

Isolation And Evaluation Of Bacteria Exhibiting Multiple Plant Growth Traits In The Rhizosphere Of Yellow Bell Pepper (*Capsicum chinense*)

ABSTRACT

AIM: The study identified and evaluated bacteria exhibiting multiple plant growth traits in the rhizosphere of Yellow Bell Pepper (*Capsicum chinense*)

Study Design; Seeds of *Capsicum chinense* were planted in a soil and allowed to grow. After five weeks of planting, soil samples from the rhizosphere were collected and the bacterial community present in the rhizosphere soil of *Capsicum chinense* was studied. The isolated organisms were assessed for their ability to produce plant growth promoting traits.

Place and duration of study: This study was carried out at an agricultural research farmland in the Federal University of Technology, Owerri, Nigeria.

Methodology: Seeds of *Capsicum chinense* were planted in the soil samples in a greenhouse. Rhizosphere soil was collected for analysis to identify the bacterial composition of the rhizosphere soil.

Results: In this study the presence of *Bacillus cereus*, *Staphylococcus aureus*, *Corynebacterium sp*, *Enterococcus faecalis* and *Bacillus polymyxa* were evident in the rhizosphere samples collected. All isolates showed multiple plant growth promoting traits except *Staphylococcus aureus* which was positive for hydrogen cyanide production only.

Conclusion: The results from this study showed that the bacterial community present in the soil can be used to effect significant vegetative crop yield and agricultural production. The isolated rhizobacteria can be formulated as bio-fertilizers or bioinoculants, etc.

1. INTRODUCTION

Peppers are an important source of nutrients in human diet [1,2]. The *Capsicum* peppers, including sweet peppers, bell peppers, hot peppers like jalapeño cayenne, serrano, cherry and many others [3] are the most worldwide cultivated [4] and are widely appreciated in the culinary as spice. These peppers are characterized by their high levels of vitamin C (ascorbic acid), provitamin A (carotene) and calcium. Ingestion of 50-100 g fresh pepper fruits can provide about 100% and 60% of recommended daily amounts of vitamin C and A, respectively. Ripe fruits of pepper are also rich in compounds with antioxidant and anticancer activity [5].

A major factor influencing plant growth and health is soil fertility which also determines the microbial population living both in rhizosphere and as endophytes within healthy plant

38 tissues. Soil fertility refers to nutrient amount in soil capable of supporting plant life [6] and
39 largely depends on micro and **macronutrients** and micro and **macroorganisms**.

40 Soil microorganisms are very important in almost every chemical transformation taking place
41 in soil. They play an active role in maintaining soil fertility as a result of their involvement in
42 nutrient synthesis and circulation. The presence of these microorganisms in **soil rhizosphere**
43 largely counts for microbial community present in soil.

44 Microbial population in and around the roots includes bacteria, fungi, yeast etc. Some are free
45 living while others form symbiotic relationships with various plants [7]. The community
46 structure of soil microorganisms in rhizosphere differs from that in non-rhizosphere soil
47 largely due to biological interactions between microorganisms and the **plant's** roots [8].

48 These biological interactions accounts for plant growth and improved soil fertility. The
49 bacterial community can be seen to synthesize nutrients and compounds that can be used to
50 enhance plant growth. Plant Growth Promoting Bacteria characterized with their fast
51 metabolism and growth are always readily colonising the root surface [7]. This makes them
52 suitable as biofertilizers, seed treatments and as biocontrol agents.

53 This study was aimed at evaluating the plant growth promoting traits of bacteria isolated in
54 the *Capsicum chinense* rhizosphere.

55 **2. MATERIALS AND METHODS**

56 **2.1 Study Area**

57 The study was carried out in the Research Farmland of School of Agriculture and
58 Agricultural Technology, Federal University of Technology, Owerri, Imo State, Nigeria.

59 **2.2 Collection of Samples.**

60 **2.2.1 Soil sample**

61 Soil samples were collected randomly from an uncultivated portion of farmland to 15-30cm
62 below the surface. The collected soil samples were bulked to form a composite sample and 5
63 kg each was measured and stored in separate polythene bags in which the pot planting
64 experiment was carried out.

