

Adsorption of Lambda Cyhalothrin onto Athi River Sediments: Apparent Thermodynamic Properties

S. S. Sherrif¹ and V. Madadi²

^{1, 2}Department of Chemistry, School of Physical Sciences, College of Biological and Physical Sciences, University of Nairobi, P.O. Box 30197-00100, Nairobi, Kenya

ABSTRACT

Lambda-Cyhalothrin is extensively used in Kenya. It is effective against many vegetable pests, rice and other disease agents. The adsorption of Lambda-Cyhalothrin onto sediments of Athi River was studied using UV-visible; the results were analyzed by fitting the data into isotherm models including Freundlich and Dubinin-Radushvich (D-R) plots to understand its environmental impacts. Different masses of sediments were spiked with different concentrations of L-Cyhalothrin. The amounts of the pesticide adsorbed were determined by analyzing the difference between the UV-visible Spectrophotometer data and the initial concentrations. Various isotherm models were used to explain the adsorption characteristics but Dubinin-Radushvich (D-R) model with average R2: 0.935, 0.938 and 0.898 fitted best in this experiment followed by Freudlich model, R2: 0.783, 0.899 and 0.812 for all sediment samples, i.e., Upstream, Midstream, and Downstream, respectively. The spontaneity of the adsorption process was also realized in ΔG (Gibb's free energy) values as predicted by Freundlich. In both models, Midstream ΔG was negative (-) showing complete spontaneous characteristics. Generally, adsorption capacities of Athi river sediments were low due to their low mineral contents and total organic Carbons including other physicochemical properties specifically, textures, Nitrogen content including temperatures and moisture contents. **Keywords :** Lambda-Cyhalothrin, Adsorption, Sediment, Isotherm Model, and Spontaneity.

I. INTRODUCTION

The use of pesticides including insecticides, herbicides, fungicides, rodenticides, and acaricides can be dated before the 16th century when chemicals such as arsenic sulphide, Paris green (Copper acetoarsenite) were used to control Malaria transmissions. Though the use of synthetic organic pesticides began around 1940 (De, 2010). Lambda-Cyhalothrin is an artificial insecticide that characterizes the biochemical effects of natural pyrethrin pesticide (He et al., 2008). It is the active ingredient in many insecticides such as Karate, Warrior, and Icon that are commonly applied to control insects and pests in public health emergency, agricultural/horticultural farming and in homes (CDPR, 2006). It is effective against many crop and disease vector insects and ticks, (He et al., 2008). The use of Pesticides including Lambda-Cyhalothrin in livestock and crop farming have led to increased food security and livelihood in many developing countries though uncontrolled excessive use is hazardous to human life

as well to the environment (FAO/WHO, 2012). Lambda-Cyhalothrin and other pesticides rapidly disappear upon application in the environment (Senoro et al., 2016). Studies have proven that approximately 2-5% of applied pesticides go to targeted organisms and subsequently the rest end up in the environment (Garrcia et al., 2011). Pesticide substrates in the environment undergo several chemical processes absorption, degradation, including oxidation, halogenations, transfer and other microbial actions (De, 2010). Transfer of pesticide residues away from the targeted sites through volatilization, spray drift, agricultural runoff, leaching, or crop removal is a major determining factor in assessing the overall environmental impacts of pesticides,(Senoro, et al., 2016).

Acute (short-term) human exposure can lead to skin irritation, burn or allergic reaction, (Sharda, 2013). It is very toxic particularly to various species of fish and other aquatic organisms (He, *et al.*, 2008).

II. MATERIALS AND METHODS

The following instruments, materials and reagents were used: UV-Visible spectrophotometer (UV-1700 Schamadzul), Mini Orbital Shaker, Analytical balance (Fischer scientific A-160), Lambda-Cyhalothrin (karate 25g/kg pure), Glass bottles, Distilled water, Stop watch, Acetone (90% pure) and River sediment from Athi River.

PROCEDURES

The samples were dried for a week and pulverized to maintaining its originality. The Calcium, Potassium and sodium contents were analyzed using Flame Photometer, whereas other elements including Phosphorus, Magnesium and Manganese were determined calorimetrically, (Mehlich, A. 1953). The available total organic carbon content was analyzed by oxidative spectrophotometer, (Gislason, E.A. et al., 2005). The total N was determined thru distillation followed by Titration, (Jan-Ake Persson et al., 2008). The PH was also determined simply by a pH meter and other trace metals (Fe, Zn and Cu) using an Atomic Absorption Spectrophotometer (AAS), (Yang, S. et al., 2005). The cation ion exchange capacity of the metallic elements at neutral pH was determined by Titration after distillation, (Carroll et al., 1959 and Tuner, R.C. et al., 1966).

