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A MATHEMATICAL ANALYSIS OF THE KINETICS OF VIRAL INACTIVATION



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A MATHEMATICAL ANALYSIS OF THE KINETICS
OF VIRAL INACTIVATION

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FOREWORD

Man and his environment must be protected from the adverse effects of pesticides, radiation, noise and other forms of pollution, and the unwise management of solid waste. Efforts to protect the environment require a focus that recognizes the interplay between the components of our physical environment--air, water, and land. The National Environmental Research Centers provide this multidisciplinary focus through programs engaged in

- studies on the effects of environmental contaminants on man and the biosphere, and
- a search for ways to prevent contamination and to recycle valuable resources.

This report describes a mathematical model which can be used to characterize the response of viruses to a disinfecting agent. Not only is the model itself presented, but a technique is described which can be used to estimate the model's parameters. Both the model and the estimation technique are being used to analyze experimental information resulting from disinfection studies.

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INTRODUCTION

Pathogenic enteric viruses transmitted via the water route present a potential hazard to public health because of their resistance to natural or artificial disinfection mechanisms. More than 100 different strains of enteric viruses, causing such diseases as poliomyelitis, meningitis, jaundice, and gastroenteritis, are excreted in human feces. The six major groups of enteroviruses responsible for these diseases are polioviruses, coxsackieviruses A and B, echoviruses, adenoviruses, infectious hepatitis, and reoviruses. Since these viruses are able to survive in sewage, natural waters, and water supplies, they may pose a health threat, particularly as wastewater reuse becomes more common.^{1, 2}

Of constant concern to public health officials is the ability of viruses to pass through water treatment plants. The chlorine levels must be adequate, not only for bacterial disinfection but for viral inactivation as well. As a result of the need for constant concern over proper disinfection levels, much research effort has been devoted to the study of the basic disinfection mechanisms.

Chick was probably the first investigator to attempt to understand the laws of disinfection by applying the principles of first-order kinetics to bacteria and spore inactivation.³ Only the experiments with anthrax spores conformed to first-order kinetics, whereas bacteria apparently followed another pattern of inactivation. Subsequent studies have obtained results that confirmed first-order kinetic inactivation for bacteria.

Many research investigations have been directed toward the study of the inactivation of viruses and enteric organisms. As a result of these studies, the process of inactivation has been found to be dependent on the time of contact between the organisms and disinfecting agent, concentration of disinfecting agent, temperature, and pH. In addition, viruses may form clumps of varying sizes and may cause aberrations due to their existence in inactivation systems.⁴ One approach to studying the interaction of these various factors is to develop a kinetic model that will systematically account for them. The development of such a model and its application are discussed in this paper.

MODEL DEVELOPMENT

One of the major features in this model is the consideration of clumping or aggregation and its effect in explaining the devitalization process and associated aberrations. For purposes of this model, it is assumed that the virions exist either as individual particles in a suspension or as aggregates or clumps made up of two or more particles.⁵ Each individual particle or aggregate will form a plaque-forming unit (PFU) before the viral suspension is subjected to a disinfecting agent. It is impossible to determine whether a PFU represents a single infective unit. If the suspension contains single particles as well as clumps of various sizes, the disinfection process will continue until the last particle in the largest clump is devitalized. When the clump is completely devitalized, a PFU is destroyed, but it is obvious that a distribution of different size clumps will lead to a non-uniform destruction of PFU's thereby causing some unusual shapes in the disinfection curve.

In this discussion, it will be assumed that this distribution of infective units represents the state of the suspension. The percentage of aggregates or clumps of all sizes which have been disinfected at any time represents the Nth state; the percentage of undisinfected single particles represents the N-1st state, etc. For illustrative purposes, let us assume a suspension in which the maximum clump size is composed of three viral particles and with clumps composed of two particles as well as single particles. Following our convention, state 1 is the percentage of undisinfected aggregates with three virions; state 2, the percentage of undisinfected aggregates with two virions; state 3, the percentage of undisinfected single particles; and state 4, the total percentage of aggregates (clumps of 1, 2, and 3 viral particles) that have been devitalized at any point in time. Obviously, under the action of a disinfectant, assuming ideal conditions, state 4 would increase as the process continues until state 4 would be 100 percent.

We can impose a frequency distribution on the various states in effect, assigning a percentage of the total plaque-forming capability to each state. The initial condition of state 4 (S_4^0) must equal 0 percent at time equal to zero or before the disinfectant acts. The percentage of undisinfected singles plus the percentages of clumps with two particles plus the percentage of clumps with three particles would equal 100 percent when time equals zero.

Associated with each state is a decay rate, k_i , that represents the probability of interaction of the destructive agent with the undisinfected singles or aggregate. The process of devitalization is assumed to take place in the following manner: The

clumps of three virions are reduced to two surviving virions, and the clumps of two are reduced to one surviving virion all the way along the chain of states until the clumps are no longer infective and are registered as a decrease in total PFU.

The set of differential equations that describes the devitalization process, where S_i ($i = 1 \dots 4$), the percent of plaque-forming capability at each state is:

$$\begin{aligned}\frac{dS_1}{dt} &= -k_1 S_1 \\ \frac{dS_2}{dt} &= k_1 S_1 - k_2 S_2 \\ \frac{dS_3}{dt} &= k_2 S_2 - k_3 S_3 \\ \frac{dS_4}{dt} &= k_3 S_3\end{aligned}\tag{1}$$

These are a set of linear first-order differential equations. The parameters k_i ($i = 1 \dots 4$) represents the devitalization rate with $k_4 = 0$, and S_i^0 is the initial condition of state i with $S_4^0 = 0$ at $t = 0$.

The solution to Equation 1 is as follows:

$$\begin{aligned}S_4 &= k_1 k_2 k_3 S_1^0 \left[\frac{e^{-k_1 t}}{(-k_1)(k_3 - k_1)(k_2 - k_1)} + \frac{e^{-k_2 t}}{(-k_2)(k_3 - k_2)(k_1 - k_2)} \right. \\ &\quad \left. + \frac{e^{-k_3 t}}{(-k_3)(k_2 - k_3)(k_1 - k_3)} + \frac{1}{(k_3)(k_2)(k_1)} \right] \\ &\quad + k_2 k_3 S_2^0 \left[\frac{e^{-k_2 t}}{(-k_2)(k_3 - k_2)} + \frac{e^{-k_3 t}}{(-k_3)(k_2 - k_3)} + \frac{1}{(k_3)(k_2)} \right] \\ &\quad - S_3^0 [e^{-k_3 t} - 1].\end{aligned}\tag{2}$$

The general closed form solution to a set of differential equations as illustrated by Equation 1 is given by the following:⁵

$$S_i = \sum_{j=1}^i \sum_{n=1}^i \frac{(k_j \cdot k_{j+1} \cdots k_{i-1}) (e^{-k_n t}) (S_j^0)}{(k_i - k_k) (k_{i-1} - k_k) \cdots (k_j - k_k)} \quad (3)$$

where $k > j$. When $j=k$, $(k_j - k_k) = 1$.

When expressed as percent survival, the equation could be written as percent survival = $100 - S_i$, where S_i is the last or final state to be considered.

Figure 1 illustrates schematically the change taking place during an experiment. Devitalized virus in an aggregate are represented by a broken circle. In a devitalization chain, the value for k_i , which indicates the rate of transition from one state into the next, differs for each state.⁶ There are also differences between chains. For example, k_3 in the first chain may be smaller than k_3 in the second chain. This might be attributed to different geometric configurations and resulting interferences. We will assume, however, that k_3 is an average reaction rate for state 3 in all of the decay chains.

Equation 2 can be reformulated in the following manner:

$$S_4 = C_0 + C_1 e^{-k_1 t} + C_2 e^{-k_2 t} + C_3 e^{-k_3 t} \quad (4)$$

where,

$$\begin{aligned} C_1 &= \frac{-k_2 k_3 S_1^0}{(k_3 - k_1) (k_2 - k_1)} \\ C_2 &= \frac{-k_1 k_3 S_1^0}{(k_3 - k_2) (k_1 - k_2)} - \frac{k_3 S_2^0}{(k_3 - k_2)} \\ C_3 &= \frac{-k_1 k_2 S_1^0}{(k_2 - k_3) (k_1 - k_3)} - \frac{k_2 S_2^0}{(k_2 - k_3)} - S_3^0 \\ C_0 &= S_1^0 + S_2^0 + S_3^0 \end{aligned} \quad (5)$$

We know that as $t \rightarrow \infty$, $S_4 \rightarrow 100$ percent; therefore, $C_0 \rightarrow 100$ percent.

Equation 4 forms the basis for the mathematical model of the kinetics of viral inactivation we wish to examine. However, to use this equation, we must be able to estimate its parameters.

