

A MATHEMATICAL MODEL FOR CONTAMINANT TRANSPORT IN AN UNCONFINED AQUIFER SYSTEM

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Abstract

The analysis of contaminant transport into an aquifer system, showing the behavior of contaminants over a time period is of paramount importance in the study of geological behavior of the aquifer system. In this paper a mathematical model for Contaminant Transport in an Unconfined Aquifer was formulated using a Laplace transform. The governing equation for solute transport given by Kumar (20014) was used for the formulation. The analysis of contaminant transport in an aquifer system, showing the behavior of contaminants for different values of Diffusive transport into the unconfined layer (α) for $0 < t \leq 7$ was modeled, the diffusion within the layer (y), and the size of the Aquifer (b) kept constant. For a uniform source of contamination at it $c^*(x)$ was observed that for different values of a , b and y the level of contamination reduces over the domain.

Keywords: Contaminant transport, unconfined aquifer, dispersion, advection, piezometric and head

Introduction

Water is therefore the most essential element for man's well-being, social and economic progress. Groundwater offers the most abundant source of water to man. It is the cheapest and the most constant in quality and quantity (Olasehinde, 2014). It is observed that in many developing countries, groundwater plays a major source of support for domestic needs and irrigation purposes (Thangarajan, 2014). Water shortages occur quite often in many areas of the world, calling for optimal management of both surface and groundwater resources (Helmut 2014, Jacques, 2014). Groundwater quality is usually better, since they are naturally more protected, once polluted, their restoration is more difficult, calling for optimal control of groundwater contamination (Amlan & Bithin, 2015; Fetter, 2014). Sagei et al. (2015) considered a fractured confined porous aquifer, and came up with a modeled solute equation, that analysed the effect of non-Fickian diffusion into surrounding rocks.

The aim of this paper to formulate a mathematical model that can be used to simulate a solute transport and analyze the movement of contaminants in an unconfined aquifer system.

Model Formulation

We considered a governing equation for solute transport given by Kumar (2014) as:

$$\frac{\partial C}{\partial t} = -\frac{\partial}{\partial x_i} (C V_i) + \frac{\partial}{\partial x_i} \left(D_{ij} \frac{\partial C}{\partial x_j} \right) + R_c \quad i, j = 1, 2, 3 \quad (2.1)$$

$$V_i = \frac{K_{ij}}{\theta} \frac{\partial h}{\partial x_j} \quad (2.2)$$

where;

- D_{ij} = Coefficient of Hydrodynamic Dispersion,
- C = Concentration of the Solute in the Source or Sink Fluid
- R_c = Sources or Sinks
- V_i = Seepage Velocity

- K_y = Hydraulic Conductivity
- h = Hydraulic Head
- x_i = Coordinate system

The initial condition (specification of the Concentration distribution of Solute at initial time $t = 0$), can be written as;

$$C(x) = C^U(x) \quad x \in \Omega \quad (2.3)$$

where;

$C^U(x)$ indicates a known Concentration distribution over the domain of interest (Ω).

$C^U = 1$ indicates a Uniform Source of Contamination at $x = 0$

Here we consider an unconfined in the formula tion of our model. Three regions were considered; the upper layer (porous layer1), middle layer (Aquifer layer) and the lower layer (porous layer 2), as shown in figure 2.1

