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PMRA Submission #	:{}	EPA MRID#: 45831104
Data Requirement:	PMRA Data Code: EPA DP Barcode: OECD Data Point: EPA MRID: EPA Guideline:	\\ D288160 \{
Test material: Common name: Chemical name:	Penoxsulam metabolite 5-Hydroxy-XDE-638 IUPAC: Not reported CAS name: Not reported CAS No.: Not reported Synonyms: Not reported	Purity: 100%
Primary Reviewer: E Staff Scientist, Dynan	•	Signature: Rhecca Pryan
QC Reviewer: Dana Staff Scientist, Dynan	Worcester	Signature: Mecca Prycon Date: 11/21/03 Signature: Dana Worcester Date: 11/21/03
Primary Reviewer: ‡ {EPA/OECD/PMRA	OHITERICKSON SOUDYEA	Date: Moodylon
Secondary Reviewer {EPA/OECD/PMRA	(s):{	Date: {}
EPA PC Code 199	[For PMRA] [For PMRA] 031 1903 ppleted: {dd-mmm-yyyy}	

CITATION: Hoberg, J.R. 2002. 5-Hydroxy-XDE-638 - Toxicity to Duckweed, *Lemna gibba*. Unpublished study performed by Springborn Laboratories, Inc., Wareham, Massachusetts. Laboratory Project Identification No. 12550.6167/Project No. 011234. Study submitted by The Dow Chemical Company for Dow AgroSciences, LLC Midland, Michigan. Experimental start date December 21, 2001 and experimental termination date January 7, 2002. The final report issued February 8, 2002.



EXECUTIVE SUMMARY:

In a 14-day acute toxicity study, freshwater aquatic vascular plants Duckweed, *Lemna gibba* G3, were exposed to Penoxsulam metabolite (5-Hydroxy-XDE-638) at mean measured concentrations <0.013-0.016 (<LOQ, negative and solvent controls), 0.081, 0.22, 0.62, 1.6, 4.6, and 11 mg/L under static conditions. Nominal concentrations were 0 (negative and solvent controls), 0.10, 0.26, 0.64, 1.6, 4.0, and 10 mg/L. After 14 days The percent reductions for frond density were significant in the 0.62, 1.6, 4.6, and 11 mg/L treatment groups. The most sensitive variable was frond numbers. The percent reductions for growth rate and dry weight were not significant in any treatment group.

The NOAEC was 0.22 mg/L, LOAEC was 0.62 mg/L/L, EC₀₅ was 0.095 mg/L, and the EC₅₀ >11 mg/L. This toxicity study is scientifically sound and satisfies the guideline $\S123-2$ for an aquatic vascular plant study with *Lemna gibba*. The study is classified as Core.

Results Synopsis

Test Organism: Lemna gibba G3

Test Type: Static

Number of fronds:

NOAEC: 0.22 mg/L LOAEC: 0.62 mg/L

 EC_{05}/IC_{05} : 0.095 mg/L 95% C.I.: 0.0014-6.4 mg/L

 EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: 0.256±0.109

Growth rates (day 7):

NOAEC: 11 mg/L LOAEC: >11 mg/L

 EC_{05}/IC_{05} : >11 mg/L 95% C.I.: N/A

 EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: N/A

Plant biomass (dry weight):

NOAEC: 11 mg/L

LOAEC: >11 mg/L

EC₀₅/IC₀₅: could not be determined 95% C.I.: N/A EC₅₀/IC₅₀: >11 mg/L 95% C.I.: N/A

Slope: N/A

Endpoint(s) Affected: Frond number

I. MATERIALS AND METHODS

GUIDELINE FOLLOWED: The test protocol was based on the following guidelines: OECD Proposed

Guideline 221 and U.S. EPA-FIFRA Pesticide Assessment Guidelines,

Subdivision J, Hazard Evaluation: Nontarget Plants Guidelines 122-2 and 123-2.

The following deviations from U.S. EPA Guideline 123-2 are noted:

1. The pretest health of the test organism was not reported.

The definitive test was conducted under static conditions and the test solution was not renewed as recommended.

These deviations do not affect the acceptability or the validity of the study.

COMPLIANCE:

Signed and dated GLP, Quality Assurance and No Data Confidentiality

statements were provided.

