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J.S.S ARTS, SCIENCE AND COMMERCE DEGREE COLLEGE

GOKAK



PROJECT WORK

ISOLATION AND IDENTIFICATION OF AZOTOBACTER FROM SOIL

NAME I VITTHAL HOSUR

CLASS: B.Sc VI Sem

REG NO: S191,7247

MENTOR: Prof. Dr. T.C. Gopal

Prof R M Mahendrakar





S.S. ARTS, SCIENCE, COMMERCE, B.B.A. COLLEGE AND P.G. DEPARTMENTS, GOKAK- 591307

DEPARTMENT OF BOTANY

CERTIFICATE

This is to certify The Project report, submitted by

Mr.

VITTHAL HOSUR Reg No. 81917247 for the Partial fulfillment of Practical in Botany, VI semester for year 2021-22 as prescribed Rani Channamma University Belgavi, is based on ISOLATION AND IDENTIFICATION OF AZOTOBACTER FROM SOIL

conducted under my supervision.

Prof. D. T.C.Gopal

M.Mahendrakar

ISOLATION AND IDENTIFICATION OF AZOTOBACTER FROM SOIL OF Kuligod region REGION

Introduction

Azotobacter in soil: In Indian soils, the population of Azotobacter is not more than 10 thousand to 1 lakh/g of soil. The population of Azotobacter is mostly influenced by other microorganisms present in soil. There are some microorganisms, which stimulate the Azotobacter population in soil thereby increasing the nitrogen fixation by Azotobacter. On the other hand there are some microorganisms, which adversely affect the Azotobacter population and hence nitrogen fixation process is hampered. For example Cephallosporium is most commonly found organisms in soil, which restricts the growth of Azotobacter. Azotobacter also produces some substances, which check the plant pathogens such as Alternaria, Fusarium and Helminthosporium. Hence Azotobacter also acts as a biological control agent.

of Azotobacter: Azotobacter Functions naturally fixes atmospheric nitrogen in the rhizosphere. There are different strains of Azotobacter each has varied chemical, biological and other characters. However, some strains have higher nitrogen fixing ability than others. Azotobacter uses carbon for its metabolism from simple or compound substances of carbonaceous in nature. Besides carbon, Azotobacter also requires calcium for nitrogen fixation. Similarly, a medium used for growth of Azotobacter is required to have presence of organic nitrogen, micronutrients and salt in order to enhance the nitrogen fixing ability of Azotobacter.

Sr. No.	Sample number	Name of the	organsms
	والمريد بمرهم المرتبة بمراجع والمعرجان بمعرمين		المرجبين فيجاور الاستحصور المرتجع المر
0	VMP1, VMP3,		vinelandii
71	VMP2,	/	paspali

Plate No. 3.1



Growth of Azotobacter spp. On Ashbys Mannitol Agar

Plate No. 3.2



Growth of Azotobacter spp. On Ashbys Mannitol Agar

<u>DISCUSSION</u>

The dominant nonsymblotic nitrogen fixing heterotrophic bacterium in Indian soils is Azotobacter chroococcum, A. vinelandii, A. insigns, A. armeniacus, A. nigricans and A. beljerinckii (Mulder and Brotonegoro, 1974). Azotobacter rarely exceeds 10⁴ to 10⁵ microorganisms per gram in Indian soils and cellulolytic microorganisms, which degrade plant residues, encourage the proliferation (Iswaran and Subba Rao, 1966).

The data showed that out of 10 soil samples collected, 3 positive for non-symblotic nitrogen fixing samples were Azotobacter. On the basis of morphological, cultural and blochemical characteristics of these 3 samples showed two of Azotobacter. The morphological different species characterization performed as gram staining, motility, cyst formation and pigment production with the help of Bergeys manual of determinative bacteriology. Differentiation of Azotobacter spp. were made and identified as Azotobacter

Azotobacter vinelandii and Azotobacter paspali The data further indicated that, in black soil of ______ Kuligod A. paspali were less in number as compared to A. vinelandii

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PRINCIPAL

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J.S.S Degree College Gokak

ACKNOWLEDGEMENT

We Sincerely express our deep sense of gratitude to our Prof. Mahindrakar sir, Dept. of Botany, J.S.S Degree College Gokak, for this valuable guidance, suggestions and encouragement throughout this project.

We are thankful to our HOD, Dr. T C Gopal, Dept. Of Botany, for providing necessary facilities to carryout the Project report on "ISOLATION AND IDENTIFICATION OF RHIZOBIUM FROM SOIL"

Our special thanks to our Principal Dr. A.S. Terdal for their valuable support in permitting this project report.

