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Vinod Thakur School of Life Science, Devi

Ahilya Vishwavidyalaya, Indore, Madhya Pradesh, India

Dinesh Thakur Deputy Director Veterinary Services Khargone, Madhya Pradesh, India Journal of Entomology and Zoology Studies

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Study of ground water quality in Barwani district with special reference to Pansemal Tehsil

Vinod Thakur and Dinesh Thakur

Abstract

The study was carried out to assess the fluoride contamination status of groundwater in Pansemal Tehsil of Barwani District M.P. For this purpose, 24 water samples collected from Hand Pumps of village of study area were analysed for Dissolve Oxygen, Biological Oxygen Demand Chemical Oxygen Demand contain.DO, BOD and COD concentration in this sampling sides varied from in groundwater samples, with DO highest range in Pansemal 6.6 mg/l and lowest range in Junapani 4.4 mg/l.BOD highest range in Piprani 0.6mg/l and lowest range in Pansemal 0.1mg/l and COD highest range in Gongwada 8.0mg/l and lowest range in Piprani 4.0 mg/l.

Keywords: Biological oxygen demand, chemical oxygen demand, Pansemal Tehsil

Introduction

Surface water and groundwater have long been considered separate entities, and have been investigated individually. Chemical, biological and physical properties of surface water and groundwater are indeed different. In the transition zone a variety of processes occur, leading to transport, degradation, transformation, precipitation, or sorption of substance ^[5].

About 50% of all the underground used in urban areas of developing countries is derived from wells, springs and bore holes and more than 1000 million inhabitants in Asia and 150 million in Latin America rely on such resources ^[8].

High concentrations of fluoride in drinking water are harmful to human health ^[3].

Water is nature's most wonderful, abundant and useful compound. Of the many essential elements for the existence of human beings, animals and plants, water is rated to be of the greatest importance. Without food, human can survive for a number of days, but water is such an essential that without it one cannot survive. Water is not only essential for the lives of animals and plants, but also occupies a unique position in industries. Groundwater is an important source of water supply throughout the world. The quantity and the suitability of groundwater for human consumption and for irrigation are determined by its physical, chemical and bacteriological properties ^[6].

Water in the atmosphere comes from evaporation from the oceans, lakes, rivers ice-fields and glaciers, moist ground transpiration from plants and animal respiration.water available in the atmosphere is carried for long distances on land from the oceans by wind and convective moment under favourable conditions, it condition and precipitates over the earth's surface as rain, snow and hail ^[7].

Groundwater is almost globally important for human consumption as well as for the support of habitat and for maintaining the river's base-flow. It is usually of excellent quality. Being naturally filtered in their passage through the ground, they are usually clear, colourless, and free from microbial contamination and require minimal treatment ^[1].

Materials and Methods

Groundwater samples from different hand pumps, bore wells were analyzed. Samples were collected in three different seasons June-May 2014-2015 Water sample could not be collected from the location during the post monsoons period due to mechanical problem in the hand pumps. The temperature pH and electrical conductivity were measured in the field using portable pH meter and EC meter. Captions and anions were analysed using ions chromatograph and standard titration method following standard procedures ^[2].

Correspondence Dinesh Thakur Deputy Director Veterinary Services Khargone, Madhya Pradesh, India **Sampling station:** Four sampling stations were selected in the present investigation-

- S1.Pansemal
- S1.Pansemal
- S2.Junapani
- S3.Gongwada
- S4.Piprani

The water samples were collected 3-3 times these selected sampling station in a precleaned and rinsed plastic container of 1litter capacity for further analysis for necessary precaution [4].

Temperature

Procedure

- 1. Immerse thermometer in the sample up to the mark specific by the manufacturer and read temp. After equilibration.
- 2. When a temp. Profile at a number of different depth is required a thermo stat with a Sufficiently long lead may be used.

pН

Reagent

- 1. Standard buffer solution of known pH.
- 2. Shake vigorously one excess (5-10g.) of finely crystalline KHC₄H₄O₆ with 100 to 300 ml distilled water at 25°c in a glass- stopper bottle.
- 3. Calcine a well washed, low alkali grade $caco_3$ in a platinum dish by igniting for 1L. At 1000° C Cool.
- 4. 0.1 N NaOH, 0.1 N Hcl, 5 N Hcl and acid potassium fluoride solution.

Procedure

- 1. In each case follows manufacturer's instructions for pH meter and for storage and preparation of electrodes for use
- 2. Sample for 1 min. Blot dry, immerse in a fresh portion of the same sample and read pH.
- 3. Take a fresh sample to measure pH.

Colour

Reagent: Dissolve 1.246 gm. Potassium chloro-platinate, K_2PtCl_6 and 1.00gm crystallised cobalt us chloride, $CaCl_{2.6}H_2O$ in distilled water with 100ml conc. Hcl and dilute to 1000ml with distilled water.

