

## APPENDIX 1

### ACTIVITY PROFILES FOR GENETIC AND RELATED EFFECTS

#### *Methods*

The x-axis of the activity profile (Waters *et al.*, 1987, 1988) represents the bioassays in phylogenetic sequence by endpoint, and the values on the y-axis represent the logarithmically transformed lowest effective doses (LED) and highest ineffective doses (HID) tested. The term 'dose', as used in this report, does not take into consideration length of treatment or exposure and may therefore be considered synonymous with concentration. In practice, the concentrations used in all the in-vitro tests were converted to  $\mu\text{g/ml}$ , and those for in-vivo tests were expressed as  $\text{mg/kg bw}$ . Because dose units are plotted on a log scale, differences in molecular weights of compounds do not, in most cases, greatly influence comparisons of their activity profiles. Conventions for dose conversions are given below.

Profile-line height (the magnitude of each bar) is a function of the LED or HID, which is associated with the characteristics of each individual test system – such as population size, cell-cycle kinetics and metabolic competence. Thus, the detection limit of each test system is different, and, across a given activity profile, responses will vary substantially. No attempt is made to adjust or relate responses in one test system to those of another.

Line heights are derived as follows: for negative test results, the highest dose tested without appreciable toxicity is defined as the HID. If there was evidence of extreme toxicity, the next highest dose is used. A single dose tested with a negative result is considered to be equivalent to the HID. Similarly, for positive results, the LED is recorded. If the original data were analysed statistically by the author, the dose recorded is that at which the response was significant ( $p < 0.05$ ). If the available data were not analysed statistically, the dose required to produce an effect is estimated as follows: when a dose-related positive response is observed with two or more doses, the lower of the doses is taken as the LED; a single dose resulting in a positive response is considered to be equivalent to the LED.

In order to accommodate both the wide range of doses encountered and positive and negative responses on a continuous scale, doses are transformed logarithmically, so that effective (LED) and ineffective (HID) doses are represented by positive and negative numbers, respectively. The response, or logarithmic dose unit ( $\text{LDU}_{ij}$ ), for a given test system  $i$  and chemical  $j$  is represented by the expressions

$$\text{LDU}_{ij} = -\log_{10} (\text{dose}), \text{ for HID values; } \text{LDU} \leq 0$$

and

$$\text{LDU}_{ij} = -\log_{10} (\text{dose} \times 10^{-5}), \text{ for LED values; } \text{LDU} \geq 0.$$

(1)

These simple relationships define a dose range of 0 to -5 logarithmic units for ineffective doses (1-100 000  $\mu\text{g/ml}$  or  $\text{mg/kg bw}$ ) and 0 to +8 logarithmic units for effective doses (100 000-0.001  $\mu\text{g/ml}$  or  $\text{mg/kg bw}$ ). A scale illustrating the LDU values is shown in Figure 1. Negative responses at doses less than 1  $\mu\text{g/ml}$  ( $\text{mg/kg bw}$ ) are set equal to 1. Effectively, an LED value  $\geq 100\ 000$  or an HID value  $\leq 1$  produces an LDU = 0; no quantitative information is gained from such extreme values. The dotted lines at the levels of log dose units 1 and -1 define a 'zone of uncertainty' in which positive results are reported at such high doses (between 10 000 and 100 000  $\mu\text{g/ml}$  or  $\text{mg/kg bw}$ ) or negative results are reported at such low dose levels (1 to 10  $\mu\text{g/ml}$  or  $\text{mg/kg bw}$ ) as to call into question the adequacy of the test.

**Fig. 1. Scale of log dose units used on the y-axis of activity profiles**

Positive ( $\mu\text{g/ml}$ or $\text{mg/kg bw}$ )	Log dose units	
0.001	8	----
0.01	7	--
0.1	6	--
1.0	5	--
10	4	--
100	3	--
1000	2	--
10 000	1	--
100 000	0	----
	10	--
	100	--
	1000	--
	10 000	--
	100 000	----
	-1	--
	-2	--
	-3	--
	-4	--
	-5	----
Negative ( $\mu\text{g/ml}$ or $\text{mg/kg bw}$ )		

LED and HID are expressed as  $\mu\text{g/ml}$  or  $\text{mg/kg bw}$ .

In practice, an activity profile is computer generated. A data entry programme is used to store abstracted data from published reports. A sequential file (in ASCII) is created for each compound, and a record within that file consists of the name and Chemical Abstracts Service number of the compound, a three-letter code for the test system (see below), the qualitative test result (with and without an exogenous metabolic system), dose (LED or HID), citation number and additional source information. An abbreviated citation for each publication is stored in a segment of a record accessing both the test data file and the citation file. During processing of the data file, an average of the logarithmic values of the data subset is calculated, and the length of the profile line represents this average value. All dose values are plotted for each profile line, regardless of whether results are positive or negative. Results obtained in the absence of an exogenous metabolic system are indicated by a bar (-), and results obtained in the presence of an exogenous metabolic system are indicated by an

upward-directed arrow ( $\uparrow$ ). When all results for a given assay are either positive or negative, the mean of the LDU values is plotted as a solid line; when conflicting data are reported for the same assay (i.e., both positive and negative results), the majority data are shown by a solid line and the minority data by a dashed line (drawn to the extreme conflicting response). In the few cases in which the numbers of positive and negative results are equal, the solid line is drawn in the positive direction and the maximal negative response is indicated with a dashed line.

Profile lines are identified by three-letter code words representing the commonly used tests. Code words for most of the test systems in current use in genetic toxicology were defined for the US Environmental Protection Agency's GENE-TOX Program (Waters, 1979; Waters & Auletta, 1981). For IARC Monographs Supplement 6, Volume 44 and subsequent volumes, including this publication, codes were redefined in a manner that should facilitate inclusion of additional tests. If a test system is not defined precisely, a general code is used that best defines the category of the test. Naming conventions are described below.

Data listings are presented with each activity profile and include endpoint and test codes, a short test code definition, results [either with (M) or without (NM) an exogenous activation system], the associated LED or HID value and a short citation. Test codes are organized phylogenetically and by endpoint from left to right across each activity profile and from top to bottom of the corresponding data listing. Endpoints are defined as follows: A, aneuploidy; C, chromosomal aberrations; D, DNA damage; F, assays of body fluids; G, gene mutation; H, host-mediated assays; I, inhibition of intercellular communication; M, micronuclei; P, sperm morphology; R, mitotic recombination or gene conversion; S, sister chromatid exchange; and T, cell transformation.

#### *Dose conversions for activity profiles*

Doses are converted to  $\mu\text{g/ml}$  for in-vitro tests and to  $\text{mg/kg bw per day}$  for in-vivo experiments.

#### 1. In-vitro test systems

- (a) Weight/volume converts directly to  $\mu\text{g/ml}$ .
- (b) Molar (M) concentration  $\times$  molecular weight =  $\text{mg/ml} = 10^3 \mu\text{g/ml}$ ; mM concentration  $\times$  molecular weight =  $\mu\text{g/ml}$ .
- (c) Soluble solids expressed as % concentration are assumed to be in units of mass per volume (i.e., 1% = 0.01 g/ml = 10 000  $\mu\text{g/ml}$ ; also, 1 ppm = 1  $\mu\text{g/ml}$ ).
- (d) Liquids and gases expressed as % concentration are assumed to be given in units of volume per volume. Liquids are converted to weight per volume using the density (D) of the solution ( $D = \text{g/ml}$ ). Gases are converted from volume to mass using the ideal gas law,  $PV = nRT$ . For exposure at 20–37°C at standard atmospheric pressure, 1% (v/v) = 0.4  $\mu\text{g/ml} \times$  molecular weight of the gas. Also, 1 ppm (v/v) =  $4 \times 10^{-5} \mu\text{g/ml} \times$  molecular weight.
- (e) In microbial plate tests, it is usual for the doses to be reported as weight/plate, whereas concentrations are required to enter data on the activity profile chart. While remaining cognisant of the errors involved in the process, it is assumed that

a 2-ml volume of top agar is delivered to each plate and that the test substance remains in solution within it; concentrations are derived from the reported weight/plate values by dividing by this arbitrary volume. For spot tests, a 1-ml volume is used in the calculation.

- (f) Conversion of particulate concentrations given in  $\mu\text{g}/\text{cm}^2$  are based on the area (A) of the dish and the volume of medium per dish; i.e., for a 100-mm dish:  $A = \pi R^2 = \pi \times (5 \text{ cm})^2 = 78.5 \text{ cm}^2$ . If the volume of medium is 10 ml, then  $78.5 \text{ cm}^2 = 10 \text{ ml}$  and  $1 \text{ cm}^2 = 0.13 \text{ ml}$ .

## 2. In-vitro systems using in-vivo activation

For the body fluid-urine (BF-) test, the concentration used is the dose (in mg/kg bw) of the compound administered to test animals or patients.

## 3. In-vivo test systems

- (a) Doses are converted to mg/kg bw per day of exposure, assuming 100% absorption. Standard values are used for each sex and species of rodent, including body weight and average intake per day, as reported by Gold *et al.* (1984). For example, in a test using male mice fed 50 ppm of the agent in the diet, the standard food intake per day is 12% of body weight, and the conversion is dose =  $50 \text{ ppm} \times 12\% = 6 \text{ mg/kg bw per day}$ .

Standard values used for humans are: weight - males, 70 kg; females, 55 kg; surface area,  $1.7 \text{ m}^2$ ; inhalation rate, 20 l/min for light work, 30 l/min for mild exercise.

- (b) When reported, the dose at the target site is used. For example, doses given in studies of lymphocytes of humans exposed *in vivo* are the measured blood concentrations in  $\mu\text{g}/\text{ml}$ .

### Codes for test systems

For specific nonmammalian test systems, the first two letters of the three-symbol code word define the test organism (e.g., SA- for *Salmonella typhimurium*, EC- for *Escherichia coli*). If the species is not known, the convention used is -S-. The third symbol may be used to define the tester strain (e.g., SA8 for *S. typhimurium* TA1538, ECW for *E. coli* WP2uvrA). When strain designation is not indicated, the third letter is used to define the specific genetic endpoint under investigation (e.g., -D for differential toxicity, -F for forward mutation, -G for gene conversion or genetic crossing-over, -N for aneuploidy, -R for reverse mutation, -U for unscheduled DNA synthesis). The third letter may also be used to define the general endpoint under investigation when a more complete definition is not possible or relevant (e.g., -M for mutation, -C for chromosomal aberration).