Date: 30/8/20 Place: Gokalc

Miss.SHRUSHTI Reg. No.S1917219





.S.S ARTS, SCIENCE, COMMERCE B.B.A COLLEGE AND P.G DEPARTMENTS, GOKAK- 591307

DEPARTMENT OF BOTANY

CERTIFICATE

This is to certify The Project report, submitted by Miss. SHRUSHTI S DAPPADHULI Reg.No: S1917219 for the Partial fulfillment of Practical in Botany, VIth semester for year 2021-22 as prescribed Rani Channamma University Belgavi, is based on "ISOLATION AND IDENTIFICATION OF RHIZOBIUM FROM SOIL OF GHATAPRABHA KARNATAKA REGION" conducted under my

supervision.

Dr. T.C. Gopal Dept. of studies in Botany Gokak L S. Science Collector GOK AK

Prof. R. M. Mahindrakar.

Dept.of studies in Botany J.S.S Degree College













J.S. S. ARTS, SCIENCE AND COMMERCE

COLLEGE, GOKAK

(AFFILIATED TO RANI CHANNAMMA UNIVERSITY, BELAGAVI)



DEPARTMENT OF BOTANY

PROJECT WORK ON WATER ANALYSIS

Topic : To Determine the Alkalinity of water

From Gokak Galls.

Class

: B.Sc Fifth Sem

Reg No

: \$3937232

Name Of The Student : Smita. A. Thorat.



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DEPARTMENT OF BOTANY

CERTIFICATE

THIS IS TO CERTIFY THAT THE PROJECT REPORT, SUBMITTED BY MAL. Smila Apparaheb. Thorat. Reg. 83917212 IN BOTANY IN 5TH SEMESTER 2021-2022 AS PRESCRIBED BY RANI CHENNAMMA UNIVERCITY BELAGAVI IS

BASED ON PROJECT REPORT.

HOD

Dr. T CGOPAL Dept of studies in botany J S S Degree College, Gokak

Jahues

Prof. Mahendrakar Sir Dept of studies in botany J S S Degree College, Gokak

Determination of water analysis

Date Tested

Tested By

: Snita A. Thorat. : To determine the alkalinity of water

Project Name

Sample Number

01 :

: 22/02/2022

Sample Location

: Crokak : River mater . Crokak falls.

Sample Description

Table -2 Total Alkalinity:

-SI No	Volume	Burette Reading (mL)		Volume of	
51.100.	of Sample (mL)	. Initial	Final	Sulphuric acid (mL)	
1.	90	0.3	4.4	4.1	
2.	20	6.0	10.2	4.2	
3.	50	10.2	14.3	4.1	

Calculation:

Volume of Sulphuric Acid: Normality of Sulphuric : Volume of Sample : Equivalent weight of CaCO2:

4.1 mL 0.02 N 50 m.L 1000

Total Alkalinity =

(volume of H2SO4(v1)* Normality * 50 * 1000) Volume of sample taken

Alkalinity as CaCO3 equivalent (mg/L)= 4.1 x 0.02 x 50 x 1000/100

= 4] mg/L as CaCO3 equivalent

T - 4.1 -> T = 41 P = 0 Sample J =

P=0. T=41 PX1/2T co3 = 2P =0 HOU3 = T-2P = 41-2×0 = 41mgl4

Determination of water analysis Data Sheet : 13/02/2023 Date Tested : Smita A Thorat : To determine the alkalingity of water

Project Name Sample Number Sample Location

Tested By

Sample Description

Table -1 Phenolphthalein Alkalinity:

: 02

: Gokak

1 and 1	Volume of Sample (mL)	Burette Reading (mL)		Volume of
SI,No.		Initial	Final	Sulphuric acid (mL)
1.	20	16	16.1	0.1
2.	90	19	19.7	0.1
3.	90	2.3	21.)	0.7

Calculation

Volume of Sulphuric Acid:	0.3 mL	
Normality of Sulphuric :	0.02 N	
Volume of Sample :	50 mL	
Equivalent weight of CaCO3: Phenolphthalein Alkalinity =	1000 (volume of H ₂ SO ₄ (v1)* Normality * 50 * 1000 Volume of sample taken	IJ
Alkalinity as CaCO3 equivalen	nt (mg/L)= x 0.02 x 50 x 1000/100	
= mg/L as CaCC	Da equivalent	

INFERENCE

Alkalinity is a measure of the capacity of water to neutralize acids. The predominant chemical system present in natural waters is one where carbonates, bicarbonates and hydroxides are present. The bicarbonate ion is usually prevalent. However, the ratio of these ions is a function of pH, mineral composition, temperature and ionic strength. Water may have a low alkalinity rating but a relatively high pH or vice versa, so alkalinity alone is not of major importance as a measure of water quality. Alkalinity is not considered detrimental to humans but is generally associated with high pH values, hardness and excess dissolved solids. High alkalinity waters may also have a distinctly flat, unpleasant taste. Based on the testing, it is found that the alkalinity of the sample is $\frac{3}{D} = 11$ wells mg/L. As per the provisional code, alkalinity should not exceed 200 mg/L for potable water. For the fresh water alkalinity ranges between 20 - 100 mg/L. Alkalinity of tested sample is within/above the limits specified in the standards.