A. MATERIALS:

1. Test Material

Penoxsulam metabolite (5-Hydroxy-XDE-638)

Description:

Not reported

Lot No./Batch No.: F0512-129A

Purity:

>99% (used as 100%)

Stability of Compound

Under Test Conditions: Day 0 measured concentrations ranged from 96 to 110% of nominal concentrations and day 14 measured concentrations ranged from 56 to 125% of nominal concentrations. The mean measured concentrations were 81 to 110% of nominal.

(OECD requires water solubility, stability in water and light, pKa, Pow, vapor pressure of test compound) OECD requirements were not reported.

Storage conditions of test chemicals: Stored at room temperature in a dark ventilated cabinet.

2. Test organism:

Name: Duckweed, Lemna gibba (EPA requires a vascular species: Lemna gibba.)

Strain, if provided: G3

Source: Laboratory cultures (original supplier: University of Toronto, Toronto, Canada)

Age of inoculum: 2 days old

Method of cultivation: 20X Algal Assay Procedure (AAP) Medium

B. STUDY DESIGN:

a) Range-finding Study: No range-finding study was conducted.

b) Definitive Study

Table 1 . Experimental Parameters	Ţ	
		Remarks
Parameter	Details	Criteria
Acclimation period:	Continuous culture	
culturing media and conditions: (same as test or not)	20X Algal Assay Procedure (AAP) Medium; same as test.	
health: (any toxicity observed)	Not reported	
Test system static/static renewal/ renewal rate for static renewal:	Static	EPA expects the test concentrations to be renewed every 3 to 4 days (one renewal for the 7 day test, 3-4 renewals for the 14 day test).
Incubation facility	Environmental chamber	
Duration of the test	14 days	EPA requires a duration of 14 days. Seven day studies will be accepted for review by the Agency.
Test vessel material: (glass/polystyrene) size: fill volume:	Sterile crystallizing dishes 270 mL 100 mL	
Details of growth medium name: pH at test initiation: pH at test termination: Chelator used: Carbon source:	20X Algal Assay Procedure (AAP) Medium 7.6-7.8 (Table 2, p. 23) 8.3-8.9 Yes NaHCO ₃	EPA recommend the following culture media: Modified hoagland's E+ or 20X-AAP.
If non-standard nutrient medium was used, detailed composition provided (Yes/No)	Not applicable	
Dilution water source/type: pH: water pretreatment (if any): Total Organic Carbon:	Sterile deionized water 7.5 ± 0.1 pH adjusted using 0.1 N hydrochloric acid 0.47-1.0 mg/L (December 2001 and January 2002 analysis)	EPA recommends a pH of ~5.0. A solution pH of 7.5 is acceptable if type 20X-AAP nutrient media is used.

		Remarks
Parameter	Details	Criteria
particulate matter: metals: pesticides: chlorine:	N/A Not detected Not detected N/A	
Indicate how the test material is added to the medium (added directly or used stock solution)	Stock solution	
Aeration or agitation	Not reported.	
Sediment used (for rooted aquatic vascular plants) origin: textural classification (% sand, silt and clay): organic carbon (%): geographic location:	Not applicable	
Number of replicates control: solvent control: treatments:	3 3 3	
Number of plants/replicate	5 plants per replicate	EPA requires 5 plants.
Number of fronds/plant	3 fronds per plant (15 total fronds per replicate)	EPA requires 3 fronds per plant.
Test concentrations nominal: measured:	0 (negative and solvent controls), 0.10, 0.26, 0.64, 1.6, 4.0, and 10 mg/L <0.013-0.016 (<loq, 0.081,="" 0.22,="" 0.62,="" 1.6,="" 11="" 4.6,="" and="" controls),="" l<="" mg="" negative="" solvent="" td=""><td>EPA requires at least 5 test concentrations with a dose range of 2X or 3X progression.</td></loq,>	EPA requires at least 5 test concentrations with a dose range of 2X or 3X progression.
Solvent (type, percentage, if used)	Dimethylformamide (DMF), 0.10 mL/L	
Method and interval of analytical verification	HPLC; days 0 and 14.	
Test conditions temperature:	23-24°C	EPA temperature: 25 °C

		Remarks
Parameter	Details	Criteria
photoperiod:	continuous light	EPA photoperiod: continuous EPA light: 5.0 Klux (±15%)
light intensity and quality:	6400-6800 lux	
Reference chemical (if used) name: concentrations:	None	
Other parameters, if any	None	

2. Observations:

Table 2: Observation parameters

Parameters	Details	Remarks/Criteria	
Parameters measured (eg: number of fronds, plant dry weight or other toxicity symptoms)	Number of fronds, toxicity symptoms, and terminal dry weights.		
Measurement technique for frond number and other end points	Direct counts and weights.		
Observation intervals	Days 7 and 14.		
Other observations, if any	None		
Indicate whether there was an exponential growth in the control	Yes		
Were raw data included?	Replicate data provided.		