For mammalian test systems, the first letter of the three-letter code word defines the genetic endpoint under investigation: A- for aneuploidy, B- for binding, C- for chromosomal aberration, D- for DNA strand breaks, G- for gene mutation, I- for inhibition of intercellular communication, M- for micronucleus formation, R- for DNA repair, S- for sister chromatid exchange, T- for cell transformation and U- for unscheduled DNA synthesis.

For animal (i.e., non-human) test systems *in vitro*, when the cell type is not specified, the code letters -IA are used. For such assays *in vivo*, when the animal species is not specified, the code letters -VA are used. Commonly used animal species are identified by the third letter (e.g., -C for Chinese hamster, -M for mouse, -R for rat, -S for Syrian hamster).

For test systems using human cells *in vitro*, when the cell type is not specified, the code letters -IH are used. For assays on humans *in vivo*, when the cell type is not specified, the code letters -VH are used. Otherwise, the second letter specifies the cell type under investigation (e.g., -BH for bone marrow, -LH for lymphocytes).

Some other specific coding conventions used for mammalian systems are as follows: BF- for body fluids, HM- for host-mediated, -L for leucocytes or lymphocytes *in vitro* (-AL, animals; -HL, humans), -L- for leucocytes *in vivo* (-LA, animals; -LH, humans), -T for transformed cells.

Note that these are examples of major conventions used to define the assay code words. The alphabetized listing of codes must be examined to confirm a specific code word. As might be expected from the limitation to three symbols, some codes do not fit the naming conventions precisely. In a few cases, test systems are defined by first-letter code words, for example: MST, mouse spot test; SLP, mouse specific locus test, postspermatogonia; SLO, mouse specific locus test, other stages; DLM, dominant lethal test in mice; DLR, dominant lethal test in rats; MHT, mouse heritable translocation test.

The genetic activity profiles and listings that follow were prepared in collaboration with Environmental Health Research and Testing Inc. (EHRT) under contract to the US Environmental Protection Agency; EHRT also determined the doses used. The references cited in each genetic activity profile listing can be found in the list of references in the appropriate monograph.

### References

- Garrett, N.E., Stack, H.F., Gross, M.R. & Waters, M.D. (1984) An analysis of the spectra of genetic activity produced by known or suspected human carcinogens. *Mutat. Res.*, 134, 89-111
- Gold, L.S., Sawyer, C.B., Magaw, R., Backman, G.M., de Veciana, M., Levinson, R., Hooper, N.K., Havender, W.R., Bernstein, L., Peto, R., Pike, M.C. & Ames, B.N. (1984) A carcinogenic potency database of the standardized results of animal bioassays. *Environ. Health Perspect.*, 58, 9-319
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- Waters, M.D., Stack, H.F., Brady, A.L., Lohman, P.H.M., Haroun, L. & Vainio, H. (1988) Use of computerized data listings and activity profiles of genetic and related effects in the review of 195 compounds. *Mutat. Res.*, 205, 295-312

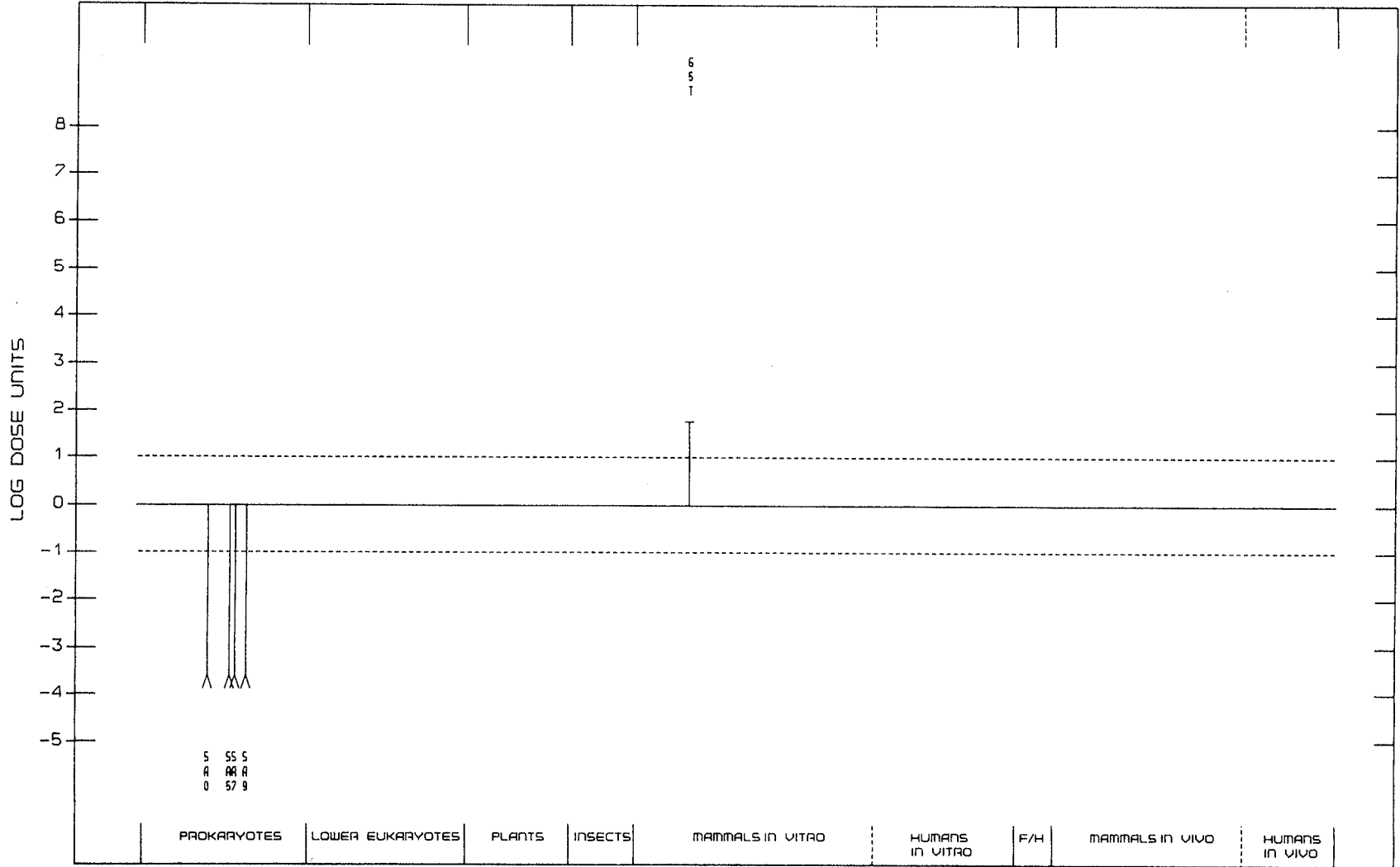
## CHLORENDIC ACID

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END POINT	TEST CODE	TEST SYSTEM	RESULTS NM M	DOSE (LED OR HID)	REFERENCE
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	3845.0000	NTP, 1987
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	3845.0000	NTP, 1987
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	3845.0000	NTP, 1987
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	3845.0000	NTP, 1987
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	+ 0	1700.0000	NTP, 1987

