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# IMPACT OF PLANT GROWTH PROMOTING ROOT ENDOPHYTES ON FLOWERING ATTRIBUTES OF CROSSANDRA (CROSSANDRA INFUNDIBULIFORMIS L.)

U. Sai Prakash\*, M. Raja Naik, K. Swarajyalakshmi, R. Lakshmipathy, M. Balakrishna and K. Arunodayam

Department of Floriculture and Landscaping, College of Horticulture (Dr. YSRHU), Anantharajupeta - 516 105, A.P., India \*Corresponding author E-mail: uppuletisaiprakash@gmail.com (Date of Receiving-12-07-2025; Date of Acceptance-17-09-2025)

ABSTRACT

Ornamental crops play a crucial role in the global floriculture industry, valued for their aesthetic appeal, socio-cultural significance and commercial importance. Among these, Crossandra (firecracker flower) is highly esteemed for its vivid blooms, utility in garlands and decorations especially in South India. But, cultivating Crossandra relies heavily on chemical fertilizers, which leads to degrading soil fertility, polluting the environment and inflating production costs. By keeping this view, a field experiment was carried out to investigate the influence of methods of inoculation, PGPRE's and their interaction on floral attributes of *Crossandra infundibuliformis* (L.) var. Arka Chenna at College of Horticulture, Anantharajupeta, Dr. Y.S.R. Horticultural University. In this experiment, various PGPRE's *viz.*, *Piriformospora indica*, *Glomus fasciculatum* and Arka Microbial Consortium were applied through different methods. Among the treatments, the interaction M<sub>1</sub>P<sub>8</sub> (Soil application with PGPRE's 10 days prior to Crossandra seedling planting x *Piriformospora indica*+ *Glomus fasciculatum*+ Arka Microbial Consortium + 25% RDF) reported superior performances among all floral parameters. These findings highlight the potential of PGPRE's as a sustainable and eco-friendly approach to improve the ornamental quality of Crossandra.

**Key words:** Crossandra infundibuliformis, Arka Chenna, Piriformospora indica, Glomus fasciculatum, Arka Microbial Consortium.

#### Introduction

Crossandra (*Crossandra infundibuliformis* L.) being a perennial plant belongs to the Acanthaceae family with a basic chromosome number of n=10. It is popularly known as "Fire cracker plant" or "Tropical Flame". Vernacularly it is called as "Kanakambaram" in Telugu (Rehana *et al.*, 2020), 'Aboli' in Marathi and 'Kanakambara' in Kannada (Sree *et al.*, 2023). It was believed to be originate from South India and Sri Lanka. The name Crossandra was derived from the two Greek words *i.e.*, 'krossoi' means fringe and 'aner' means male, combinedly known as fringed stamens (Das *et al.*, 2022). Despite lacking any fragrance, it gained significant popularity owing to its remarkable colour, light weight and strong market demand (Vinod and Kannan, 2020).

In India, Karnataka, Andhra Pradesh, Tamil Nadu and Telangana are the regions where Crossandra is cultivating in a commercial scale. The flowers serve various purposes, including religious offerings, hair adornments, garlands, venis and are also suitable for garden display (Prasanth *et al.*, 2020). This serves as a significant ornamental pot flower in Sweden, Denmark and Hungary (Bharath *et al.*, 2018).

Piriformospora indica, formerly known as Serendipita indica is a root endosymbiotic fungus having a wide host range, belongs to the Sebacinales in the Basidiomycota (Kaboosi et al., 2022). It was isolated by Prof. Dr. Ajit Varma and group from the rhizosphere of two woody shrubs, Prosopis juliflora and Zizyphus nummularia, in Rajasthan's Thar desert (Varma et al.,

2013). Upon root colonization with the host plant, *P. inidca* produces acid phosphatases and nitrate reductases which helps the host plant to access the insoluble, condensed or complex forms of phosphorus and nitrogen present in the soil (Gill *et al.*, 2016). It forms pear shaped chlamydospores in the root region.

