

Surface Initiated Toxicity in Water and analysis in Sagar District

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Abstract - Groundwater is an important source of water for people all around the globe. A bore or a well is used to obtain groundwater. The reconstituted reagent was balanced at + 4 ° C for half an hour. The reagent was then stabilised at + 15 ° C for 30 minutes before pipetting into the cuvettes. A ten-level 1:2 dilution sequence was laid. The Sample Diluent 1243-552 was used to create the sample dilution sequence. All samples and dilutions were tempered to +15 ° C for 15 minutes. All samples and dilutions were held at + 15 ° C during the measurement. Duplicate measurements were taken. The programme Ascent Software, developed by Aboatox Co. in Finland, assisted in evaluating the sample's toxicity. TOC was measured using the Apollo 9000 TOC Combustion Analyzer. It calculates sample TOC levels in milligrammes per litre. It is concluded that the water is likely to have a range of pathogenic bacteria, as well as Bacterial pathogens of water grasping, indicating a water-specific risk of re-use for vulnerable staff.

Keywords - Water, Toxicity, Analysis

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INTRODUCTION

In addition to being the most abundant substance on our planet, water also has the distinction of being the only clear liquid. A water molecule consists of two hydrogen and one oxygen atom. The absence of water would kill every other living thing on the planet. Water is clearly vital for the existence of all organisms on our watery earth. No person can survive for any appreciable length of time without water, since it is required for the proper functioning of every single cell and organ system in the body. Water is essential for every bodily process. Unreliable and polluted water supplies are a major issue in many rural and urban areas in developing countries (1). In rural areas, groundwater is a finite but renewable resource. The two most common sources of water for irrigation and other purposes are surface water and groundwater. Undersea and surface water systems are interconnected. Above ground, in the form of oceans, rivers, lakes, ponds, and streams; and below ground, in the form of permeable soils and rocks, are the two primary sites where water may be found on Earth.

As a result, groundwater is sometimes the sole source of potable water in many regions of the world. To access groundwater, a well or bore must be bored into the earth. Groundwater is an important aspect of the water cycle because of its many uses in agriculture, industry, and human consumption. Runoff, animal waste, leaking underground storage tanks, and industrial chemical waste have all contributed to the pollution of groundwater caused by the overuse of pesticides and fertilisers. Contaminated groundwater

may make many activities useless, and removing the contamination is a time-consuming and expensive task. It's possible that might have negative effects on both humans and the planet. Health. When pollutants are present in groundwater in such amounts to make the water unusable, we say that there is groundwater pollution. Groundwater contamination may come from both natural and manmade causes. Human activities change the chemical composition of groundwater. Foreign compounds and other contaminants introduced into the earth may contaminate the water supply. Polluted groundwater poses serious health risks to humans, therefore it's important to keep an eye on the processes that affect the water table. Drinking water from the ground is common in rural places (2).

Water quality land use impacts

The structure and function of aquatic ecosystems are affected by a wide variety of landscape factors. These factors include riparian zone condition, channel slope and aspect, local geology, vegetation, and hydrography. Water quality is affected by several factors, but land use and land cover are two of the most significant. Water balance, water chemistry, and the variety of aquatic life in streams that receive runoff are all affected by land use and management practises. Most of the area around Old Wives Lake is used for agriculture, mostly raising annual crops and animals. The Wood River is an important water supply in the watershed, but it might be harmed by runoff from these areas. Discharging treated sewage into the Wood River is a common

practise in Gravelbourg, and it poses a serious risk to the river's health. Twice a year, in the spring and autumn, there is a release. An elaborate system of shallow basins called a lagoon is utilised to treat the city's wastewater via biological processes. This method is widely used in remote areas since its effluent is on par with that of a secondary treatment system (3-6).

