#### **3-6 Mechanism of water softening**

Clay minerals may attain a net negative charge by ion replacement in which Si(IV) and Al(III) ions are replaced by metal ions at similar size but lesser charge. This negative charge must be compensated by association of cations with the clay layer surfaces. Because of their structure and high surface area per unit weight, clays have a strong tendency to absorb cations(Ca<sup>++</sup>, Mg<sup>++</sup>) from water. One noticed this in sample 1 which has 17% clay (page 62) and it is always the best for removing water total hardness. But in samples 2, 3, 4 and 5 when hard water is passed through a bed of modified clays, the calcium and magnesium are taken up and held by the sodium cation exchanger (modified clays) which simultaneously gives up equivalent amount of sodium ion exchange for them.

Reverse Osmosis, this is an applied pressure that can remove many types of molecules and ions and bacteria from water i.e. isolation of large molecules (retaining of ions through pores with small molecules like H<sub>2</sub>O can pass) the membrane used, is of special type of polymeric matrix e.g. West Soba, Blue Nile, East Nile, could be used for earths of existence as solutes.

Ion exchange: replacing ofCa<sup>+2</sup>and Mg<sup>+2</sup>by Na<sup>+</sup> and K<sup>+</sup>. Zeolites can be used for softening water (Metuchen 1961). Cation exchangers have adsorption power for divalent cations more than monovalent cations. Adsorption power for all cations increases with increasing atomic weight, table (19), on page 67 sample 3,4 and 5 are the best because the high content of easy removing ions of Na<sup>+</sup> and K<sup>+</sup>. But sample 1and 2 are the least.

Molecular sieves: species with a kinetic diameter which makes this too large to pass through a modified clays pore are effectively "sieved" this "sieved" effect can be utilized to produce sharp separation of molecules by size and shape.

Aluminosilicates can be used as molecular sieves also zeolites can be used as sieves, those having pore of definite sizes ranging from 4Åto13Å as stated by (Bulut and Baysal 2006). It is noticed that the task was done as volume isolation of cations more than electrostatic or exchange or even adsorption, the best samples are 1, 2, and 3.

#### Adsorption of Ca<sup>+2</sup> and Mg<sup>+2</sup> on clays:

It could be visualized as adsorption of  $Ca^{+2}$  and  $Mg^{+2}$  on clays as physical adsorption that need only contact between cations and clays as "physisorption" i.e. involving electronic, electrical, electrostatic attachment on surfaces that could be classified by easy desorption, and for further attachment as chemisorption like any chemical reaction between the cations and earths.

One can involve XRD and EM to measure the degree of the strong (binding) attachment between  $Ca^{+2}$  and  $Mg^{+2}$  on the earths. Sample 1 is hematite. However sample 2 contains enstatite and albite. Sample 3 is labrodorite. But samples 4 and 5 are diopside and plagioclase feldspar respectively.

The X-Ray Diffraction showed that all earths were aluminosilicates with different orientations and the bonding strength between the constituent varieties of compounds to give a cluster of specific nature.

Reverse osmosis mechanism depends upon the physical and mechanical force applied to enforce the membrane, while the bigger particles are retained on the membrane and then discarded.

Ion exchange mechanism that uses the concept of solvation power of  $H^+$  and  $OH^-$  i.e. by exchanging the Na<sup>+</sup> and K<sup>+</sup> by Ca<sup>+2</sup> and Mg<sup>+2</sup>. As a result the ions of Ca<sup>+2</sup> and Mg<sup>+2</sup> are isolated by attachment to the clay precipitates while Na<sup>+</sup> and K<sup>+</sup> are passing through with the water to the boiler without making hardness. Therefore samples 4 and 5 are the best suit for ion exchange mechanism.

The molecular sieves of earths are working as inert volume isolation of substrates, regardless of ionic or even covalent or dative bonds.

