

Laboratory Experiment of the Evaporation Level of Surface Water Salinity

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ABSTRACT:Evaporation causes а very significant mass loss in any kind of surface water body and is significant for attempts at resource recovery (Uche and Amgbara, 2020). Evaporation from surface water depends on the water body's salinity. This investigation is on direct measurement at the scale of natural water body collected from a free surface water body in AfikpoEbonyi State. Five different salts such as NaCL, NaOH, MgSO₄, NH₄CL, and KNO₃ were selected based on the fact that saline water bodies contain these salt compounds. These salts applied to the 500g of water varied as follows 0g, 15g, 25g, 35, 45g, and 55g. These samples after mixing were kept at room temperature in the water laboratory of AkanuIbiam Federal Polytechnic Unwana and evaporation allowed to take place within 24hours interval for 15 days. The results show that the higher the concentration of salt in the water samples the less evaporation occurred. Ammonium Chloride (NH₄CL) has the highest rate of evaporation followed by Potassium Nitrate (KNO₃), Magnesium Sulphate (MgSO₄), Sodium Hydoxide (NaOH) and Sodium Chloride (NaCL) has the least rate of evaporation.

Keywords: Salinity, surface water, evaporation, salts

I. INTRODUCTION

Evaporation from a free surface water body occur when water from liquid or solid state passes into the vapour state and diffuse into the atmosphere. Evaporation from open water bodies depends on many factors which include water temperature, air temperature, air humidity and air velocity above the water surface. The effect of surface water salinity on evaporation increases the complication of the challenge of quantifying evaporation (Salhotra et al., 1987). It has been difficult to ascertain the rate of evaporation of a saline water by direct measurement at the scale of natural water bodiesMor et,.al (1995). Salinity can cause unfavourable environment and hydrological situation that restrict the production of crops. Certain factors contribute immensely to the development of saline water upon evaporation on a free surface water. Salinity desiccates the soil as air pores in the spoil act like a sponge and the salt water is drown up to the surface of soil as a result of capillary rise buildings, roads and other structures can be damaged. Salinity is one of the serious environmental problems that cause osmotic stress and reduction in plant growth and crop production as when enough salt accumulate in the root zone. It also affects soil physical properties by causing fine particles to bind together into coarse aggregates. The factors contributing significantly to the development of saline soil are tidal flooding during raining seasons (June - October). When there is water logging within two meters of the surface, there is high concentration of salt surrounding compounds within the environmentscausing damages to building, roads and other structures. Soil riches vavels (pH) in coastal regions range from 6.0 - 8.4, the organic matter content of the soil is also pretty low 1.0 -1.5 %. The mechanism that induces salinity can work very quickly to affect vegetation In drainage lines and on sporting ovals and to damage buildings, roads and pipe systems. Salinity shortens the life of infrastructures and increase building cost due to the need for protective works and use of the higher specification materials. There are different types of salinity such as dryland salinity, irrigation salinity, urban salinity, river salinity and industrial salinity.

II. SALINITY

• DRYLAND SALINITY. Dryland salinity occurs when removal or loss of native vegetation and its replacement with crops and pastures that has shallower roots and different



water use requirements result in more water reaching the groundwater system. The groundwater rises to near the ground surface in low-lying areas or on the break of slope called discharge. Groundwater can also flow underground directly into streams. The groundwater carries dissolved salts from the underlying soil and bedrock materials through which it travels. As saline groundwater comes close within two meters to the soil surface, salt enters the plant root zone. Even where groundwater does not bring much salt with it, the water-logging to the root zone alone can damage or kill vegetation. The impact of water-logging and salinization will vary depending on soil type, climate and land use.

- IRRIGATION SALINITY. The main reason for this type of salinity is the application of large volumes of irrigation water, equivalent to as much as four times the average naturally occurring rainfall, compounded by the replacement of native vegetation by plants with different water sue patterns.
- URBAN SALINITY. Urban salinity is a combination of dryland and irrigation salinity. It is caused both by rising water tables due to clearing, and by the application of additional quantities of water from watering garden and parks; leaking water, sewage and drainage pipes; and obstruction or modification of natural surface and sub-surface drainage paths.
- RIVER SALINITY. River salinity is caused by saline discharges from dryland, irrigation and urban salinity into creeks and rivers. Overtime, as salinity within catchments worsens, the quality of river water declines, becoming more saline.
- INDUSTRIAL SALINITY. Industries, towns, intensive agriculture, coal mines effluents influence salinity.