65 **2.2.2. Collection of Yellow bell pepper seeds**

66 The yellow pepper seeds were obtained in sealed plastic bags from Imo Agricultural
67 Development Program [ADP] Centre, Owerri, Imo state, Nigeria.

68 **2.3 Planting of Seed**

69 *Capsicum chinense* seeds (5 seeds per bag) were planted in bags containing soil samples
70 collected at random from farmland and allowed to grow for five weeks.

71 **2.4. Isolation of microorganisms**

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73 The rhizospheric soil samples of growing yellow bell pepper seeds were aseptically collected
74 and introduced into different sterile test tubes, properly labelled and taken to the laboratory

75 for microbiological investigation. **Microorganisms were isolated** by using **the** spread plate
76 method according to Cheesbrough [8]. The nutrient agar plates were incubated at 37°C for 24
77 hours. The culture plates were observed for microorganism growth.

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79 **2.5. Identification of Microorganisms**

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81 The bacterial isolates were identified by using cultural, morphological and biochemical
82 characteristics as described by Cheesbrough [9]

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84 **2.6. Evaluation of plant growth promoting traits**

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86 **IAA production:** IAA production by the isolates was estimated by using Salkowaskis
87 reagent. Appearance of pink color indicated IAA production which was read at 535nm [10].

88 **Phosphate solubilization activity:** All bacterial isolates were screened for inorganic
89 phosphate solubilization. Qualitative estimation was done by using Pikovskaya medium
90 containing tri-calcium phosphate, iron phosphate. Positive results **were** recorded by formation
91 of **clear halo zone** around the culture [11].

92 **Hydrogen cyanide production**

93 The production of HCN was detected by spreading 1 ml of 24 h old broth culture on King's B
94 medium supplemented with 4.4g/l glycine and incubated with Whatmann filter paper flooded
95 with solution containing 0.5% picric acid in 2% sodium carbonate. After 24-48 h, yellow to
96 brown change in color of the filter paper was observed [12].

97 **Ammonia production**

98 All the bacterial isolates were tested for the ammonia production of using Nessler's reagent.

99 Ammonia production **was** detected by formation of faint yellow to dark brown color [13].

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101 **3. RESULTS AND DISCUSSION**

102 The increasing importance of beneficial bacteria in agriculture has resulted in many efforts
103 to isolate and identify bacteria associated with the soil and rhizosphere of plants, in order to
104 identify their roles in plant growth promotion and protection against pathogens. The
105 application of PGPR has a potentially attractive approach to disease management and
106 improved crop productivity in sustainable agriculture.

107 Bacterial analysis of rhizosphere soil showed the presence of mostly gram-positive
108 organisms. Results in Tables 1 and 2 reveals the morphological and biochemical
109 characteristics of the bacterial isolates. The bacterial isolates were *Bacillus cereus*, *Bacillus*
110 *polymyxa*, *Enterococcus faecalis*, *Corynebacterium* sp. and *Staphylococcus aureus*.

