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### Influence of biofertilizers and media on germination parameters and chlorophyll content in aonla

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#### Abstract

Aonla is one of the highly remunerative crops usually found in dry deciduous forests of India. For raising the rootstock, it is mainly propagated by sexual methods. Findings regarding effect of biofertilizers and media on germination parameters and chlorophyll content is less hence the present experiment was conducted to study the effect of biofertilizers and media on germination parameters and chlorophyll content of aonla. Treatment combinations has showed the significant difference. Among the different treatment combinations used media containing of soil + cocopeat + vermicompost enriched with the biofertilizer mixture of *Azatobacter* + PSB + KRB has recorded minimum days taken for initiation of seed germination, days taken to 50 per cent germination, maximum germination percent at 20 and 30 DAS and highest chlorophyll content.

Keywords: biofertilizers, GA3, germination parameters, vermicompost, FYM, aonla

#### Introduction

Aonla botanically *Emblica officinalis* L. bearing the chromosome number of 2n=28. In India it is also called as Indian gooseberry, fruit of 21st century, amritphal fruit, aonal, usuri which belongs to Euphorbiaceae family, native to tropical South-East Asia. It is the deep-rooted tree having the wide range of adoptability to grow in any types of soil (Lilabati and Sahoo, 2016)<sup>[4]</sup>. It is a rich source of ascorbic acid (vitamin C) next to barbados cherry and has high nutritional and medicinal properties (Vishen et al., 2017)<sup>[13]</sup>. Dried fruits are used to treat diarrhoea, chronic dysentery, haemorrhage, diabetes, cough, jaundice etc. (Vishen et al., 2017)<sup>[13]</sup>. Uttara Pradesh is the leading state in India in terms of both area and production of aonla (NHB annonymus 2019) <sup>[7]</sup>. Aonla can be propagated by both sexual and asexual methods. Asexually it is propagated by patch budding, shield budding, chip budding, forket budding and wedge grafting (Negi et al., 2010) <sup>[6]</sup>. Seed propagation is essential to raise the rootstocks for asexual propagation. Growing media is the important factor for the plants that gives anchorage to the plants and provide essential nutrients required by the plants. The growing media enriched with biofertilizers possess the advantages like add nutrients through natural process like nitrogen fixing, phosphorus solubilizing and stimulate plant growth through the synthesis of growth promoting substances. They help to build up soil microflora and there by maintain soil health. Vermicompost application is one of the effective methods to improve the depleted soil fertility and enrich the nutrients availability, conserve more water and maintain soil quality (Makode, 2015) <sup>[5]</sup>. So, the present investigation was undertaken to study the effect of different growing media, biofertilizers and their interaction on seedling vigour of aonla.

#### **Materials and Methods**

The present investigation was carried out at College of Horticulture, Dr. Y.S.R Horticultural University, Venkataramannagudem, West Godavari District, Andhra Pradesh during the year 2020-2021.

Experiment was laid out in the factorial randomized block design (FRBD) with 2 factors, biofertilizers (8 levels:  $B_0$ : Control,  $B_1$ : Nitrogen fixing bacteria (*Azatobactor* @ 2%),  $B_2$ : Phosphorous solubilizing bacteria (PSB @ 2%),  $B_3$ : Potassium releasing bacteria (KRB @ 2%),  $B_4$ : *Azatobactor* @ 2% + PSB @ 2%,  $B_5$ : *Azatobactor* @ 2% + KRB @ 2%,  $B_6$ : PSB @ 2% + KRB @ 2%,  $B_7$ : *Azatobactor* @ 2% + PSB @ 2% + KRB @ 2%) and media [2 levels:  $M_1$ : Soil + Cocopeat + FYM (1:1:1),  $M_2$ : Soil + Cocopeat + Vermicompost (1:1:1)] with 16 treatment combinations replicated twice. For experimental purpose seeds were collected from the reliable source and treated with GA<sub>3</sub> @ 500 ppm for 24 hr to break the dormancy. Polybags of  $20 \times$ 