III. IMPACTS OF SALINITY

Increasing river salinity levels will put pressure on the provision of municipal water supplies and increases treatment and infrastructure costs. Where the water for municipal supplies is ''hard'' (the salinity being made up of high levels of calcium and magnesium), water may require extensive and expensive treatment before it is suitable for consumption. Hard water can result I n damage to hot water system and household appliances, and increase use of soap, detergents and water conditioners. The implications of river salinity are also costly for agricultures. The yields of some particular sensitive plants such as rice and horticultural crops, are affected by saline water at levels as low as 700EC (Electrical Conductivity). The impact of river salinity is more costly on high value enterprises such as irrigated crops and horticultural crops. Dryland salinity can lead to lower agricultural production, lower profitability due to costs of mitigation, reduced yield, changed land use, and in extreme case to the total removal of land from agricultural production. Recreational and tourism opportunities may also be reduced, and cultural heritage including aboriginal cultural heritage may be damaged.

✤ Effect of Salinity on Evaporation

Salinity is the saltiness or dissolved salt content of water in soil such as sodium chloride, magnesium, and calcium sulfates, and bicarbonates of a water body. In oceanography it has been traditional to express salinity not as percent but as per million (parts per thousand) which is approximately grams of salt per kilogram of solution. Salinity is the sum weight of many different elements within a given volume of water (DeCosmo et al., 1996). The factors significantly contributing to the development of saline soil are tidal flooding during wet seasons (June – October) direct inundation by saline water, and upward or lateral movement of saline ground water during dry season (November – March) the severity of salinity problems increase with the desiccation of the soil. It affects crops depending on degree of salinity at the graphical stages of growth, which reduces yield and in severe case cause total vield lost.

* Factors Affecting Evaporation

There are many factors affecting the rate of evaporation from a free surface water. These include:

• Climate Effects

For evaporation to take place, energy is required to change the water from a liquid form to a vapour form. The air above the evaporation pond becomes saturated with water vapour and this moist air must be removed in order to allow the process to affect the rate of evaporation.

• Temperature

Heating the water molecules to the required temperature for vaporization will change the water from liquid form to vapour form. The higher the temperature, the higher the rate of evaporation. When the temperature of the water increases, it makes the water molecules to gain energy, move faster and escape at a faster rate.

Evaporation Rate

The rate at which water evaporates depends on the temperature of the liquid as how compact the molecules bonds to each other, the surface area of the liquid and the temperature, pressure, and air movements in the surrounding



into which evaporation occurs. The warmer the liquid, the faster the evaporation rate. The stronger the molecules bond to each together the slower the evaporation rate.

• Properties of the Liquid being evaporated

Dissolved salts in the water result in a lower saturation vapour pressure due to the decreased chemical potential of the water and thus, lower the evaporation rate. The second law of thermodynamics implies that an increase in ion activity as a result of the presence of solute reduce the chemical potential of a liquid solvent and also the rate of spontaneous transformation of the liquid phase into the vapour phase (Kokya and Kokya, 2006).

• Surface Area

The surface area of the water body will have an effect on the evaporation rates for some areas of similar climate and evaporation rates from a maximum value for small water bodies, as determined for a standard evaporation pan, to a fraction of that value for larger of that bodies. The larger the exposed surface the faster the evaporation rate. Procedures for calculating evaporation rates indicate that an evaporation rate is directly proportional to vapour pressure. The vapour pressure of saline water is lower than that of fresh water resulting in a reduction in evaporation cohesive forces between the dissolved ions. Evaporation rate decreases exponentially with increasing salinity. The evaporation rates of saline solutions are calculated by multiplying the rates of evaporation of water by a salinity factor.

• Humidity

Composition will have an effect on the humidity level at which evaporation will cease, for example, for a water body saturated with sodium chloride there will be no evaporation during periods when the humidity is above 70% for other salt evaporation may cease at lower humidity levels. If the humidity level is high, less evaporation occurs as saturation levels are quickly reached when humidity is high. It is more difficult for water to evaporate. If the space above the water becomes completely filled with water vapour, (100% relative humidity) then evaporation is balanced by the opposite process i.e. condensation.

IV. METHODS

Sampling Techniques

The sample of these salts include: Sodium Chloride, Magnesium Sulphate, Sodium Hydroxide, Ammonium Chloride and Potassium Nitrate.

The concentration of the salts were varied ie (0g, 15g, 25g, 35g, 45, and 55g)

Laboratory Procedure

500grams of water was measured and put into an evaporation can which served for the control sample. ,then another 500grams of water with the salts while the salts varies with 0g, 15g, 25g, 35g, 45g and 55g. The mixtures were thorough.The entire set-up were kept at room temperature, evaporation allowed to take place at 24hours interval for 15days; to check the effect of salinity on evaporation process at free water surface. Electronic weighing balance was used to know the amount of water that has been evaporated in the concentrations every day for 15 days. Graphs of cumulative evaporation were plotted against time in days. After the experiments, data were collected on evaporation for the different salts.