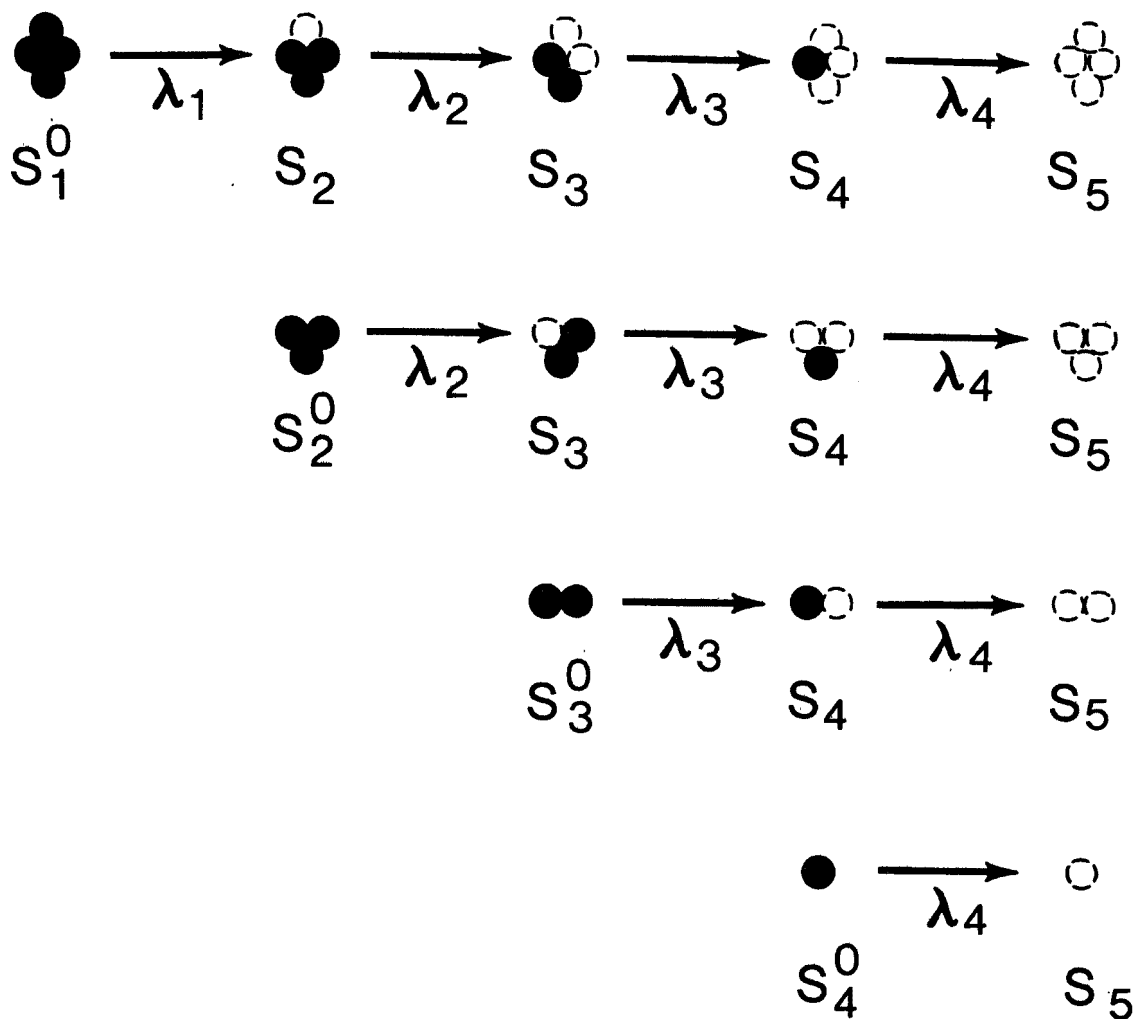


Figure 1. Schematic illustration of the physical change taking place in a suspension of virions under the influence of a devitalizing agent. Devitalized virus in a clump is represented by a broken circle. The values for λ indicate the rates of transition from one state into the next in a devitalization chain. S_i^0 is the initial percent of plaque-forming capability in state i .

ESTIMATION OF PARAMETERS

If we were to rewrite Equation 4 in terms of percent survival, we would have the following:

$$100\% - S_4^0 = -C_1 e^{-k_1 t} - C_2 e^{-k_2 t} - C_3 e^{-k_3 t} \quad (6)$$

or

$$\bar{y} = -C_1 e^{-k_1 t} - C_2 e^{-k_2 t} - C_3 e^{-k_3 t} \quad (7)$$

where $\bar{y} = 100\% - S_4^0$.

For simplicity, we shall assume that our observations are equidistant, as in Figure 2, and that the difference in the successive abscissa values is h . With the use of our three-term example, we find the value of the i th ordinate at $t_0 + (i-1)h$, where t_0 is the value of y_0 at t_0 , is then:⁷

$$\begin{aligned} \bar{y}_i = & -C_1 \exp[-k_1 t_0 + (i-1)h] - C_2 \exp[-k_2 t_0 + (i-1)h] \\ & - C_3 \exp[-k_3 t_0 + (i-1)h] \end{aligned} \quad (8)$$

or, if we make the following substitutions:

$$e^{-k_1 h} = u_1;$$

$$e^{-k_2 h} = u_2;$$

$$e^{-k_3 h} = u_3;$$

$$-C_1 \exp[-k_1 t_0 + (i-1)h] = f_1;$$

$$-C_2 \exp[-k_2 t_0 + (i-1)h] = f_2;$$

$$-C_3 \exp[-k_3 t_0 + (i-1)h] = f_3;$$

then, for five equidistant measurements, we have:

$$f_1 + f_2 + f_3 = y_i$$

$$f_1 u_1 + f_2 u_2 + f_3 u_3 = y_{i+1}$$

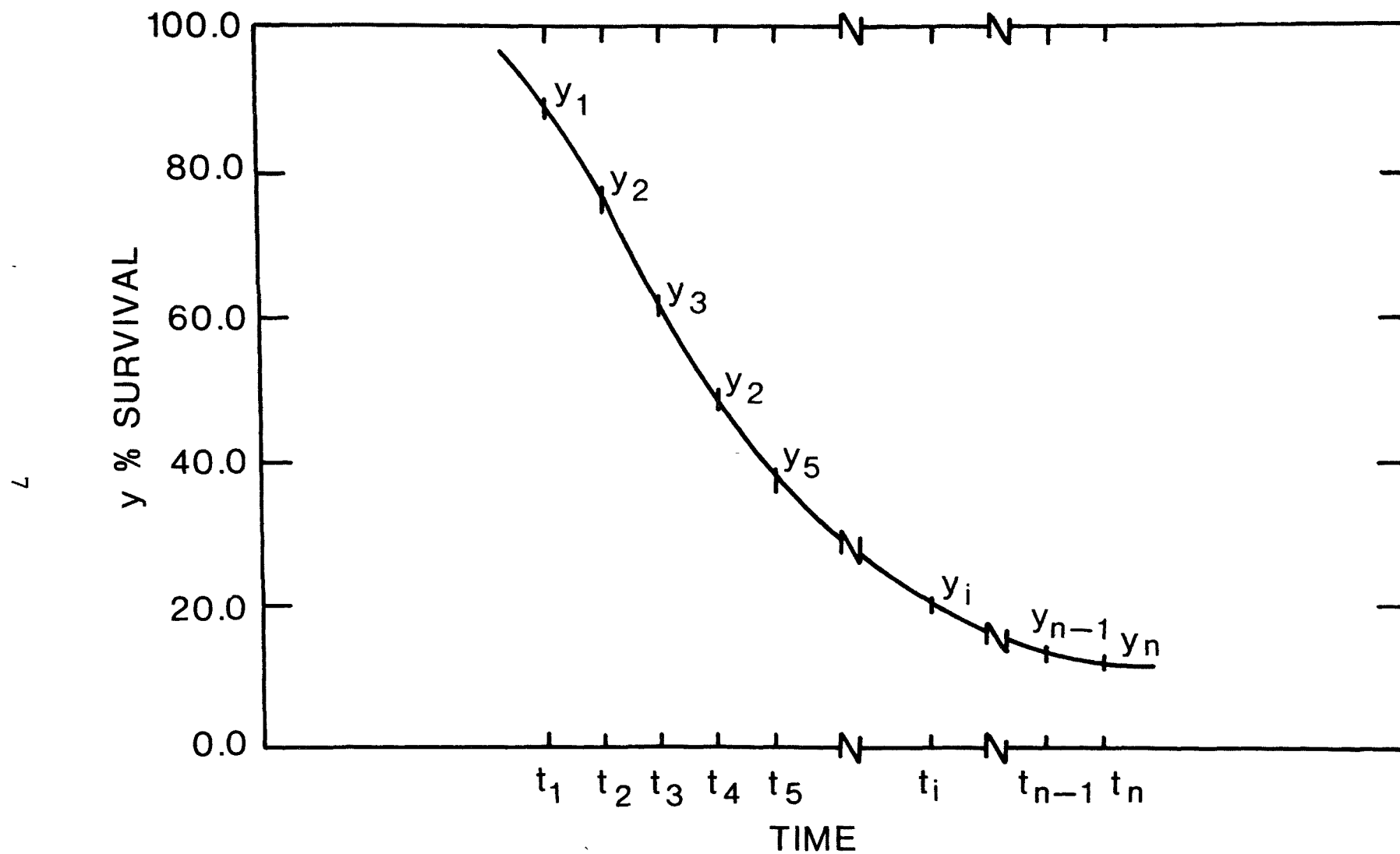


Figure 2. Equally distant values for percent survival versus time.

$$\begin{aligned}
f_1 u_1^2 + f_2 u_2^2 + f_3 u_3^2 &= y_{i+2} \\
f_1 u_1^3 + f_2 u_2^3 + f_3 u_3^3 &= y_{i+3} \\
f_1 u_1^4 + f_2 u_2^4 + f_3 u_3^4 &= y_{i+4}
\end{aligned} \tag{9}$$

In general, the set of equations for N observations would be as shown below:

$$\begin{aligned}
f_1 + f_2 + f_3 &= y_0 \\
f_1 u_1 + f_2 u_2 + f_3 u_3 &= y_1 \\
f_1 u_1^2 + f_2 u_2^2 + f_3 u_3^2 &= y_2 \\
&\dots \dots \dots \\
f_1 u_1^{N-1} + f_2 u_2^{N-1} + f_3 u_3^{N-1} &= y_{N-1}
\end{aligned} \tag{10}$$

which would necessarily be satisfied identically. If the constants u_1 , u_2 , and u_3 were known (or preassigned), Equations 10 would comprise N linear equations in the three unknowns f_1 , f_2 , and f_3 , and be solved exactly if $N=3$ or approximately by least squares if $N>3$.