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CHLORENDIC ACID  
115-28-6

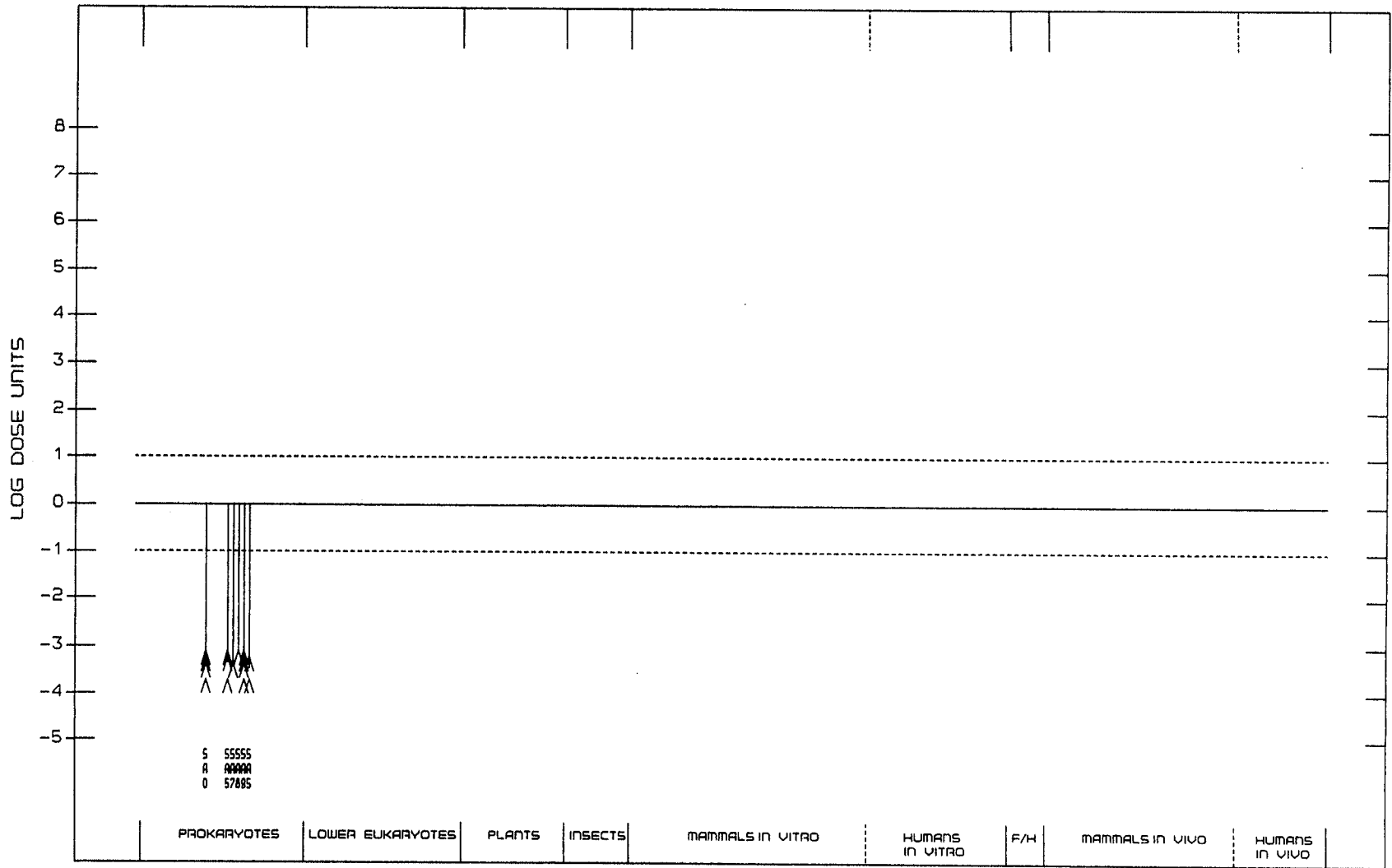


## CHLORINATED PARAFFINS

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G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	1250.0000	BIRTLEY ET AL., 1980
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	2300.0000	MEIJER ET AL., 1981
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	1667.0000	NTPa, 1986
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	5000.0000	NTPb, 1986
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	1250.0000	BIRTLEY ET AL., 1980
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	1667.0000	NTPa, 1986
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	5000.0000	NTPb, 1986
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	2300.0000	MEIJER ET AL., 1981
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	- -	1250.0000	BIRTLEY ET AL., 1980
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	1250.0000	BIRTLEY ET AL., 1980
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	2300.0000	MEIJER ET AL., 1981
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	1667.0000	NTPa, 1986
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	5000.0000	NTPb, 1986
G	SAS	S. TYPHIMURIUM (OTHER), REVERSE MUTATION	- -	1667.0000	NTPa, 1986
G	SAS	S. TYPHIMURIUM (OTHER), REVERSE MUTATION	- -	5000.0000	NTPb, 1986



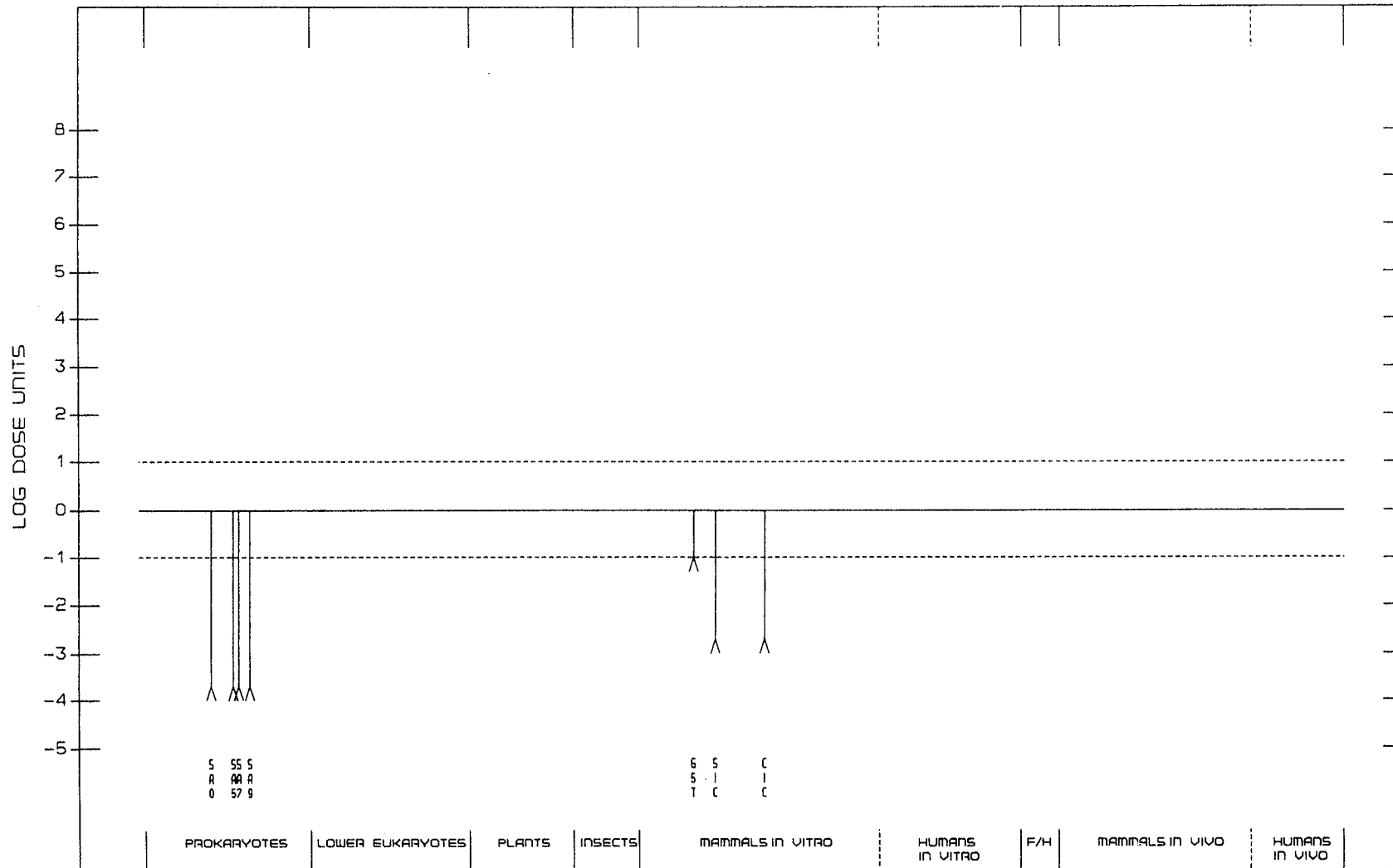
CHLORINATED PARAFFINS  
63449-39-8



## DECABROMODIPHENYL OXIDE

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G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	5000.0000	NTP, 1986
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	5000.0000	NTP, 1986
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	5000.0000	NTP, 1986
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	5000.0000	NTP, 1986
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	-	-	10.0000	NTP, 1986
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	-	-	500.0000	NTP, 1986
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	-	-	500.0000	NTP, 1986

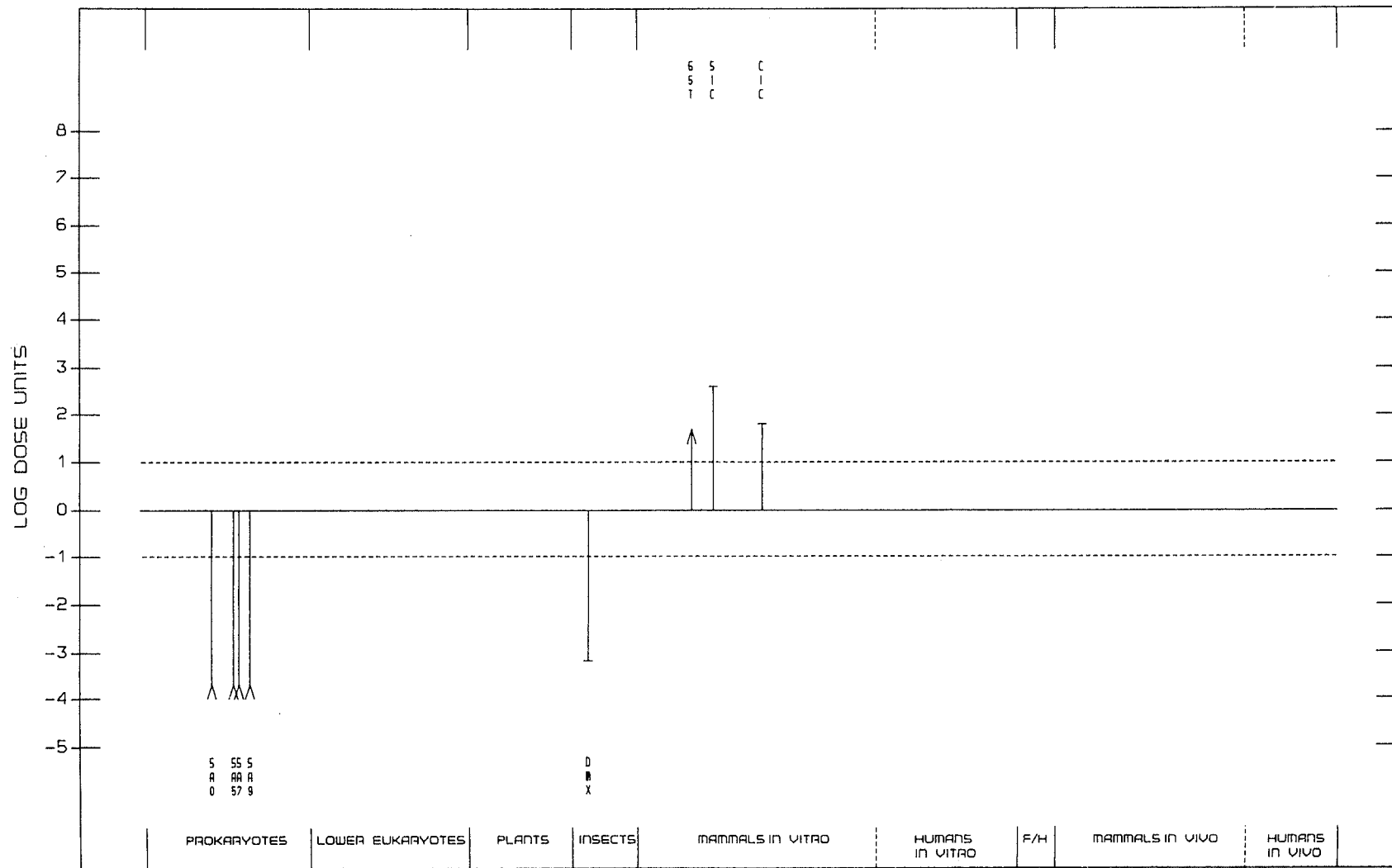
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## DIMETHYL HYDROGEN PHOSPHITE