V. RESULTS

The rate of evaporation was determined by using electronic weighing balance to know the amount of water that has been evaporated in the concentrations every day for 15 days Fig. 4.1 Variation of Cumulative Evaporation with time for control experiment of all salts



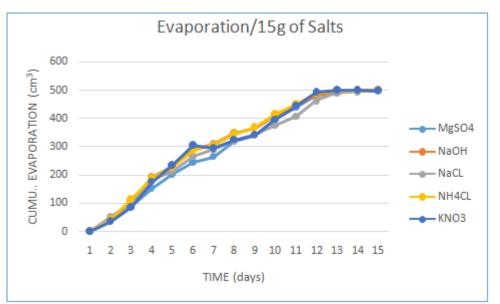


Fig. 4.1 Variation of Cumulative Evaporation with Time at 15gConcentration experiment of all salts

Figure 4.1 shows variation of cumulative evaporation with time at 15gconcentration. At this concentration, evaporation is highest with NH_4CL because on the 12^{th} day, all the moisture in the can

has evaporated. Next is KNO_3 , $MgSO_4$, NaOH, while the least is NaCL on day 12. This shows that inhibition of evaporation is highest with NaCl while NH_4CL has the least inhibition.

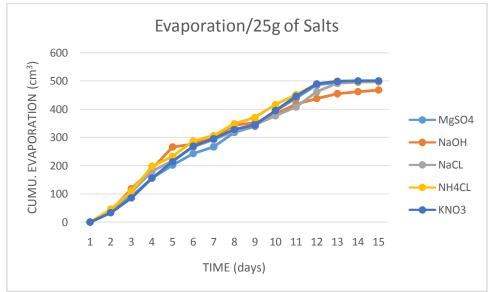


Fig. 4.2 Variation of Cumulative Evaporation with Time at 25gConcentration experiment of all salts

Figure 4.2 shows variation of cumulative evaporation with time at 25g concentration. At this concentration, evaporation is highest with NH4CL, KNO_3 and $MgSO_4$ because on the 13th day, all the moisture in the cans had evaporated. Next is NaCL,

while the least is NaOH on day 13. This shows that inhibition of evaporation is highest with NH_4CL , KNO_3 and $MgSO_4$ while NaOH has the least inhibition.



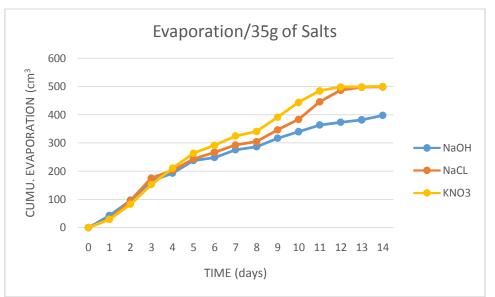


Fig. 4.3 Variation of Cumulative Evaporation with Time at 35g Concentration experiment of all salts

Figure 4.3 shows variation of cumulative evaporation with time at 35g concentration. At this concentration, evaporation is highest with NH_4CL and $MgSO_4$ because on the 12^{th} day, all the moisture in the cans had evaporated. Next is

 KNO_3 , NaCL, while the least is NaOH on day 12. This shows that inhibition of evaporation is highest with NH₄CL, and MgSO₄ while NaOH has the least inhibition.

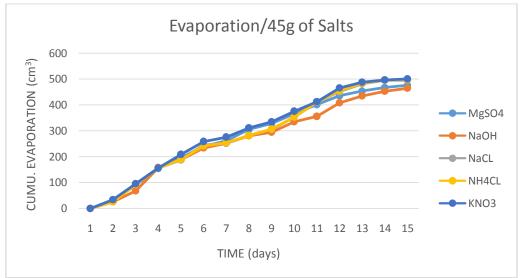


Fig. 4.4 Variation of Cumulative Evaporation with Time at 45gConcentration experiment of all salts

Figure 4.4 shows variation of cumulative evaporation with time at 45g concentration. At this concentration, evaporation is highest with NH_4CL because on the 12^{th} day, all the moisture in the cans had evaporated. Next is KNO_3 and $MgSO_4$,

followed by NaCL, while the least is NaOH on day 12. This shows that inhibition of evaporation is highest with NH_4CL while NaOH has the least inhibition.



