

Monitoring of Pollutant Levels in Stagnant and Flowing Water Bodies in and around Ahmedabad, India



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Abstract: *There are various water bodies in and around Ahmedabad, India which serve as source of water for irrigation and other recreational purposes. The stagnant water bodies available are lakes such as Kankaria lake and Vastrapur lake whereas flowing water can be seen in form of Kharicut canal, or Sabarmati river which flow across the city. With increasing anthropogenic activities, pollutants levels have increased in these water bodies. Nitrogen and phosphorus are known to be the primary causes of eutrophication (i.e., nutrient enrichment due to human activities) in those surface waters. Around 110 years ago Kharicut canal was built for providing potable water as well as for irrigation on peripheries of Ahmedabad district. Uncontrolled release of wastewater from various sources including municipal treatment plants, industrial treatment plants and common effluent treatment plants led to pollution of Kharicut canal and further adversely affected the farms and their crops. Apart from the orders given by High Court in 2016 for cleaning of canal and recent reports of the leading newspaper, Times of India also represented still worst conditions of canal. So providing adequate sanitation and clean water to villages and lakes through the Kharicut canal is a burning issue and economic challenge for the Gujarat Government. The water from various water bodies were collected and analysed for BoD, COD, TDS, Phosphate, pH, salinity etc. The water samples revealed alkaline water in all the locations. Most of the nutrients were in the range however COD in Kharicut Canal was alarming.*

Keywords: *Water purity, BOD, COD, Pollution.*

I. INTRODUCTION

There are various water bodies in and around Ahmedabad, India which serve as source of water for irrigation and other recreational. Approximately 25% of all water contamination are due to nutrient-related causes (e.g., nutrient inputs, oxygen depletion, algal growth, ammonia, harmful algal blooms affecting biological integrity and turbidity) [1]. There have been many chemical and biological approaches for removal of nutrients from the water. Neutralization and application of flocculants and coagulants like, alum and FeSO_4 have been employed as traditional processes in effluent treatment plants for phosphorus removal [2].

During wastewater treatment, even after secondary biological treatment by activated sludge system where degradable organics are removed, much amount of non-biodegradable organic and inorganic pollutants remain in effluents contributing to high chemical oxygen demand (COD) levels. These high COD and BOD values in treated effluent usually exceed the maximum permissible limit. Hence this water cannot be used even for irrigation purpose. Usually according to Gujarat Pollution Control Board (GPCB) guidelines, the permissible limit of COD and BOD in industrial wastewater is 250 mg/ml and 30 mg/ml respectively [3].

The various studies on river contamination in India are reported. Gopal et al., 2021 collected and analysed water samples from 35 locations in the Arkavathi river, its tributaries and lakes spread across the Bengaluru metropolitan area in Southern India during monsoon. They reported that Ibuprofen was present in high concentrations (1834 ng/L) followed by triclosan (1761 ng/L) and diclofenac (1130 ng/L) in the river [4]. Williams et al., 2019 carried out longitudinal survey of a small Indian city's river receiving untreated wastewater wherein emerging contaminants were detected and concluded that minimal attenuation of contaminants beyond city limits indicated the river was largely composed of untreated wastewater. They recommend that a wastewater treatment facility would substantially reduce most contaminant loads within the receiving environment [5]. Bhanu and Vellanki, 2021 employed the studies on detection of 16 emerging contaminants (ECs), including pharmaceuticals, personal care products and hormones in river Yamuna [6]. Nandeshwar et al., 2016 conducted iron contamination study for the Nag River (India). They treated water with various adsorbents like orange peels, coconut shells, sawdust etc. out of which the most promising green adsorbents were found to be orange peels and the best activating agent was HCl [7]. Birol and Das, 2010 suggested improvements in the capacity and technology of a sewage treatment plant in Chandernagore municipality, located on the banks of the River Ganga in India [8]. The similar studies of river and local water body contamination are reported for the various parts of the world [9]. Kassa, 2021 evaluated the hazardous effects of untreated municipal wastewater discharge from the city of Bahir Dar to the head of the Blue Nile River. This was done by collecting wastewater samples from the upper, middle and lower sites of the stream and characterizing its physicochemical qualities. In all cases, tap water was taken as a control and the wastewater qualities were significantly varied among the studied sites.

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The highest values of EC, TDS, COD, etc. were found at the middle site [10]. Nieto-Juárez et al., 2021 studied removal of the pharmaceutical substances in municipal wastewater treatment plants (MWWTPs) from Peru and the impact of these compounds in surface waters receiving treated wastewater. Acetaminophen was the drug found at the highest concentration, and it was present in all the treated wastewater samples reaching average values above 100 µg/L in the department of Puno [11].