Procedure

- 1. Pour sample in a nessler tube of 50 ml mark. Similarly fill three to four tube with colour standard which appear to correspond to the colour the of the sample
- 2. Compare colour of the sample with that of the standard by viewing vertically down words while the tube are placed on a white surface calculation.

Calculation: for dilute sample calculate colour units as:

 $A \!\!\times 50$

Where: -

- Colour units =
- A= estimation colour dilute sample
- B= ml sample in 50 ml dilute sample

Odour Procedure

- Full of sample, insert stopper and shake vigorously for 2-3 sec. And then quickly observe the odour. The sample should be at ambient temperature.
- 2. Odour free, rotten egg. Burnt sugar, soapy, fishy, septic aromatic, chlorines, alcoholic odour or any other specific odour.

DO (Dissolve Oxygen)

- 1. Manganous sulphate solution: Dissolve 480 gm MnSo₄.4 H₂O, 400gm MnSo₄.2H₂O or 364 gm MnSo4 H₂O in distilled water, filter and dilute to 1 litter.
- 2. Alkali iodine Azid reagent: Dissolve 500gm NaOH or 700 gm KOH and 135 gm NaI or 150 gm KI in dissolved water and dilute to 1 litter.Add 10 gm sodium azide (NaN₃) dissolved in 40 ml distilled water.
- **3.** Sulphuric acid concentrated: 1 ml is equivalent to about 3 ml alkali iodide azide reagent.
- **4. Standard sodium thiosulphate 0.025N:** Dissolve 6.205 gm sodium thiosulphate (Na₂S₂O₃.5H₂O) in freshly boiled and cooled distilled water and dilute to 1 litter. Add 5 ml chloroform or 0.4 gm NaOH /L or 4 gm borax and 510 mg HgI₂/L. 0.025 N potassium dichromate solution dissolved 1.226 gm potassium dichromate in distilled water and dilute to 1 L.
- 5. Standard potassium dichromate: A solution potassium dichromate equivalent to 0.025 N sodium thiosulphate contains 1.226 gm /L.K₂Cr₂O₇ at 103° C for 2 hrs before making the solution.
- 6. Standard disation of 0.025 N sodium thiosulphate solutions: Dissolved approximately 2 gm KI in an Erlenmeyer flask with 100-150 ml distilled water. Add 10 ml of H₂So₄, followed by exactly 20 ml 0.25 N potassium dichromate solution.
- 7. Starch Indicator: Add cold water suspension of 5 gm soluble starch to approximately 800ml boiling water with stirring. Dilute to 1 L allow to boil for a few minute and let settle overnight.

BOD (Bio Chemical Oxygen Demand)

Procedure

Fill BOD Bottle sample + 1ml MnSo₄ + 1ml Azide solution. And shake well

add 1ml H₂So₄ to dissolved brown p <u>titrate 200ml</u> Against

 $0.025N\ Na_2S_2O_3$ using starch as indicator till the solution become colourless.

A. Fill BOD Bottle with sample (incubate) at 27° C for three days (then) follow procedure as per 'A'.

Calculation

BOD % mg /L = D1 -D2 D1 = Ini DO of 1^{st} day D2 = final DO of 3^{rd} day

Reagents

- 1. Monogamous sulphate: 480g MnSo₄ 4 H₂O dilute in 1 litter water.
- Alkali iodide Azide: 500gm NaOH o 700 gm KOH and 135 g NaI on 150 gm KI in Distilled water.
- 3. Sulphuric acid : $conc.H_2So_4$
- 4. Starch solution :

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5. Standard $Na_2S_2O_3$ 0.025 N: Dissolve 6.205 gm $Na_2S_2O_3.5H_2O$ in one litter distilled water.

COD (Chemical Oxygen Demand) Procedure

20 ml sample add \rightarrow 1gm HgSo_4 + Glass beads add & mix \rightarrow 30 ml H_2So_4

Cool drops

10 ml of 0.25 N K_2Cr_2O_7 + \rightarrow reflux for 2 hrs at 150 ° C \rightarrow cool to room

While mixing temperatures wash condenser with 25 ml distilled water

Add 2-3 drops triturate against FAS till Blue Green to just reddish

Ferroin indicator Brown.

"Reflux & Triturate a blank simultaneously."

Calculation:
$$C = (A-B) \times N \times 8000$$

ml of sample

Where,

C = COD as mg O_2/L

A = ml of FAS used for sample

B = ml of FAS used for blank

N = Normality of FAS

Reagent

- Std. Potassium Dichromate solution : 0.25 N Dry K₂Cr₂O₇ at 103 ° C for 2 hrs and dissolved 12.259 gm of dried k₂Cr₂O₇ in distilled water dilute to 1 litter.
- Ferro in Indicator : Dissolved 1.485 gm 1:10 Phenanthrolein monohydrate + 695 mg.