15 cm pinched with 16 holes in each were filled with the media enriched with the different biofertilizers used in the experiment. Treatment combinations comprised of T<sub>1</sub>: B<sub>0</sub>M<sub>1</sub>: Soil: Cocopeat: FYM (1:1:1), T<sub>2</sub>: B<sub>1</sub>M<sub>1</sub>: Azatobactor @ 2% +Soil: Cocopeat: FYM (1:1:1), T<sub>3</sub>: B<sub>2</sub>M<sub>1</sub>: PSB @ 2% + Soil: Cocopeat: FYM (1:1:1), T<sub>4</sub>: B<sub>3</sub>M<sub>1</sub>: KRB @ 2% + Soil: Cocopeat: FYM (1:1:1), T<sub>5</sub>: B<sub>4</sub>M<sub>1</sub>: Azatobactor @ 2% + PSB @ 2% + Soil: Cocopeat: FYM (1:1:1), T<sub>6</sub>: B<sub>5</sub>M<sub>1</sub>: Azatobactor @ 2% + KRB @ 2% + Soil: Cocopeat: FYM (1:1:1), T<sub>7</sub>: B<sub>6</sub>M<sub>1</sub>: PSB @ 2%+ KRB @ 2% +Soil: Cocopeat: FYM (1:1:1), T<sub>8</sub>: B<sub>7</sub>M<sub>1</sub>: *Azatobactor* @ 2% + PSB @ 2% + KRB @ 2% + Soil: Cocopeat: FYM (1:1:1), T<sub>9</sub>: B<sub>0</sub>M<sub>2</sub>: Soil: Cocopeat: Vermicompost (1:1:1), T<sub>10</sub>: B<sub>1</sub>M<sub>2</sub>: Azatobactor @ 2% +Soil: Cocopeat: Vermicompost (1:1:1), T<sub>11</sub>: B<sub>2</sub>M<sub>2</sub>: PSB @ 2% + Soil: Cocopeat: Vermicompost (1:1:1), T<sub>12</sub>: B<sub>3</sub>M<sub>2</sub>: KRB @ 2% + Soil: Cocopeat: Vermicompost (1:1:1), T<sub>13</sub>: B<sub>4</sub>M<sub>2</sub>: Azatobactor @ 2% + PSB @ 2% + Soil: Cocopeat: Vermicompost (1:1:1), T<sub>14</sub>: B<sub>5</sub>M<sub>2</sub>: Azatobactor @ 2% + KRB @ 2% + Soil: Cocopeat: Vermicompost (1:1:1),  $T_{15}$ :  $B_6M_2$ : PSB @ 2% + KRB @ 2% +Soil: Cocopeat: Vermicompost (1:1:1), T<sub>16</sub>: B<sub>7</sub>M<sub>2</sub>: Azatobactor @ 2% + PSB @ 2% + KRB @ 2% + Soil: Cocopeat: Vermicompost (1:1:1). During February 2-3 seeds were sown in each polybag according to the treatment combinations. Watering was done regularly using rose can and different plant protection measures was taken at all the growth stages. From each replication representative 5 seedlings were taken to record the different parameters at monthly interval.

#### Statistical analysis

The data on various characters studied during the course of investigation were analysed statistically by applying the procedures of analysis of variance as outlined by Panse and Sukhatme (1985) <sup>[8]</sup>. The critical difference among the treatment means was compared at 5 per cent level of significance.

#### **Results and Discussion**

Less number of days taken for initiation of seed germination (8.50 days), days taken to 50 per cent germination (20.50 days), maximum germination percentage at 20 DAS (88.82%) and 30 DAS (97.38%) and total chlorophyll content (61.70) was recorded in the treatment  $T_{16}$  containing the media of soil + cocopeat + vermicompost inoculated with the biofertilizers of *Azatobacter* + PSB + KRB against lowest in the uninoculated seedlings (Table 1).

The plants grown in media enriched with vermicompost exhibit the positive effects on various germination parameters (Khare *et al*, 2018)<sup>[2]</sup>. Kumar *et al.* (2015)<sup>[3]</sup> showed that, media containing vermicompost showed highest seed germination of *Pinus gerardiana*. Media inoculated with biofertilizers like *Azospirillum, Azatobacter* and phosphobacteria showed earliness and highest seed germination of cardamom seeds showed by Paulraj and Raj in 2018<sup>[9]</sup>.

Biofertilizers helps in the synthesis of beneficial hormones and growth factors that will leads to the increased cell division and multiplication of cell also increases the assimilation and accumulation of food materials (Raman, 2012)<sup>[12]</sup> in apple that will helps in better growth of the seedlings. Similar results were observed in shea tree by Abdullahi *et al.*, 2012<sup>[11]</sup>, Verma *et al.*, 2019<sup>[12]</sup> in dragon fruit and in sweet orange by Rana *et al.*, 2020<sup>[11]</sup>

 Table 1: Influence of biofertilizers and growing media on days taken to initiation of m germination, days to 50 % germination, germination % at 20 and 30 DAS and total chlorophyll content.