II. RESULTS and DISCUSSION:

A. INHIBITORY EFFECTS:

After 14 days, the mean frond number percent inhibitions compared to the pooled controls were 3, 3, 11, 9, 13, and 11% in the 0.081, 0.22, 0.62, 1.6, 4.6, and 11 mg/L treatment groups, respectively. The percent reductions for frond density were significant in the 0.62, 1.6, 4.6, and 11 mg/L treatment groups. The mean growth rate percent inhibitions compared to the pooled controls were -2, -2, 0, 2, -2, and 4% in the 0.081, 0.22, 0.62, 1.6, 4.6, and 11 mg/L treatment groups, respectively. The mean dry weight percent inhibitions compared to the pooled controls were 17, -15, -16, -5, -16, and 5% in the 0.081, 0.22, 0.62, 1.6, 4.6, and 11 mg/L treatment groups, respectively. The percent reductions for growth rate and dry weight were not significant in any treatment group.

Table 3: Effect of Penoxsulam metabolite (5-Hydroxy-XDE-638) on frond number and dry weight of Duckweed, Lemna gibba

Treatment ¹	Me	Mean frond number at			Mean Biomass	
(estimated measured and nominal concentration) mg/L	number/test solution	7 days	14 days	% inhibition at 14 days	Growth Rate (days ⁻¹)	(dry weights, g)
Negative control (dilution water)	15	338	806		0.44	0.1049
Solvent control	15	363	812		0.45	0.1298
0.081 (0.10)	15	375	781	3	0.46	0.0971
0.22 (0.26)	15	394	786	3	0.46	0.1350
0.62 (0.64)	15	358	720	11*	0.45	0.1364
1.6 (1.6)	15	331	733	9*	0.44	0.1234
4.6 (4.0)	15	376	707	13*	0.46	0.1361
11 (10)	15	305	719	11*	0.43	0.1112
Reference chemical (if used)	Not applicable					

¹ Nominal concentrations are in parentheses.

Table 4: Statistical endpoint values.

Statistical Endpoint ^a	frond No.	growth rate (day 7)	dry weight
NOAEC or EC ₀₅ (mg/L)	0.22	4.6	11
LOAEC (mg/L)	0.62	.62	
EC ₅₀ (mg/L) (95% C.I.)	>11	>11	>11
EC ₂₅ (mg/L) (95% C.I.)	>11	Not reported	>11
Reference chemical NOAEC IC ₅₀ /EC ₅₀	Not applicable	Not applicable	Not reported

^a Statistical data based on measured test concentrations.

B. REPORTED STATISTICS: A t-test was used to compare the dilution water (negative) and solvent controls. The controls were pooled for all statistical analyses. The data was analyzed for normality using the Shapiro-Wilk's Test and homogeneity of variance using Bartlett's Test. The Williams' test was used to compare the treatment groups to the pooled control. The NOAEC and LOAEC were determined from significance data. The EC₅₀ was empirically estimated to be greater than the highest concentration tested (no concentrations with >50% inhibition). The reported statistics were based on the mean measured test concentrations..

^{*} Significantly reduced compared to the pooled control (Williams' Test).

C. VERIFICATION OF STATISTICAL RESULTS:

Statistical method: Frond number, growth rate, and dry weight data satisfied the assumptions of ANOVA (i.e., normal distribution and variance homogeneity); the NOAEC and LOAEC values were determined using ANOVA (growth rate and dry weight), followed by Bonferroni's test (frond number) via TOXSTAT statistical software. For all endpoints, the solvent control was compared to the negative control using a Student's t-test and no difference was found, so the two were pooled for comparison to treatment. The EC_{05} for frond number was determined using the Probit method via Nuthatch statistical software. The EC_{05} could not be determined for dry weight because the response was not monotonic, so the Probit method could not be used; reductions in growth rate did not exceed 4%, so the EC_{05} for this endpoint was visually determined. Reductions did not exceed 50% for any endpoint, so the EC_{05} could be visually determined for all endpoints.