On the other hand, Glomus fasciculatum belongs to the Glomeromycota taxonomy, which includes 110 species, makes the genus important in the order Glomerales (Dalpe et al., 2005). Arbuscles present in AMF helps in secretion of organic acids, thereby promoting nutrient uptake by the colonised plants (Gong and Tian, 2019). It produces extramatrical hyphae which acts as root extensions and thereby increasing the root surface area, making it easy for absorption of water and diffusion limited nutrients, especially in P-deficient soils (Mamatha et al., 2002). Arka Microbial Consortium is a carrier-based product developed by ICAR-IIHR, Bengaluru, conveniently applied through soil, seed, water and growing media. It contains three types of bacterial strains i.e., Bacillus aryabhattai, Pseudomonas taiwanensis and Azotobacter tropicalis (Aswathi et al., 2020).

Among the ornamental crops, Crossandra is one of the perennial crop which requires continuous feed of fertilizers throughout its life. Prolonged use of chemical fertilizers causes serious consequences for soil and environmental health viz., soil acidification, leaching salinization, disruption of soil microbial ecology, nutrient imbalance, reducing soil fertility over the time etc. Besides, application of chemical fertilizers is unsafe to humans also. Meanwhile, Plant Growth Promoting Root Endophytes (PGPRE's) are the rhizosphere biota which makes a symbiotic association with the host plant and promotes the growth of host plant. This PGPRE's solubilize the insoluble form of nutrients present in the soil and make them available for the plant. As these PGPRE's are cheaper than chemical fertilizers, the cost of cultivation can also be reduced. The existing research gap pertains to the lack of clear and systematic assessment of various application methods and the combinations of PGPRE's on overall plant performance in Crossandra. Moreover, no prior studies have examined the combined application of Piriformospora indica, Glomus fasciculatum and Arka Microbial Consortium in this crop, which makes the work novel with increased yield by reducing fertilizer regimes indicates the potentiality of creating a sustainable floriculture in the future. By keeping this in view, the present investigation was carried out to study the influence of PGPRE's on the floral attributes of Crossandra var. Arka Chenna.

#### **Materials and Methods**

The field experiment was conducted at College of Horticulture, Anantharajupeta, Dr. Y.S.R. Horticultural University (latitude 13.99'N and longitude 79.32'E) during the Rabi season, 2023 under open field conditions. Uniform and healthy rooted cuttings of Crossandra variety Arka Chenna were used for the planting. Before planting, the soil was sterilised with 40% formalin solution to eliminate all the existing microbial population in the soil. After 3 days of covering the soil with the tarpaulin sheet, the soil was exposed to the sunlight for 3 days continuously to get rid of the fumes. Then, the soil was used for planting in the polybags of size  $13 \times 13$  inches. The experiment followed a statistical design of 2 factorial RBD, with 60 treatment combinations and 3 replications. Each replication consisted of five polybags containing single rooted cutting per polybag. The details regarding the factors were listed below:

#### Factor-1: Method of application

- $\rm M_{\scriptscriptstyle 1}$  Soil application with PGPRE's 10 days prior to Crossandra seedling planting
- $\mathrm{M}_{2}$  Inoculation of PGPRE's to cuttings prior to planting
- M<sub>3</sub>- Seedling root inoculation with PGPRE's through root treatment just prior to planting
- ${
  m M_4} ext{-}$  Seedling root inoculation with PGPRE's through soil drenching on  $10^{\rm th}$  day after transplanting
- ${
  m M_5}$  Seedling root inoculation with PGPRE's through soil drenching on  $20^{\rm th}$  day after transplanting
- $M_6$  Seedling root inoculation with PGPRE's through soil drenching on  $30^{th}$  day after transplanting

## Factor-2: Plant Growth Promoting Root Endophytes (Treatments)

- T<sub>1</sub>- Piriformospora indica
- T<sub>2</sub>- Glomus fasciculatum
- T<sub>3</sub>- Arka Microbial Consortium
- T<sub>4</sub>- *Piriformospora indica* + *Glomus fasciculatum* + Arka Microbial Consortium
  - T<sub>5</sub>- Piriformospora indica + 50% RDF
  - T<sub>6</sub>- Glomus fasciculatum + 50% RDF
  - T<sub>7</sub>- Arka Microbial Consortium + 50% RDF
- T<sub>8</sub>- *Piriformospora indica* + *Glomus fasciculatum* + Arka Microbial Consortium + 25% RDF
- T<sub>9</sub>- RDF alone (Dr. YSRHU recommendation for Crossandra)
  - T<sub>10</sub>- Absolute control

#### **Procurement of materials**

The rooted cuttings of Crossandra var. Arka Chenna and Arka Microbial Consortium were procured from ICAR-IIHR, Bengaluru. While other PGPRE's *viz.*, *Piriformospora indica* was procured from AMITY University, Noida and *Glomus fasciculatum* was procured from Agricultural Research Station, Amaravathi, Andhra Pradesh.