The sieving mechanism is working for isolation of calcium and magnesium large ions by the surfaces of the earths and even further by fractures and inner pores of earths.

The XRD and Electron Microscopy studies for earths can give a clear picture for that mechanism.

The adsorption is the best mechanism to apply for  $Ca^{+2}$  and  $Mg^{+2}$ onto earths using many concepts, but the most suitable one is that of Freundlich solid /liquid adsorption i.e. the amount of the adsorbates( $Ca^{+2}$  and $Mg^{+2}$ ) on the earths as adsorbents could give a clear view about the attachment nature, this is known as physical adsorption,(physiosorption).

The physiosorption is usually followed by chemical adsorption (chemisorption) i.e. like a chemical reaction with strong bonds between the  $Ca^{+2}$  and  $Mg^{+2}$  and the earths.

The chemisorption follows  $\sigma$ - bond attachment accompanied by  $\pi$ - back donation resulting in a very strong and stable attachment.

Sample 5, 4 and 3 may be good examples of hardness removal by the adsorption mechanism.

One can classify the 5 earths according to the previous mechanism for the best removing action of the hardness (scales) from the industrial waters. The adsorption is the best mechanism followed by ion –exchange and the finally molecular sieves. For the revers osmosis, one can conduct studies shortly.

## Conclusion

This study shows the possibility to use some local soil as a synthetic resin to remove hardness from industrial water. The following results were obtained according to the efficiency of hardness removal by different alkali concentration, for NaOH the efficiencies were in the range 50% to80% for Jurdiga sample, the West Nile clay in the range 18% to 70%, East Nile clay 13% to 70%, Blue Nile 8% to 62% and West Soba 18% to 48%. The pilot results gave efficiency of scale removal of 40% from (250g earth treated with 0.1M NaOH), gave 18 liters of soft water while the calcined samples gave 8 liters. The SPSS results showed that there were no immaterial differences in using NaOH or KOH. The differences that in concentration of the base as given by ANOVA.

The chemical analysis and XRF showed that the samples were aluminum silicates. While XRD showed the dominant of hematite, of West Nile clay while feldspar of East Nile clay, Blue Nile and Jurdiga.But West Soba was diopside and microcline mineral. Grain size distribution range from silty clay to fine sand.

# Recommendation

The used earths contain amounts of  $Ca^{+2}$  and  $Mg^{+2}$  as waste products, then for further research work, the waste could be moulded, calcined and material impact tested as ceramic bricks. Also for further work should be conducted to investigate the recovery efficiency of made resin as well as be examined other local clays to be a potential polymer resin to remove hardness in water.

## REFERENCES

- Ahmed, S. and Igbal, Y Ghani F. (2008) "Phase and micro structure of brick- clay soil and fired clay- bricks from some areas in Peshawar Pakistan", *Journal of the Pakistan Material Society*, vol. 2, pp.33-39.
- Alkan, M., Demirbas, O., Celikcapa, S. and Dogan, M. (2004), "Hazardous matter' vol. 116, pp.135.
- Alley, BES. and Robert, E. (2007), Water quality control hand book. Mc Grow-Hill, second edition .
- American Public Health Association, American Water Work Association, Water Pollution Control Federation (1966) *Standard and Methods of the Examination of Water and Wastewater. Including Buttom Sediments and Sledges*, 70, 146.
- Arden, U., T. (1968).Water purification by ion exchange. London: Butter Worth, P. 1.
- Aruidhas, G. (2004). Molecular of Structure and Spectroscopy. New Delhi: Prentice, Hall of India.
- Asheoft, W N. and Mermin, D N. (1967). Solid State physics. United State of America: Harcourt college publishers.
- Barmatl, CT. and Patgiri, K D. (1996). Physics and chemistry of soils . New age international publishers, p. 105.
- Brady, N. (1974). The nature and Properties of Soils. Macmillan publishing Co, mc,8<sup>th</sup> edition.
- Brown, T L IRH. and LE, May. (1977). Chemistry the central Science. Prentice Hall.