However, if the u 's are also to be determined, at least six equations are needed, and a difficulty occurs because the equations are non-linear in the u 's. This difficulty can be minimized by the following method.

Let u_1 , u_2 , and u_3 be the roots of the algebraic equation:

$$u^3 - a_1 u^2 - a_2 u - a_3 = 0 \tag{11}$$

so that the left-hand member of Equation 11 is identified with the product $(u-u_1)(u-u_2)(u-u_3)$. To determine the coefficients a_1 , a_2 , a_3 , we multiply the first line in Equation 10 by a_3 , and the second line by a_2 , and the third line by a_1 , and the fourth line by -1 , and add the results. If use is made of the fact that each u satisfies Equation 11, the result is seen to be of the form:

$$y_3 - a_1 y_2 - a_2 y_1 - a_3 y_0 = 0 \tag{12}$$

A set of N-4 additional equations of similar type is obtained in the same way by starting instead successively with the second, third . . . (N-3)th equations. In this way, we find that Equations 10 and 11 imply the N-3 linear equations:⁸

$$\begin{aligned}
 y_2 a_1 + y_1 a_2 + y_0 a_3 &= y_3 \\
 y_3 a_1 + y_2 a_2 + y_1 a_3 &= y_4 \\
 &\dots \\
 y_{N-2} a_1 + y_{N-3} a_2 + \dots y_{N-4} a_3 &= y_{N-1}
 \end{aligned} \tag{13}$$

Since the ordinates y_k are known if $N=6$, this set generally can be solved directly for a_1 , a_2 , and a_3 , or it can be solved approximately by the method of least squares if $N>6$.

In theory, after the a 's are determined, the u 's are found as the roots of Equation 11 and may be real or complex. Equation 10 then becomes linear and the f 's can be determined from the first n of these equations or preferably by applying a least-squares technique applied to the entire set.

We have examined the situations in which there are only three terms to analyze in Equation 2. However, most often the situation will occur when there are n terms in the equation to be solved. This would take the form as follows:

$$S_n = C_0 + C_1 e^{-k_1 t} + C_2 e^{-k_2 t} + \dots C_{n-1} e^{-k_{n-1} t} \tag{14}$$

Assuming that there are N points equally spaced at $t=0, 1, 2, 3 \dots N-1$, and following the logic described in this paper, we get a set of equations similar to Equations 8:

$$\begin{aligned}
 f_1 + f_2 + f_3 + \dots + f_n &= y_0 \\
 f_1 u_1 + f_2 u_2 + f_3 u_3 + \dots + f_n u_n &= y_1 \\
 f_1 u_1^2 + f_2 u_2^2 + f_3 u_3^2 + \dots + f_n u_n^2 &= y_2 \\
 &\dots \\
 f_1 u_1^{N-1} + f_2 u_2^{N-1} + f_3 u_3^{N-1} + \dots + f_n u_n^{N-1} &= y_{N-1}
 \end{aligned} \tag{15}$$

Again, following the logic described earlier, we have the following $N-n$ linear equations where the columns of data are labeled 1 through $n+1$.

(1)	(2)	(3)	(n)	(n+1)	
$y_{n-1}a_1$	$+ y_{n-2}a_2$	$+ y_{n-2}a_3$	$+ \dots$	$+ y_0a_n$	$= y_n$
y_na_1	$+ y_{n-1}a_2$	$+ y_{n-2}a_3$	$+ \dots$	$+ y_1a_n$	$= y_{n+1}$
\dots	\dots	\dots	\dots	\dots	\dots
$y_{N-2}a_1$	$+ y_{N-3}a_2$	$+ y_{N-4}a_3$	$+ \dots$	$+ y_{N-n-1}a_n$	$= y_{N-1} \quad (16)$

After the a's have been determined by least squares, the values for the c's can be found as roots of the following equation:

$$u^n - a_1u^{n-1} - a_2u^{n-2} - \dots - a_{n-2}u - a_n = 0 \quad (17)$$

And once the u's have been found, the f's can be found from Equation 15.

The application of this approach presumes that the number of terms that make up the model is known. Generally this number is unknown, and a major part of the analysis becomes the estimation of the optimum number of terms describing the disinfection process. Even if the number of terms is known, the solution to Equation 17 is often complex because of estimation errors in determining the coefficients. To make this analysis usable, we must be able to determine the number of terms (number of states) that make up the inactivation process. The following section describes a technique for estimating the number of components that "best" describe the inactivation process.

OPTIMAL NUMBER OF TERMS

To determine the proper number of terms that will describe the inactivation process, we would formulate the set of linear equations shown in Equation 16. In this set, the column labeled n+1 is the response or dependent variable, and the columns 1 through n are the independent variables. Using step-wise regression, we regress the independent variables (1 through n) against the n+1st or dependent variable.⁹ As each variable is forced into the equation, a value for its coefficient is calculated. Each coefficient has an associated sign. When the signed coefficient is substituted into Equation 17, it is possible that an equation with alternating signs may result; for example, Equation 17 might look as follows:

$$a_0u^n - a_1u^{n-1} + a_2u^{n-2} - \dots + a_{n-2}u - a_n = 0 \quad (18)$$

According to Des Cartes' rule of signs:

The number of positive real roots of a real algebraic equation either is equal to the number N_a of sign changes in the sequence $a_0, a_1, a_2, \dots, a_n$ of coefficients where vanishing terms are disregarded or it is less than N_a by a positive even integer.

Since the decay coefficients in Equation 14 are the positive real roots in Equation 18, we can use Des Cartes' rule to give us an indication as to the number of terms which optimally describes the inactivation process. We will assume that when the number of terms in the regression equation is one more than the number of sign changes, the optimal number of terms has been identified, and the variables in the regression equation are to be used in calculating k_n . The approach will be discussed beginning with the identification of the optimal number of terms.

We can illustrate this approach by assuming a model of three terms as follows:

$$y = 20.00e^{-0.10t} + 30.00e^{-0.30t} + 50.00e^{-0.50t} \quad (19)$$

Table 1 (Page 12) contains values for Equation 19 which have been generated at intervals of $t=0.50$ to simulate a disinfection curve. Table 2 illustrates the way in which these data are organized to solve for the coefficients in Equation 17. As shown in Equation 16, a matrix of data points is established with n dependent variables. In this case, 27 independent variables have been constructed. The value of $y_1 = 100.00$ is the first value in the upper left-hand corner of the matrix, and the value $y_{27} = 5.7660$ is the first value for the dependent variable. The second value for the first independent value is $y_1 = 83.7858$, and the second value for $y_{28} = 5.4274$. This same pattern is repeated throughout the matrix.

Table 2. MATRIX OF DATA FOR REGRESSION ANALYSIS

Var 1	Var 2	. . .	Var n	. . .	Var 28
100.00	83.786				5.7660
83.786	70.648				5.4274
.
2.0359	1.9338				0.5203
1.9338	1.8370				0.4949