#### Floral studies

Flowering parameters like early spike initiation, number of days to first floret bud opening, early spike formation, number of days taken to full flowering, longevity of basal florets and duration of flowering were recorded per each treatment in each replication until the end of the flowering cycle.

#### Statistical analysis

All the data recorded for floral parameters were analyzed using two-way ANOVA under a two-factorial RBD. Comparison was done at 5% level of probability (p<0.05) to determine significant differences among the treatments. The critical difference (CD) values were determined to differentiate the means of substantially different treatments.

#### **Results**

All the floral parameters were recorded throughout the cropping period. Analysis of the data revealed that there was significant difference among the methods of inoculation, PGPRE's and the interactions ( $p \le 0.05$ ).

From the study, it was clear that among the methods of inoculation, M<sub>1</sub> (Soil application with PGPRE's 10 days prior to Crossandra seedling planting) was shown significant response and recorded early spike initiation (122.81 days), early opening of first floret bud (139.98)

days), quick spike formation (154.81 days), complete flowering (165.38 days), longevity of basal florets (7.61 days) and more duration of flowering (187.66 days). While,  ${\rm M_2}$  (Inoculation of PGPRE's to cuttings prior to planting) took more number of days for first spike initiation (135.98 days), number of days for first floret bud opening (152.36 days), complete spike formation (165.33 days), full flowering (180.23 days), less longevity of basal florets (4.84 days) and less duration of flowering (158.67 days).

Among the PGPRE's, the combinational application of PGPRE's i.e., P. (Piriformospora indica+ Glomus fasciculatum+ Arka Microbial Consortium + 25% RDF) elicited a statistically significant impact on early spike initiation (122.30 days), early opening of first floret bud (138.20 days), longevity of basal florets (8.94 days) and more duration of flowering (218.81 days) except days taken to full spike formation and complete flowering. Early full spike formation (150.69 days) was found in P (Piriformospora indica+ Glomus fasciculatum+ Arka Microbial Consortium), while early days to complete flowering (162.54 days) was exhibited in P<sub>7</sub> (Arka Microbial Consortium + 50% RDF). Meanwhile, P<sub>10</sub> (Absolute control) took long time for first spike initiation (145.09 days), number of days for first floret bud opening (162.07 days), complete spike formation (176.76 days), full flowering (197.00 days), less longevity of basal florets (2.96 days) and less duration of flowering (110.98 days).

The interactional treatment of  $\rm M_1P_8$  (Soil application with PGPRE's 10 days prior to Crossandra seedling planting x *Piriformospora indica+ Glomus fasciculatum+* Arka Microbial Consortium + 25% RDF) recorded a consistent and pronounced enhancement in almost all the floral parameters viz., early spike initiation (109.67 days), early opening of first floret bud (126.22 days), longevity of basal florets (11.89 days) and more

**Table 1:** Effect of methods of inoculation, Plant Growth Promoting Root Endophytes (PGPRE's) and their interaction on number of days taken to early spike initiation in Crossandra var. Arka Chenna.

Method of				Plant G	owth Pro	vth Promoting Root Endophytes (P)						
inoculation (M)	P <sub>1</sub>	P <sub>2</sub>	$\mathbf{P}_{3}$	P <sub>4</sub>	P <sub>5</sub>	P <sub>6</sub>	P <sub>7</sub>	P <sub>8</sub>	P <sub>9</sub>	P <sub>10</sub>	Mean M	
$M_{_1}$	118.67	123.67	118.44	119.56	119.78	127.56	113.78	109.67	139.89	137.11	122.81	
$M_2$	137.78	127.00	127.78	132.22	137.33	134.44	129.67	129.56	149.11	154.89	135.98	
$M_3$	124.33	123.00	125.89	120.44	120.33	130.11	118.78	118.22	138.67	148.11	126.79	
$M_4$	119.22	126.56	134.00	124.89	129.00	130.89	121.00	120.78	137.22	153.44	129.70	
$M_{5}$	130.67	123.00	122.11	126.44	122.22	138.00	138.56	130.33	133.22	134.00	129.86	
$M_{6}$	132.11	131.44	132.11	121.00	123.67	139.00	123.44	125.22	132.22	143.00	130.32	
Mean P	127.13	125.78	126.72	124.09	125.39	133.33	124.20	122.30	138.39	145.09		
Source	Met	thod of inc	culation (	(M)	PGPRE's (P)			MxP				
S.Em±		2.2	31			2.8	880			7.054		
CD (P=0.05)		6.2	47			8.0	065			19.755	5	