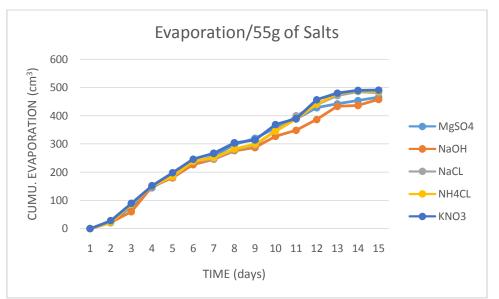


Fig. 4.5 Variation of Cumulative Evaporation with Time at 55gConcentration experiment of all salts

Figure 4.5 shows variation of cumulative evaporation with time at 55g concentration. At this concentration, evaporation is highest with KNO_3 because on the 12^{th} day, all the moisture in the cans had evaporated. Next is NH_4CL which have values 498cm^3 followed by MgSO₄, and NaCL while the least is NaOH on day 12. This shows that inhibition of evaporation is highest with KNO_3 while NaOH has the least inhibition.

VI. CONCLUSION

From the experiment, it was observed that evaporation decreases the mass per unit volume of water with time. The higher temperature and surface area, the higher the rate of evaporation. It was observed that the higher the quantity of salt, the lesser the evaporation that is 15g of salt has the highest rate of evaporation followed by 25g, 35, 45, and 55. On comparison between the salts, Sodium Chloride (NaCL) has the least rate of evaporation, followed by sodium hydroxide (NaOH), Magnesium sulphate (MgSO₄), ammonium chloride (NH₄CL) and potassium nitrate (KNO₃) which has the highest rate of evaporation. Salinity decreases as evaporation takes places in the present of various salt. Precipitation will be limited in areas prone to salinity and this will invariably affect the hydrological balance.

REFERENCES

 Adams, T. C., (1934). Evaporation from Great Salt Lake. Bull Am. Meteorol, Soc. V. 15 P. 35-39.

- [2]. Ahmed, A. b., Omoogun, G. A. &Shaida, S. S., (2000). Typical Salinity factor produced from treatment.
- [3]. Akridge, D. G., (2008). Methods of calculating evaporation rates during salt production, journal of Archaeological Science American Water Works Association, (1999). Water Quality and Treatment. A hand book of Community water supplied (5th edition ed.). R. O. Letterman, ED.
- [4]. Bowen, I. S., (1926). The ratio of heat losses by conduction and by evaporation from any water Surface; Phys. Rev. Scr. 2, V. 27, P. 779-787
- [5]. DECosmo, J., K. B. Katsaros, S. D. Smith, R. J. Anderson, W. A. Oost, K. Bumke, and Chadwick, H. (1996). Effect of Salinity on Evaporation from water surfaces. Journal of Physical Oceanography, Vol.40.
- [6]. Du Plessis, J. Burger, R. Swart and Museven (2006). A desalination guide for South Africa Municipal Engineers Pectoris Water Research Commission.
- [7]. EPA (United State Environmental Protection Agency, (1987). Salinity principal for water treatment.
- [8]. Kokya B. A. and Kokya T. A., (2006). Proposing a formular for the evaporation measurement from salt water resources hydrological processes.
- [9]. Lee, C. H., (1927). Discussion of "Evaporation on Reclamation Projects". Trans. Am. Soc. Civ. Engineers, V. 90, P. 340 – 343.



- [10]. Mickey, M., (2001). Membrane concentrate disposal practices and regulations. US Department of Interior
- [11]. National Research Council (1928). International Critical Table: New York, McGraw-Hill, !st ed., V.3.
- [12]. Saxton K. E., Rawls W. J., (2006). Soil Water Characteristic Estimates by Texture and Organic Matter
- [13]. for Hydrologic Solutions. Soil Sci. Society of America.
- [14]. Schulte, F., (2006). Handbook for the operation of water treatment works, Petroria. Water Institute of South Africa.
- [15]. Sverdrup H.U., Johnson M. W., and Fleming R. H., (1942). The Oceans: New York, Prentrice-Hall.

- [16]. Sverdrup H. U. (1937). On the evaporation from the ocean. Jour. Marine Research V. 1, No. 1 P.3 -14
- [17]. Swinbank, W. C., (1951). The measurement of vertical transfer of heat and water vapour by eddies in the lower atmosphere: Jour. Meteorology. V.8, No. 3, p. 135-145.
- [18]. U. S. Geological Survey, (1954). Water-loss investigation, Lake Hefner studies, technical report: U. S. Geol Survey Prof. Paper 269 (Also published as U. S. Navy electronics laboratory report
- [19]. 327, 1952)
- [20]. Young, A. A. , (1947). Some recent evaporation investigations: trans. Am. Geophys. Union V. 28. No. 2 p. 279-284.