Water pollution from agriculture, municipal and industrial waste

A Look at the Consequences of Not Having Access to Clean Water Pollution might be caused by many different types of human activity, such as farming or the incorrect disposal of municipal waste water. Wastewater treatment plants and factories are examples of point sources since they produce waste only at certain locations. Point sources of pollution are simple to identify, meaning that they may be monitored and regulated, and in some cases even treated at the source. Agricultural practises are a major contributor to nonpoint source pollution, often known as long-distance pollution. Agricultural activities and weather extremes are two examples of transient drivers of nonpoint source pollution. Nonpoint sources contribute significantly to water pollution, and their contamination may start from large land regions and travel via the air, the ground, and the water to reach the ocean. Because of this, keeping tabs on and controlling emissions from diffuse and mobile sources is no easy task (7).

Various toxic substances, including human waste, suspended particles, rubbish, and chemicals from households, businesses, and industries, may be found in municipal wastewater (8). The cellular and organ levels, the organismal and community levels, and even the trophic level may all be affected by urban sewage. Nutrients like as nitrogen and phosphate, viruses such as *Cryptosporidium*, and endocrine disrupting compounds such as medicines and hormones from birth control pills are just some of the many contaminants that may be found in municipal waste water effluents (9).

To a large extent, agricultural practises are blamed for the deterioration and contamination of aquatic environments. Scientific research indicates that agriculture is too responsible for the deteriorating water quality of as much as 77% of the rivers and streams in the Great Plains. The agricultural sector is a major user of freshwater. Over ninety-five percent of western Canada's fertile grassland has been converted for the ecologically destructive purposes of cereal crop and animal production. When natural grasslands and riparian zones are converted into agricultural land, the stream's chemistry may be significantly affected by changes in the stream's flow, temperature, channel features, bed disturbance regime, and organic matter intake. Therefore, the stream biota's species composition shifts and their habitat quality decreases as a result of these changes

to the physical environment. The amount of agricultural production seems to correlate negatively with the severity of these outcomes. One study of fish populations in many U.S. rivers found that the health of fish communities decreased fairly linearly with increasing agricultural intensity. Nutrients like nitrogen and phosphate, together with pesticides, silt, viruses, and hormone-affecting compounds, are among the most pervasive contaminants originating from farms. Drinking water contaminated with these substances raises concerns about potential health effects to humans and ecological damage (10-16).

MATERIAL AND METHODS

Sample collection

Groundwater samples were obtained from five different regions of Sagar and analysed using traditional procedures for toxicity were measured.

Eco toxicity testing

- **ToxAlert test**

Using the reconstitution solution provided by the kits, the bacterial suspension was prepared to reconstitute the freeze-dried bacteria (*Vibrio fischeri*). The freeze-dried bacteria and the resolution for 26 reconstitution are kept in the freezer at -18°C before the test. The ToxAlert® 100 luminometer contains a separate well for reagent vials in order to maintain the proper temperature (15°C). To begin, 12.5 mL of well-shacked reconstitution solution (to ensure that enough oxygen is dissolved) was placed in the microquant vial and maintained in the liquid dried reagent for at least 15 minutes. The bacterial vial was then added to the mix, along with 0.5 mL of reconstitution solution. After 15 minutes, the bacterial suspension was transferred to the Microquant vial, which contained the balance of the reconstitution solution and was utilised as the bacterial experiment's suspension. A fifteen-minute preincubation time has been established. During that time, 500 l of the coral sample suspension was pumped into all cuvettes, along with control cuvettes (A1 and B1). Cuvette A1 was installed in the turret just before the contact time began. At contact moment $t=0$, the RLU of the alternative was determined. 500 l of NaCl solution was then gently mixed into cuvette A1. Because the inter-cuvette duration was set to 30 seconds, the identical procedure was used for cuvette B1 at $t=30$ times. At $t=60$ and 90 seconds, a somewhat different procedure for the sample cuvettes was used. After measuring RLU, 500 l of diluted water was added and gently mixed for the test. For each experiment, the WET technique manuals recommended a dilution sequence of 6.25 percent, 12.5 percent, 25 percent, 50 percent, and 100 percent samples (USEPA, 1994; USEPA, 1993). As a diluent, a 2 percent NaCl solution was utilised to prepare the sample dilution sequence. At $t=30$ minutes, the

exposure time was completed. The RLU of all cuvettes was then assessed once again using a 30-second inter-cuvette instant in the same series.