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G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	5000.0000	NTP, 1985
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	5000.0000	NTP, 1985
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	5000.0000	NTP, 1985
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	5000.0000	NTP, 1985
G	DMX	D. MELANOGASTER, SEX-LINKED RECESSIVES	- 0	1500.0000	NTP, 1985
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	- +	2100.0000	MCGREGOR ET AL., 1988
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	+ +	250.0000	TENNANT ET AL., 1987b
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	+ +	1600.0000	TENNANT ET AL., 1987b

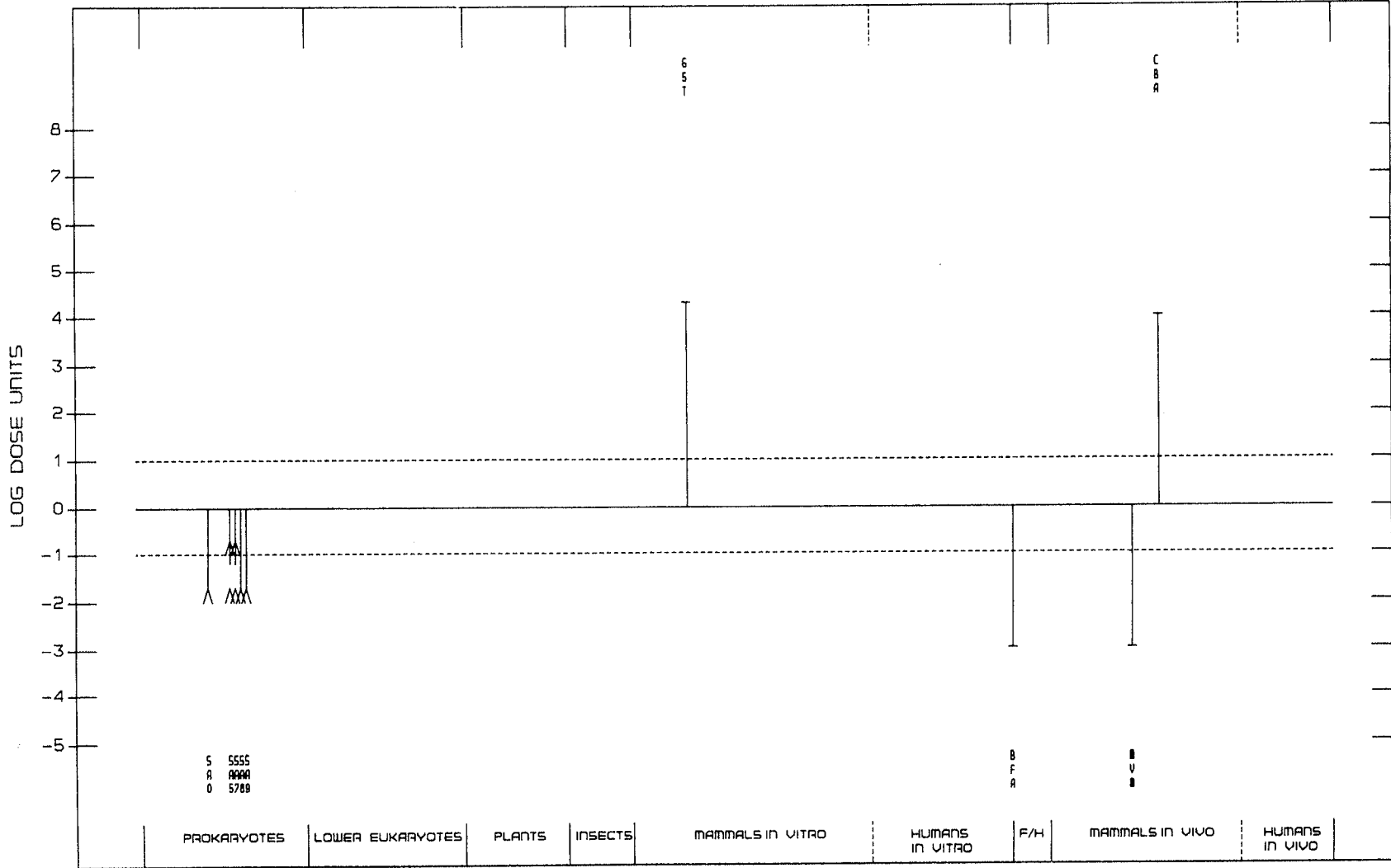
DIMETHYL HYDROGEN PHOSPHITE  
868-85-9



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END POINT	TEST CODE	TEST SYSTEM	RESULTS NM M	DOSE (LED OR HID)	REFERENCE
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	50.0000	CONNOR ET AL., 1980
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	50.0000	MACGREGOR ET AL., 1980
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	50.0000	CONNOR ET AL., 1980
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	5.0000	MACGREGOR ET AL., 1980
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	50.0000	CONNOR ET AL., 1980
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	5.0000	MACGREGOR ET AL., 1980
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	- -	50.0000	CONNOR ET AL., 1980
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	50.0000	CONNOR ET AL., 1980
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	50.0000	MACGREGOR ET AL., 1980
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	+ 0	5.0000	NTP, 1987
F	BFA	ANIMAL BODY FLUIDS, MICROBIAL MUTAGENICITY	- 0	1000.0000	CONNOR ET AL., 1980
M	MVM	MICRONUCLEUS TEST, MICE IN VIVO	- 0	1000.0000	CONNOR ET AL., 1980
C	CBA	CHROM ABERR, ANIMAL BONE MARROW IN VIVO	(+) 0	10.0000	CONNOR ET AL., 1980

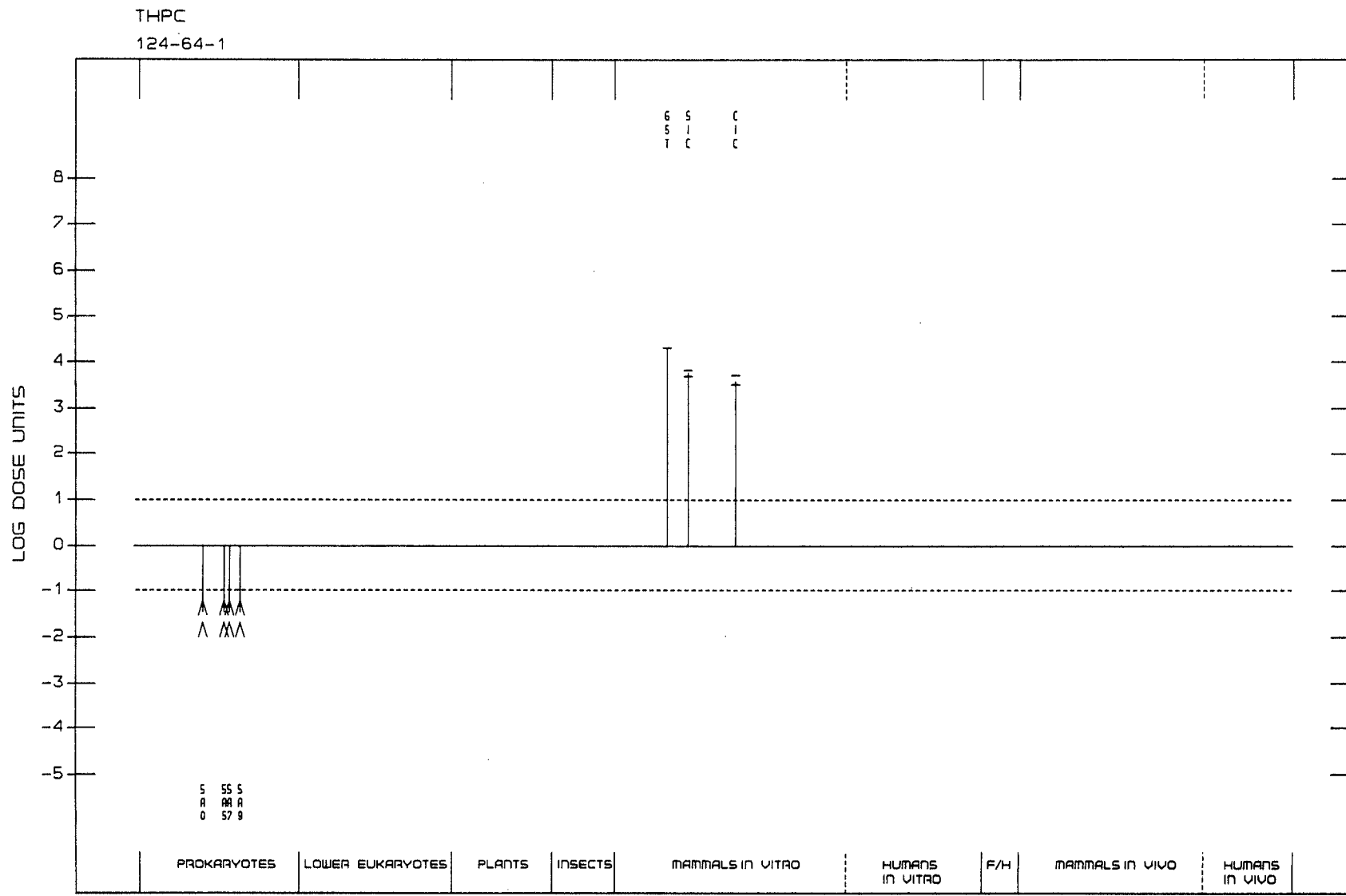
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## THPC