Hence the water sample is fit / unfit for drinking. So,

Sample I 95 unfet for drenking Sample IT is fet for dranking.

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CERTIFICATE

This is to certify that the study tour report, Submitted by Sachen, Rejanagel (Reg No: S1817197), for the partial fulfillment of practical in Botany, 3rd semester for year 2019-20 as prescribed by Rani Channamma University Belagavi, is based on study tour conducted under my supervision.

HOD

Dr. T C Gopal

prof. H S Dasar

Dept. of studies in Botany

J.S.S. Degree College, Gokak



Dept. of studies in Botany

J.S.S. Degree college , Gokak

JSS ARTS SCIENCE COMMERCE DEGREE COLLEGE GOKAK



BOTANY PROJECT

METHODS USED IN EFFLUENT TREATMENT BY INDUSTRIES

1

CLASS : BSC It

SUBJECT : Botany - II

REG.NO : 51716161

Prema. V. Meti

SIGNATURE OF TEACHER

yopal 218/10/19 **SIGNATUE OF H.O.D**

APPL



J. S. S. ARTS, SCIENCE, COMMERCE, B.B.A. COLLEGE AND P.G. DEPARTMENTS, GOKAK - 591 307

DEPARTMENT OF BOTANY

PROJECT WORK

NAME: SHREYA, S. BANAKARI

REG :51616223

Staff:

INTEGRATED FARMING

TOPIC -

Prof:Mahendrakar & Sudharshan

25/20/38

Dept of studies in Botany

J.S.S Degree college Gokak

HOD:

Dr.T.C.Gopal

Dept of studies in Botany

J.S.S Degree college Gokak

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Problems of present agriculture

- Decline in agriculture growth rate
- Decline in factor productivity
- Static or decline in food production
- Increasing malnutrition
- Shrinkage in net cultivable area
- Increasing environmental pollution
- Depleting ground water table
- Increasing cost of production
- Low farm income
- Problems of Farm labours due to large scale migration

What is the solution?

"Integrated Farming System"

Source: http://planningcommission.nic.in/plans/planrel/fiveyr/1st/1planch18.html

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EETTING MORE OUT OF EVERY ACRE





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(NAAC Re-accreditation : "A" Grade Institution) Department of Botany

PROJECT REPORT ON

"TREATEMENT OF INDUSTRIAL EFFLUENTS IN A BIOREACTOR"

SUBMITTED BY:-

Savita Hosapeti REG NO. S1616194

UNDER THE GUIDANCE OF

xan

Mr. Mahendrakar

CERTIFICATE

This is to certify that the project entitled, "TREATEMENT OF INDUSTRIAL EFFLUENTS IN A BIOREACTOR" submitted by Miss Savita Hosapeti Reg No. S1616194 in partial fulfillments for the botany project, V Sem for year 2018-19 as prescribed by the Rani Channamma University Belagavi is based on "TREATEMENT OF INDUSTRIAL EFFLUENTS

HOD J Dr. T. C Gopal

Dept of studies in botany J S S Degree College Gok

Prof. Mahendrakar Dept of studies in botany J S S Degree College Gokak

Analytical methods for water quality parameters BOD-The quantity of oxygen required by the microorganisms for the stabilization of the biological decomposable organic matter. BOD tests measure the molecular oxygen utilized during a specified incubation duration for the biochemical degradation of organic material and the oxygen used to oxidize inorganic material such as ferrous iron and sulfides. The most common BOD test consists of a 5 day period in which a sample is placed in an airtight bottle under controlled conditions temperature (20°C \pm 1°C), keeping any light from penetrating the sample to prevent photosynthesis. The Dissolved Oxygen (DO) in the sample is measured before and after the 5 day incubation period, and BOD is then calculated as the difference between initial and final DO measurements. BOD can be considered a more "natural" test in determining the oxygen required to oxidize organic

COD

Chemical oxygen demand test determines the oxygen required for chemical oxydation of organic component as well as the no. of inorganic component with the help of strong oxidant i.e. potassium dichromate.

Once oxidation, is complete, the excess potassium dichromate is titrated with ferrous ammonium sulfate (FAS) until all of the excess oxidizing agent has been reduced to Cr^{3+} . Typically, the oxidation-reduction indicator Ferroin is added during this titration step as well. Once all the excess dichromate has been reduced, the Ferroin indicator changes from blue-green to reddish-brown.

COD is often preferred for daily analysis since it is inherently more reproducible, accounts for changing conditions and takes a short time to complete.

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