- **Flash assay (kinetic determination)**

The reconstituted reagent was balanced at + 4 ° C for half an hour. The reagent was then stabilised at + 15 ° C for 30 minutes before pipetting into the cuvettes. A ten-level 1:2 dilution sequence was laid. The Sample Diluent 1243-552 was used to create the sample dilution sequence. All samples and dilutions were tempered to +15 ° C for 15 minutes. All samples and dilutions were held at + 15 ° C during the measurement. Duplicate measurements were taken. The programme Ascent Software, developed by Aboatox Co. in Finland, assisted in evaluating the sample's toxicity.

- **Estimation of eco-toxicity results:**

The Environmental Protection Agency's (EPA, USA) Probit software is used to determine EC₅₀ values for the ToxAlert testing. The EC₅₀ and EC₂₀ principles for Flash tests were calculated using the ABOATOX software that included with the equipment.

RESULT AND DISCUSSION

Toxicity Test and Reduce the toxicity of GROUND WATER

Determination of toxic effects of Ground water for recipient freshwater systems with the help of different inocula

As stated in the Methods and Materials part, the samples were composed from the ground water healing plant of Sagar. 2 samples were taken: (1) raw and (2) treated Ground water. Together kinds were than varied an inoculum taken from a usual, intact water body, Bhakra canal (A) and from the receiver stream (B).

Figure 1 indicates the toxicity changes of sample 1 (raw sewage water) and the inoculate raw sewage water sample (1A and 1B). In seven days, the toxicity augmented realization app. 99% inhibition till the 3rd week. Subsequent to the 3rd week, there was an arresting decline in toxicity and the toxicity ongoing to dwindle in stable manner.

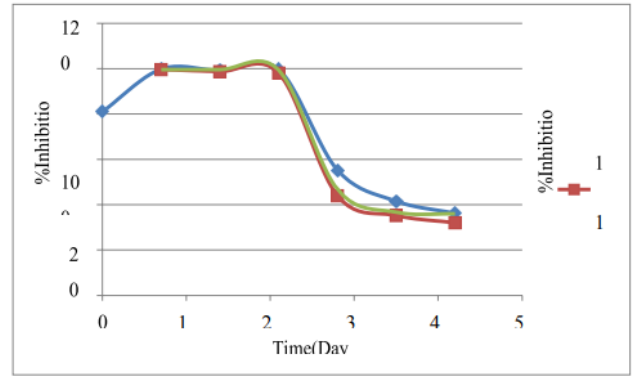


Figure 1: Toxicity changes of the sample 1 in comparison with the presence of different inocula

Figure 4.6 depicts that the toxicity of sample 2 (treated sewage water) and the inoculate treat sewage water samples (2A & 2B). There is an augment till the ending of 1st week and after the town expected decline can be seen. By the finish of the test, the toxicity of the sample behave in a stable manner.

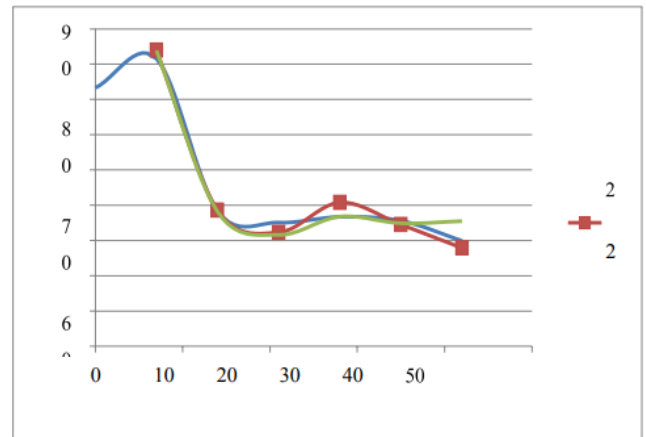


Figure 2: Toxicity changes of Sample 2 in comparison with the presence of different inocula