- Bulut, Y. and Baysal, Z. (2006). "removal of Pb<sup>2+</sup>from waste water using wheat bran" *Journal of Environmental Management*. Vol. 78, pp.107-113.
- Deer, A. Howie, A R. and Zussman, J. (1992). Introduction to the rock forming minerals. Hong Kong: Long man group, second edition, pp. 105-120.
- Duggen, M J. and Inskip ,M J. (1985). "Childhood exposure to lead in surface dust". *Public Health Review*. Vol. 13, pp2-54.
- Filtration facts (http: // www.epa. Gov/ ogwdw/ faq/ pdfs/ fs- health seriesfiltration – pdf), sept 2005, U. S. Environmental Protection Administration , pp. 6-7. Accessed 6 January 2013.
- Frye, Keith. (1993). Mineral Science and Introduction Survey. Macmillan publishing company.
- Gill, R. (1996). Chemical Fundamental of Geology. Chapman and Hall, Second edition, p. 95.
- Grim, E Ralph. (1962). Applied Clay mineralogy. New York: McGraw-Hill Book Company, pp. 166-204.
- Guggenheium, Stephen. and Martin, RT. (1995). "Definition of clay and clay mineral" *Journal report of the AIP EA nomenclature and CMS nomenclature* .PP. 255-256.
- Guglielmo, Ventura. and Lara, Risegari. (2012). The Art of CryogEnics: low-temperature, Experimental Techniques. SurendraKum.
- Harvey, D. (2000). Modern analytical chemistry. McGraw Hill.
- Ibrahim, S W. (2014). Removal of naphthenic acid from Sudanese crude oil by zeolites and clays. Sudan.

- Jana, Dypayan. (2007). "Clinoptilolite-apromsingpozzolan in concrete". "A New look at an Old pozzolan".29<sup>th</sup> ICMA Conference. Quebec City, Canada: Construction Materials Consultants, Inc .Retrieved 7 October 2013.
- John, stone S J. and John, stone M G. (1961). Minerals for the chemical and allied industries. Second edition. Chapman and hail.
- Klein, cornelis. (2001). Manual of Mineral science, 22<sup>nd</sup> edition. New York: Wiley, pp.245-250.
- Klein, C. and Hurlbut, Ir C S. (1999). Manual of mineralogy. 21 edition. New York: John Wiley and Sons, INC, p.70.
- Konta, J. (1995) "Clay and Man: clay raw materials in the service of man". *Appliedclayscience*, vol. 10, pp.257-335.
- The Associated press (via NY times), (2011). Levels of Radioactive Materials Rise Near Japanese Plant.
- Lewis, D W. and M c Conchie, D. (1994). practical Sedimentology. second edition. Chapman and hall.
- "Lime softening" (http:// water. me. VCCS.Edu / concepts / softening lime .html). Retrieved 4 Nov 2001.
- Lower, Stephen. (2007). Hard water and water softening.
- Malakootian, M., Nouri, J. and Hossaini, H. (2009). "Removal of heavy metals from paint industry's waste water using leca as an available adsorbent". *International Journal Environment Science and Technology*, vol. 6, pp.183-190.
- Mellor, JW. (1941). Intermediate inorganic chemistry, long mans.
- Metuchen, NJ. (1961). Water treatment for industrial and other uses. second edition, Permutit Company.