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G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	17.0000	NTP, 1987
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	50.0000	MACGREGOR ET AL., 1980
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	17.0000	NTP, 1987
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	50.0000	MACGREGOR ET AL., 1980
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	17.0000	NTP, 1987
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	50.0000	MACGREGOR ET AL., 1980
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	17.0000	NTP, 1987
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	+	0	5.0000	NTP, 1987
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	+	+	15.0000	NTP, 1987
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	+	+	20.0000	LOVEDAY ET AL., 1989
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	+	+	30.0000	NTP, 1987
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	+	0	19.0000	SASAKI ET AL., 1980
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	+	?	30.0000	LOVEDAY ET AL., 1989
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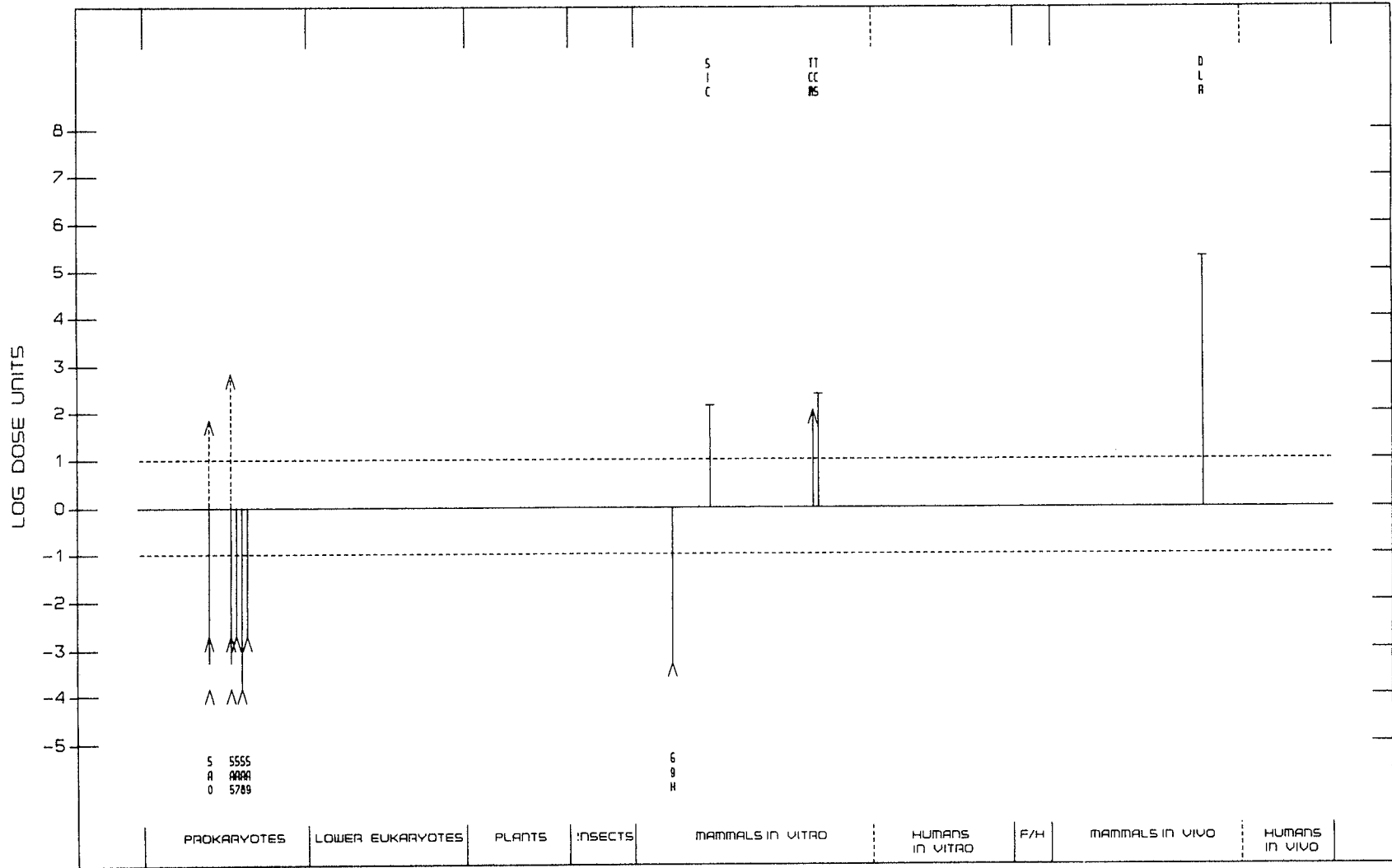




## TRIS(2-CHLOROETHYL) PHOSPHATE

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			NM	M		
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	6950.0000	PRIVAL ET AL., 1977
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	(+)	1427.0000	NAKAMURA ET AL., 1979
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	500.0000	HAWORTH ET AL., 1983
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	6950.0000	PRIVAL ET AL., 1977
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	+	143.0000	NAKAMURA ET AL., 1979
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	500.0000	HAWORTH ET AL., 1983
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	500.0000	HAWORTH ET AL., 1983
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	-	-	6950.0000	PRIVAL ET AL., 1977
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	500.0000	HAWORTH ET AL., 1983
G	G9H	MUTATION, CHL V79 CELLS, HPRT	-	-	2000.0000	SALA ET AL., 1982
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	+	+	700.0000	SALA ET AL., 1982
T	TCM	CELL TRANSFORMATION, C3H10T1/2 CELLS	-	(+)	900.0000	SALA ET AL., 1982
T	TCS	CELL TRANSFORMATION, SHE, CLONAL ASSAY	+	0	400.0000	SALA ET AL., 1982
M	MVM	MICRONUCLEUS TEST, MICE IN VIVO	?	0	250.0000	SALA ET AL., 1982
C	DLR	DOMINANT LETHAL TEST, RATS	+	0	0.5000	SHEPEL'SKAIA & DYSHGINEVICH, 1981

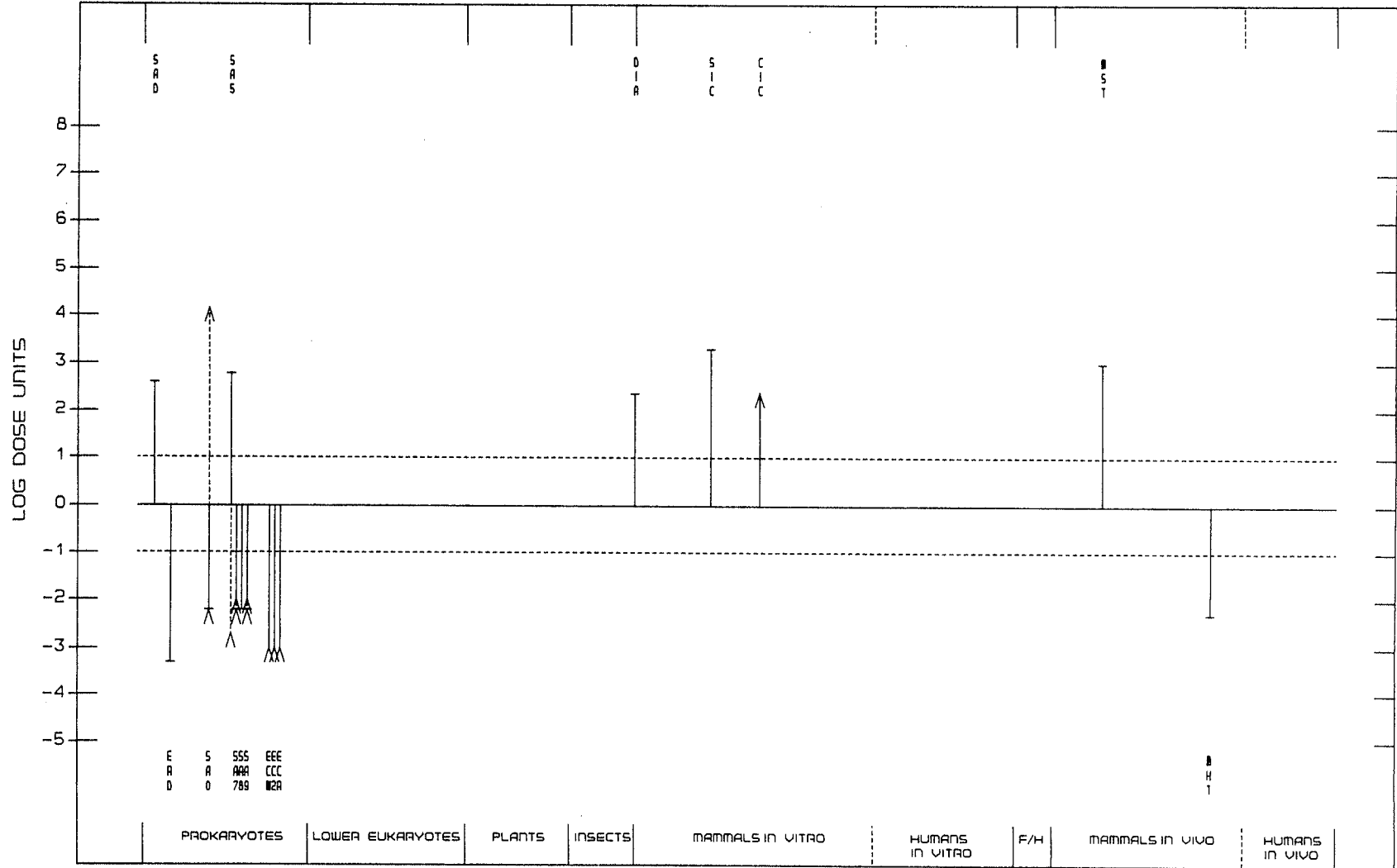
TRIS(2-CHLOROETHYL) PHOSPHATE  
115-96-8