Number of fronds:

NOAEC: 0.22 mg/L LOAEC: 0.62 mg/L

EC₀₅/IC₀₅: 0.095 mg/L 95% C.I.: 0.0014-6.4 mg/L

 EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: 0.256±0.109

Growth rates (day 7):

NOAEC: 11 mg/L LOAEC: >11 mg/L

 EC_{05}/IC_{05} : >11 mg/L 95% C.I.: N/A EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: N/A

Plant biomass (dry weight):

NOAEC: 11 mg/L LOAEC: >11 mg/L

 EC_{05}/IC_{05} : could not be determined 95% C.I.: N/A EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: N/A

Endpoint(s) Affected: Frond number

D. STUDY DEFICIENCIES:

The deviations did not affect the acceptability or the validity of the study.

E. REVIEWER'S COMMENTS:

With the exception of the growth rate NOAEC, the reviewer's conclusions agreed with the study author's; the reviewer's analysis detected no effect on growth rate at any treatment level, while the study author's detected a significant reduction at the highest treatment level. No endpoints were reduced 50%, so the EC_{50} for this study is 11 mg/L. According to the reviewer's analysis, frond number was the only affected endpoint; the reviewer determined the EC_{05} value for this endpoint to be 0.095 mg/L.

The amount of test substance was limited, so further tests to determine EC₅₀ were not performed. The study author reported these test results define the toxicity of the metabolite relative to the parent compound.

The test was conducted according to U.S. EPA Good Laboratory Practice Regulations with the following exception: The data for routine water contaminant screening analysis was not collected in accordance to GLP procedures. A GLP statement was provided.

F. CONCLUSIONS: This toxicity study is scientifically sound and satisfies the U.S. EPA Guideline Subdivision J, §123-2 for an aquatic vascular plant study with *Lemna gibba*. As a result, this study is classified as Core.

Number of fronds:

NOAEC: 0.22 mg/L LOAEC: 0.62 mg/L

EC₀₅/IC₀₅: 0.095 mg/L 95% C.I.: 0.0014-6.4 mg/L

 EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: 0.256±0.109

Growth rates (day 7):

NOAEC: 11 mg/L LOAEC: >11 mg/L

 EC_{05}/IC_{05} : >11 mg/L 95% C.I.: N/A EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: N/A

Plant biomass (dry weight):

NOAEC: 11 mg/L LOAEC: >11 mg/L

 EC_{05}/IC_{05} : could not be determined 95% C.I.: N/A EC_{50}/IC_{50} : >11 mg/L 95% C.I.: N/A

Slope: N/A

Endpoint(s) Affected: Frond number

III. REFERENCES:

ASTM. 2000. Standard practice for conducting acute toxicity tests with fishes, macroinvertebrates, and amphibians. Standard E729-88a, American Society for Testing and Materials, 100 Barr Harbor Drive, West Conshohocken, Pennsylvania.

Horning, W.B. and C.I. Weber, 1985. Short-term methods for estimating the chronic toxicity of effluents and receiving waters to freshwater organisms. EPA/600/4-89/014. Environmental Monitoring and Support Laboratory, U.S. Environmental Protection Agency, Cincinnati, Ohio.

Hillman, W.S. 1961. The Lemnacea, or duckweeds. Bot. Rev. 27:221-287.

Miller, W.E., J.C. Green and T. Shiroyama. 1978. The *Selenastrum capricornutum* Printz algal assay bottle test. EPA 600/9-78-018. U.S. Environmental Protection Agency, Corvallis, Oregon.

OECD. 1997. Good Laboratory Practices as acknowledged in the EEC Council Directive 88/320/EEC of 9 June 1988.