To conduct the adsorption study, 2, 4, 6, 8, 10, 20, 40,

60, 80, and 100g/L were prepared and adsorption values determined at the UV-vis between 200 to 900nm. The existence of the adsorption/desorption equilibrium of 2.0g, 1.0g, 0.5g, 0.2g and 0.1g of the soil samples were investigated. The dried sediments were placed in glass bottles and shaken with the pesticide for an hour. The aqueous parts were poured and filtered with a Whitman filter paper to obtain a clear concentrated L-Cyhalothrin solution. The absorbance of Ce and Qe was obtained at the UV-Visible spectrophotometer at 218 nm. ΔG , n and K were found using 0.5g of each sample mixed with deionized water. The samples were spiked with 50, 40, 30, 20, and 10ppm of L-Cyhalothrin and shaken for 15min, 30min, 45min and 60min using an orbital shaker. Similarly, concentration study of L-Cyhalothrin was carried out using UV visible Spectrophotometer.

RESULTS AND DISCUSSION

The soils properties have significant impacts on its adsorption capacity. The profile of the soils according to Table 1.1 on Athi River soil showed that the organic carbon content is low for all three sampling sites, (0.54, 0.91 and 0.75 respectively). This means the adsorption capacity of these samples is generally low thus not proportionately. Another factor that influences the adsorption is the physical dipole-dipole attraction (Van der Waals force) that improves the adsorption capacity for the soil. Below in Table 1.1, are the sediment profiles of the Athi River.

| Field | Upstream (A) | Midstream (B) | Downstream (C) | |
|---------------------|--------------|---------------|----------------|--|
| Sediment depth | 2-5cm | 2-5cm | 2-5cm | |
| Parameters | Value | Value | Value | |
| Total Nitrogen % | 0.08 | 0.11 | 0.10 | |
| Total Org. Carbon % | 0.54 | 0.91 | 0.75 | |
| Phosphorus ppm | 8.00 | 1100 | 40.00 | |
| Potassium me % | 0.82 | 1.06 | 0.60 | |
| Calcium me% | 23.90 | 30.20 | 5.10 | |
| Magnesium me% | 3.28 | 3.36 | 1.76 | |
| Manganese me% | 2.12 | 2.68 | 0.61 | |
| Copper ppm | 1.56 | 1.55 | 1.71 | |
| Iron ppm | 191.00 | 174.00 | 39.30 | |
| Zinc ppm | 5.00 | 5.94 | 2.04 | |
| Sodium me% | 1.00 | 1.29 | 0.50 | |
| Elect. Cond. Ms/cm | 0.93 | 0.86 | N/A | |

Table 1.1 sediment profile of Athi River.

CONCENTRATION STUDY

The prepared standard solution of Lambda-Cyhalothrin behaved well at λ max 218 nm at UV-Visible spectra as shown below in *figures 1.1*. The plot of absorbance against concentration at 218 nm shaped a straight line. This means Beer's Law is obeyed:

Α= έ CL.....1.1

In the above equation, A is the absorbance, L is the path length of the light and C is concentration. The symbol $\hat{\epsilon}$, is the molar absorption coefficient for the chemical species of interest.

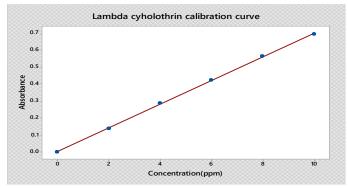


Figure 1.1 Plot of Concentration from 0-12 ppm of Lambda-Cyhalothrin.

ADSORPTION ISOTHERM

The process of adsorption is defined as the summative accumulation of a chemical species at the interface of an aqueous solution and a solid phase (Sposito, 1989). The chemical species is called adsorbate and the surface at which it accumulates is called adsorbent. The adsorption kinetics of Lambda-Cyhalothrin onto Athi River sediments was carried out by preparing 10ml of Lambda-Cyhalothrin in various concentrations (10ppm-50ppm) and shaken at varying times (15 minutes, 30

Table 1.2 Freudlich adsorption data.

minutes, 45 minutes and 60 minutes) on an orbital shaker. The mixture was filtered and the concentrated residual solution was analyzed using UV-spectrophotometer (SHEMAZU 1700). The amount of the pesticide adsorbed (mg/g) was determined using the Vanderburgh and Van Griekenm's formula (Igwe and Abia, 2006):

$$Q =$$

Where Q is the amount of solute adsorbed, V is the volume of the absorbate, C_i and C_f are the initial and final concentrations, respectively. W is the weight in gram of the adsorbent and data fitted into freudlich and Dubinin-Radushkevich isotherm models.