## P-CHLORO-O-TOLUIDINE

END POINT	TEST CODE	TEST SYSTEM	RESULTS NM M	DOSE (LED OR HID)	REFERENCE
D	SAD	S. TYPHIMURIUM, DIFFERENTIAL TOXICITY	+ 0	250.0000	RASHID ET AL., 1984
D	ERD	E. COLI REC, DIFFERENTIAL TOXICITY	- 0	2000.0000	RASHID ET AL., 1984
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- +	7.0000	ZIMMER ET AL., 1980
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	167.0000	HAWORTH ET AL., 1983
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- 0	163.0000	RASHID ET AL., 1984
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	500.0000	HAWORTH ET AL., 1983
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	+ 0	163.0000	RASHID ET AL., 1984
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	100.0000	ZIMMER ET AL., 1980
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- -	167.0000	HAWORTH ET AL., 1983
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	- 0	163.0000	RASHID ET AL., 1984
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	- 0	163.0000	RASHID ET AL., 1984
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	100.0000	ZIMMER ET AL., 1980
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	167.0000	HAWORTH ET AL., 1983
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- 0	163.0000	RASHID ET AL., 1984
G	ECW	E. COLI WP2 UVRA, REVERSE MUTATION	- -	1000.0000	RASHID ET AL., 1984
G	EC2	E. COLI WP2, REVERSE MUTATION	- -	1000.0000	RASHID ET AL., 1984
G	ECR	E. COLI (OTHER), REVERSE MUTATION	- -	1000.0000	RASHID ET AL., 1984
D	DIA	STRAND BREAKS/X-LINKS, ANIMAL CELLS IN VITRO	(+) 0	425.0000	ZIMMER ET AL., 1980
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	+ +	50.0000	GALLOWAY ET AL., 1987
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	- +	400.0000	GALLOWAY ET AL., 1987
G	MST	MOUSE SPOT TEST	+ 0	100.0000	LANG, 1984
C	MHT	MOUSE HERITABLE TRANSLOCATION TEST	- 0	200.0000	LANG & ADLER, 1982
D	BVD	BINDING TO DNA, ANIMALS IN VIVO	+ 0	25.0000	BENTLEY ET AL., 1986
D	BVP	BINDING TO RNA/PROTEIN, ANIMALS IN VIVO	+ 0	25.0000	BENTLEY ET AL., 1986

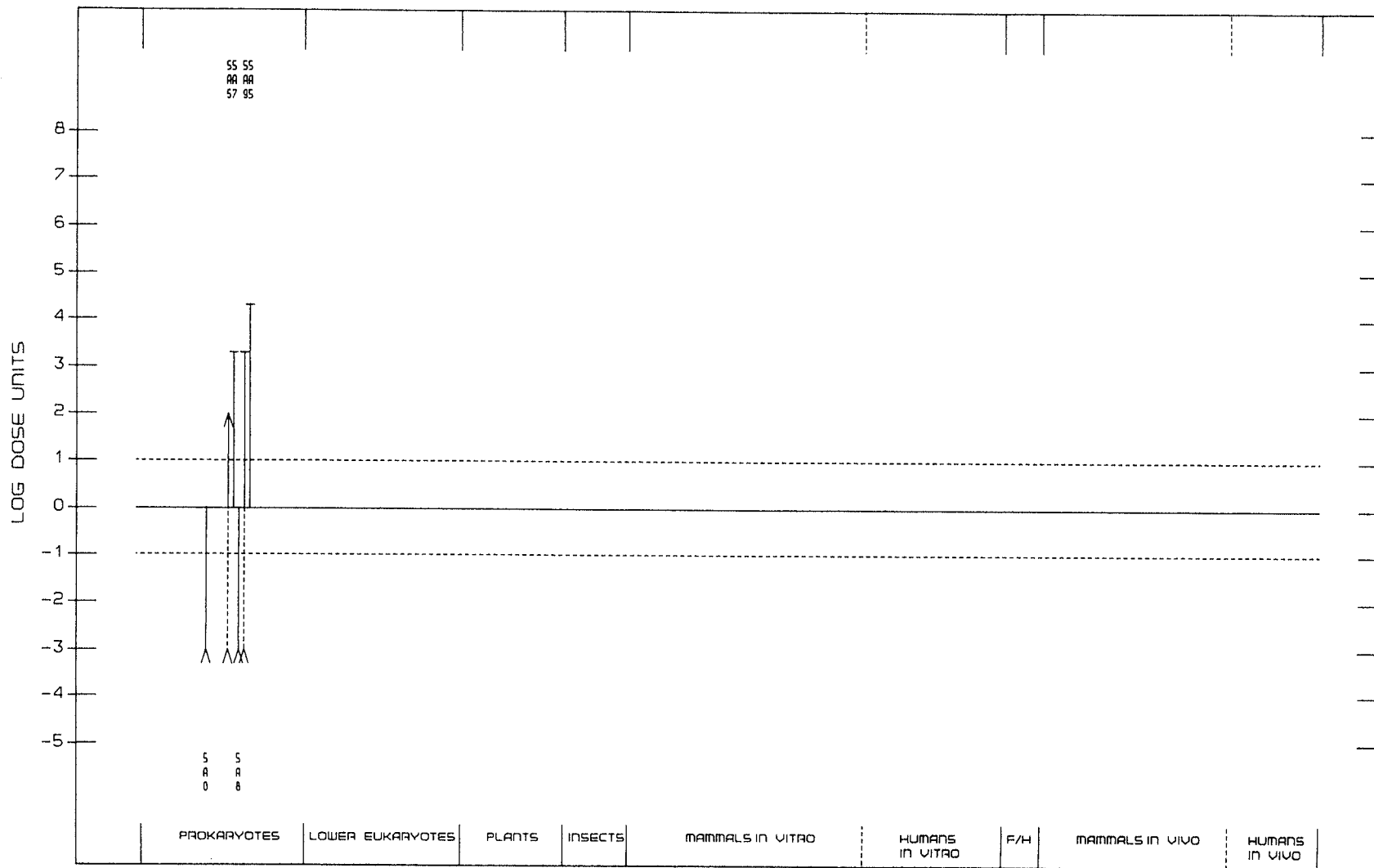
P-CHLORO-O-TOLUIDINE  
95-69-2



## DISPERSE BLUE 1

END POINT	TEST CODE	TEST SYSTEM	RESULTS NM M	DOSE (LED OR HID)	REFERENCE
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	1000.0000	BROWN & BROWN, 1976
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	- -	1000.0000	NTP, 1986
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- -	1000.0000	BROWN & BROWN, 1976
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	- +	1000.0000	NTP, 1986
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	(+) (+)	50.0000	BROWN & BROWN, 1976
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	- -	1000.0000	BROWN & BROWN, 1976
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	- -	1000.0000	BROWN & BROWN, 1976
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	+ +	50.0000	NTP, 1986
G	SAS	S. TYPHIMURIUM (OTHER), REVERSE MUTATION	+ -	5.0000	NTP, 1986

DISPERSE BLUE 1  
2475-45-8

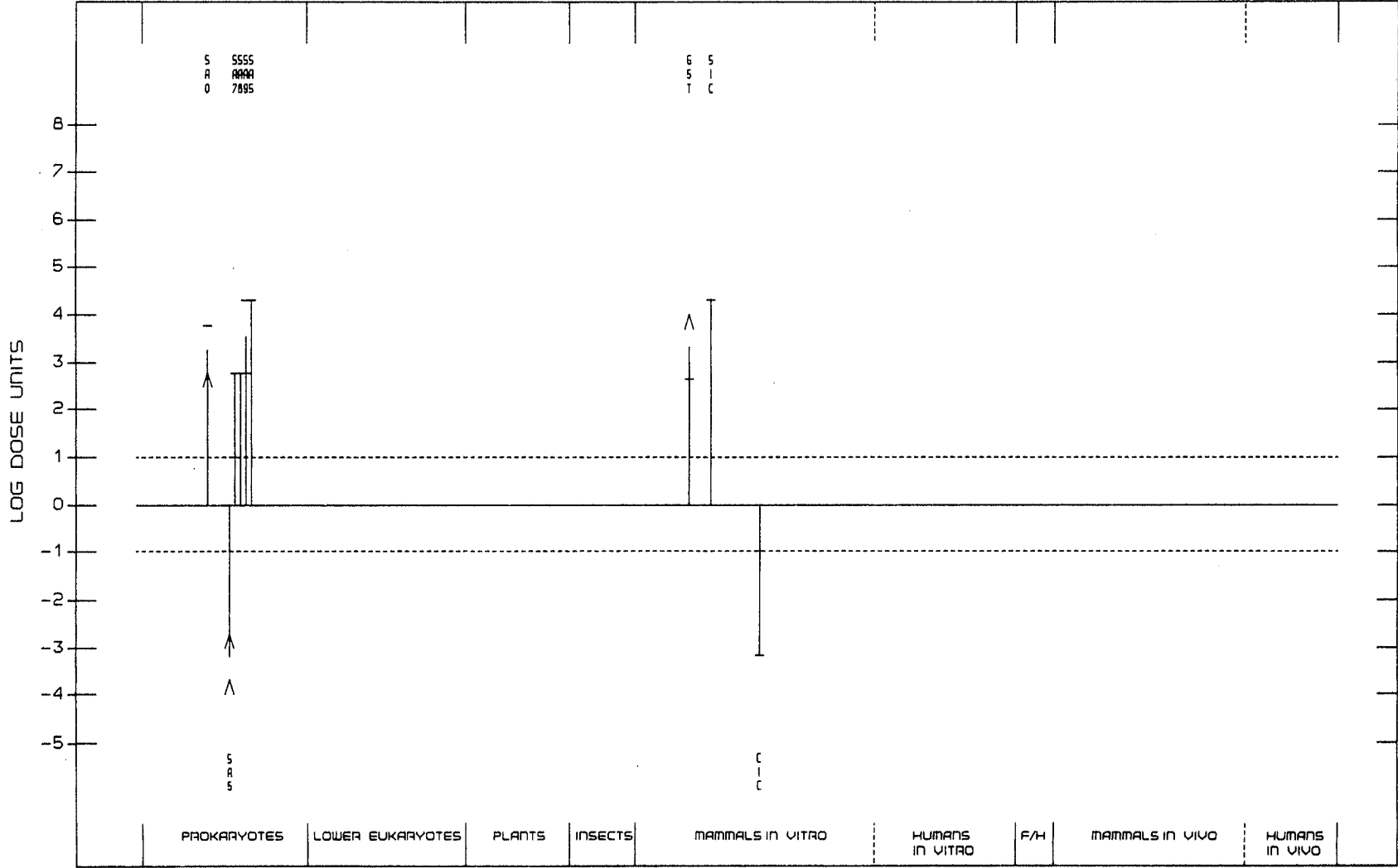


## DISPERSE YELLOW 3

END POINT	TEST CODE	TEST SYSTEM	RESULTS		DOSE (LED OR HID)	REFERENCE
			NM	M		
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	+	167.0000	CAMERON ET AL., 1987
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	+	+	17.0000	ZEIGER ET AL., 1988
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	5000.0000	CAMERON ET AL., 1987
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	500.0000	ZEIGER ET AL., 1988
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	+	+	167.0000	CAMERON ET AL., 1987
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	+	+	167.0000	CAMERON ET AL., 1987
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	+	+	167.0000	CAMERON ET AL., 1987
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	+	+	5.0000	ZEIGER ET AL., 1988
G	SAS	S. TYPHIMURIUM (OTHER), REVERSE MUTATION	+	+	5.0000	ZEIGER ET AL., 1988
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	-	+	10.0000	MCGREGOR ET AL., 1988
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	(+)	-	229.0000	CAMERON ET AL., 1987
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	+	0	5.0000	TENNANT ET AL., 1987b
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	-	0	1500.0000	TENNANT ET AL., 1987b