Figs 3 display shifts in harmfulness from day 0 to day 153. A elevated toxic content (transmitted as 80.9 percent bioluminescence restriction) and risk was introduced to the world by crude mutual wastewater water. With no inoculum included, this toxic quality even greater than before during the original 12 days, depicting to some extent different instances in the three initiation processes: at room temperature (experiment R22) hindrance at that stage was as large as 91.8 percent, extending to Day 12, attainment its maximum limit of 97.8 percent. Under the other two schemes of temperature, 10 ° C (test R10) and 30 ° C (test R30) initially reported a small decline from the maximum harmfulness of 94.9% and 97.3% respectively by day 12. Thereafter, a speedy decline could be seen between Day 12 and Day 19 initially, than from Day 19 on, an ongoing, faster decline, finally reaching an "average" point of implementation. 30% of the punishment by day 26 for R22 (34.55%), by day 40 for test R10 (34.15% restraint) and by day 54 for test R30 (32.45% restraint). It be supposed to be

observed, however, that this instance show an uneven low limit of 26.1% on Day 19, which can be in all probability ascribed to test error.

Within the presence of the inoculum, for RS10 and RS22 trials on Day 12 the maximum toxic output was smaller than the scenario with inoculum-free trials. Shortly thereafter, a considerably quicker decrease was began, resulting in restraint even by Day 19 below 30 percent.

(There was marginally greater hindrance appreciation for RS22 on Day 54.) Sample RS30, nevertheless, showed a rather exceptional instance: toxic performance extended to 94.3 percent of the restraint by daylight hours 12 than a fast and persistent reduce could be seen, but in conclusion the risk started to augment again, indicating 40.7 percent of the restraint by Day153.

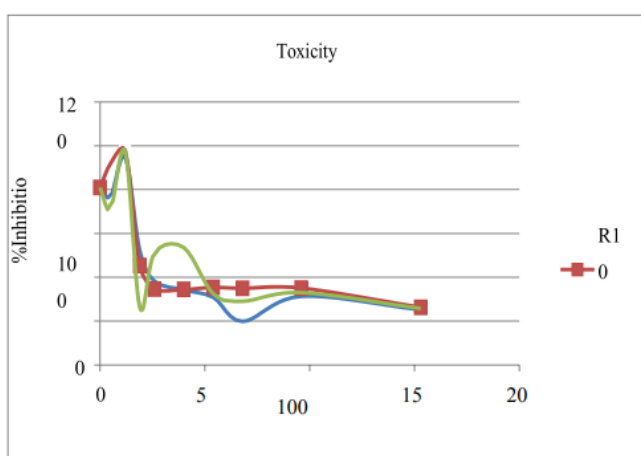


Figure 3: Toxicity changes of the raw sewage water sample without inoculums, at various temperature regimes

Figures 4 indicate changes in the lethality of the sans inoculum and vaccinated sewage water from day 0 to day 153. The treated collective sewage water showed a bearable harmfulness (transmitted as a 35.8 percent restriction of bioluminescence). With no inoculum included, the hindrance shifts at room temperature (T22 experiment) between a small augment and a reduce attainment its utmost (41 percent restriction in the Day40) follow by a quick reduce attainment 25.1 percent in the Day 54. 10 ° C (e.g. T10) and 30 ° C (e.g. T30) under the other two heat schemes: T10 and T30 toxic quality stated a radical decline on Day 19 (arriving at 16.5% and 15.9% correspondingly), demonstrating to some degree unexpected example in comparison to T22 (36.9%). The examples demonstrated lessening in poisonous quality in Day 54 beneath the three warmth systems. T22 and T30 demonstrated an unflinching more slow decline, at last coming to a "bearable" height of request. 30% of hindrance by Day 153, while, T10 demonstrated an expansion reaching53.45%.

In view of the inoculum, toxic modifications in the quality of the specimens showed a comparison

instance under the three warmth schemes up to Day 68. The instances reach a hindrance below 30 percent by the implementation of Day 19 at that stage. By Day 26.A, 40 percent showed an unusual decline in Day 54, reaching a 22 percent hindrance. Tests TS22 and TS30 showed an unwavering slower decline, finally achieving "tolerable" level of app.26 percent of inhibition by Day153, however, experiment TS10 showed a variety of examples to some extent: harm extended to 49.8 percent of inhibition by Day 153.