- OECD. 2000. OECD Guideline for Testing of Chemicals. *Lemna* sp. Growth Inhibition Test. Proposed Guideline #221. Revised Draft, October 2000.
- Sokal, R.R. and F.J. Rohlf. 1981. Biometry. 2nd Edition. W.H. Freeman and Co. New York, NY. 859 pp.
- U.S. EPA. 1982. Pesticide Assessment Guidelines, Subdivision J, Hazard Evaluation: Nontarget Plants. EPA 540/9-82-020, 27 October 1982. U.S. EPA, Washington, D.C.
- U.S. EPA. Federal Insecticide, Fungicide and Rodenticide Act (FIFRA); Good Laboratory Practice Standards; Final Rule (40 CFR, Part 160). Federal Register, 48 (230); 34052-34074. U.S. Environmental Protection Agency, Washington, DC.
- Weber, C.I., W.H. Peltier, T.J. Norberg-King, W.B. Horning II, F.A. Kessier, J.R. Menkedick, T.W. Neiheisel, P.A. Lewis, D.J. Kiemm, Q.H. Pickering, E.L. Robinson, J.M. Lazorchak, L.J. Wymer and R.W. Freyberg (eds.). 1989. Short-term methods for estimating the chronic toxicity of effluents and receiving waters to freshwater organisms. 2nd ed. EPA/600/4/89/001. Environmental Monitoring Systems Laboratory, U.S. Environmental Protection Agency, Cincinnati, OH.
- Williams, D.A. 1971. A test for differences between treatment means when several dose levels are compared with a zero dose control. *Biometrics* 27: 103-117.

Williams, D.A. 1972. A comparison of several dose levels with a zero control. Biometrics 28: 519-531.

APPENDIX I. OUTPUT OF REVIEWER'S STATISTICAL RESULTS:

frond production

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SOURCE	DF	SS	MS	F
Between	6	38167.458	6361.243	4.814
Within (Error) 17	22465.500	1321.500	
Total	23	60632.958		

Critical F value = 2.70 (0.05,6,17) Since F > Critical F REJECT Ho:All groups equal

frond production

File: 1104f Transform: NO TRANSFORMATION

BONFERRONI T-TEST - TABLE 1 OF 2 Ho:Control<Treatment

TRANSFORMED MEAN CALCULATED IN
GROUP IDENTIFICATION MEAN ORIGINAL UNITS T STAT SIG

1 GRPS 1&2 POOLED 809.167 809.167

2 0.081 781.000 781.000 1.096

3	0.22	786.000	786.000	0.901
4	0.62	719.667	719.667	3.482 *
5	1.6	732.667	732.667	2.976 *
6	4.6	707.333	707.333	3.962 *
7	11	719.333	719.333	3.495 *

Bonferroni T table value = 2.65 (1 Tailed Value, P=0.05, df=17,6)

frond production

File: 1104f Transform: NO TRANSFORMATION

BONFERRONI T-TEST - TABLE 2 OF 2 Ho:Control<Treatment

NUM OF Minimum Sig Diff % of DIFFERENCE
GROUP IDENTIFICATION REPS (IN ORIG. UNITS) CONTROL FROM CONTROL

1 GRPS 1&2 POOLED 6
2 0.081 3 68.247 8.4 28.167

	UKFS INZ PC		U		
2	0.081	3	68.247	8.4	28.167
3	0.22	3	68.247	8.4	23.167
4	0.62	3	68.247	8.4	89.500
5	1.6	3	68.247	8.4	76.500
6	4.6	3	68.247	8.4	101.833
7	11	3	68.247	8.4	89.833

frond production

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WILLIAMS TEST (Isotonic regression model) TABLE 1 OF 2

GROL	JP		ORIGI	NAL	TRANS	FORMED	ISOTONIZED
	IDENTIFICATI	ON	N	MEAN	1	MEAN	MEAN
1	GRPS 1&2	P00	LED 6	809.1	67	809.167	809.167
2	0.081	3	781.00	00 7	781.000	783.50	0
3	0.22	3	786.00	0 7	86.000	783.500)
4	0.62	3	719.66	7 7	19.667	726,167	7
5	1.6	3	732.667	7 73	32.667	726.167	
6	4.6	3	707.333	3 70	7.333	713.333	
7	11	3	719.333	7	19.333	713.333	

frond production

File: 1104f Transform: NO TRANSFORMATION

WILLIAMS TEST (Isotonic regression model) TABLE 2 OF 2

ISOTONIZED CALC. SIG TABLE DEGREES OF IDENTIFICATION MEAN WILLIAMS P=.05 WILLIAMS FREEDOM

GRP	S 1&2 P	OOLED 8	09.167			
	0.081	783.500	0.999		1.74	k= 1, v=17
	0.22	783.500	0.999		1.82	k= 2, v=17
	0.62	726.167	3.229	*	1.85	k= 3, v=17
	1.6	726.167	3.229	*	1.87	k= 4, v=17
	4.6	713.333	3.728	*	1.87	k= 5, V=17
	11	713.333	3.728	*	1.88	k= 6, V=17

s = 36.352

Note: df used for table values are approximate when v > 20.