In this present study, the aim was to study the pollutant levels in terms of nutrient concentrations and COD and BOD values in various stagnant and flowing water bodies in and around Ahmedabad.

II. MATERIALS AND METHODS

A. Physicochemical characterization

All the samples collected were investigated for pH, temperature, salinity, electrical conductivity, TDS etc. using automated multimeter. Samples were centrifuged at 7500 rpm at room temperature. Biochemical oxygen demand (BOD) and chemical oxygen demand (COD) were determined by standard methods [12] (APHA, 2005) for the selected samples. Total phosphate level was also determined for nutrient estimation.

B. Biochemical oxygen demand (BOD) determination

Bacteria utilize dissolved oxygen (DO) present in wastewater for oxidation of organic matters present. Biochemical oxygen demand (BOD) is the amount of oxygen consumed by bacteria while they decompose organic matter under aerobic conditions at 25 °C. The amount of reduced DO is directly proportional to the amount of organic matter present in the sample. Higher the BOD, higher is the concentration of organic matter present in the water. BOD is used often in wastewater-treatment plants as an index of the degree of organic pollution in water. BOD was estimated by measuring dissolved oxygen. Two parallel sets of BOD bottles were filled with sample water. One was used for determination of DO value (D_0) and another was incubated at 20 °C for 3 days, following which DO was determined (D_3) from this water.

Chemicals and Reagents:

- 0.025 N $\text{Na}_2\text{S}_2\text{O}_3$: (0.1 N Stock): Dissolve 24.82 g $\text{Na}_2\text{S}_2\text{O}_3$ in distilled water and make the final volume to 1 litre. Prepare in volumetric flask: 250 ml of 0.1 N $\text{Na}_2\text{S}_2\text{O}_3$ and add 750 ml distilled water (d/w).
- MnSO_4 : Dissolve 480 g of $\text{MnSO}_4 \cdot 4\text{H}_2\text{O}$ in distilled water and make the final volume to 1 litre.
- Concentrated H_2SO_4
- Alkaline iodide azide: Dissolve 500 g of NaOH, 150 g of KI in little amount of distilled water, add 10 g of NaNO_3 in 40 ml d/w. Mix both the reagent and make the final volume to 1 litre.
- 0.5% Starch: Dissolve 0.5 g of starch in 100 ml d/w by heating.

Procedure:

- Sample was filled till the top in BOD bottle. Two parallel sets of BOD bottles were filled with each sample water. As the strength of the waste (organic load) was not known in various water samples, various

dilutions were made arbitrarily. So, 1:2, 1:4, 1:6, 1:8, 1:10, and 1:20 dilutions were made from the collected water samples. Undiluted water sample was also taken. Hence total 14 bottles were taken for each water sample. Aerated d/w was used for preparing dilutions. One bottle from each dilution was incubated at 20°C for 3 days. The other bottle from each dilution was immediately used for determination of D_0 values of DO.

- For estimation of DO, 2 ml of MnSO_4 and 2 ml of alkaline iodide azide reagents were added in the bottle.
- One ml concentrated H_2SO_4 was added. The excess water was allowed to spill.
- One hundred ml of sample was taken from bottle and one ml starch was added. Color changed to deep blue.
- The mixture was titrated with 0.025N $\text{Na}_2\text{S}_2\text{O}_3$ till the color changed to colorless.
- The DO was calculated using the following formula:
- Dissolved O_2 in mg/l = Burette reading \times N of $\text{Na}_2\text{S}_2\text{O}_3 \times 8000$ / ml of sample
- After 3 days, DO readings were taken from the second bottle and BOD was calculated using the following formula.

C. $\text{BOD} = (D_0 - D_3) \times \text{Dilution Factor}$

where,

D_0 = dissolve oxygen at day 0

D_3 = dissolve oxygen at day 3

The BOD value of a sample was calculated by taking average of all the possible BOD values obtained from different dilutions.

D. Chemical oxygen demand (COD)

COD is the indicative measurement of amount of the oxygen that can be consumed by organisms in wastewater for oxidation of the organics present. Specifically, it measures the equivalent amount of oxygen required to chemically oxidize organic compounds in water. The COD test is often used to monitor water treatment plant efficiency and use to keep a check on residual organic material in wastewater. This test is based on the fact that a strong oxidizing agent, under acidic conditions, can fully oxidize almost any organic compound to carbon dioxide and water. It is an important and rapidly measured variable for characterizing water bodies, sewage, industrial wastes, and treatment plant effluents. Higher COD levels mean that a greater amount of oxidizable organic material is present in the sample, and indicate organic pollution of the water body. Higher amount of degradable or oxidizable organics present will lead to depletion of DO of that water body due to microbial and algal activity which may prove deleterious to aquatic life forms.