111 This implies that microorganisms that were isolated from the plant rhizosphere are
112 pathogenic and potentially toxin-producing microorganisms which can lower the quality of
113 Yellow pepper plants and can also be responsible for causing pepper diseases.
114 Similar work by Hanna et al. [14] also revealed the isolation of *Pseudomonas* and *Bacillus*
115 spp. *Bacillus* and *Pseudomonas* spp. are the most frequently reported genera of PGPR
116 [15,16,17].
117 The plant growth promoting characteristics were examined with ten selected PGPR isolates.
118 Table 3 shows the plant growth promoting potentials of the bacterial isolates. All isolates
119 showed multiple Plant Growth Promoting (PGP) trait except *Staphylococcus aureus* which
120 was positive for **Hydrogen Cyanide production** only.
121 Hydrogen cyanide production was found to be the most frequent trait exhibited by *Bacillus*
122 *cereus* and *Enterococcus faecalis* while Ammonia production was exhibited mostly by
123 *Bacillus cereus*, *Bacillus polymyxa* and then *Corynebacterium* sp. Bacterial plant growth
124 promotion is a well-established and complex phenomenon that often is achieved by activities
125 of more than one PGP trait exhibited by plant isolated bacteria [18]. In this study, 80% of
126 isolates exhibited more than two PGP traits **that** may promote plant growth directly, indirectly
127 or synergistically. Similar to these findings, multiple PGP activities among Plant Growth
128 Promoting Rhizobacteria has been found in some bacteria including species of
129 *Pseudomonas*, *Azospirillum* sp., *Azotobacter* sp and *Serratia* sp etc. and they have been
130 reported to enhance plant growth [19]. Hartmann et al. [18] had reported that some studies
131 suggest that PGPR enhances the growth, seed emergence, crop yield and contribute to the
132 plant protection against certain pests and pathogens as well as nutrient availability.
133 Indole Acetic Acid is effective in root growth and development, fruit growth and
134 development, apical dominance and flowering [7]. Similar studies have shown that IAA
135 production is very common among PGPR [20, 21, 22, 23, 24, 17]. The production of
136 producing hydrogen cyanide by some of the **rhizobacteria isolated**, which several studies
137 have attributed a disease protective effect, is a very strong indication of biocontrol potentials
138 of these organisms. This is similar to how **phosphorus-solubilizing** bacteria like *Bacillus*
139 *cereus* and *Enterococcus faecalis* are effective in increasing the plant available phosphorus
140 in the soil as well as the growth yield of crops [25]. A review by Kucey et al. [26] had
141 emphasized on the ability of some phosphorus-solubilizing microbes to stimulate
142 phytopathogen biocontrol that affect plant growth via production of siderophores, hydrolytic
143 enzymes and HCN. Most isolates from this study tested was positive to the **ammonia**

144 production which has been reported as another key trait that significantly increases the crop
 145 vegetative growth and yield [27].

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147 **Table 1 Showing cell morphology and microscopic characteristics of bacterial isolates.**

Colony code	Cell morphology	Mot	Gram Stain	Spore	Flagellum	Capsule	Probable identity
YPB1	Dull, dry serrated cream colonies	+	+R	+	+	-	<i>Bacillus cereus</i>
YPB2	Serrated with medusa head	-	+R	+	-	-	<i>Bacillus polymyxa</i>
YPB3	Moist and shiny cream colonies	-	+S	-	-	-	<i>Enterococcus sp.</i>
YPB4	Dull, dry umbonate cream colonies	-	+R	-	-	-	<i>Corynebacterium sp.</i>
YPB5	Golden yellow colonies	-	+S	-	-	-	<i>Staphylococcus sp.</i>

148 **Key:** Mot= Motility, - = Negative, += Positive, +R= Positive Rod, +S= Positive Spherical,
 149 YPB= Yellow Pepper Bacterial Isolate

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160 **Table 2 :Biochemical and carbohydrate fermentation test of bacterial isolates**

Colony Code	Cat	Oxi	Coag	Ind	MR	VP	Cit	TSI	NO ₃	Ure	Glu	Suc	Lac	Fru	Mal	Mann	Identity Of Isolates
YPB1	+	-	-	-	-	+	+	-	+	+	+	-	-	-	-	-	<i>Bacillus cereus</i>
YPB2	+	-	-	-	-	-	+	-	-	-	+	+	-	-	+	-	<i>Bacillus polymyxa</i>
YPB3	-	-	-	-	+	-	+	-	-	-	+	+	+	+	-	+	<i>Enterococcus faecalis</i>
YPB4	+	-	-	-	-	+	+	-	+	-	+	-	-	-	+	-	<i>Corynebacterium sp.</i>
YPB5	+	-	+	-	-	+	-	-	+	+	+	+	+	+	+	+	<i>Staphylococcus aureus</i>

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162 Key; Cat=Catalase, Coag= Coagulase, Oxi=Oxidase, Ind=Indole, MR=Methyl Red,
 163 VP=VogesProskauer,Cit= Citrate Utilization, Ure= Urease Production, NO₃=Nitrate
 164 Production, TSI= Triple Sugar Iron Test, Glu= Glucose, Suc=Sucrose, Mal=Maltose,
 165 Lac=Lactose, Mann=Mannitol, Fru=Fructose.