FeSo₄.7H₂O in distilled water dilute to 1 litter.

- 3. Td. Ferrous Amm. Sulphate Solution :0.25 N (FAS)
- Dissolved 98.0gm FAS in distilled water. And 20 ml come H₂SO₄ cool and dilute to 1 litre standardise against 0.25 N K₂Cr₂O₇ solutions.

Total solid

Procedure

A. Dry evaporating Dish /Biker at $103\pm 1^{\circ}$ C cool and store in a desiccators, weigh immediately before use.

B. While stirring pipette a measured volume in to the pre – weighed evaporating dish / Beaker using a wide pore pipette. Choose a sample volume to yield between 10 and 200 mg dried residue

Calculation - mg Total solids/ $L = (A-B) \times 10^6$ /ml sample)

Where,

A= weight of dish /Beaker + residue, gm.

B = weight of dish / Beaker, g

Observation Tables and Results

 Table 1: Comparison of D O (dissolved oxygen) at different depth on Northern and southern Hand Pump and Bore well.

Depth	Pansemal		Junapani		Gongwada		Piprani	
in feet	H.P	B.W	H.P	B.W	H.P	B.W	H.P	B.W
70-80	5.4	5.6	5.4	5.3	5.2	5.3	5.2	5.4
80-90	5.6	6.6	4.4	5.7	5.6	6.6	6.1	6.6
90-100	5.8	5.4	5.6	5.8	4.8	5.6	5.4	4.8



Fig 1: Hand Pump



Fig 2: Bore well

DO: The Dissolved oxygen in the sampling station was ranged between 4.4 to 6.6 mg/L in the present study period.

Table 2: Comparison of B O D (Biological oxygen Demand) at

 different depth on Northern and southern Hand Pump and Bore well.

Depth	Pansemal		Junapani		Gongwada		Piprani	
in feet	H.P	B.W	H.P	B.W	H.P	B.W	H.P	B.W
70-80	0.2	0.3	0.3	0.4	0.6	0.4	0.2	0.4
80-90	0.6	0.2	0.2	0.4	0.1	0.6	0.2	0.5
90-100	0.4	0.1	0.3	0.2	0.4	0.2	0.1	0.6



Fig 3: Hand Pump





Fig 4: Bore well

BOD: Biological oxygen demand level of sampling station range between 0.1 to 0.6 mg/L. The value of B O D in different groundwater sample was also under the permissible limit.

Table 3: Comparison of C O D (Chemical oxygen Demand) at different depth on Northern and southern Hand Pump and Bore well.

Depth	Pansemal		Junapani		Gongwada		Piprani	
in feet	H.P	B.W	H.P	B.W	H.P	B.W	H.P	B.W
70-80	4.0	4.0	4.0	6.0	8.0	6.0	4.0	6.0
80-90	5.0	4.0	6.0	5.0	7.0	4.0	5.0	6.0
90-100	6.0	5.0	7.0	4.0	6.0	5.0	4.0	5.0



Fig 5: Hand Pump



Fig 6: Bore well

COD: The C O D value of sampling station was range between 4.0 to 7.0 mg/L in competition to the low D O value, the C O D value always observed higher than B O D value.

 Table 4: Comparison of Total solid at different depth on Northern and southern Hand Pump and Bore well.

Depth in	Pansemal		Junapani		Gongwada		Piprani	
feet	H.P	B.W	H.P	B.W	H.P	B.W	H.P	B.W
70-80	300	356	368	442	546	632	670	478
80-90	340	450	348	452	560	432	375	573
90-100	298	385	476	460	548	562	480	580



Fig 7: Hand Pump



Fig 8: Bore well

Total solid: The total solid content of sampling station was ranged between 298 to 670 mg/L the observed result of total solid indicate that all the sampling station were suitable for drinking purpose after necessary treatment.

Conclusion

The present study investigates the hydrochemistry of water and DO, BOD, COD and Total Solid in ground water.

DO: The Dissolved oxygen in the sampling station was ranged between 4.4 to 6.6 mg/L in the present study period.

BOD: Biological oxygen demand level of sampling station range between 0.1 to 0.6 mg/L. The value of BOD in different groundwater sample was also under the permissible limit.

COD: The COD value of sampling station was range between 4.0 to 7.0 mg/L in competition to the low DO value, the C O D value always observed higher than BOD value.

Total solid: The total solid content of sampling station was ranged between 298 to 670 mg/L the observed result of total solid indicate that all the sampling station were suitable for drinking purpose after necessary treatment.

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