<b>Table 2:</b> Effect of methods of i	inoculation, Plant Growth Prom	noting Root Endophytes (PGPRE's) and their interaction on	on
number of days taken	to early floret bud initiation in Cr	Crossandra var. Arka Chenna.	

Method of				Plant G	rowth Pro	moting R	oot Endop	hytes (P)				
inoculation (M)	P <sub>1</sub>	$\mathbf{P}_{2}$	$\mathbf{P}_{3}$	P <sub>4</sub>	$\mathbf{P}_{_{5}}$	P <sub>6</sub>	P <sub>7</sub>	P <sub>8</sub>	P <sub>9</sub>	P <sub>10</sub>	MeanM	
M <sub>1</sub>	118.67	123.67	118.44	119.56	119.78	127.56	113.78	109.67	139.89	137.11	122.81	
$M_2$	137.78	127.00	127.78	132.22	137.33	134.44	129.67	129.56	149.11	154.89	135.98	
$M_3$	124.33	123.00	125.89	120.44	120.33	130.11	118.78	118.22	138.67	148.11	126.79	
$M_4$	119.22	126.56	134.00	124.89	129.00	130.89	121.00	120.78	137.22	153.44	129.70	
$M_{5}$	130.67	123.00	122.11	126.44	122.22	138.00	138.56	130.33	133.22	134.00	129.86	
$M_{_6}$	132.11	131.44	132.11	121.00	123.67	139.00	123.44	125.22	132.22	143.00	130.32	
Mean P	127.13	125.78	126.72	124.09	125.39	133.33	124.20	122.30	138.39	145.09		
Source	Met	thod of inc	culation (	(M)	PGPRE's (P)				MxP			
S. Em±		2.2	31			2.8	80			7.054		
CD (P=0.05)		6.2	47			8.0	65			19.755	5	

**Table 3:** Effect of methods of inoculation, Plant Growth Promoting Root Endophytes (PGPRE's) and their interaction on number of days taken to full spike initiation in Crossandra var. Arka Chenna.

Method of	Plant Growth Promoting Root Endophytes (P)												
inoculation (M)	P <sub>1</sub>	P <sub>2</sub>	$\mathbf{P}_{3}$	$\mathbf{P}_{_{4}}$	$\mathbf{P}_{_{5}}$	$P_6$	P <sub>7</sub>	P <sub>8</sub> P <sub>9</sub> P <sub>10</sub> 4         152.56         175.22         178.44           0         159.67         177.67         185.67           6         151.33         169.22         180.44           2         165.33         173.78         171.11           8         153.78         166.89         171.00           2         149.22         160.00         173.89           4         155.31         170.46         176.76           MxP	MeanM				
$\mathbf{M}_{_{1}}$	146.78	149.78	146.56	143.33	152.44	161.56	141.44	152.56	175.22		154.81		
$M_2$	165.22	153.33	166.11	159.33	165.78	161.56	159.00	159.67	177.67	185.67	165.33		
$M_3$	153.44	152.56	151.22	145.89	154.67	159.44	147.56	151.33	169.22	180.44	156.58		
$M_4$	150.44	163.44	161.33	147.11	158.56	169.67	150.22	165.33	173.78	171.11	161.10		
$M_{5}$	161.11	148.56	163.44	158.44	157.22	159.22	164.78	153.78	166.89	171.00	160.44		
$M_6$	164.78	158.44	155.33	150.00	151.44	158.00	152.22	149.22	160.00	173.89	157.33		
Mean P	156.96	154.35	157.33	150.69	156.69	161.57	152.54	155.31	170.46	176.76			
Source	Met	thod of inc	culation (	(M)		PGPRE's (P)				MxP			
S. Em±		2.2	51			2.9	06			7.117			
CD (P=0.05)		6.3	03			8.1	37			19.932	2		

**Table 4:** Effect of methods of inoculation, Plant Growth Promoting Root Endophytes (PGPRE's) and their interaction on number of days taken to full flowering in Crossandra var. Arka Chenna.

