## Chapter IV

## Results

Under non-steady state aeration of a deoxygenated water, an overall transfer coefficient (K ) can be La determined by:-

1. the graphical method and

2. the method of least squares

By graphical method, K is determined from the La slope of a semilogarithmic plot of the concentration deficit  $(C_s-C_1)$  versus time as inllustrated in Apendix B.

$$K_{La} = 2.3 \log_{10} \frac{(c_s - c_1)_{t_1}}{(c_s - c_1)_{t_2}} / t_2 - t_1 \text{ min.}^{-1}$$

The method of least squares as suggested be Fair and Geyer (1954) is also illustrated in Apendix B, where K is determined by the equations stated below:-

n.A +  $\leq x.B$  =  $\leq y$  ..... (1)  $\leq x.A$  +  $\leq x^2.B$  =  $\leq xy$  ..... (2) where n = number of samples taken x = elapsed time in minutes y = log (C<sub>s</sub>-C<sub>1</sub>) B = overall oxygen transfer rate constant with base 10

$$K_{La} = \frac{B}{0.4343} = overall transfer coefficient$$

K is corrected to standard conditions at 20°C tem-La perature and one atmospheric pressure.

$$K_{La(20^{\circ}C)} = K_{La(T)^{\circ}\Theta}^{(20-T)}$$

On bubble aeration the value of  $\Theta$  varies from 1.016 to 1.037 (Eckenfelder and Barnhart, 1960)

A value of 1.024 was applied to this study.

A straightline relationship between log  $(C_s-C_1)$ and time (t) was tested by Pearson (r) and the significance of Pearson r, as suggested by Downie and Heath (1970).

Pearson (r) = 
$$\frac{n \cdot \xi x y - (\xi x) (\xi y)}{\sqrt{\left[n \cdot \xi x^2 - (\xi x)^2\right] \left[n \cdot \xi y^2 - (\xi y)^2\right]}}$$

t, the significance of Pearson (r) is equal to  $\frac{r}{(1-r)^{\frac{1}{2}}} \times (n-2)^{\frac{1}{2}} \text{ with } (n-2) \text{ degrees of freedom.}$ The presented values of (r) for different is the presented values of (r) for different is a significance and degrees of freedom to simplify significance test for Pearson (r).

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Since the present of chloride in tap water observed, varied from 18 to 38 mg/l, the correction for chloride of 0.008 mg/l at 25°C per 100 mg/l chloride as suggested by Fair & Geyer (1954) determined in this study is very small (0.003 mg/l), it may be negligible when compare to the accurate reading of only one decimal observed from the oxygen meter.

All the so obtained results are shown in Appendix C.