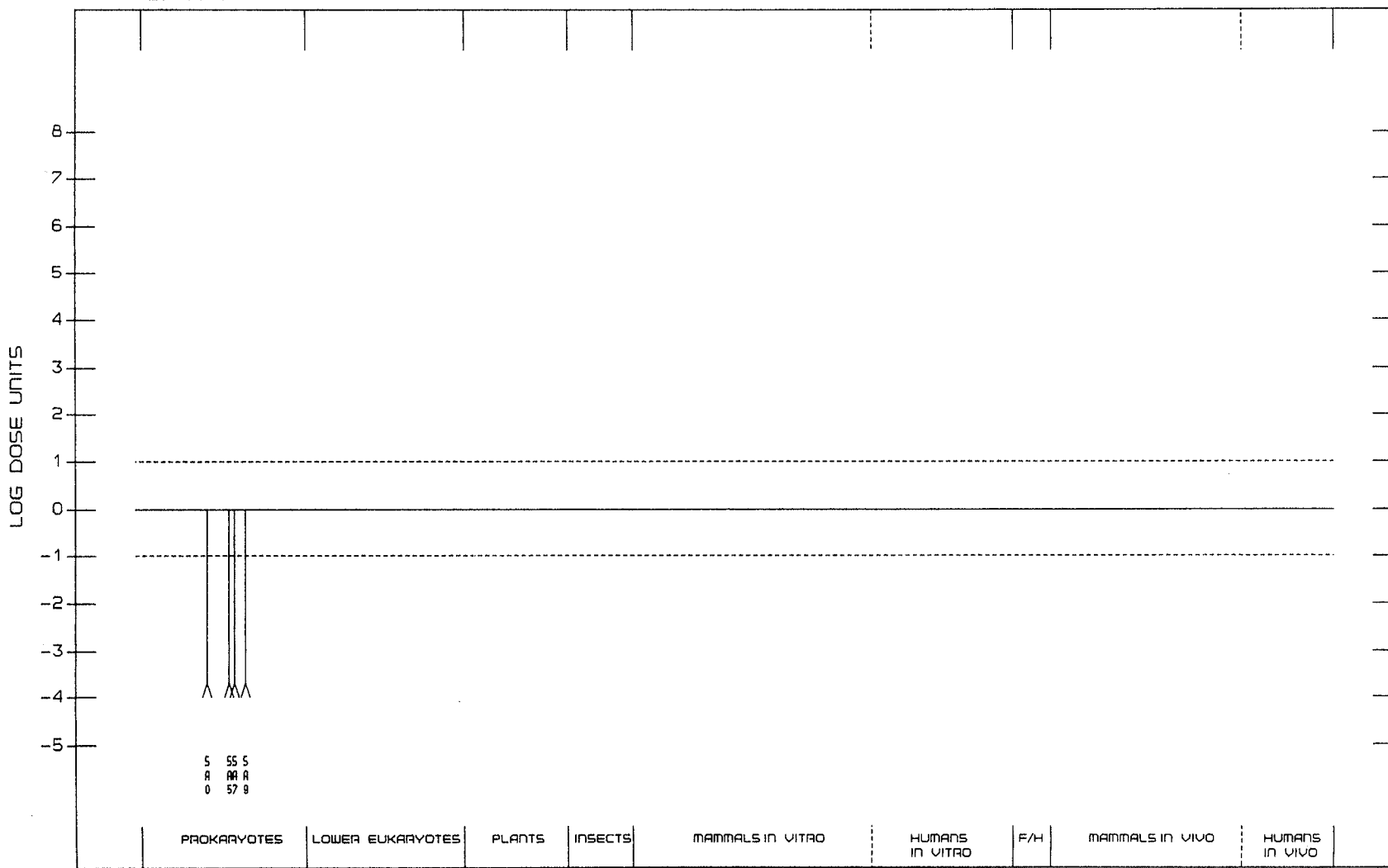
DISPERSE YELLOW 3  
2832-40-B



## VAT YELLOW 4

END POINT	TEST CODE	TEST SYSTEM	RESULTS		DOSE	REFERENCE
			NM	M (LED OR HID)		
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	5000.0000	ZEIGER ET AL., 1987
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	5000.0000	ZEIGER ET AL., 1987
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	5000.0000	ZEIGER ET AL., 1987
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	5000.0000	ZEIGER ET AL., 1987

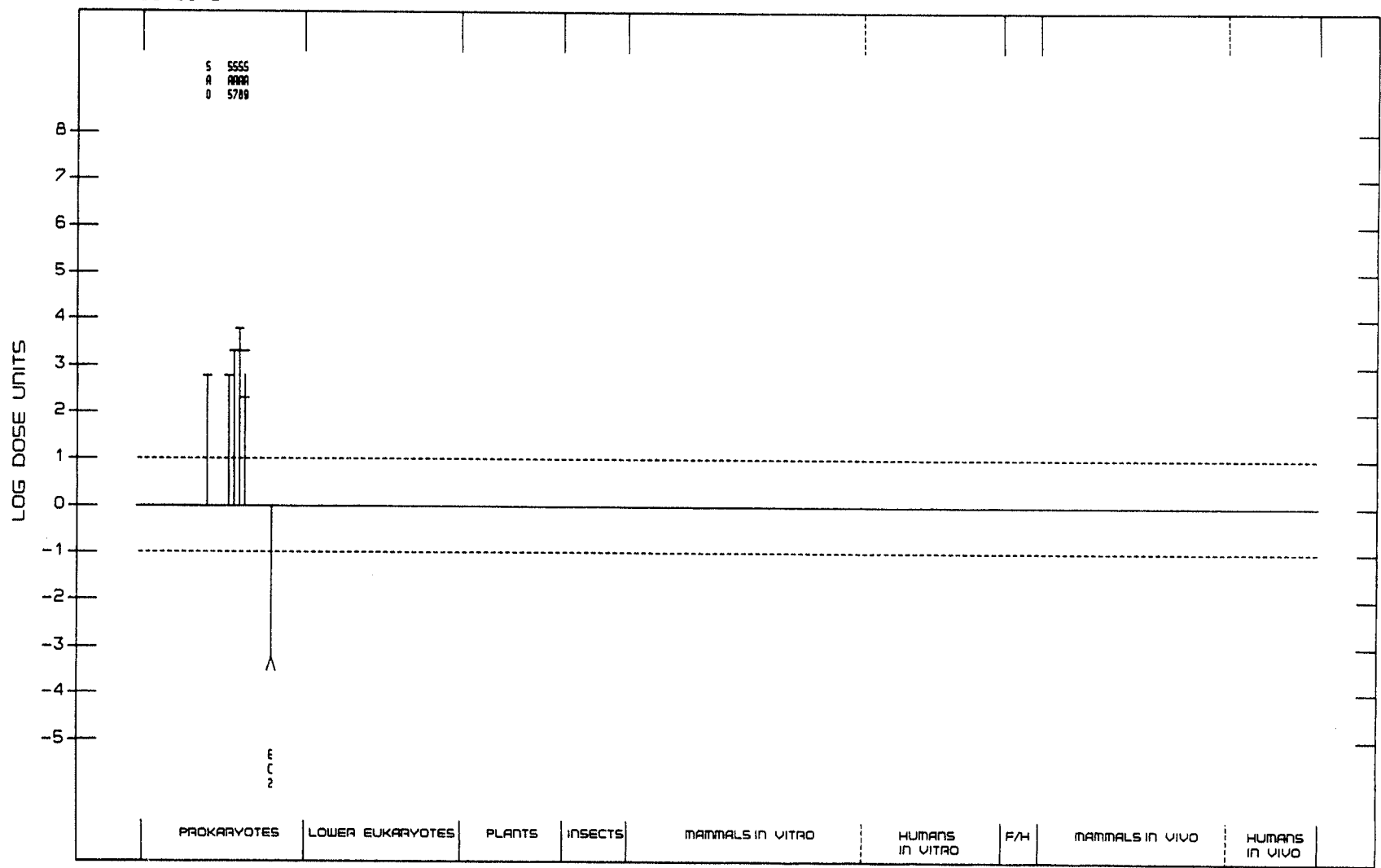
VAT YELLOW 4  
128-66-5



## 5-NITRO-O-TOLUIDINE

END POINT	TEST CODE	TEST SYSTEM	RESULTS		DOSE (LED OR HID)	REFERENCE
			NM	M		
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	+	+	0.0000	SPANGGORD ET AL., 1982b
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	+	+	167.0000	DUNKEL ET AL., 1985
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	+	-	0.0000	SPANGGORD ET AL., 1982b
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	+	+	167.0000	DUNKEL ET AL., 1985
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	+	+	0.0000	SPANGGORD ET AL., 1982b
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	+	+	50.0000	DUNKEL ET AL., 1985
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	+	+	0.0000	SPANGGORD ET AL., 1982b
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	+	+	17.0000	DUNKEL ET AL., 1985
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	+	+	0.0000	SPANGGORD ET AL., 1982b
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	+	+	50.0000	DUNKEL ET AL., 1985
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	+	+	500.0000	COUCH ET AL., 1987
G	EC2	E. COLI WP2, REVERSE MUTATION	-	-	1667.0000	DUNKEL ET AL., 1985

