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APPENDIX A

Slope of the stopped-FIgrams was used to plot versus the concentration of standard phosphate solutions to make a calibration graph. Procedure for evaluation of the slope from the stopped-FIgram was illustrated in Figure A.1.

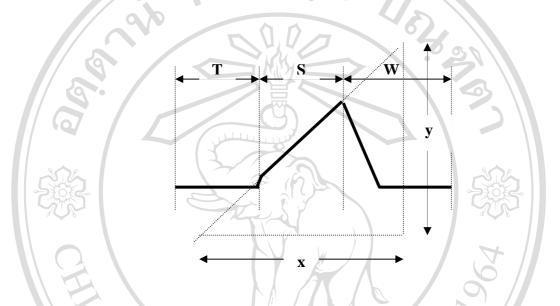


Figure A.1 Procedure of measuring slope of the stopped-FIgram: T = Travelling time , S = Stopping time, W = Washing time, y = detector signal (mV) and x = time(s)

In this work, full scale sensitivity range of recorder was 2 V while is corresponded to x cm on chart.

So, 1 cm length on chart recorder = $\underline{2} = 0.02$ V = 20 mV

Then, from Figure A.1;

١.,

y-value = height y (cm) * 20 mV

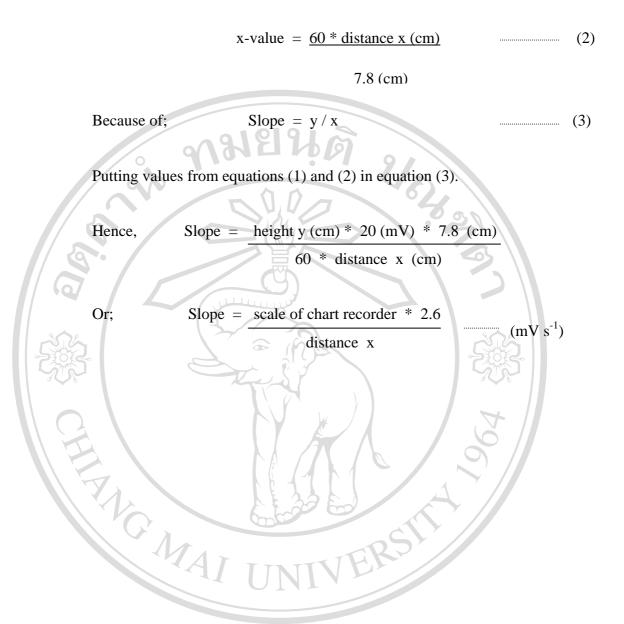
Chart speed of chart recorder was 7.8 cm min⁻¹

Thus, distance 7.8 cm equivalent to the time of 60

So; distance x cm equivalent to the time for 60 * x s

S

niv



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APPENDIX B

1. Batch method for determination of phosphate [38]

Pipette 15.00 ml of standard phosphate solution into a 25.00 ml volumetric flask. Added 0.02 M sodium molybdate, 0.25 % w/v ascorbic acid and 0.15 M nitric acid at 2.0, 2.0 and 1.0 ml, respectively into the flask. Adjusted the volume to the mark by water and let the mixture at room temperature for 10-15 minutes before measuring absorbance at 660 nm.

2. Titrimetric standard method for determination of chlorate [12] The procedure is based upon the reaction between chlorate and iodide in the presence of concentrated hydrochloric acid :

 $ClO_3^- + 6I^- + 6H^+ \rightarrow Cl^- + 3I_2 + 3H_2O$

The liberated iodine is titrate with standard sodium thiosulphate solution.

Place 25.00 ml of the chlorate solution in a glass-stoppered conical flask and added 3 ml of concentrated hydrochloric acid followed by two portions of about 0.3 g each of pure sodium hydrogen carbonate to remove air. Added immediately about 1.0 g of potassium iodide and 22 ml of concentrated hydrochloric acid. Stopper the flask, shaked the contents and allowed it to stand for 5-10 min. Titrate the solution with a standard solution of 0.1 M sodium thiosulphate.

3. Batch method for determination of chlorate [39,41] liding

-Making a calibration graph

Pipetted 25 mg Γ^1 of standard potassium chlorate solutions: 0.00, 1.00, 2.00, 3.00 and 4.00 ml into a 25.00 ml volumetric flask. Added 3 x 10⁻⁴ M 5.00 ml indigo carmine solution for each flask and adjusted the volume to the mark by 1:1 hydrochloric acid. Heat the solution in water bath at 50 °C for 10 minute before measuring absorbance at 610 nm.

-Determination of soil samples

The extracted chlorate that was prepared as mentioned in 2.4.2 was analysed using the same procedure as above for the calibration graph.



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APPENDIX C

1. Sampling of soil samples [42]

Sampling of soil samples from same area was collected throughout at least 15 points and mixed together. This sample was called "composite sample". Soil was dug at the depth of 6 inches (15 cm).

2. Preparation of samples [42]

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Samples were dried in clean room and without dust. After drying, matrix in soil was abandoned. Soil was grinded, sifted with a sieve and collected in plastic bag before analysis.

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2/02/03/15

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List of Publication:

- Somnam, S., Sangthong, W. and Saiyasombat, W., "Mulberry pulp preparing", 27th Congress on Science and Technology of Thailand, Songkla, 2001.
 - Somnam, S., Jayavasti, S., Jakmunee, J. and Grudpan, K., "Stopped-flow injection analyzer for determination of soluble phosphorus in fertilizer", 28th Congress on Science and Technology of Thailand, Bangkok, 2002.

 Somnam, S., Jakmunee, J. and Grudpan, K., "Development of stopped-flow injection system for determination of phosphate in soil samples", 29th Congress on Science and Technology of Thailand, Khon Kaen, 2003.



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