Table 1. VALUES FOR EQUATION 19 AT INTERVALS OF $t=0.5$

t	$e^{-0.100t}$	$e^{-0.300t}$	$e^{-0.500t}$	y	t	$e^{-0.100t}$	$e^{-0.300t}$	$e^{-0.500t}$	y	t	$e^{-0.100t}$	$e^{-0.300t}$	$e^{-0.500t}$	y
0.00	20.0000	30.0000	50.0000	100.0000	13.00	5.4506	0.6072	0.0751	6.1330	26.00	1.4854	0.0122	0.0000	1.4978
0.50	19.0245	25.8212	38.9400	83.7858	13.50	5.1848	0.5226	0.0585	5.7660	26.50	1.4130	0.0105	0.0000	1.4236
1.00	18.0967	22.2245	30.3265	70.6478	14.00	4.9319	0.4498	0.0455	5.4274	27.00	1.3441	0.0091	0.0000	1.3532
1.50	17.2141	19.1288	23.6183	59.9613	14.50	4.6914	0.3872	0.0355	5.1141	27.50	1.3785	0.0078	0.0000	1.2864
2.00	16.3746	16.4643	18.3939	51.2329	15.00	4.4626	0.3332	0.0276	4.8235	28.00	1.2162	0.0067	0.0000	1.2229
2.50	15.5760	14.1710	14.3252	44.0722	15.50	4.2449	0.2868	0.0215	4.5533	28.50	1.1568	0.0058	0.0000	1.1627
3.00	14.8163	12.1970	11.1565	38.1699	16.00	4.0379	0.2468	0.0167	4.3016	29.00	1.1004	0.0049	0.0000	1.1054
3.50	14.0937	10.4981	8.6887	33.2806	16.50	3.8410	0.2125	0.0130	4.0665	29.50	1.0467	0.0043	0.0000	1.0511
4.00	13.4064	9.0358	6.7667	29.2090	17.00	3.6536	0.1829	0.0101	3.8467	30.00	0.9957	0.0037	0.0000	0.9994
4.50	12.7525	7.7772	5.2699	25.7997	17.50	3.4754	0.1574	0.0079	3.6408	30.50	0.9471	0.0031	0.0000	0.9503
5.00	12.1306	6.6939	4.1042	22.9287	18.00	3.3059	0.1354	0.0061	3.4476	31.00	0.9009	0.0027	0.0000	0.9037
5.50	11.5390	5.7615	3.1963	20.4969	18.50	3.1447	0.1166	0.0048	3.2661	31.50	0.8570	0.0023	0.0000	0.8594
6.00	10.9762	4.9589	2.4893	18.4245	19.00	2.9913	0.1003	0.0037	3.0954	32.00	0.8152	0.0020	0.0000	0.8172
6.50	10.4409	4.2682	1.9387	16.6478	19.50	2.8454	0.0863	0.0029	2.9347	32.50	0.7754	0.0017	0.0000	0.7772
7.00	9.9317	3.6736	1.5098	15.1152	20.00	2.7067	0.0743	0.0022	2.7833	33.00	0.7376	0.0015	0.0000	0.7391
7.50	9.4473	3.1619	1.1758	13.7852	20.50	2.5747	0.0640	0.0017	2.6404	33.50	0.7016	0.0012	0.0000	0.7029
8.00	8.9865	2.7215	0.9157	12.6239	21.00	2.4491	0.0550	0.0013	2.5055	34.00	0.6674	0.0011	0.0000	0.6585
8.50	8.5483	2.3424	0.7132	11.6039	21.50	2.3296	0.0474	0.0010	2.3781	34.50	0.6349	0.0009	0.0000	0.6358
9.00	8.1313	2.0161	0.5554	10.7030	22.00	2.2160	0.0408	0.0008	2.2577	35.00	0.6039	0.0008	0.0000	0.6047
9.50	7.7348	1.7353	0.4325	9.9027	22.50	2.1079	0.0351	0.0006	2.1437	35.50	0.5744	0.0007	0.0000	0.5752
10.00	7.3575	1.4936	0.3368	9.1881	23.00	2.0051	0.0202	0.0005	2.0359	36.00	0.5464	0.0006	0.0000	0.5470
10.50	6.9987	1.2855	0.2623	8.5467	23.50	1.9073	0.0260	0.0003	1.9338	36.50	0.5198	0.0005	0.0000	0.5203
11.00	6.6574	1.1064	0.2043	7.9682	24.00	1.8143	0.0223	0.0003	1.8370	37.00	0.4944	0.0004	0.0000	0.4949
11.50	6.3327	0.9523	0.1591	7.4443	24.50	1.7258	0.0192	0.0002	1.7453					
12.00	6.0238	0.8197	0.1239	6.9675	25.00	1.6417	0.0165	0.0001	1.6584					
12.50	5.7301	0.7055	0.0965	6.5321	25.50	1.5616	0.0142	0.0001	1.5760					

Table 3 contains the results of the application of the step-wise regression program to the matrix of data in Table 2. The equations resulting from each step are as follows:

$$x^{28} - 0.9418x^{27} = 0 \quad (20)$$

$$x^{28} - 1.1223x^{27} + 0.1402x^{23} = 0 \quad (21)$$

$$x^{28} - 1.1420x^{27} + 0.1565x^{23} - 0.000089x^1 = 0 \quad (22)$$

$$x^{28} + 0.0318x^{27} - 0.9398x^{23} + 0.07968x^8 - 0.01124x^1 = 0 \quad (23)$$

Equation 22 combines the maximum number of sign changes with the minimum number of variables in the equation and is, therefore, selected as the equation governing the number of terms in the disinfection equation. This matches identically with the three terms used in the simulated data. After the best estimate has been made of the number of terms which makes up the data, the next step in the analysis is to estimate the decay coefficients in the equation. This step is described in the following section.

Table 3. RESULTS OF REGRESSION ANALYSIS
USING DATA FROM TABLE 2

Step	Var	Coefficient
1	27	0.94179753
2	23	-0.14022224
	27	1.12226027
3	1	0.00008941
	23	-0.15649791
	27	1.14201651
4	1	0.01124259
	8	-0.07967716
	23	0.93985016
	27	-0.03183525

ESTIMATION OF PARAMETERS

Decay Rates

Based on the data in Table 1, the parameters for Equation 19 can be estimated using the techniques outlined in Appendix A. The first decay coefficient to be calculated will be that associated with variable x^{23} and the calculation is as follows:

$$\frac{[(x^{24} - r_1 x^{23}) - (x^2 - r_1 x^1)]}{(23 - 1)} =$$

$$- \frac{[(x^{24} - r_1 x^{23}) - (x^{28} - r_1 x^{27})]}{(23 - 27)} \quad (24)$$

Substituting the average values for x^{28} , x^{27} , x^{24} , x^{23} , x^2 , and x^1 , into Equation 24 yields the following values for r :

$$r_1 = 0.85 \quad (25)$$

where

$$r_1 = e^{-k_1 h} \quad (26)$$

From Equation 26, we can calculate k_1 as follows:

$$k_1 = -\ln(r_1)/h \quad (27)$$

$$k_1 = -[\ln(0.85)]/0.50$$

$$k_1 = 0.32$$

The decay coefficient associated with variable x^1 is calculated as follows:

$$\ln(r_2) = \ln(x^2) - \ln(2x^1 - x^2) \quad (28)$$

$$\ln(r_1) = -0.24$$

Substituting into Equation 27 for k_2 we get the following:

$$k_2 = (0.24)/0.50 \quad (29)$$

$$k_2 = 0.48$$

The decay coefficient associated with variable x^{27} is calculated as follows:

$$\ln(r_3) = \ln(x^{28}/x^{27}) \quad (30)$$

$$\ln(r_3) = \ln(0.95)$$

Substituting into Equation 27 for k_3 we get the following:

$$k_3 = (0.05)/0.50$$

$$k_3 = 0.10$$

Coefficients

Once the decay rates in Equation 19 have been estimated, the values for the coefficients are relatively easy to obtain. Values for each exponential term can be calculated at the appropriate time interval and these values regressed against the values of y in Table 1. Stepwise regression can then be used to estimate the coefficients (Appendix D).

EXAMPLE INACTIVATION PROBLEM

To illustrate the utilization of this technique, it will be applied to experimental data collected from a series of electromicroscopy investigations conducted by Gordon Sharp at the University of North Carolina.¹⁰ Sharp prepared electron micrographs of dilute preparations of T7 virus that had been subjected to a devitalizing agent.

Figure 3 shows the inactivation curve, and Table 4 contains the distribution of T7 coliphage particles resulting from these experiments. Column 1 of Table 4 lists the group size of the aggregates, that is, the number of particles in each clump of virus. Column 2 lists the number of groups in the suspension, and Column 3 lists the number of particles in each group. Column 4 lists the percent of plaque-forming capability that each group represents in the suspension. For example, there are 770 groups in the suspension, but 610/770 or 79.1 percent of them are groups of single viral particles, and 116/770 or approximately 15.1 percent of them are groups of two viral particles, etc.

Table 4. T7 VIRUS DATA

Group size	Number of groups	Number of particles	Plaque-forming capability (%)
1	610	610	79.22
2	116	232	15.06
3	24	72	3.12
4	12	48	1.56
5	6	30	0.78
6	1	6	0.13
18	1	18	0.13
Total	770	1,016	100.00