Chemicals and Reagents:

- $\text{K}_2\text{Cr}_2\text{O}_7$: Dissolve 12.258 g $\text{K}_2\text{Cr}_2\text{O}_7$ in 1000 ml distilled water.

- b. FAS (Ferrous Ammonium Sulfate): Dissolve 39.29 g FAS into some amount of distilled water then add 20 ml concentrated H₂SO₄ and make up the final volume up to 1000 ml with distilled water.
- c. HgSO₄ Crystals
- d. Ferroin indicator: Dissolve 695 mg Ferrous sulfate and 1.458 g 1, 10- Phenanthroline monohydrate in 100 ml distilled water.
- e. H₂SO₄ + AgSO₄ reagent: Dissolve 10.1 g AgSO₄ in 1000 ml H₂SO₄.

Procedure:

1. Total of 25 ml sample was taken in a reflux flask for COD estimation with the ratio of 24 ml of double distilled water + 1 ml water sample.
2. Following reagents were added into flask sequentially -
 - 0.5 g HgSO₄ (1 pinch),
 - 2.5 ml H₂SO₄ + AgSO₄ reagent,
 - 12.5 ml K₂Cr₂O₇ solution, and
 - 35 ml H₂SO₄ + AgSO₄ reagent
3. After cooling the sample, 5-6 porcelain beads were added and flask was placed on heater for 2 h at 70°C temperature.
4. Flasks were joined with condenser. The tap water was allowed to get filled with water.
5. After 2 h, condenser was disconnected and sample was allowed to cool down.
6. Total volume was made up to 100 ml by using double distilled water in 250 ml flask.
7. Ferroin indicator of 3-5 drops was added and sample was titrated with (0.1 N) FAS.
8. The colour change from pale yellow to green to blue to red was recorded.

E. Phosphate estimation (SnCl₂ method)

Chemicals and Reagents

- a. Phenolphthalein Indicator
500 mg Phenolphthalein was dissolved in 50 ml ethyl/isopropyl alcohol and then 50 ml distilled water was added. 0.02 N NaOH solution was added drop wise to this solution until faint pink colour appears.
- b. Sulfuric acid – Nitric acid reagent
75 ml concentrated H₂SO₄ is added to about 150 ml of distilled water. The solution was kept for sometimes to cool down and 1 ml concentrated HNO₃ was added followed by diluting it to 250 ml distilled water.
- c. Ammonium Molybdate reagent
Part 1: Dissolve 25 g of Ammonium Molybdate in 200 ml distilled water.
Part 2: Add 280 ml concentrated H₂SO₄ to 400 ml distilled water. Add molybdate solution to part 2 and make up to total of 1000 ml.
- d. SnCl₂ solution
2.5 g of fresh SnCl₂ was added in 100 ml glycerol and heated in water bath to dissolve.
- e. KH₂PO₄ stock solution
439 mg of KH₂PO₄ was dissolved in 1000 ml distilled water in which further two drops of toluene was added as preservative.
- f. KH₂PO₄ working solution

Freshly dilute the stock solution 1:100 times to give final phosphorus concentration of 1 ml = 1 µg P = 3.06 µg PO₄.

F. Procedure

1. Suitable aliquot of the centrifuged sample (containing not more than 20 µg of phosphate) was taken and one drop of phenolphthalein indicator was added. If pink colour appeared, two to three drops of sulfuric acid nitric acid solution was added to neutralize the solution.
2. Four micro liter of Ammonium molybdate reagent and 0.5 ml of stannous chloride solution were added to the sample. The contents were mixed well after each addition.
3. After 10 min but before 15 minutes absorption was read at 690 nm.
4. Calibration curve was prepared using Potassium dihydrogen phosphate as standard.

III. RESULTS

Though the Biological Nutrient Removal (BNR) has been practiced in wastewater treatment plants since long, lot of nutrients are still released in the treated water.

All types of nutrient present in the water originate from a variety of sources, and amongst them nitrogen (N) and phosphorus (P) have been regarded as the most important due to their impact on the aquatic life.

Being the limiting nutrients for algal growth in fresh water bodies they directly influence the acceptability of water in recreational bodies like lakes. Figure 1 depicts the calibration curve for phosphorous estimation by SnCl₂ method.

Concentration of PO₄ in samples collected from various time intervals was determined using the equation. The biochemical characteristics of the water samples collected from different lakes are shown in Table 1. The DO levels in these water samples ranged between 1.2 to 1.8 mg/L.