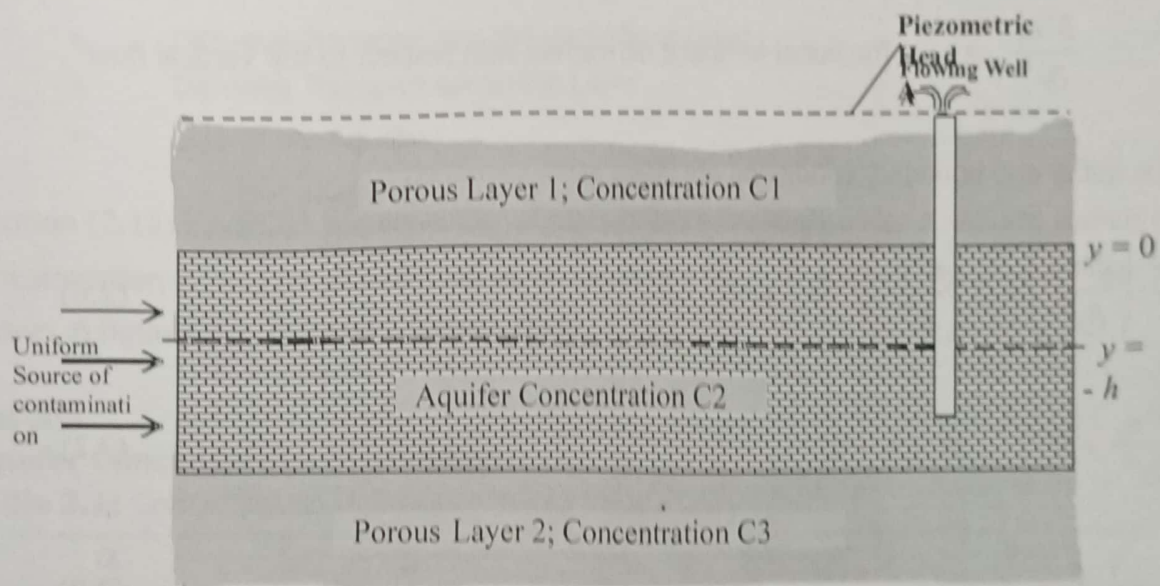


Figure 2.1 Diagram of an unconfined Aquifer for the model formulation

The partial differential equation describing the contaminant transport in the upper layer is given as;

$$\frac{\partial c_1}{\partial \tau} = D_1 \frac{\partial}{\partial y} \left[\frac{\partial^\alpha c_1}{\partial y^\alpha} \right] \quad 0 < y < \infty, \tau > 0 \quad (2.4)$$

where;

D_1 = Effective Diffusivity of Porous layer

c = contaminant concentration

Similarly, the partial differential equation describing the contaminant transport in the aquifer is given as;

$$\frac{\partial c_2}{\partial \tau} + \beta \frac{\partial^\gamma c_2}{\partial \tau^\gamma} = \bar{D}_2 \frac{\partial}{\partial y} \left(\frac{\partial^\alpha c_2}{\partial y^\alpha} \right) + D_2 \frac{\partial}{\partial x} \left(\frac{\partial^\alpha c_2}{\partial x^\alpha} \right) - v \frac{\partial c_2}{\partial x} \quad (2.5)$$

$$-h < y < 0, 0 < x < \infty, \tau > 0$$

D_2 and \bar{D}_2 = Effective Diffusivities, in the aquifer in the x and y -direction, respectively;

v = Fluid Velocity

τ = Time

β = Capacity Coefficient

$\frac{\partial^\gamma c_2}{\partial \tau^\gamma}$ = Fractional-in-time Derivatives

$\frac{\partial^\alpha c_2}{\partial x^\alpha}$ = fractional-in-Space derivatives with respect to the horizontal flow

$\frac{\partial^\alpha c_2}{\partial y^\alpha}$ = fractional-in-Space derivatives with respect to the vertical flow