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167 **Table 3 showing plant growth promoting potentials of the bacterial isolates**

Isolates	HCN	NH ₃	IAA	PO ₄
<i>Bacillus cereus</i>	++	++	+	++
<i>Bacillus polymyxa</i>	+	++	+	+
<i>Enterococcus faecalis</i>	++	+	+	+
<i>Corynebacterium sp.</i>	+	++	-	+
<i>Staphylococcus aureus</i>	+	-	-	-

168 **Key;** HCN= Hydrogen Cyanide Production, NH₃= Ammonia Production, IAA= Production
 169 of Indole Acetic Acid, PO₄= Phosphate Solubilisation,

170 += Positive, - = Negative

171 **Note:**The positive reaction intensity is indicated by the number of (+) symbols.

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175 5. CONCLUSION

176 The use of PGPR inoculants to improve agricultural production is a dynamic process and one
 177 with a wide range of capabilities. This study isolated bacterial isolates that demonstrated

178 PGPR traits. These soil microbes are active elements for soil development and in long run

179 pushes for sustainable agricultural practices. Taken together; these results suggest that PGPR
180 are able of inducing IAA production, phosphate solubilization and resistance to fungal
181 pathogens, thereby improving plant growth. The potentials of these strains may be applied to
182 enhance the growth and yield of yellow bell pepper. Due to the diverse nature of PGPR
183 strains, instead of one strain, two or more strains with multiple PGP traits can be used as
184 biofertilizer which is an efficient approach to replace chemical fertilizers and pesticides for
185 sustainable pepper cultivation. Further investigations, including tests under field conditions,
186 are needed to ascertain the role of PGPR as biofertilizers that exert beneficial effects on plant
187 growth and development.

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189 6. COMPETING INTERESTS. There are no competing interests

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193 7. REFERENCES

194

- 195 1. Bossland PW, Votava EJ. Peppers: Vegetable and Spice. Capiscums. 2000; 1:204.
- 196
- 197 2. Shetty AA, Magadam S, Managanvi K. Vegetables as sources of antioxidants. Journal
198 of Food & Nutritional Disorders. 2013; 2(1):1-5.
- 199 3. Alvarez-Parrilla E, De La Rosa LA, Amarowicz R, Shahidi F. Protective effect of
200 fresh and processed Jalapeño and Serrano peppers against food lipid and
201 human LDL cholesterol oxidation. Food Chemistry. 2012; 133(3):827-834
- 202 4. Hwang D G, Park JH, Lim, JY, Kim D, Choi Y, Kim S, Reeves G, Yeom SI, Lee JS,
203 Park M, Kim S, Choi IY, Choi D, Shin, C. The hot pepper (*Capsicum*
204 *annuum*) micro RNA transcriptome reveals novel and conserved targets: a
205 foundation for understanding Micro RNA functional roles in hot pepper. PLoS
206 One. 2013; 8(5)
- 207 5. Mateos RM, Jiménez A, Román P, Romojaro F, Bacarizo S, Leterrier M, Gómez M,
208 Sevilla F, Del Río LA, Corpas FJ, Palma JM. Antioxidant systems
209 from pepper (*Capsicum annuum* L.): Involvement in the response to
210 temperature changes in ripe fruits. International Journal of Molecular
211 Sciences. 2013; 14(5):9556-9580