Method of				Plant Gr	owth Pro	moting R	oot Endop	hytes (P)			
inoculation (M)	P <sub>1</sub>	P <sub>2</sub>	$\mathbf{P}_{_{3}}$	$\mathbf{P}_{_{4}}$	P <sub>5</sub>	P <sub>6</sub>	P <sub>7</sub>	P <sub>8</sub>	P <sub>9</sub>	P <sub>10</sub>	MeanM
$\mathbf{M}_{_{1}}$	156.78	159.78	156.56	155.33	162.44	171.56	151.44	162.56	193.22	184.11	165.38
$M_2$	175.22	163.33	179.00	181.56	175.78	171.56	169.00	169.67	196.00	221.22	180.23
$M_3$	163.44	162.56	161.22	157.44	164.67	169.44	157.56	161.33	187.22	194.89	167.98
$M_4$	160.44	173.44	173.44	160.33	168.56	179.67	160.22	175.33	192.22	192.33	173.60
$M_{5}$	171.11	158.56	174.22	171.56	167.22	169.22	174.78	163.78	184.44	195.11	173.00
$M_6$	174.78	168.44	165.33	177.78	161.44	168.00	162.22	159.22	177.44	194.33	170.90
Mean P	166.96	164.35	168.30	167.33	166.69	171.57	162.54	165.31	188.43	197.00	
Source	Met	thod of inc	culation (	<b>M</b> )	PGPRE's (P)				MxP	•	
S.Em±		2.6	10	·	·	3.3	669			8.253	
CD (P=0.05)		7.3	09			9.4	35			23.112	2

duration of flowering (241.11 days) except days taken to full spike formation and complete flowering. While, days taken to full spike formation (141.44 days) and early days to complete flowering (151.44 days) was exhibited in

 $\rm M_{_1}P_{_7}$  (Soil application with PGPRE's 10 days prior to Crossandra seedling planting x Arka Microbial Consortium + 50% RDF). However,  $\rm M_{_2}P_{_{10}}$  (Inoculation of PGPRE's to cuttings prior to planting x Absolute control) took

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Method of	Plant Growth Promoting Root Endophytes (P)												
inoculation (M)	P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>	P <sub>4</sub>	$\mathbf{P}_{_{5}}$	$P_6$	P <sub>7</sub>	P <sub>8</sub>	P <sub>9</sub>	P <sub>10</sub>	MeanM		
$M_{_1}$	175.78	183.44	185.56	228.89	202.00	196.78	222.67	241.11	134.00	106.33	187.66		
$M_2$	195.00	178.67	153.11	189.78	179.33	125.78	150.33	209.56	103.33	101.78	158.67		
$M_3$	157.22	161.67	181.78	201.56	195.78	182.67	214.56	228.44	137.78	107.11	176.86		
$M_4$	147.44	163.56	121.56	198.67	153.22	175.33	172.56	200.11	136.44	127.44	159.63		
$M_{5}$	151.00	169.78	143.00	201.67	152.78	159.78	183.89	218.89	133.33	105.89	162.00		
$M_6$	173.44	200.11	155.78	199.67	182.22	172.89	160.56	214.78	130.56	117.33	170.73		
Mean P	166.65	176.20	156.80	203.37	177.56	168.87	184.09	218.81	129.24	110.98			
Source	Met	thod of inc	oculation (	(M)	PGPRE's (P)			MxP	)				
S.Em±		0.4	62			0.5	97			1.462			
CD (P=0.05)		1.2	95			1.6	71			4.094			

Table 5: Effect of methods of inoculation, Plant Growth Promoting Root Endophytes (PGPRE's) and their interaction on flowering duration in Crossandra var. Arka Chenna.

Table 6: Effect of methods of inoculation, Plant Growth Promoting Root Endophytes (PGPRE's) and their interaction on longevity of basal florets on the spike (days) in Crossandra var. Arka Chenna.