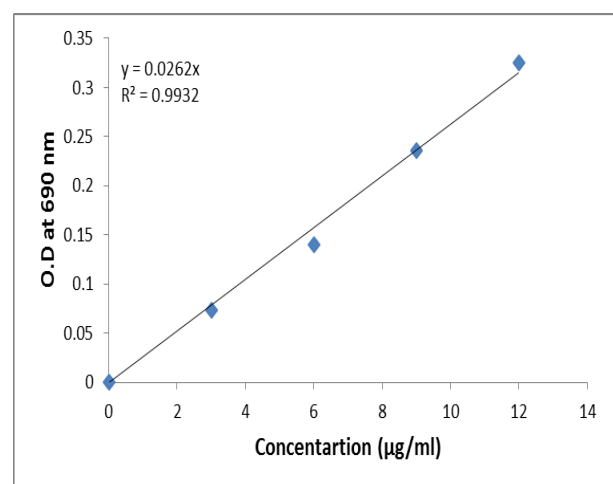


Figure 1: Standard curve for phosphorous estimation by SnCl₂ method



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Table 1: Comparative data of physic-chemical analysis of lake water samples

Lake	pH	Conductivity (µs)	salinity	TDS	Phosphate (ppm)
Kankaria	9.0	1961	1.09 ppt	1.38 ppt	2.850
Chiloda	9.4	468	250 ppm	333 ppm	2.693
Nikol	9.5	1519	832 ppm	1.07 ppt	2.857
Manipur	9.6	420	223 ppm	292 ppm	ND

Note: 'ND'-Not determined

All the samples were found to have different properties. All the samples showed alkaline pH ($\text{pH} \geq 9.0$). Electrical conductivity, salinity and TDS were high in Kankaria lake water sample as compared to the water samples collected from other lakes. Water sample collected from Nikol lake also had high TDS demonstrating higher pollutant level in the water body. Amount of phosphate concentration was almost same for all the lake water samples. Four water samples were collected from different sites of Kharicut canal are shown in Figure 2. Their physic-chemical parameters are shown in Table 2.



(a)



(b)



(c)



(d)

Figure 2: water sample collection site from Kharicut canal. (a) Site 1: Near Jay Chemical Industries and OEPL, (b) Site 2: Near Odhav, (c) Site 3: Near RTO, (d) Site 4: Near CETP Vatva GIDC.

Table 2: Physico-chemical parameters of water from four different sites of Kharicut Canal

Water Sample	pH	Salinity (ppm)	Conductivity (µ/s)	TDS (ppm)	BOD (mg/L)	COD (mg/L)
S-1	9.3	210	384	277	85	320
S-2	8.78	211	396	280	121	480
S-3	8.5	241	454	322	39	163
S-4	8.6	434	824	570	121	480

From the above results, it can be concluded that samples S-2 and S-4 contained higher organic load as compared to samples S-1 and S-3 as they have higher COD and BOD values. Although presence of maximum pollutant can be predicted in Sample 4 (S-4) as it contained maximum TDS,

and conductivity along with COD. The reason of higher TDS and COD of sample 4 might be due to the collection site which is near GIDC, Vatva, and a most well-known industrial zone in Ahmedabad.

IV. CONCLUSION

Out of total 39 small, medium sized and large lakes in the Ahmedabd city of India, samples from 4 lakes were collected on the basis of the diversity of the area and exposure to human activities. All the water samples were found to be alkaline in nature having more than 9 pH.

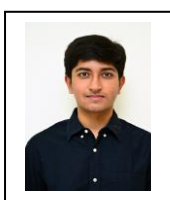


The dissolved oxygen level of all the lake water samples were ranging in between 1.2 to 1.8 mg/l. Nitrogen and phosphorus are the primary causes of cultural eutrophication (i.e., nutrient enrichment due to human activities) in surface waters. Though the phosphate content was almost same for all water samples collected from lake and hence this nutrient was present in negligible concentration. Kharicut canal which was primarily built for irrigation purpose became polluted in the industrial zone and it is an alarming sign as the COD exceeds the permissible limit (250 mg/L) detected by Gujarat Pollution Control Board (GPCB). This indicates the requirement of proper treatment of wastewater to reduce COD levels before being released in the canal water.

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AUTHOR PROFILE



Hemil H Patel is a 12th grade student of Sattva Vikas School in Ahmedabad, India. He is a passionate research student and this paper is one of his few research works heralding his entry into the research field. He is an energetic and effective leader of the School Science Club who carried out many innovative initiatives like conducting of webinars and organizing industry visits during the covid-induced lockdown, for benefit of his fellow students. Hemil plans to pursue his undergraduate

studies in Engineering in USA and would like to take up research assignments in the field of Engineering.