- 212 6. Jilani G, Akram, A, Ali RM, Hafeez FY, Shamsi IH, Chaudhry AN, Chaudhry AG.
213 Enhancing crop growth, nutrients availability, economics and beneficial
214 rhizosphere microflora through organic and biofertilizers. *Ann. Microbiol.*
215 2007; 57(2): 177-183.
- 216 7. Wiley JM, Sherwood LM, Woolverton CJ. Isolation of pure culture. *Prescott's*
217 *Microbiology*. McGraw Hill, New York. 2011; 149-152.
- 218 8. Van LLC. Plant responses to plant growth-promoting rhizobacteria. *Eur J Plant*
219 *Pathol.* 2007; 119: 243-254.
- 220 9. Cheesbrough M. Microbiological tests. *District Laboratory Practice in Tropical*
221 *Countries*. Part 2. University press, Cambridge. 2000; 37-71.
- 222 10. Kumar K, Amaresan N, Bhagat S, Madhuri K, Srivastava RC. Isolation and
223 characterization of rhizobacteria associated with coastal agricultural
224 ecosystem of rhizosphere soils of cultivated vegetable crops. *World Journal*
225 *of Microbiology and Biotechnology*, 2011;27: 1625-1632.
- 226 11. Pikovskaya RE. Mobilization of phosphorus in soil in connection with vital activity
227 of some microbial species. *Microbiology* 1948;17: 362-370
- 228 12. Castric PA. Hydrogen cyanide, a secondary metabolite of *Pseudomonas aeruginosa*.
229 *Journal of Microbiology*. 1975;16: 793-796.
- 230 13. Cappuccino JG, Sherman N. Biochemical activities of microorganisms: a Lab
231 manual. *Journal of Microbiology*, 1992;14:1123-1245.
- 232 14. Hanna R, Melgorzata B, Agata GS. Cultivable microorganisms inhabiting the aerial
233 parts of *Hypericum perforatum*. *ACTA Sci.Pol., Hortum Cultum*.
234 2014;13(5):117-129
- 235 15. Laguerre G, Attard M R, Revoy F, Amarger N. Rapid identification of Rhizobia by
236 restriction fragment length polymorphism analysis of PCR amplified 16S
237 rRNA genes. *Appl. Environ. Microbiol.* 1994; 60: 56-63.
- 238 16. Hallmann J, Berg G. Spectrum and population dynamics of bacterial root endophytes.
239 In: *Microbial Root Endophytes*. eds Schulz B, Boyle C, Sieber T. Springer,
240 Heidelberg. 2006; 15-31.
- 241 17. Zahid M, Abbasi MK, Hameed S, Rahim N. Isolation and identification of indigenous
242 plant growth promoting rhizobacteria from Himalayan region of Kashmir and
243 their effect on improving growth and nutrient contents of maize (*Zea mays L.*). *Front.*
244 *Microbiol.* 2015; 6:207

- 245 18. Hartmann A, Schmid M, Van TD, Berg G. Plant-driven selection of microbes. Plant
246 Soil. 2009; 321:235–257.
- 247 19. Kloepper JW, Leong, J, Teintze M, Schroth MN. Enhanced plant growth promoting
248 rhizobacteria. Nature.1990; 286:885-886.
- 249 20. Yasmin S, Bakar MAR, Malik KA, Hafeez FY. Isolation, characterization and
250 beneficial effects of rice associated plant growth promoting bacteria from
251 Zanzibar soils. J. Basic Microbiol.2004; 44:241–252
- 252 21. Ahad F, Ahmad I, Khan M. Screening of free-living rhizospheric bacteria for their
253 multiple plant growth promoting activities. Microbial Res, 2008;165(2):173-
254 181.
- 255 22. Nihorimbere V, Ongena M, Smargiassi M, Thonart P. Beneficial effect of the
256 rhizosphere microbial community for plant growth and health. Biotechnol.
257 Agron. Soc. 2011). 15:327–337.
- 258 23. Ng LC, Sariah M, Sariam O, Radziah O, Abi MAZ. Rice seed bacterization for
259 promoting germination and seedling growth under aerobic cultivation system.
260 Aust. J. Crop. Sci. 2012; 6 170–6175.
- 261 24. Dalal J, Kulkarni N. Antagonistic and plant growth promoting potentials of
262 indigenous endophytic bacteria of soybean (*Glycine max*(L). Current Research
263 in Microbiology and Biotechnology 2013;1(2): 62-69
- 264 25. Arshad M, Frankkeberger Jr. WT. Microbial production of plant growth regulators.
265 Soil Microbial Ecology. 2005; 307:34-37
- 266 26. Kucey RMN, Jenzen HH, Leggett ME. Microbial mediated increases in plant
267 available phosphorus. *AdvAgron*, 2003; 42:199-228.
- 268 27. Kennedy AC, Smith KL. Soil microbial diversity and sustainable agricultural soils.
269 Plant soil.1995;170:75-86
- 270
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