1.1 FREUDLICH ISOTHERM MODEL

The same data obtained from the statistics of the research were exposed to Freundlich isotherm which gave positive (+) ΔG values for sample C (Downstream) and negative (-) ΔG for samples A and B (Up and Mid Streams, respectively). This means that samples A and B underwent spontaneous adsorption using Freudlich isotherm while sample C was nonspontaneous. Thus the Freundlich Isotherm is characteristic of heterogeneous surface, (Freundlich, 1906). It can be expressed as:

From equation 1.2, qe (mg/g) is amount of pesticide adsorbed and Ce (mg/L) is the equilibrium concentration and n is a constant that measures the adsorption of non-linearity between the solute concentration in the solution and the adsorption. n is characteristic of the quasi-Gaussian heterogeneity related to the adsorption surface. Kf (L/g) is the Freudlich isotherm constant.

| Time | Sample | 1/n | LnK ^F | KF | _R 2 | ΔG |
|----------|--------|--------|------------------|--------|----------------|-----------|
| (minute) | | | | | | |
| | | | | | | |
| 15 | А | 0.8957 | 1.883 | 6.5732 | 75.4 | -4,665.27 |
| | | | | | | |
| | _ | | | | | |
| | В | 1.002 | 1.691 | 5.4249 | 82.9 | -1,921.94 |
| | | | | | | |

| | C | 2.136 | -0.322 | 0.7247 | 78.1 | 797.78 |
|----|---|--------|---------|---------|------|-----------|
| 30 | А | 0.9518 | 1.834 | 6.2589 | 86.1 | -4,543.88 |
| | В | 1.841 | 0.5828 | 1.7915 | 99.2 | -1,444.56 |
| | С | 1.966 | -0.2400 | 0.7867 | 90.4 | 594.62 |
| 45 | А | 0.8852 | 1.852 | 6.3726 | 75 | -4,588.48 |
| | В | 1.174 | 1.549 | 4.7066 | 84.4 | -3,837.67 |
| | С | 3.228 | -2.2880 | 0.1015 | 74.2 | 5,668.69 |
| 60 | А | 1.053 | 1.635 | 5.1295 | 76.6 | -4,050.85 |
| | В | 1.249 | 1.484 | 4.41055 | 93.3 | -3,676.74 |
| | С | 2.930 | -2.151 | 0.1164 | 82 | 5,329.27 |

The linearized equation for Freudlich isotherm is expressed as follows:

From this expression, plots of lnqe versus ln C_e as presented in this linear form of Freundlich Isotherm have slopes at 1/n and intercepts at lnKF. Graphs (in figures 1.2, 1.3, and 1.4) below were generated using the data highlighted in Table 1.2 above. The data fitted strongly in the adsorption of Lambda-Cyhalothrin unto Athi River sediments of the three (3) sampling areas. This can be validated from the regression (r²) values ranging from 0.742 to 0.992 and the linearity of the plot in figs 1.2, 1.3, and 1.4 below:

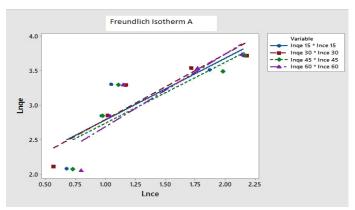


Figure 1.2 : Freundlich Isotherm for Sample

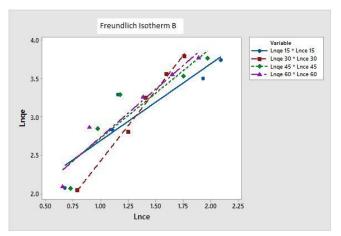


Figure 1.3: Freundlich Isotherm for Sample B

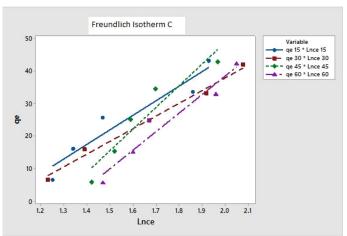


Figure 1.4: Freundlich Isotherm for Sample C

1.2 DUBININ-RADUSHKEVICH (D-R) ISOTHERM MODEL

This model was used to show the apparent adsorption energy heterogeneity at sites of adsorption. This model has fitted well with high solute activities including intermediate concentration range data. This approach was originally applied to differentiate between the physical and chemical adsorptions of metal ions with it main free energy (E) per molecule (Dubinin, M.M. 1960). The equation and its linear forms are expressed below:

From the above expressions, q_D is adsorption strength of soil/sediment (mg/g), B_D is Dubinin-Radushkevich constant (mol²/KJ²), qe is amount of adsorbate at equilibrium (mg/g), and E is Polanyi potential (Dubinin isotherm constant). T is the absolute temperature (k), R is the universal gas constant (j/molk) and Ce is the adsorbate equilibrium concentration (mg/L).One unique characteristic of this model is that it is temperature dependent. This allows for all suitable data to lie in the same curve called characteristic curve when adsorption data at separate temperatures are plotted as function of the amount adsorbed (lnqe) against the energy (E). Table 1.3 below summarizes D-R data used to generate the graphs in figures 1.5, 1.6, and 1.7:

| Time | Sample | Ln qD | BD | qD | \mathbf{R}^2 | E (Kj/mol) |
|----------|--------|-------|-----------|----------|----------------|-------------|
| (minute) | | | | | | |
| 15 | А | 3.854 | -0.000002 | 47.1814 | 91.8 | -353,553.39 |
| | В | 3.844 | -0.000002 | 46.7120 | 93.8 | -353,553.39 |
| | С | 4.421 | -0.000006 | 83.1794 | 86.4 | -117,851.13 |
| 30 | А | 3.795 | -0.000001 | 44.4782 | 98 | -707,106.78 |
| | В | 3.839 | -0.000002 | 46.4790 | 89.6 | -353,553.39 |
| | С | 4.203 | -0.000005 | 66.8867 | 96.9 | -141,421.36 |
| 45 | А | 3.841 | -0.000002 | 46.5720 | 91.5 | -353,553.39 |
| | В | 3.995 | -0.000002 | 54.3258 | 94.1 | -353,553.39 |
| | С | 5.139 | -0.000010 | 170.5451 | 85.7 | -70,710.68 |
| 60 | А | 3.952 | -0.000002 | 52.0393 | 92.6 | -353,553.39 |
| | В | 3.888 | -0.000002 | 48.8132 | 97.5 | -353,553.39 |
| | С | 4.754 | -0.000010 | 116.0475 | 90.2 | -70,710.68 |

From Table 1.3 above, the potential energy (E) can be calculated from the value of B_D using the formula: Inqe = $\ln q_D - B_D E^2$. For example, in 15 minutes of shaking time, $\ln qe 15 = 3.844 - 0.000002 E^2$, $r^2 = 0.938$. $Lnq_D = 3.844$, $B_D = 0.000002 q_D = e^{3.844}$ and E = -353,553.39 kj/mol.

Generally, when the adsorption energy value falls below 8Kj/mol, the adsorption process is said to be characterized by physiosorption, i.e., physical binding of the pesticide to the surface of the soil/sediment. If the value is higher than 8kj/mol but less than 20kj/mol, it is said to be characterized by ion exchange and similarly, its values beginning from 20kj/mol and above are said to be characterized by particle diffusion. From Table 1.3 E (kj/mol) values are lesser than 8kj/mol, which deduces that the adsorption process of Lamda-Cyhalothrin was predominantly a physiosorption process. From fig 1.5 to fig 1.7, the data fitted well in Dubinin-Radushkevich isotherm at the various shaking times for samples A (Upstream) and sample B (Midstream) whereas sample C (Downstream) has some deviations, though the regression (r^2) values range from 0.857 --- 0.980.