*INACTIVATION OF COLIPHAGE T₇
BY ULTRAVIOLET RAYS*

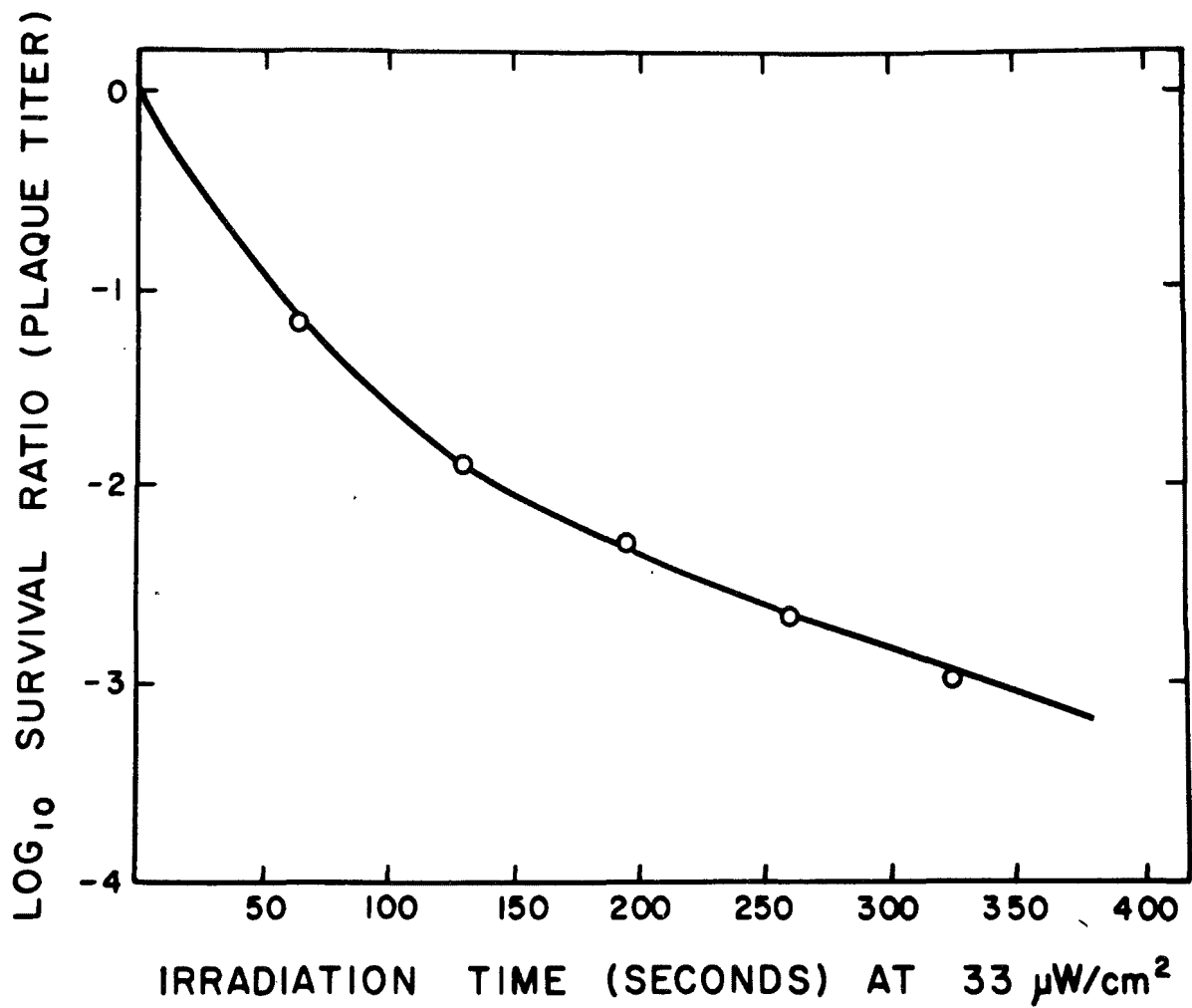


Figure 3. Inactivation of coliphage

Table 5 contains the data from Figure 3, at intervals of 5 seconds, arranged in 10 columns of data. Table 6 contains the coefficients associated with each set of variables as they enter the stepwise regression equation. It is obvious from the alternating signs that six variables will describe the inactivation process. The decay coefficients per minute calculated from the techniques outlined in Appendix A are as follows:

$$r_1 = 1.56 \quad (32)$$

$$r_2 = 2.12$$

$$r_3 = 2.32$$

$$r_4 = 2.68$$

$$r_5 = 2.83$$

$$r_6 = 4.68$$

Each of these values represents a k_i in Equation 14, and each value of $e^{-k_i t}$ can be generated at various intervals of t by the program in Appendix B. If all six values of $e^{-k_i t}$ represented by Equation 32 are regressed against the values for y as obtained from the graph in Figure 3 then Table 7 contains the values for their coefficients. Using the program in Appendix C, the values for each predicted S_i (percentage of plaque-forming capability) can be calculated. The predicted and actual values are shown in Table 8.

When the regression is performed, the values shown in Table 7 result. At the fourth step of the regression, the corrected R^2 begins to decrease which is an indicator that the regression should be terminated at that point, and step 3 is, therefore, used as the last step in the regression analysis. Equation 5 and the program contained in Appendix C, where $k_1 = 1.56$, $k_2 = 2.83$, and $k_3 = 4.68$, yields the following values for S_i : $S_3 = 73.97\%$, $S_2 = 15.51\%$, and $S_1 = 11.30\%$. Physically, this means that there are 73.97% singles, 15.51% doubles, and the rest of the particles amount to approximately 11.30%. The comparison between the results obtained from the model and the electron micrographs is shown in Table 8. The agreement seems reasonable.

Table 5. DATA FROM DISINFECTION CURVE

Var 1	Var 2	Var 3	Var 4	Var 5	Var 6	Var 7	Var 8	Var 9	Var 10
100.00000	79.00000	58.00000	45.00000	36.50000	28.50000	25.00000	19.80000	17.50000	15.00000
79.00000	58.00000	45.00000	36.00000	28.50000	25.00000	19.80000	17.50000	15.00000	13.00000
.
.
.
.
.
.
.
000.16500	00.16000	00.15500	00.15000	00.14000	00.13500	00.13000	00.12000	00.11000	00.10500
000.16000	00.15500	00.15000	00.14000	00.13500	00.13000	00.12000	00.11000	00.10500	00.10000

Table 6. RESULTS OF REGRESSION ANALYSIS
USING DATA FROM TABLE 5

Step	Var	Coefficient
1	9	0.84449432
2	6	0.16137199
	9	0.57866191
3	6	0.22635578
	8	-0.22446799
	9	0.73339338
4	5	-0.14587300
	6	0.36096394
	8	-0.35799450
	9	0.95901767
5	4	0.12430537
	5	-0.26913616
	6	0.29413744
	8	-0.36433149
	9	1.01961917
6	1	-0.03634866
	4	0.29008852
	5	-0.30158431
	6	0.22952950
	8	-0.50623862
	9	1.13438561
7	1	-0.03743081
	4	0.32109643
	5	-0.43943162
	6	0.20853501
	7	0.25010067
	8	-0.67970749
	9	1.22741919

Table 7. RESULTS OF REGRESSION
USING DR. SHARP'S INACTIVATION DATA

Step	Var	Coefficient	Corrected R ² as a percent	t value
1	2	96.77	99.501	117.26
2	2	89.08	99.549	32.05
	6	6.64		2.88
3	1	58.89	99.933	19.67
	2	4.12		0.92
	6	37.77		20.82
4	1	61.68	99.932	6.69
	2	-10.82		-0.23
	4	15.24		0.32
	6	34.70		3.55

Table 8. COMPARISON BETWEEN MODEL OUTPUT
AND ELECTRON MICROGRAPHS

Group size	Percent plaque-forming capability (counted)	Percent plaque-forming capability (predicted)
1	79.22	73.97
2	15.06	15.51
3 + 4 + 5 + 6 + 18	5.82	11.30

SUMMARY AND CONCLUSIONS

The kinetics of viral inactivation have been examined from a rational point of view. A mathematical model based on the radionuclide chain decay concept was formulated and a solution technique developed that allows for estimations of the optimal number of terms in the equation and for estimating the equation's parameters. With the use of data derived from electron microscopy, the model was tested and achieved reasonable results.

Based on this discussion, it is obvious that the postulated mathematical model and its solution techniques are superior to others that have been formulated. The approach outlined in this report not only determines the number of aggregate groups in the suspension, but the values for decay coefficients as well. There are some deficiencies in this approach, however, and it is important that these be considered. The approach suggested here is statistical in nature and is, therefore, subject to experimental error in the various estimations made. More importantly, the estimates of aggregate size and concentration are blind. That is, if this approach estimates three terms as optimal, there is no way to provide information on the make-up of these aggregate groupings. They might be clumps of single, double, and triple particles, or clumps of 20, 21, and 25 particles. The values for the decay coefficients may give some insight as to clump size, but these insights are hardly sufficient. This technique must be coupled with a physical assay approach incorporating electron microscopy. A research project that combines the elements of mathematical analysis with electron microscopy is currently underway.

APPENDIX A

In this appendix, the mathematical justification for the techniques used in the section entitled "Estimation of Parameters" is developed. Table 9 contains the first three values for the individual terms which make up the variables 1, 2, 23, 24, 27, and 28, as shown in Table 2. The first variable to enter the stepwise regression equation is x^{27} , as shown in Equation 20. Looking at variables 27 and 28 in Table 9, it is obvious that the term labeled f_3 dominates variable 27 and is most highly correlated with variable 28, while the terms f_1 and f_2 in variable 27 are relatively insignificant. The next variable to enter the stepwise regression equation is variable 23, and it can be seen that terms f_2 and f_3 in variable 23 are significant but that term f_1 is insignificant, and finally in variable 1, terms f_1 , f_2 , and f_3 , are all significant. It can be concluded from this that a variable enters the regression equation when one of the terms which comprise it is significant enough to alter the rate of change of the entering variable. Therefore, we would expect that variables would enter the regression equation with alternating signs associated with their coefficients, since the entrance of each variable into the equation signifies a significant change in the functions slope. Moreover, we would expect that the variables entering the equation with alternating signs represents the maximum change in the slope of the function with respect to the other variables in the regression equation. Using Equation 19 as an example, we would, therefore, attempt to find a u_i such that b_n in the following equation is a maximum relative to its adjacent variables:

$$f_1(u_1^{n+1} - u_i u_1^n) + f_2(u_2^{n+1} - u_i u_2^n) + f_3(u_3^{n+1} - u_i u_3^n) = b_n \quad (33)$$

or in a more simple form, we would attempt to find a u_i such that b_{23} is a maximum relative to b_1 and b_{27} in the following set of equations:

$$\begin{aligned} x^2 - u_i x^1 &= b_1 \\ x^{24} - u_i x^{23} &= b_{23} \\ x^{28} - u_i x^{27} &= b_{27} \end{aligned} \quad (34)$$

Table 9. FIRST THREE VALUES FOR SELECTED VARIABLES

Var	Time	$c_1 e^{-0.500t_*}$	$c_2 e^{-0.300t_+}$	$c_3 e^{-0.100t_+}$	y
1	0	50.0000	30.0000	20.0000	100.0000
	0.5	38.9400	25.8212	19.0245	83.7858
	1.0	30.3265	22.2245	18.0967	70.6478
2	0.5	38.9400	25.8212	19.0245	83.7858
	1.0	30.3265	22.2245	18.0967	70.6478
	1.5	23.6183	19.1288	17.2141	59.9613
23	11.0	0.2043	1.1064	6.6574	7.9682
	11.5	0.1591	0.9523	6.3327	7.4443
	12.0	0.1239	0.8197	6.0238	6.9675
24	11.5	0.1591	0.9523	6.3327	7.4443
	12.0	0.1239	0.8197	6.0238	6.9675
	12.5	0.0965	0.7055	5.7301	6.5321
27	13.0	0.0751	0.6072	5.4506	6.1330
	13.5	0.0585	0.5226	5.1848	5.7660
	14.0	0.0455	0.4498	4.9139	5.4274
28	13.5	0.0585	0.5226	5.1848	5.7660
	14.0	0.0455	0.4498	4.9139	5.4274
	14.5	0.0355	0.3872	4.6914	5.1141

*f₁.†f₂.‡f₃.