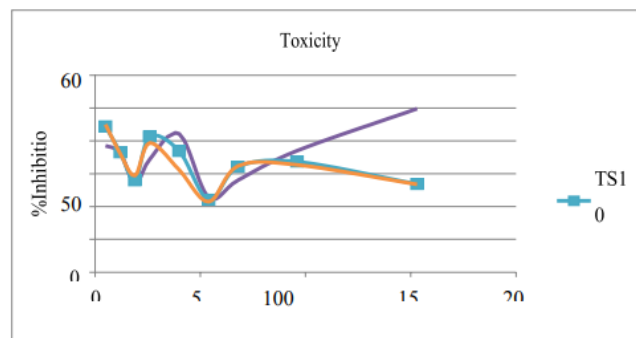


Figure-4: Toxicity changes of the treated sewage water sample with inoculum, at various temperature regimes

Biodegradation evaluation of liquid manure disposal

Two years earlier, one of the lakes was deserted and no mechanical isolation was constructed here. Groundwater was evaluated to evaluate the environmental impact in the area of the lake (Table 1), but no toxicity was assessed. Furthermore, the farm has numerous stabilization ponds with adequate mechanical security, which are motionless in use.

Table 1: Results of groundwater analysis

Components	Measured vlues (mg/l)(µg/l)			Limit values	
	F1	F2	F3	A	B
pH	6.72	6.56	6.67	-	6.5-9.0
HCO ₃	420.13	490.16	437.64		
CO ₃	0.0	0.0	0.0		
Cl	10.97	52.85	30.91		
COD _{ps}	0.72	0.96	0.72		
NO ₃	12.48	23.45	18.44	10	25
NO ₂	0.05	0.03	0.03		
NH ₄ -N	360	360	320	250µg/l	500µg/l
SO ₄	1.76	7.68	3.08	200	250
Na	9.85	7.05	6.6		
K	0.92	0.98	0.88		
Ca	53.0	61.0	42.0		
Mg	18.3	31.7	21.7		
PO ₄	310	270	240	200 µg/l	500 µg/l
e. conductivity	0.79 mS/cm	0.86 mS/cm	0.95 mS/cm		

Different inocula were used to evaluate the modifications induced by biodegradation, one

gathered from a safe (reference) setting and one gathered at the surface.

Sagar one (the sample of new manure) from the pond presently in use, while the other (the sample of old manure) from the reservoir left two years earlier. On the site and from Sagar (orientation uncontaminated soil) the soil inocula was collected.

The six sample samples have been designed and labeled as follows: 1: liquid compost from the pond presently in use

- **1T1:** clean soil mixed with liquid manure (1). (200 cm³ soil + 200 cm³ liquid manure)
- **1T2:** soil collected from the site mixed with the liquid manure (1). (200 cm³ soil + 200cm³ liquid manure)
- **2:** Liquid manure from the deserted tank
- **2T1:** clean soil mixed with liquid manure (2). (200 cm³ soil + 200cm³ liquid manure)
- **2T2:** soil collected from the site mixed with liquid manure (2) (200 cm³ soil + 200 cm³ liquid manure) All the sample were incubate at 20°C for 12 weeks.

Figures indicate the distinction in sample 1 toxicity (new fluid manure), sample 2 (old liquid dung) and sub-sample toxicity. Sample 2 toxicity is obviously much smaller than sample 1 since the deserted pond has more time (almost 2 years) to restore without incorporating any fresh pollutant and the microbial community residing there have been adjusted to these pollutant.

As the sample 1 and 2 were dissolved with the soil inocula, the 1T1, 1T2, 2T1 and 2T2 subsamples indicated more unreliable patterns in toxicity change through the process of squalor than the raw samples. The subsample toxicity is smaller than the initial.

CONCLUSION

Toxicity has reached a kind of equilibrium for both samples by the end of the experiment. 90% By the 3rd month (Day 84), demonstrating that the sample duration be not sufficient to achieve a substantial toxicity decrease. Sample 2, however, showed a marked decline in toxicity as of the original 65% to the rather steady 45% inhibition.

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