Method of				Plant G	rowth Pro	moting R	oot Endop	hytes (P)				
inoculation (M)	P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>	P <sub>4</sub>	$\mathbf{P}_{_{5}}$	P <sub>6</sub>	P <sub>7</sub>	P <sub>8</sub>	P <sub>9</sub>	P <sub>10</sub>	MeanM	
$M_{_1}$	6.89	7.56	6.22	10.44	7.33	10.11	8.67	11.89	3.56	3.44	7.61	
$M_2$	4.22	6.22	3.22	7.44	4.56	3.56	4.11	8.44	4.56	2.11	4.84	
$M_3$	6.78	8.78	5.78	10.11	6.56	5.11	6.56	10.22	2.89	2.67	6.54	
$M_4$	5.33	4.56	4.22	8.11	4.56	4.89	8.56	9.44	4.56	3.33	5.76	
$M_{5}$	4.33	4.44	3.22	5.56	6.11	4.78	8.67	4.56	4.56	3.00	4.92	
$M_{_6}$	5.78	6.11	4.11	9.78	5.44	4.89	4.89	9.11	3.78	3.22	5.71	
Mean P	5.56	6.28	4.46	8.57	5.76	5.56	6.91	8.94	3.98	2.96		
Source	Met	thod of inc	culation (	(M)	PGPRE's (P)					MxP		
S. Em±		0.0	92			0.1	.19			0.292		
CD (P=0.05)		0.2	59			0.3	334			0.818		

maximum number of days took for first spike initiation (154.89 days), number of days for first floret bud opening (177.44 days), complete spike formation (185.67 days), full flowering (221.22 days), less longevity of basal florets (2.11 days) and less duration of flowering (101.78 days).

#### **Discussion**

Perusal of data from the study clearly showed that all of the floral parameters were performed superiorly under the interaction treatment of M<sub>1</sub>P<sub>8</sub> (Soil application with PGPRE's 10 days prior to Crossandra seedling planting x Piriformospora indica+ Glomus fasciculatum+ Arka Microbial Consortium + 25% RDF) except days taken to full spike formation and complete flowering. Due to pre-planting microbial application, bioavailability of phosphorus- a key nutrient for the development of floral primordia, nitrogen, zinc and boron was enhanced and ready for uptake when the plant was introduced. This enables in early transformation of plants from vegetative phase to reproductive phase and early spike initiation and early first floret bud opening (Bohra and Kumar, 2014). This result was in favour with Pathak and Kumar (2009) in gladiolus. According to Goswami et al. (2016), better establishment of microbes and higher microbial activity was observed when the microbes were applied early, which primes the plant for early reproductive transition. Patil and Narayana (2018) and Kusagur et al. (2022) found that combined application of microbes along with fertilizers resulted in early flowering. More duration of flowering might be due to application of Piriformospora indica which helps in altering C:N ratio and balancing the management of vegetative as well as reproductive phase and enhances flowering days (Noorjahan et al., 2018). Similar results were found by Naik (2015) in marigold. Bohra and Kumar (2014) reported that the extension of blooming length is related to enhanced protein synthesis, accelerated nutrient mobilisation and the inhibition of chlorophyll degradation, resulting from adequate nutrient availability in soils supplemented with VAM. The improved longevity of basal florets was due to presence of ethylene inhibitors or cytokinins (Pathak and Kumar, 2009) or due to delayed senescence, induced by microbial regulation of ethylene biosynthesis and antioxidant activity as exhibited by Tongon *et al.* (2015) in snapdragon flowers.

#### Conclusion

Till now, no research had done with the combination of Piriformospora indica, Glomus fasciculatum and Arka Microbial Consortium in Crossandra. Based on the findings, the plants treated with the combination of these three PGPRE's along with supplemented nutrients particularly through soil application of 10 days prior to planting enhances all the floral attributes than control. This study suggested that early application of PGPRE's will result in early microbial colonisation in the rhizosphere, united with the synergistic generation of plant growth promoting chemicals like auxins and cytokinins, making the environment favourable for vigorous plant growth and development. These experimental findings highlighted the interesting utilisation of microbial consortia in sustainable floriculture. Using these microbial inoculants not only improves the crop quality and production, but also eliminates the need of chemical fertilizers, thereby making it a feasible and eco-friendly approach for commercial producers. Furthermore, the incorporation of bioinoculants with diminished fertiliser application is a sustainable alternative to traditional high-input floricultural methods.

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