Therefore, u_i can be calculated from the following equation:

$$\frac{(x^2 - u_i x^1) - (x^{24} - u_i x^{23})}{(1 - 23)} = \frac{(x^{24} - u_i x^{23}) - (x^{28} - u_i x^{22})}{(23 - 27)} \quad (35)$$

since we know that b_n is a maximum round the point x^{23} .
Table 10 confirms that this is in fact the case for variable 23.

Table 10. b_n FOR SELECTED VALUES OF u_i

$x^2 - u_2 x^1$	$x^{24} - u_2 x^{23}$	$x^{28} - u_2 x^{27}$
-2.2800	0.5839	0.4857
-1.4613	0.5583	0.4449
-0.7931	0.5000	0.4569

For variables x^1 and x^{27} , this computation is impossible since there are no variables which can be used to make a computation similar to Equation 2.

However, for the decay coefficient associated with x^1 , the following equation can be developed using the properties of infinite series:

$$u_i^n (x^2 - u_i x^1) \rightarrow u_i^{n+1} (x^1 - x^2) \text{ as } n \rightarrow \infty \quad (36)$$

Therefore,

$$\ln(u_i) = \ln(x^2) - \ln(2x^1 - x^2) \quad (37)$$

For the decay coefficient associated with x^{27} , the following relationship can be developed:

$$u_i^n x^{28} - u_i^{n+1} x^{27} \rightarrow 0 \text{ as } n \rightarrow \infty \quad (38)$$

Therefore,

$$\ln(u_i) = \ln(x^{28}) - \ln(x^{27}) \quad (39)$$

APPENDIX B

PROGRAM FOR GENERATING VALUES OF $e^{-k_i t}$

AT GIVEN INTERVALS OF t

```

        DIMENSION A(15),B(15),C(15)
        MR = 1
        MW = 15
    5  READ(MR,10)T,N
    10 FORMAT(F5.2,I2)
        IF (N)70,70,15
    15 READ(MR,20)(A(J),J=1,N)
        READ(MR,20)(C(J),J=1,N)
    20 FORMAT(10F8.0)
        E=2.71828
        S=0.0
        Y=0.0
        WRITE(MW,25)
    25 FORMAT('1'//26X,'-XT')
        WRITE(MW,30)
    30 FORMAT(25X,'E      TABLE')
        WRITE(MW,35)
    35 FORMAT(25X,'-----')
        WRITE(MW,40)(A(J),J=1,N)
    40 FORMAT(/10X, 3(3X,'-',F5.3,'T'))
        WRITE(MW,45)
    45 FORMAT(5X,'T',6X,'E', 2(9X,'E'),12X,'Y')
    50 DO 60 I=1,N
    55 B(I)=C(I)*E**(-A(I)*S)
    60 Y=Y+B(I)
        WRITE(MW,65) S,(B(K),K=1,N),Y
    65 FORMAT(/3X,F5.2,2X,11F10.4)
        S=S+T
        Y=0.0
        IF (B(1)-.0001)      5,50,50
    70 STOP
        END

```

APPENDIX C

PROGRAM FOR CALCULATION S_i

DIMENSION A(16),B(16),S(10),TEMP(10),STOR(10)	001
2 READ (1,5)N	002
5 FORMAT (I2)	003
IF(N)8,42,8	
8 READ (1,10)A,8	004
10 FORMAT (8F10.3/8F10.3)	005
DC 41 I=1,N	006
TEMP(1)=1.0	006A
STOR(1)=1.0	006B
DO 30 J=1,N	007
IF(I-J)15,30,15	008
15 TEMP(1)=TEMP(1)*(A(J)-A(I))	009
STOR(1)=STOR(1)*A(J)	010
30 CONTINUE	011
U1=TEMP(1)/STOR(1)	012
IF(I-1)45,35,45	013
35 S(1)=B(1)*U1	014
WRITE (3,40) S(1)	015
40 FORMAT ('1'////' S(1) = ',F12.2///)	
GO TO 41	017
45 TEMP(2)=1.0	017A
STOR(2)=1.0	017B
DO 65 J=1,N	018
IF(I-J)50,65,50	019
50 STOR(2)=STOR(2)*A(J)	020
TEMP(2)=TEMP(2)*(A(J)-A(I))	021
65 CONTINUE	022
U2=STOR(2)*S(1)/TEMP(2)	023
IF(I-2)80,70,80	024
70 S(2)=(B(2)-U2)*U1	025
WRITE (3,75) S(2)	026
75 FORMAT (' S(2) = ',F12.2///)	027
GO TO 41	028
80 TEMP(3)=1.0	028A
STOR(3)=1.00	028B
DO 100 J=2,N	029
IF(I-J)85,100,85	030
85 STOR(3)=STOR(3)*A(J)	031
TEMP(3)=TEMP(3)*(A(J)-A(I))	032
100 CONTINUE	033
U3=STOR(3)*S(2)/TEMP(3)	034
IF(I-3)115,105,115	035
105 S(3)=(B(3)-U2-U3)*U1	036
WRITE (3,110) S(3)	037
110 FORMAT (' S(3) = ',F12.2///)	038
GO TO 41	039
115 TEMP(4)=1.0	039A
STOR(4)=1.0	039B
DO 135 J=3,N	040
IF(I-J)120,135,120	041
120 STOR(4)=STOR(4)*A(J)	042
TEMP(4)=TEMP(4)*(A(J)-A(I))	043
135 CONTINUE	044
U4=STOR(4)*S(3)/TEMP(4)	045
IF(I-4)150,140,150	046
140 S(4)=(B(4)-U2-U3-U4)*U1	047
WRITE (3,145) S(4)	048
145 FORMAT (' S(4) = ',F12.2///)	049
GO TO 41	050
150 TEMP(5)=1.0	050A
STOR(5)=1.0	050B

DO 170 J=4,N	051
IF(I-J)155,170,155	052
155 STOR(5)=STOR(5)*A(J)	053
TEMP(5)=TEMP(5)*(A(J)-A(I))	054
170 CONTINUE	055
U5=STOR(5)*S(4)/TEMP(5)	056
IF(I-5)185,175,185	057
175 S(5)=(B(5)-U2-U3-U4-U5)*U1	058
WRITE (3,180) S(5)	059
180 FCRMAT (' S(5) = ',F12.2///)	060
GO TO 41	061
185 TEMP(6)=1.0	061A
STOR(6)=1.0	061B
DO 205 J=5,N	062
IF(I-J)190,205,190	063
190 STOR(6)=STOR(6)*A(J)	064
TEMP(6)=TEMP(6)*(A(J)-A(I))	065
205 CONTINUE	066
U6=STOR(6)*S(5)/TEMP(6)	067
IF(I-6)220,210,220	068
210 S(6)=(B(6)-U2-U3-U4-U5-U6)*U1	069
WRITE (3,215) S(6)	070
215 FCRMAT (' S(6) = ',F12.2///)	071
GO TO 41	072
220 TEMP(7)=1.0	072A
STOR(7)=1.0	072B
DO 240 J=6,N	073
IF(I-J)225,240,225	074
225 STOR(7)=STOR(7)*A(J)	075
TEMP(7)=TEMP(7)*(A(J)-A(I))	076
240 CONTINUE	077
U7=STOR(7)*S(6)/TEMP(7)	078
IF(I-7)255,245,255	079
245 S(7)=(B(7)-U2-U3-U4-U5-U6-U7)*U1	080
WRITE (3,250) S(7)	081
250 FORMAT (' S(7) = ',F12.2///)	082
GO TO 41	083
255 TEMP(8)=1.0	083A
STOR(8)=1.0	083B
DO 275 J=7,N	084
IF(I-J)260,275,260	085
260 STOR(8)=STOR(8)*A(J)	086
TEMP(8)=TEMP(8)*(A(J)-A(I))	087
275 CONTINUE	088
U8=STOR(8)*S(7)/TEMP(8)	089
IF(I-8)290,280,290	090
280 S(8)=(B(8)-U2-U3-U4-U5-U6-U7-U8)*U1	091
WRITE (3,285) S(8)	092
285 FORMAT (' S(8) = ',F12.2///)	093
GO TO 41	094
290 TEMP(9)=1.0	094A
STOR(9)=1.0	094B
DO 300 J=8,N	095
IF(I-J)295,300,295	096
295 STOR(9)=STOR(9)*A(J)	097
TEMP(9)=TEMP(9)*(A(J)-A(I))	098
300 CONTINUE	099
U9=STOR(9)*S(8)/TEMP(9)	100
IF(I-9)315,305,315	101
305 S(9)=(B(9)-U2-U3-U4-U5-U6-U7-U8-U9)*U1	102
WRITE (3,310) S(9)	103