The initial and boundary conditions are given as;

$$\bar{D}_2 \left(\frac{\partial^\alpha c_2}{\partial y^\alpha} \right) = D_1 \left(\frac{\partial^\alpha c_1}{\partial y^\alpha} \right) \quad \text{at } y=0 \quad (2.6)$$

$$\frac{\partial^\alpha c_2}{\partial y^\alpha} = 0 \quad \text{at } y = -h \quad (2.7)$$

$$c = \frac{1}{h} \int_{-h}^0 c_2 dy \quad (2.8)$$

Method of Solution

Applying Duhamel's Theorem (Randall and Leveque, 2005), and for a uniform source of contamination at $x=0$, we obtain the solution of problem (2.5) as;

$$\psi(\tau, x) = 1 - \left(\frac{-1 + e^{-\tau}}{\tau} - \frac{-1 + e^{x(b\gamma \cos(\pi\gamma) + \pi \cos \beta^2)}}{x(b\gamma \cos(\pi\gamma) + \pi \cos \beta^2)} \right) \left(\frac{\sin x b \sin \pi\gamma}{\gamma + 1} - \frac{x \sin \pi\beta}{\beta + 1} \right) \quad \text{and} \quad (2.9)$$

$$C(t, x) = \frac{\partial}{\partial t} \int_0^t C_0(t-\tau) \varphi(\tau - x, x) d\tau \tag{2.10}$$

where;

φ is defined by equation (2.9)

and

$$C = 1 - \left(\left(\frac{-1 + e^{-t}}{t} \right) - \left(\frac{xb\gamma + xt^\beta \left(\frac{1}{2}\gamma \right) + \frac{1}{2}}{1-\gamma} \right) + \left(\frac{xt^\gamma \left(\frac{1}{2}\beta \right) + \frac{1}{2}}{1+\beta} \right) \right) \tag{2.11}$$

That is;

$$C = \frac{-\Gamma(-\beta - 1)\Gamma(-\gamma + 1) + xt^\beta \Gamma(-\gamma + 1) + bxt^\gamma \Gamma(-\beta - 1)}{\Gamma(-\beta - 1)\Gamma(-\gamma + 1)} \tag{2.12}$$

and,

$$\beta = \frac{\alpha}{\alpha + 1} \tag{2.13}$$

where;

$$\Gamma(\beta) = \int_0^1 t^{\beta-1} e^{-t} dt; \quad \text{and} \quad \Gamma(\beta + 1) = \int_0^1 t^\beta e^{-t} dt; \tag{2.14}$$

- α = Diffusive Transport into the unconfined Layer
- γ = Diffusive Transport within the Layer
- b = Size of the Aquifer

Solution (2.12) describes the behaviour of contaminants transport under a uniform source of contamination $C^U(x)$ for different values of values of α , b and γ for $0 < t \leq 7$ as shown in figures 3.1, 3.2 and 3.3 respectively.