- Naddafi, K., AM, Mahvi., Nasseri, S., M, Mokhtati. and H, Zeraati. (2005).
  "Evaluation of the Efficiency of clay pot in removal of water impurities", *Iranian JEnv Health Sci Eng*, vol. 2, pp.12-16.
- Orumwense, E. (1996). "Removal of lead from water by adsorption on kaolintic clay". *Journal of chemical and biotechnology*, vol 58, pp. 363-369.
- Petti, J E J. (1975). Sedimentary rocks. Third edition, Harper and kow publisher.
- Pitty, AF. (1979). Geography and Soil properties. First Published, Methuen and Co Ltd.
- Potts, PJ. (1992). Handbook of silicate rock analysis. Blackie.
- Powell, T Sheppard. (1954). Water conditioning for industry. first edition, McGraw Hill, company, INC.
- Ppcoetzee, (1998). Scale reduction and scale modification effects induced by Zn. [PDF].
- Prapat, Pentamwa., Wipasinum, Thipthara. and Suparat, Nuangon. (2001)."Hardness removal from ground water by synthetic resin from waste plastics "*international conference on environmental and computer science* vol.19.pp. 59-63.
- Putnis, A. (1995). Introduction to mineral science. BA spriters Ltd, Over Wallop, Hamp-shir.
- Rob, D., Xanat, F., Melanie, P. and Georges. (2003). "T.point- of use water treatment technology investigations in Nepal. Clean water for Nepal" MA02139.
- Scott, M Auerbach., Kathleen, ACarrado. and Prabirk, Dutta. (2003).
  Handbook of Zeolite Science and technology.CRC press.

- Selley, R C. (1982). An Introduction of Sedimentology. Second edition, Academic press mc.
- Skoog, A D., Holler, FJ. and Nieman, AT. (1997). Principle of Instrumental analysis. fifth edition, United state of America.
- Skooge, A Douglas. (1995). principle of Instrumental analysis. Third edition, United State of America: Saunders College publishing.
- Smith, P RJ. (2000). "Use of Glauconitic clay to remove copper from effluents produced during manufacture of printed circuit boards", *on line* Available at <a href="http://www.aecuk.com/PCB">http://www.aecuk.com/PCB</a>. Html>. vol. 50,pp. 393-399.
- Sorg, Thomas J., Schock, Michael R. and Lytle, Darren A. (1999)."Ion exchange softening : effects on metal concentration", *JournalAWWA91* vol. 8,pp. 85 -97.
- USGS- US. Geological Survey Office of Water Quality. "USGS Water Quality information: water hardness and alkalinity" .usgs.gov.
- Vjvnov, Aleksei., .Fulton, John L. Huthwelker, Thomas, Pin, Sonia. Mei. Donghai-Schenter, Gregory K., Govind, Niranjan. Camaioni., Donald M. Hu., JianZhi, Lercher. and Johannes, A. (2014). "Quantitatively Probing the AL Distribution in Zeolites" *Journal of American Chemical Society* vol. 136, n. 23, pp. 8296-8306.
- Vogel's. (1989) Textbook of quantitative chemical analysis. fifth edition, New York: John Wiley and Sons.
- Wainerda, E R. and Uken, A E. (1971). Modern Methods of geochemical analysis. NewYork: Plenum press.
- Weaver, Charlese. and Pollard, Lind. (1973). The chemistry of clay Minerals. New York: Elsevier and Scientific Publishing Company, pp.2-4, 133.

- WHO, Hardness in drinking water, Background document for development of WHO guid lines for drinking water quality world health organization (2003).
- Worden, H R. and Morad, S. (2003). Clay Minerals Cements in sand stones. Black well publishing.
- Xu, Ruren., Pang, Wengin., Yu, Jihon., Huo, Qisheng. and Chen, Jiesheng. (2007). Chemistry of zeolites and related materials: synthesis and structure, John Wiley and Sons.
- Yoder, HS. (1957). Experimental studies on micas, as synthesis, clay and clay minerals, proc. Natl 16<sup>th</sup> conf. clay.42-61.
- Zhang, G., Wasyliuk, K. and Pan, Y. (2004). the characterization and quantitative analysis of clay minerals, application of software infrared reflectance spectroscopy, vol. 39, pp. 1348.
- Zoltai, T. and Stout, J H. (1984). Mineralogy Concepts and Principles .Burgess.