310	FORMAT (' S(9) = ',F12.2///)	104
	GC TO 41	105
315	TEMP(10)=1.0	105A
	STOR(10)=1.0	105B
	DO 325 J=9,N	106
	IF(I-J)320,325,320	107
320	STOR(10)=STOR(10)*A(J)	108
	TEMP(10)=TEMP(10)*(A(J)-A(I))	109
325	CONTINUE	110
	U10=STOR(10)*S(9)/TEMP(10)	111
	IF(I-10)41,330,41	112
330	S(10)=(B(10)-U2-U3-U4-U5-U6-U7-U8-U9-U10)*U1	113
	WRITE (3,335) S(10)	114
335	FORMAT (' S(10) = ',F12.2///)	115
41	CONTINUE	116
	GO TO 2	116A
42	STOP	117
	END	118

APPENDIX D

STEPWISE REGRESSION PROGRAM

```

C      1130 STEPWISE MULTIPLE REGRESSION PROGRAM, 3/14/66
C      PHASES 1 AND 2 CAN BE OVERLAID TO CONSERVE CORE. THE STEPS TO
C      READY PHASES 1 AND 2 FOR OVERLAY ARE
C          1. SET UP A COMMON AREA CONSISTING OF RIJ,XBAR,SIGMA,FIN,
C          FOUT,DBS,NVAR,NOBS,NINDV,IRES,IFA.
C          2. SET SIGMA AND DATA EQUIVALENT IN PHASE 2.
C          3. REPEAT PHASE 1 DEFINE FILE STATEMENT IN PHASE 2.
C          4. REMOVE STATEMENT 101-3 FROM PHASE 1 AND INSERT IT
C          BEHIND DIMENSION COMMENTS CARD IN PHASE 2.
C      PHASE 1. TRANSFORM ORIGINAL DATA, COMPUTE AND PRINT MEANS,
C      STANDARD DEVIATIONS, AND SIMPLE CORRELATION COEFFICIENTS.
C      DIMENSIONS
C      IMPLICIT REAL*8(A-H,C-Z)
C      DIMENSION DATA(30),CONST(12),ITRAN(30),JTRAN(30),KTRAN(30),LTRAN(3
010)
C      DIMENSION RIJ(30,30),XBAR(30), SIGMA(30),AID(18)
C      DIMENSION SIGB(30),B(30),ID(30)
C      EQUIVALENCES
C      EQUIVALENCE (SIGMA(1),DATA(1))
C      DEFINE DATA FILE
C      DEFINE FILE 1011000,60,U,IFA)
C      STATEMENT LABEL 101 IS NOT REFERENCED. IT MARKS THE FIRST
C      EXECUTABLE STATEMENT OF THE SOURCE PROGRAM.
C      ICOM IS FIXED DECIMAL REPRESENTATION OF ALPHABETIC COMMA.
101 ICOM=27456
C      INITIALIZE DATA FILE
C      IFA=1
C      READ I.D.
C      READ(1,1, END=999) (AID(I),I=1,18)
C      1 FORMAT(18A4)
C      READ CONTROL CARD
C      READ(1,2)NVIN,NVAR,NOBS,NTRAN,NCONS,FIN,FOUT,IRES
C      2 FORMAT(2I2,14,2I2,2F6.3,11)
C      IF(FIN-FOUT)1020,690,690
690 IF(NTRAN)1000,730,700
C      READ TRANSFORMATION CARDS
700 READ(1,71)((ITRAN(I),JTRAN(I),KTRAN(I),LTRAN(I),I=1,NTRAN)
71 FORMAT(36I2)
C      IF(NCONS)1000,730,720
720 READ(1,72)(CONST(I),I=1,NCONS)
72 FORMAT(12F6.3)
C      INITIALIZE.
730 OBS=NOBS
C      NINDV=NVAR-1
C      DO 90 I=1,NVIN
C      XBAR(I)=0.0
C      DO 90 J=1,NVIN
90 RIJ(I,J)=0.0
C      READ DATA, FORM SUMS VECTOR, SUMS OF SQUARES MATRIX
C      DO 110 I=1,NOBS
C      READ(1,3)(DATA(J),J=1,NVIN)
C      3 FORMAT(12F6.0)
C      IF(NTRAN)1000,860,750
C      TRANSFORMATION OF RAW DATA
750 DO 850 M=1,NTRAN
C      II=ITRAN(M)
C      JJ=JTRAN(M)
C      KK=KTRAN(M)
C      LL=LTRAN(M)
C      GO TO (760,770,780,790,800,810,820,830,840),II

```

C	X(J)=X(K)	0610
760	DATA(JJ)=DATA(KK)	0620
	GO TO 850	0630
C	X(J)=-X(K)	0640
770	DATA(JJ)=-DATA(KK)	0650
	GO TO 850	0660
C	X(J)=LOG X(K)	0670
780	DATA(JJ)=DLOG(DATA(KK))	0680
	GO TO 850	0690
C	X(J)=1/X(K)	0700
790	DATA(JJ)=1.0/DATA(KK)	0710
	GO TO 850	0720
C	X(J)=X(K)+X(L)	0730
800	DATA(JJ)=DATA(KK)+DATA(LL)	0740
	GO TO 850	0750
C	X(J)=X(K)*X(L)	0760
810	DATA(JJ)=DATA(KK)*DATA(LL)	0770
	GO TO 850	0780
C	X(J)=X(K)/X(L)	0790
820	DATA(JJ)=DATA(KK)/DATA(LL)	0800
	GO TO 850	0810
C	X(J)=X(K)+C(L)	0820
830	DATA(JJ)=DATA(KK)+CCNST(LL)	0830
	GO TO 850	0840
C	X(J)=X(K)*C(L)	0850
840	DATA(JJ)=DATA(KK)*CCNST(LL)	0860
850	CONTINUE	0870
860	IF(IRES)870,880,870	0880
C	WRITE DATA FILE	0890
870	WRITE(10,'IFA')(DATA(J),J=1,NVAR)	
880	DO 100 J=1,NVAR	0910
	XBAR(J)=XBAR(J)+DATA(J)	0920
	DO 100 K=1,NVAR	0930
100	RIJ(J,K)=RIJ(J,K)+DATA(J)*DATA(K)	0940
110	CONTINUE	0950
C	COMPUTE STANDARD DEVIATIONS*SQR ROUTE (OBS-1)	0960
	DO 120 I=1,NVAR	0970
120	SIGMA(I)=(RIJ(I,I)-XBAR(I)*XBAR(I)/OBS)**.5	0980
C	COMPUTE CORRELATION MATRIX	0990
	DO 130 I=1,NVAR	1000
	DO 130 J=1,NVAR	1010
130	RIJ(I,J)=(RIJ(I,J)-XBAR(I)*XBAR(J)/OBS)/(SIGMA(I)*SIGMA(J))	1020
C	COMPUTE MEANS AND STANDARD DEVIATIONS	1030
	DO 140 I=1,NVAR	1040
	XBAR(I)=XBAR(I)/OBS	1050
140	SIGMA(I)=SIGMA(I)/(OBS-1.0)**.5	1060
C	SKIP TO NEW PAGE, WRITE I.D., AVERAGES, STANDARD DEVIATIONS,	1070
C	AND SIMPLE CORRELATION MATRIX.	1080
	WRITE(3,65)(AID(I),I=1,18)	1090
65	FORMAT('1',18A4)	1100
	WRITE(3,51)	1110
51	FORMAT('O AVERAGES')	1120
	WRITE(3,52)(I,XBAR(I),ICOM,I=1,NINDV),NVAR,XBAR(NVAR)	1130
52	FORMAT(4(' VAR(' ,I2,')=' ,F13.2,A1))	1140
	WRITE(3,53)	1150
53	FORMAT('O STANDARD DEVIATIONS')	1160
	WRITE(3,52)(I,SIGMA(I),ICOM,I=1,NINDV),NVAR,SIGMA(NVAR)	1170
	WRITE(3,55)	1180
55	FORMAT('O SIMPLE CORRELATION COEFFICIENTS')	1190
	DO 150 I=1,NINDV	1200
	WRITE(3,56)(I,J,RIJ(I,J),ICOM,J=1,NINDV),I,NVAR,RIJ(I,NVAR)	1210