Results

Aquifer Concentration Distribution

Table 3.1: Concentration Distribution Values for $\alpha, b,$ and γ

α	b	γ	t
1.00	0.50	0.50	$0 < t \leq 7$

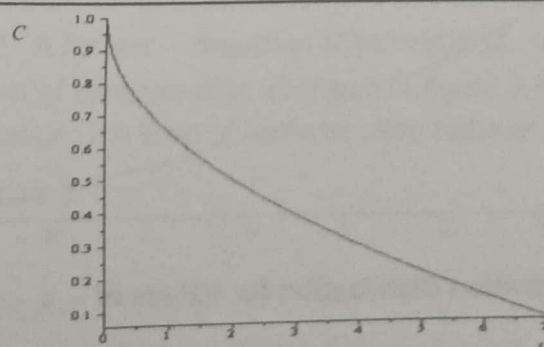


Figure 3.1: Aquifer Concentration Distribution for Values of $\alpha, b,$ and γ

Table 3.2: Concentration Distribution Values for $\alpha, b,$ and γ

α	b	γ	t
0.50	0.50	0.50	$0 < t \leq 7$

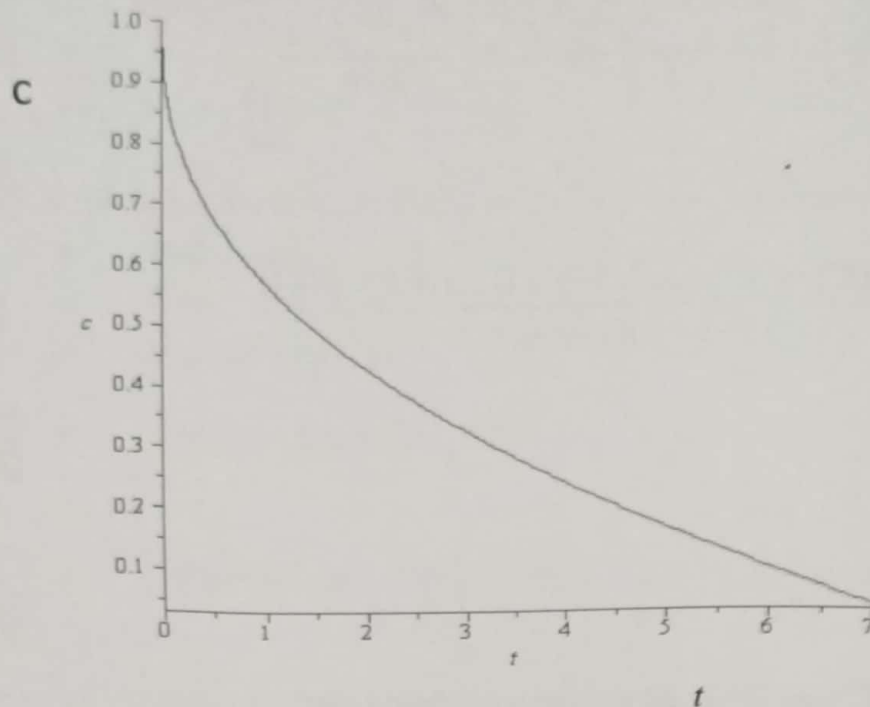


Figure 3.2: Aquifer Concentration Distribution for Values of $\alpha, b,$ and γ

Table 3.3: Concentration Distribution Values for $\alpha, b,$ and γ

α	b	γ	t
0.20	0.50	0.50	$0 < t \leq 7$

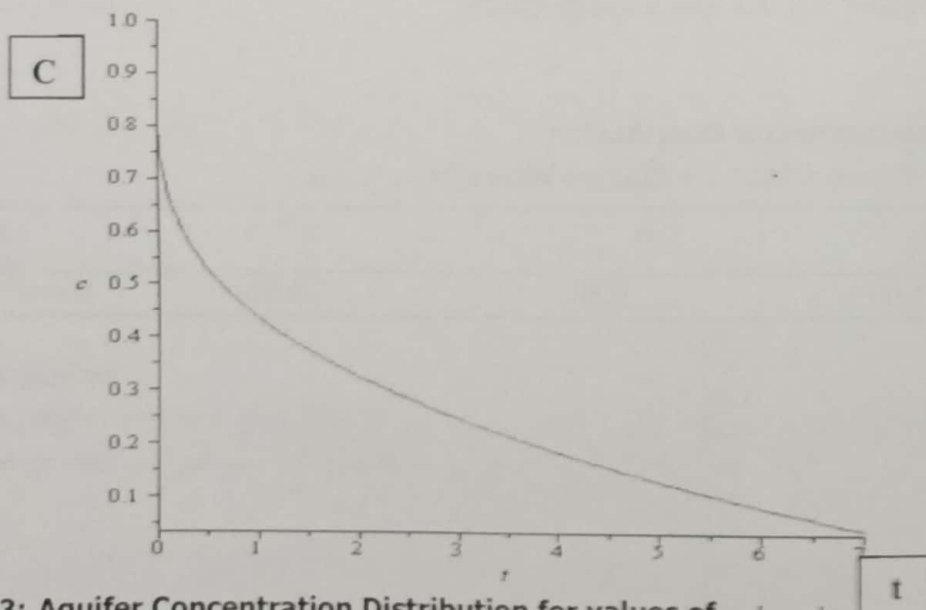
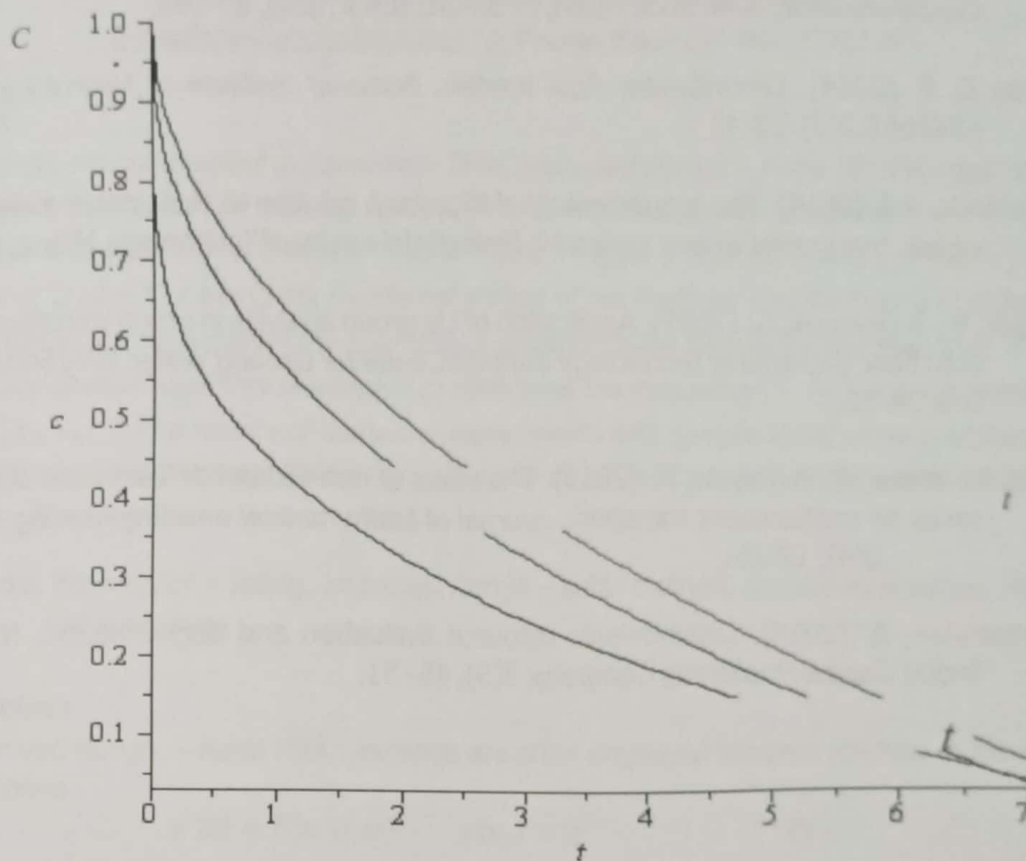


Figure 3.3: Aquifer Concentration Distribution for values of $\alpha, b,$ and γ

Table 3.4: Concentration Distribution for Different Values of $\alpha, b,$ and γ

α	b	γ	t
1.00	0.50	0.50	$0 < t \leq 7$
0.50	0.50	0.50	$0 < t \leq 7$
0.20	0.50	0.50	$0 < t \leq 7$

**Figure 3.4: Aquifer Concentration Distribution for Different Values of $\alpha, b,$ and γ** **Conclusion**

The results indicate that for a uniform source of contamination, $C^U(x)$, for values of $\alpha = 1.00$, $b = 0.5$, and $\gamma = 0.5$ the level of contamination is at its peak as shown in figure 3.1. A reduction of the value α to 0.50, we noticed a reduction in the level of concentration as shown in figure 3.2. A further reduction of the value of α to 0.20, we noticed a further reduction in the level of concentration as shown in figure 3.3. This shows that for a uniform source of contamination, the level of contamination reduces over time depending on the values of α , b and γ .

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