



Abstract

Aquatic environments differ in their chemical composition, which, in turn, affects the organisms living in the water. Among the dissolved materials that affect the chemistry of water and as a consequence, the aquatic organisms, are salinity and the dissolved oxygen (DO) and carbon dioxide. Here we present the results of our research into the salinity of a surface-water sample from East River and the effect of different salinity on vacuoles of cells normally adapted to fresh water. We also determined the amount of DO in our water samples, and studied the anaerobic process of cell respiration. Finally, we determined the presence of coliform bacteria by inoculating media which specifically allows coliform bacteria growth. We developed experiments and questions to determine the properties of our water sample taken from East River, and developed tables and graphs that allowed us to analyze our results.

Introduction

Aquatic life is affected by three important factors, salinity, dissolved oxygen and dissolved carbon dioxide. In order to measure the effects caused by these factors we conducted experiments using water samples from the East River. Our water sample came frame francisisperistic father East River River Jahred Island Island Dirp. Deeps The bampte owle ctollected/37/9/27/1835 and bad hachapter appranted for 3 for 3 letail Dissolved so system (Dx) gin (Dx) en thay ged is sal is diss the dist the through this onghe his sach it the twater aquatia quarits plants apro by protocort photo photos sister as a set of the organisos danisuse bei a use is is a me for star i four their a life iver a Furlex a compte fisbase us state ir gists their gials this sectra ed dis segme dot a yberbioted the bloode an tot the eds lof the ed lu for restirtationspitationinglesselled bacteria buckeria fungi alefungi dependiber Desetterause up there use in in the regression of the state wateren Withoutstetheaperes Without dissplacehoe vogedissed tendaoxygeno bangeriabeafed of longen the step hantsughdtheide events all did lefte Gardlyn die offle (arbon) is axialth (CQ23) tis at is ther diaschatchisteringthentsater Rhintalandkaninallsrikes condicrees faceather that dean in the viate indepenventer depend our GOA to harry southesistasych the sistawith for Herverthr, How edierg tocEbedOrgeton PloetOlcEentRoldal Sestate20thest statist that ano indiscost and Gissolved at 62 in invatersin the asei they avidity vesicits results icade crease at no fints with a karbo watch a wheteine dyganinenorganische kindkletbeinskrektet osiste Halsingneadebomater (paranoff)er sAmothen sourflet that in one one satisfies is satisfies a satisfies of the source of badtyin favboudy. Aftwatchin Tilyei nabinaity ish bracki sla wate framges 5 from 50 ppto (Mapiphal diffected different by by the syntian solution they care placed in. Attimpth cell in la oed inc sistions subvicons start adostant planeas place oche flacoide Claraid. Contain aon ditions uld affect whe latifiest the satinity siprecipitation precipitation reappointed and the second of the se would tibers expected electrease the overal healtion location bocadis into would involve an f firstraageof Trishwater This would have an impactiffe baring different rerganistiffecentite differental aviels infordenity inverter sortinor this important background of the akigs of wed of the relates of wed meterials in water to observe fiter influence of a reaching are esseause they are essential and lafkey element in life.

Materials and Methods

Our water sample, Long Island City Deep, was tested for dissolved O2, dissolved CO2, and salinity. Our instructor took the pH and temperature at the time of collection. Salinity was measured using the evaporation method. Also, our water sample and NaCl was used to determine the percentage of change in mass through a standard curve graph. Dissolved Oxygen was measured with a LaMotte oxygen titration kit. We had to add two chemical reagents which will fix the oxygen present in our water sample. This will allow us to titrate the amount of dissolved oxygen in our water. Dissolved Carbon Dioxide was measured using the LaMotte Titration kit. We add two drops of Phenolphthalein Indicator and filling the direct reading titrator with Carbon Dioxide Reagent and inserting into the test tube. Also, with the use of our water sample and bicarbonate concentrations to determine the rate of photosynthesis. Lastly, we determine if our water was a hypotonic or hypertonic by using our microscope. We observe and measure the vacuoles placed in each solution as well in our water sample.

LIFE IN THE WATERS OF EAST RIVER: EFFECTS OF SALINITY AND DISSOLVED OXYGEN ON CELLS



Figure 1. The graph above illustrates the final percent change in mass.

Length of onion cell vacuoles in different solutions			tions
Measurements	Length vacuales hypotonic solution	Length vacuoles hypertonic solution	Length vacuales water sample
1	108 µm	250 µm	350 µm
2	100 µm	160 µm	107 µm
3	306 µm	300 µm	208 µm
4	207 µm	220 µm	102 µm
5	304 µm	310 µm	200 µm
Average	205µm	248 µm	193. 4 µm

Figure 3. This table shows the measurements of onion cell vacuoles placed in tonicity solutions.



Figure A: Onion cell vacuoles in hypertonic solution



Figure B: Onion cell vacuoles in isotonic solution



Figure 2. The graph above show how Glucose Concentration can affect the fermentation rate in yeast. None of the four trials show consistent pattern.

ange in Mass / NaCI Concentration			
Μ	%Change		
0	2.89		
0	7.04		
0.1	0.78		
0.2	0.68		
0.3	2.48		
0.4	3.74		
0.5	6.2		
0.6	4.49		

Figure 2. The table depicts the change of mass in percentage of each NaCI solutions that has change within time.

Pictures of Onion Cells Vacuoles in Tonicity Solutions

Figure C: Onion cell vacuoles placed in hypotonic solution

Before performing both experiments, we predicted that the enzyme concentration and the enzymes will increase. Also, we predicted that our water sample will have the results of being a hypotonic solution. In figure 1 the fermentation rate in yeast is affected by glucose concentrations. The results made in the photosynthesis lab contributes to our hypothesis because in the second graph from figure 2, there is a consistent pattern on the amount of Bicarbonate concentration on the rate of photosynthesis. The initial mass of our water sample was 7.1g and after 45 minutes the mass was 7.6g, resulting in a mass difference of 0.5g with a 7.04% change in mass. Based on the results from figure 3, our water sample Long Island City Deep, is a hypertonic solution. Some possible errors that might have skewed our data: Not making the appropriate dilutions would not allow us to determine what effect enzymes and substrate concentration have on enzyme activity. In the Osmosis and Diffusion activity is not tying the string on the bag tight enough

Conclusion and Future work

Based on our studies we can conclude that the quality of aquatic life depends heavily on the levels of salinity, dissolved oxygen and carbon dioxide. By conducting these experiments, we were able to have a better understanding on how each individual factor affect one another. When the levels of dissolved oxygen and carbon dioxide are out of their optimum range processes such as photosynthesis cannot be carried out normally. The amount of salinity affects aquatic life because certain life organisms acquire a certain amount in order to have a balanced homeostasis. Throughout this research I've learned the heavy impact mankind has on the environment as well as aquatic life. I have gained knowledge of the East River through these experiments because it allowed me to study the effects these factors have on our water quality. Based on what I have learned I can use this advantage to educate others in order to secure a better future to our surroundings. With this data, I learned that the dissolved materials played an important role on aquatic organisms. In the future I can conduct other experiments that would allow me to observe more chemical and physical properties of other water bodies.

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Discussion

References