56	FORMAT (4(' VARS(' ,I2,' ,',I2,')=' ,F10.3,A1))	1220
150	CONTINUE	
C	PHASE 2. PERFORM STEPWISE CALCULATIONS AND PRINT RESULTS.	1230
C	DIMENSIONS	1240
C	INITIALIZE	1250
	DO 190 I=1,NVAR	1260
	SIGB(I)=0.0	1270
190	B(I)=0.0	1280
	NENT=0	1290
	DF=OBS-1.0	1300
	NSTEP=-1	1310
C	TRANSFORM SIGMA VECTOR FROM STANDARD DEVIATIONS TO SQUARE	1320
C	ROOTS OF SUMS CF SQUARES.	1330
	DO 310 I=1,NVAR	1340
310	SIGMA(I)=SIGMA(I)*(OBS-1.0)**.5	1350
C	BEGIN STEP NUMBER NSTEP.	1360
200	NSTEP=NSTEP+1	1370
	STDEE=((RIJ(NVAR,NVAR)/DF)**.5)*SIGMA(NVAR)	1380
	DF=DF-1.0	1390
	IF(DF)1010,1010,205	1400
205	VMIN=0.0	1410
	VMAX=0.0	1420
	NIN=0	1430
C	FIND MINIMUM VARIANCE CONTRIBUTION OF VARIABLES IN REGRESSION	1440
C	EQUATION. FIND MAXIMUM VARIANCE CONTRIBUTION OF VARIABLES	1450
C	NOT IN REGRESSION EQUATION.	1460
	DO 300 I=1,NINDV	1470
210	VI=RIJ(I,NVAR)*RIJ(NVAR,I)/RIJ(I,I)	1490
	IF(VI)240,300,220	1500
220	IF(VI-VMAX)300,300,230	1510
230	VMAX=VI	1520
	NMAX=I	1530
	GO TO 300	1540
240	NIN=NIN+1	1550
	ID(NIN)=I	1560
C	COMPUTE REGRESSION COEFFICIENT AND ITS STANDARD DEVIATION.	1570
	B(NIN)=RIJ(I,NVAR)*SIGMA(NVAR)/SIGMA(I)	1580
	SIGB(NIN)=(STDEE*RIJ(I,I)**.5)/SIGMA(I)	1590
	IF(VMIN)250,260,1000	1600
250	IF(VI-VMIN)300,300,260	1610
260	VMIN=VI	1620
	NMIN=I	1630
300	CONTINUE	1640
	IF(NIN)1000,460,400	1650
C	COMPUTE CONSTANT TERM.	1660
400	BSUBO=XBAR(NVAR)	1670
	DO 410 I=1,NIN	1680
	J=ID(I)	1690
410	BSUBO=BSUBO-B(I)*XBAR(J)	1700
	IF(NENT)1000,480,420	1710
C	OUTPUT FOR VARIABLE ADDED	1720
420	WRITE(3,57)NSTEP,K	1730
57	FORMAT('STEP NUMBER ',I2,10X,'ENTER VARIABLE ',I2)	1740
	DEPV = NSTEP	
425	WRITE(3,58)STDEE	1750
58	FORMAT(' STANCARD DEVIATION OF RESIDUALS=' ,F16.3)	1760
	SDPRM=(STDEE/XBAR(NVAR))*100.	1761
	WRITE(3,49)SDPRM	1762
49	FORMAT(' STD. DEV. AS PERCENT OF RESPONSE MEAN=' ,F10.3)	1763
	R=(1.-RIJ(NVAR,NVAR))**.5	1770
	RSQ = R**2.	

RSQP = RSQ * 100.	
WRITE(3,59) RSQP	
59 FORMAT(' PERCENT VARIATION EXPLAINED R-SQ=',F15.3)	1773
CRSQ = 1.-((1.-RSQ)*(CHS-1.))/(OBS-DEPV-1.)	
CRSQP = CRSQ * 100.	
WRITE(3,84) CRSQP	
84 FORMAT(' CORRECTED R-SQ AS A PERCENT=',F20.3)	
IDFN=OBS-DF-2.0	1800
IDFD=DF+1.0	1810
F=(SIGMA(NVAR)**2-(STDEE**2)*(DF+1.0))/((OBS-DF-2.0)*STDEE**2)	1820
WRITE(3,66)IDFN,IDFD,F	1830
66 FORMAT(' GOODNESS OF FIT OR OVERALL F,F(' ,I3,' ,',I3,')=',F8.3)	1840
WRITE(3,60)BSUBC	1850
60 FORMAT(' CONSTANT TERM=',18X,F16.8)	1860
WRITE(3,61)	1870
61 FORMAT('OVAR COEFF STD DEV T VALUE'	1880
1)	1881
WRITE(3,62)	1890
62 FORMAT(' COEFF')	1900
DC 430 I=1,NIN	1910
J=ID(I)	1920
T=B(I)/SIGB(I)	1930
WRITE(3,63)ID(I),B(I),SIGB(I),T	1940
63 FORMAT(' ',I3,F18.8,F20.8,F18.8)	1950
430 CONTINUE	
C COMPUTE F LEVEL FOR MINIMUM VARIANCE CONTRIBUTION VARIABLE	1960
C IN REGRESSION EQUATION.	1970
FLEVL=VMIN*DF/RIJ(NVAR,NVAR)	1980
IF(FOUT+FLEVL)460,460,450	1990
C INITIALIZE FOR REMOVAL OF VARIABLE K FROM EQUATION.	2000
450 K=NMIN	2010
NENT=0	2020
DF=DF+2.0	2030
GO TO 500	2040
C COMPUTE F LEVEL FOR MAXIMUM VARIANCE CONTRIBUTION VARIABLE	2050
C NOT IN EQUATION.	2060
460 FLEVL=VMAX*DF/(RIJ(NVAR,NVAR)-VMAX)	2070
IF(FLEVL-FIN)600,600,470	2080
C INITIALIZE FOR ENTRY OF VARIABLE K INTO EQUATION.	2090
470 K=NMAX	2100
NENT=K	2110
GO TO 500	2120
C OUTPUT FOR VARIABLE DELETED	2130
480 WRITE(3,64)NSTEP,K	2140
64 FORMAT('OSTEP NUMBER ',I2,10X,'DELETE VARIABLE ',I2)	2150
GO TO 425	2160
C UPDATE MATRIX	2170
500 DC 540 I=1,NVAR	2180
IF(I-K)510,540,510	2190
510 DO 530 J=1,NVAR	2200
IF(J-K)520,530,520	2210
520 RIJ(I,J)=RIJ(I,J)-RIJ(I,K)*RIJ(K,J)/RIJ(K,K)	2220
530 CONTINUE	2230
540 CONTINUE	2240
DO 560 J=1,NVAR	2250
IF(J-K)550,560,550	2260
550 RIJ(K,J)=RIJ(K,J)/RIJ(K,K)	2270
560 CONTINUE	2280
DO 580 I=1,NVAR	2290
IF(I-K)570,580,570	2300
570 RIJ(I,K)=-RIJ(I,K)/RIJ(K,K)	2310

	580	CONTINUE	2320
		RIJ(K,K)=1.0/RIJ(K,K)	2330
		GO TO 200	2340
	600	IF(IRES)610,640,610	2350
C		PRINT RESIDUALS	2360
	610	IFA=1	2370
		WRITE(3,67)	2380
	67	FORMAT('O OBS ACTUAL ESTIMATE RESIDUAL NORMAL')	2390
		WRITE(3,69)	2391
	69	FORMAT(' DEVIATE')	2392
		DO 630 K=1,NGBS	2400
		READ(10' IFA)(DATA(I),I=1,NVAR)	2410
		EST=BSUBD	2420
		DO 620 I=1,NIN	2430
		J>ID(I)	2440
	620	EST=EST+B(I)*DATA(J)	2450
		RESID = DATA(NVAR)-EST	2460
		XNORD = RESID/STDEE	2461
		IF(DABS(XNORD)-3.)91,92,92	2470
	91	IF(DABS(XNORD)-2.)93,94,94	2471
	92	WRITE(3,30)K,DATA(NVAR),EST,RESID,XNORD	2480
	30	FORMAT(' ',I4,4F12.2,' **')	2481
		GO TO 630	2482
	94	WRITE(3,31)K,DATA(NVAR),EST,RESID,XNORD	2483
	31	FORMAT(' ',I4,4F12.2,' **')	2484
		GO TO 630	2485
	93	WRITE(3,68)K,DATA(NVAR),EST,RESID,XNORD	2486
	68	FORMAT(' ',I4,4F12.2)	2487
	630	CONTINUE	2490
C		NORMAL END OF JOB	2500
	640	GO TO 101	2501
	999	CALL EXIT	
C		ERRGR. NIN, NENT, VMIN, NCONS, OR NTRANS IS NEGATIVE. CHECK	2520
C		FOR CONTROL CARD ERROR.	2530
	1000	STOP1	
C		ERROR DEGREES OF FREEDOM =0. EITHER ADD MORE DATA OBSERVATIONS OR	2550
C		DELETE ONE OR MCRE INDEPENDENT VARIABLES. SAMPLE SIZE MUST EXCEED	2560
C		NUMBER OF INDEPENDENT VARIABLES BY AT LEAST 2.	2570
	1010	STOP2	2580
C		ERROR. F LEVEL FOR INCOMING VARIABLE IS LESS THAN F LEVEL FOR	2590
C		OUTGOING VARIABLE.	2600
	1020	STOP4	2610
		END	

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16. ABSTRACT Pathogenic enteric viruses transmitted via the water route present a potential hazard to public health because of their resistance to natural or artificial disinfection mechanisms. Of constant concern to public health officials is the ability of viruses to pass through water treatment plants. Therefore, many research investigations have been directed toward the study of the inactivation of viruses and enteric organisms. This report describes a mathematical model which can be used to characterize the response of viruses to a disinfecting agent. Not only is the model presented, but a technique is described which can be used to estimate the model's parameters. Both the model and the estimation technique are being used to analyze experimental information resulting from disinfection studies.				
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