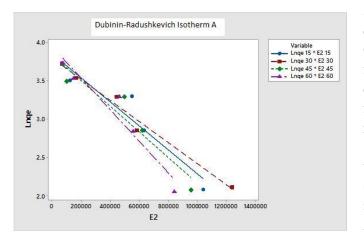


Figure 1.5: D-R Isotherm for Sample A

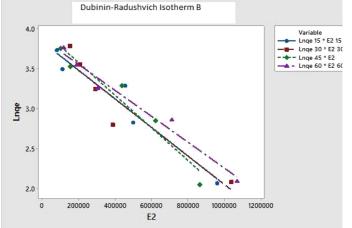


Figure 1.6: D-R Isotherm for Sample B

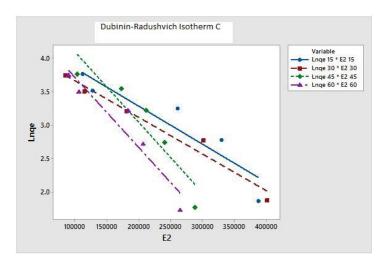


Figure 1.7: D-R Isotherm for Sample C

III. CONCLUSION

Adsorption of Lambda-Cyhalothrin decreases with increase in mass of loam sediments. The longer the contact time the higher the adsorption. Increase in concentration results in decreased proportion of adsorption. This happens because at high initial concentration, the number of moles of Lambda-Cyhalothrin available to the surface area is high, so functional adsorption becomes dependent on initial concentration. Adsorption of lambda-Cyhalothrin followed Fredlich-peterson isotherm model with regression values ranging from 0.954 to 0.992. As shaking time increased from 15 minutes to 60 minutes, the value of n increased from 0.1998 to 0.2914. The ΔG (Gibb's free energy) was 11.7946 \pm 0.3Kjol/mol which indicated that adsorption of Lambda-Cyhalothrin by Athi River sediments was spontaneous, indicating high affinity to the sediments.

IV. REFERENCES

- [1]. CDPR (2011). California Air Resource Board, Cdpr.ca.gov. Retrieved in November 20, 2011.
- [2]. Carroll, Dorothy (1959). Ion exchange in clays and other minerals, Geological Society of America Bulletin, **70** (6): 749-780.
- [3]. De, A.K. (2010). *Environmental Chemistry* 7th edition. New Age International Publishers. New Delhi, India, 198-261.
- [4]. Dubinin, M. M., and Radushkevich, L. V. 1974). Equation of the Characteristic Curve of Activated

Charcoal, Proceedings of the Academy of sciences of the USSR, Chemistry Section **55**: 331-333.

- [5]. FAO/WHO (2012). Evaluation Report based on submission of data from BharaRasayan Limited (TC). In: Food and Agriculture Organization, ed. FAO specifications and evaluations for agricultural pesticides- Lambda-Cyhalothrin. Rome: FAO, 15-23.
- [6]. Freundlich, H.M.F. (1906). Over the Adsorption in Solution. J. Phys. Chem. **57:**385–471.
- [7]. Garcia, S.G., Scheiben, D. and Binder, C.R. (2011). The weight method: a new screening method for estimating pesticide deposition from knapsack sprayers in developing countries, *Chemosphere*, 82, 1571-1577.
- [8]. Gislason, E.A. and Craig, N.C. (2005), cementing the foundations of thermodynamics: comparison of system-based and surroundings-based definitions of work and heat, *J. Chem. Thermodynamics***37**: 954-966.
- [9]. He, L-M., Troiano, J., Wang, A., Kean Goh, K. (2008) Environmental Chemistry, Ecotoxicity and Fate of Lambda-Cyhalothrin: *Reviews of Environmental Contamination and Toxicicolgy*, Vol.195, 71-89, Springer New York, NY, USA.
- [10]. Igwe, J.C and Abia, A.A. (2006). A Bioseparation process for removing heavy metals from waste water using biosorbents, African journal of Biochemistry, 5 (12), 1167-1179.
- [11]. Jan-Ake Persson, Marten Wennerholm, Stephen O'Halloran, (2008). Handbook for Kjeldahl Digestion: 11-42.
- [12]. Mehlich, A. (1953). Determination of P, Ca, Mg, K, Na, and NH4. North Carolina Soil Test Division (Mimeo 1953), 23-89.
- [13]. Senoro, D.B., Maravillas, S.L., Ghafari, N., Rivera, C., Quiambao, E. and Lorenzo, M.C.M. (2016). Modeling of the residues Transport of Lambda-Cyhalothrin, Cypermithrin, Malathion and Endosulfan in three different Environmental compartments in the Philippines. *Sustainable Environmental Research*. <u>http://dx.doi.org/10.1016/j.serj.2016.04.010</u>.
- [14]. Sharda, USA LLC (2013). Sharda Lambda-Cyhalothrin 1EC Insecticide, 7217 Lancaster Pike, suite A, Hockessin, DE 19707.
- [15]. Simard, R.R. and Trans, T.S (1993). Mehlich III-Extractable Elements. In: M.R. carter, Ed. Soil

Sampling and Methods of Analysis: 43-49.

- [16]. Sposito, G. (1989). The Chemistry of Soils. Oxford Univ. Press, Oxford, England.
- [17]. Tuner, R.C. and Clark J.S. (1966). Lime potential in acid clay and soil suspensions. Trans. Comm. II and IV Int. Soc. Soil Science, pp.208-215.
- [18]. Yang, S.Y. and W.L. Chang, (2005).Soil Sci., 170, 55.