5-NITRO-O-TOLUIDINE  
99-55-8

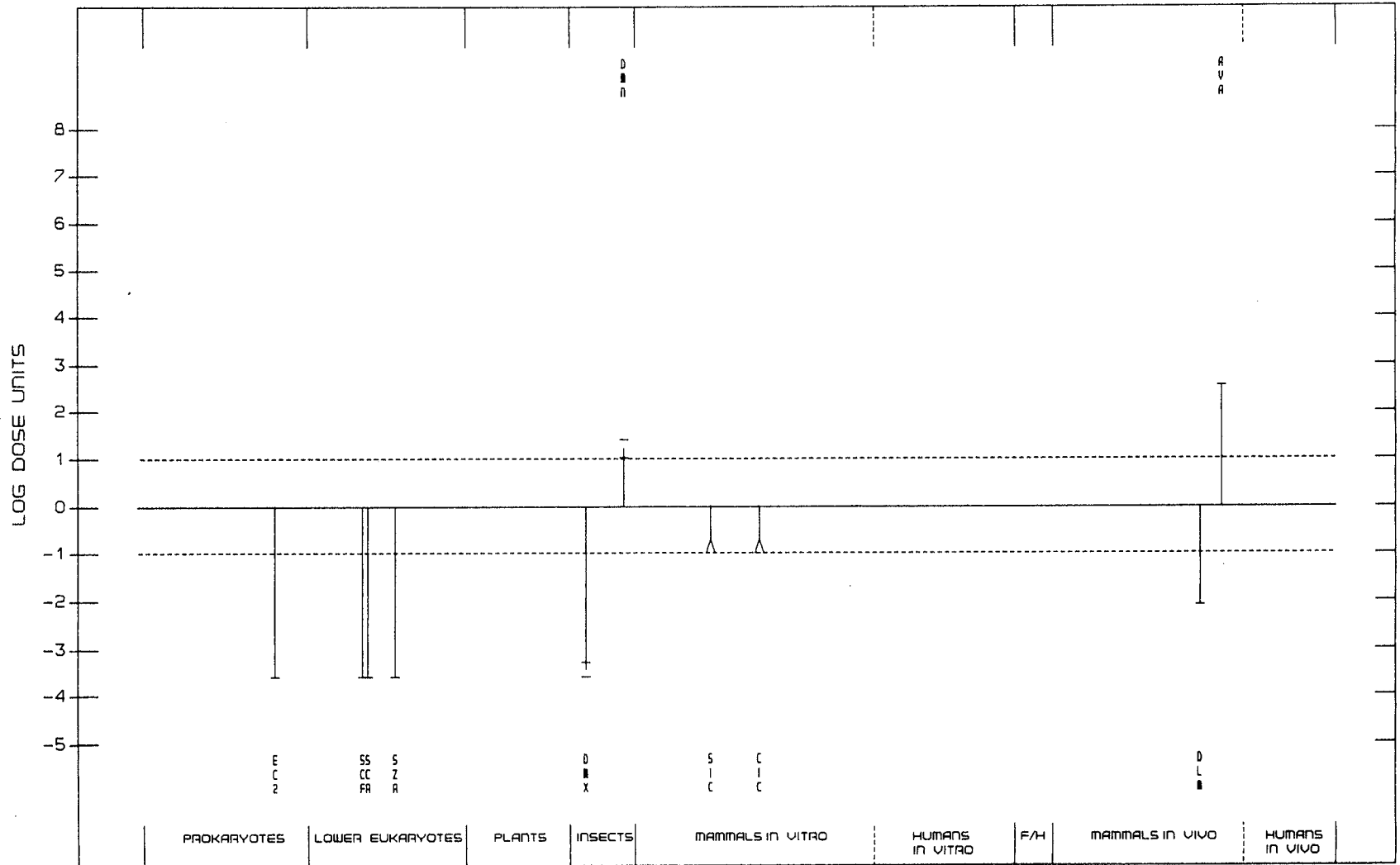


## NITRILOTRIACETIC ACID

END POINT	TEST CODE	TEST SYSTEM	RESULTS NM M	DOSE (LED OR HID)	REFERENCE
G	EC2	E. COLI WP2, REVERSE MUTATION	- 0	4000.0000	ZETTERBERG, 1970
G	SCF	S. CEREVISIAE, FORWARD MUTATION	- 0	4000.0000	ZETTERBERG, 1970
G	SCR	S. CEREVISIAE, REVERSE MUTATION	- 0	4000.0000	ZETTERBERG, 1970
G	SZR	S. POMBE, REVERSE MUTATION	- 0	4000.0000	ZETTERBERG, 1970
G	DMX	D. MELANOGASTER, SEX-LINKED RECESSIVES	- 0	1911.0000	KRAMERS, 1976
G	DMX	D. MELANOGASTER, SEX-LINKED RECESSIVES	- 0	4000.0000	WOODRUFF ET AL., 1985
C	DML	D. MELANOGASTER, DOMINANT LETHALS	- 0	0.0000	KRAMERS, 1976
A	DMN	D. MELANOGASTER, ANEUPLOIDY	+ 0	9557.0000	COSTA ET AL., 1988a
A	DMN	D. MELANOGASTER, ANEUPLOIDY	+ 0	4000.0000	RAMEL & MAGNUSSON, 1979
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	- -	5.0000	LOVEDAY ET AL., 1989
C	CIC	CHROM ABERR, CHINESE HAMSTER CELLS IN VITRO	- -	5.0000	LOVEDAY ET AL., 1989
C	DLM	DOMINANT LETHAL TEST, MICE	- 0	125.0000	EPSTEIN ET AL., 1972
A	AVA	ANEUPLOIDY, ANIMAL CELLS IN VIVO	+ 0	275.0000	COSTA ET AL., 1988a

NITRILOTRIACETIC ACID

139-13-9



END POINT	TEST CODE	TEST SYSTEM	RESULTS		DOSE (LED OR HID)	REFERENCE
			NM	M		
D	PRB	PROPHAGE, INDUCT/SOS/STRAND BREAKS/X-LINKS	-	-	0.0000	VENIER ET AL., 1987
D	ERD	E. COLI REC, DIFFERENTIAL TOXICITY	+	+	250.0000	VENIER ET AL., 1987
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	435.0000	LOPRIENO ET AL., 1985
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	5000.0000	DUNKEL ET AL., 1985
G	SA0	S. TYPHIMURIUM TA100, REVERSE MUTATION	-	-	435.0000	VENIER ET AL., 1987
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	435.0000	LOPRIENO ET AL., 1985
G	SA5	S. TYPHIMURIUM TA1535, REVERSE MUTATION	-	-	5000.0000	DUNKEL ET AL., 1985
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	435.0000	LOPRIENO ET AL., 1985
G	SA7	S. TYPHIMURIUM TA1537, REVERSE MUTATION	-	-	5000.0000	DUNKEL ET AL., 1985
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	-	-	435.0000	LOPRIENO ET AL., 1985
G	SA8	S. TYPHIMURIUM TA1538, REVERSE MUTATION	-	-	5000.0000	DUNKEL ET AL., 1985
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	435.0000	LOPRIENO ET AL., 1985
G	SA9	S. TYPHIMURIUM TA98, REVERSE MUTATION	-	-	5000.0000	DUNKEL ET AL., 1985
G	ECW	E. COLI WP2 UVRA, REVERSE MUTATION	-	-	100000.0000	VENIER ET AL., 1987
R	SCG	S. CEREVISIAE, GENE CONVERSION	-	-	40.0000	LOPRIENO ET AL., 1985
R	ANG	A. NIDULANS, CROSSING-OVER	-	0	10930.0000	CREBELLI ET AL., 1986
G	SZF	S. POMBE, FORWARD MUTATION	-	-	40.0000	LOPRIENO ET AL., 1985
G	ANF	A. NIDULANS, FORWARD MUTATION	-	0	18510.0000	CREBELLI ET AL., 1986
A	ANN	A. NIDULANS, ANEUPLOIDY	-	0	10930.0000	CREBELLI ET AL., 1986
M	PLI	PLANTS (OTHER), MICRONUCLEI	+	0	550.0000	DE MARCO ET AL., 1986
C	VFC	VICIA FABAE, CHROM ABERR	+	0	1375.0000	KIHLMAN & STURELID, 1970
D	URP	UDS, RAT PRIMARY HEPATOCYTES	-	0	1000.0000	WILLIAMS ET AL., 1982
G	G9H	MUTATION, CHL V79 CELLS, HPRT	-	0	1.5000	CELOTTI ET AL., 1987
G	G5T	MUTATION, L5178Y CELLS, TK LOCUS	-	-	2350.0000	MITCHELL ET AL., 1985
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	-	0	1.9000	LOPRIENO ET AL., 1985
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	-	0	1.0000	VENIER ET AL., 1985
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	-	0	275.0000	VED BRAT & WILLIAMS, 1984
S	SIC	SCE, CHINESE HAMSTER CELLS IN VITRO	-	0	514.0000	MONTALDI ET AL., 1985
S	SIM	SCE, MOUSE CELLS IN VITRO	-	0	257.0000	MONTALDI ET AL., 1985



C	CIA	CHROM ABERR, OTHER ANIMAL CELLS IN VITRO	+	0	688.0000	KIHLMAN & STURELID, 1970
G	GIH	MUTATION, HUMAN CELLS IN VITRO	+	0	3.0000	GRILLI & CAPUCCI, 1985
S	SHL	SCE, HUMAN LYMPHOCYTES IN VITRO	-	0	275.0000	VED BRAT & WILLIAMS, 1984
C	CHL	CHROM ABERR, HUMAN LYMPHOCYTES IN VITRO	-	0	2063.0000	MONTALDI ET AL., 1988
M	MVM	MICRONUCLEUS TEST, MICE IN VIVO	-	0	400.0000	MONTALDI ET AL., 1988

NITRILOTRIACETIC ACID 3NA SALT  
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