Treatment	Days taken to initiation of seed	Days taken to 50%	Germination % at	Germination % at	Total chlorophyll
	germination	germination	20 DAS	30 DAS	content (SPAD)
Biofertilizers	5				
$B_0$	17.25	30.50	43.43	45.80	40.44
B1	12.75	24.75	67.72	68.25	46.93
<b>B</b> <sub>2</sub>	13.50	25.25	63.99	64.94	45.94
<b>B</b> 3	14.00	26.75	56.63	56.74	44.66
<b>B</b> 4	9.50	23.25	85.72	87.84	52.30
<b>B</b> 5	11.25	23.75	76.77	80.10	50.88
<b>B</b> <sub>6</sub>	11.50	24.00	74.45	78.09	49.81
<b>B</b> <sub>7</sub>	8.50	21.00	87.66	93.96	57.74
SEm±	0.19	0.27	0.82	0.77	0.24
CD at 5%	0.58	0.83	2.48	2.34	0.74
Media					
M1	12.69	25.38	67.17	69.32	47.46
M <sub>2</sub>	11.88	24.44	71.92	74.61	49.71
SEm±	0.10	0.14	0.41	0.38	0.12
CD at 5%	0.29	0.42	1.24	1.17	0.37
Interaction					
$B_0M_1$	18.50	32.50	35.77	37.20	38.94
$B_1M_1$	13.00	25.00	65.93	67.01	46.27
$B_2M_1$	14.00	25.50	63.76	64.72	45.44
B <sub>3</sub> M <sub>1</sub>	14.00	27.00	55.83	55.83	44.39
B <sub>4</sub> M <sub>1</sub>	10.00	23.50	85.44	87.11	51.87
B5M1	11.50	24.00	72.58	76.51	50.37
$B_6M_1$	12.00	24.00	71.58	75.68	48.61
B7M1	8.50	21.50	86.50	90.55	53.78
$B_0M_2$	16.00	28.50	51.09	54.41	41.94
$B_1M_2$	12.50	24.50	69.50	69.50	47.59
$B_2M_2$	13.00	25.00	64.21	65.16	46.43
$B_3M_2$	14.00	26.50	57.44	57.66	44.92
B <sub>4</sub> M <sub>2</sub>	9.00	23.00	86.00	88.57	52.73

B5M2	11.00	23.50	80.96	83.70	51.38
$B_6M_2$	11.00	24.00	77.33	80.50	51.00
$B_7M_2$	8.50	20.50	88.82	97.38	61.70
SEm±	0.27	0.39	1.15	1.09	0.35
CD at 5%	0.83	1.17	3.51	3.31	1.05

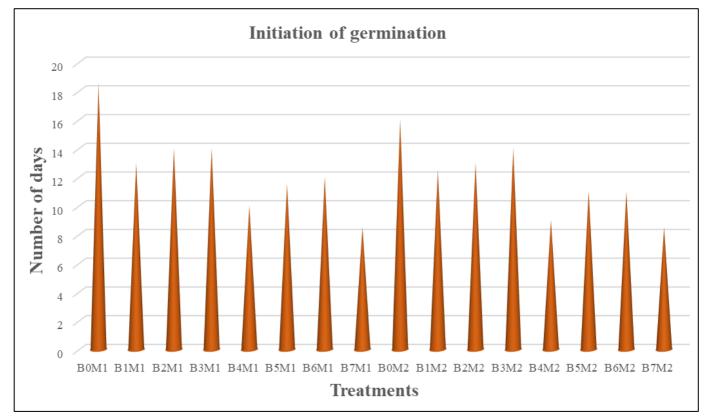


Fig 1: Influence of media and biofertilizers on initiation of seed germination in Aonla

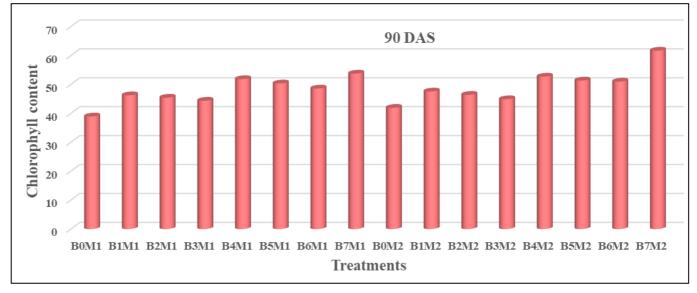


Fig 2: Influence of media and biofertilizers on chlorophyll content (SPAD) in aonla

#### Conclusion

Experiment revealed that the treatment  $T_{16}$  containing media of soil + cocopeat + vermicompost with biofertilizers of *Azatobacter* + PSB + KRB recoded less number of days taken for initiation of seed germination (8.50 days), days taken to 50 per cent germination (20.50 days), maximum germination percentage at 20 DAS (88.82%) and 30 DAS (97.38%) and total chlorophyll content (61.70).

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