# The Presence of Iron in Surface Waters 

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Phelan, James, "The Presence of Iron in Surface Waters" (1967). Honors Theses. 357.
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A

> Report
> for

Dr. Joe Nix

Special Studies 491

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\text { May } 10,1967
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THE PRESENCE OF IRON IN SURFACE WATERS

Iron is found in most natural surface waters, although in widely varying amounts, depending on several factors relating to the water and its environment. This report is desioned primarily to gain an understanding of these factors contributing to the stability and solubility of iron in these natural waters.

It has been found that iron is present in rather large amounts in aquatic vegetation and that this same vegetation is important is adding iron to surface water and in removing dissolved or syspended iron from the water. When iron is combined with organic ions or molecules, it may form comnlexes that are both stable and soluble. When larser water nlants are present, they may produce rather drastic chances in iron content of natural waters. When dense prowths of a auatic plants cover all or nearly all the open water surface, the exchance of air and water is either completely stonped or greatly reduced. Too, aquatic lants such as algae, funæi, and bacteria greatly reduce the oxysen supply already present. ${ }^{1}$

One may ask, "Why does a change in oxysen supply affect the stability and solubility of iron?" This question may

[^0]perhaps be best answered by the following discussion concerning the relation of redox notential, pH , and iron content: The measurement of redox potentials in ground water is very difficult, for the concentrations of the ions in solution that produce such a potential are usually very small. The reserve ions that have the ability to stabilize the redox potential is similar to the solute supply which stabilizes the oH . When related to redox potential, this effect is called poising, and when related to pH , it is called buffering. A laboratory determination of pH is possible, for the pH of a ground samrle may not closely show the hydrosen-ion activity of the water in the aquifer.

Upon reaching the atmosphere, ground water is generally only weakly poise. When the water reaches the zone near or above the water table where oxygen is found, some oxygen immediately diffuses into the solution, the result being a weter-oxyren couple which stabilizes the redox potential
as lons as a contact with air is available. This occurs because the laroe available supply of oxypen found in the atmosnhere sives a poisinf, effect.

If the ground water has only a few parts per million of iron, it will have a low redox potential in the aquifer, but when oxygenation takes place the redox potential rises. At saturation, there is a possibility of the water containing oxyoen in a molarity 10 to 100 times srester than that of iron. The subsequent adjustment of iron to this new and stable redox notential ultimately results in the oxidation of
nearly all the iron to the ferric state.

$$
\mathrm{Fe}^{++}+3 \mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{Fe}(\mathrm{OH})_{3} \quad 3 \mathrm{H}^{+}+\mathrm{le}^{-}
$$

The above oxidation and hydrolysis to ferric hydroxide exolains the precipitation of iron from the water and the subsequent incorrect laboratory determinations of iron in water. Thus, the iron content should be determined in the field. ${ }^{2}$

The following data taken from the Caddo River and DeGray Creek provides excellent examples for the above discussion.

| LOCATION | SAMPLINS PERIOD | pH | IRON |  |
| :--- | :---: | :--- | :--- | :--- |
| Caddo River <br> (Station 5) | 6 | $(10 / 22 / 66)$ | 7.2 | 0.19 |
| DeGray Creek | 6 | $(10 / 22 / 66)$ | 6.8 | 1.90 |

Carbon dioxide-bicarbonate-carbonate equilibria occurs in most natursl waters. Thus, the type of rock(such es those rich in calcite) over which the water runs may be very imrortant to the iron content, for it is this equilibrium thet detrrmines the linits for solubility of iron under reducinc conditions. This occurs beceuse of its effect on pH , which in turn affects the solubility of iron.

[^1]
[^0]:    $1_{\text {Hem, John D., and Oborn, Eugene T., 1961, Microbiologic }}$ Factors in the Solution and Transport of Iron: U.S. Geolorical Survey water-Supply Paper $1459-\mathrm{H}$.

[^1]:    $=2_{\text {Hem, John D., 1960, Restraints on Dissolved Ferrous Iron }}$ Imoosed by Bicarbonate Redox potential, ana pH: U.B. Geolo्यc ai survey nater-suply paper 1459-B.

