforts.

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