Estimates of EC%

Parameter Estimate 95% Bounds Std.Err. Lower Bound Lower Upper /Estimate EC5 0.095 0.0014 6.4 0.88 0.015

EC10 2.5 0.26 24. 0.47 0.10 EC25 5.8E+02 5.7 6.0E+04 0.97 0.0097 EC50 2.5E+05 15. 4.1E+09 2.0 6.1E-05

Slope = 0.256 Std.Err. = 0.109

Goodness of fit: p = 0.37 based on DF= 4.0 17.

1104F: frond production

Observed vs. Predicted Treatment Group Means

Dose #Reps. Obs. Pred. Obs. Pred. %Change Mean Mean -Pred. %Control 0.00 6.00 809. 811. -1.60 100. 0.00 0.0810 3.00 *7*81. *772*. 9.29 95.2 4.82 0.220 3.00 786. 762. 24.2 94.0 6.04 0.620 3.00 720. 750. -29.9 92.5 7.54 1.60 3.00 733. 736. -3.82 90.8 9.16 4.60 3.00 707. 720. -12,2 88.8 11.2 11.0 3.00 719. 704. 15.7 86.8

!!!Warning: EC25 not bracketed by doses evaluated.

!!!Warning: EC50 not bracketed by doses evaluated.

growth rate

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ANOVA TABLE

SOURCE	DF	SS	MS	F
Between	6	0.0026	0.0004	2.000

Within (Error) 17 0.0027 0.0002

Total 23 0.0053

Critical F value = 2.70 (0.05,6,17)

Since F < Critical F FAIL TO REJECT Ho:All groups equal

growth rate

File: 1104g Transform: NO TRANSFORMATION

BONFERRONI T-TEST - TABLE 1 OF 2 Ho:Control<Treatment

TRANSFORMED MEAN CALCULATED IN GROUP IDENTIFICATION MEAN ORIGINAL UNITS T STAT SIG GRPS 1&2 POOLED 0.448 0.448 0.081 0.457 -0.833 0.457 2 3 0.22 0.467 0.467 -1.833 4 0.450 0.450 0.62 -0.1675 1.6 0.440 0.440 0.833 4.6 0.457 6 0.457 -0.83311 0.430 0.430 1.833

Bonferroni T table value = 2.65 (1 Tailed Value, P=0.05, df=17,6)

growth rate

File: 1104g Transform: NO TRANSFORMATION

BONFERRONI T-TEST - TABLE 2 OF 2 Ho:Control<Treatment

NUM OF Minimum Sig Diff % of DIFFERENCE GROUP IDENTIFICATION REPS (IN ORIG. UNITS) CONTROL FROM CONTROL

1	GRPS 1&2 POOLED	6			
2	0.081 3	0.027	5.9	-0.008	
3	0.22 3	0.027	5.9	-0.018	
4	0.62 3	0.027	5.9	-0.002	
5	1.6 3	0.027	5.9	0.008	
6	4.6 3	0.027	5.9	-0.008	
7	11 3	0.027	5.9	0.018	

growth rate

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WILLIAMS TEST (Isotonic regression model) TABLE 1 OF 2

GROUP ORIGINAL TRANSFORMED ISOTONIZED

Παγε 13 οφ 16

	IDENTIFIC	ATI	ON	N	MEAN	MEAN	MEAN
1	GRPS 1	&2 F	2001	ED 6	0.448	0.448	0.455
2	0.	.081	3	0.45	7 0.457	0.455	
3	C	.22	3	0.467	0.467	0.455	
4	C	.62	3	0.450	0.450	0.450	
5	•	1.6	3	0.440	0.440	0.448	
6	4	4.6	3	0.457	0.457	0.448	
7		11	3	0.430	0.430	0.430	

growth rate

File: 1104g Transform: NO TRANSFORMATION

WILLIAMS TEST (Isotonic regression model) TABLE 2 OF 2

	IS	OTONIZE	D CALC	c. s	IG -	TABLE	DEGREES	OF
	IDENTIFICATI	ON M	IEAN	WILL	IAMS	P=.05	WILLIAMS	FREEDOM
٠								
	GRPS 1&2 P	OOLED	0.455					
	0.081	0.455	0.745		1.74	4 k=	: 1, V=17	
	0.22	0.455	0.745		1.82	! k =	2, v=17	
	0.62	0.450	0.186		1.85	k=	3, v=17	
	1.6	0.448	0.000		1.87	k= 4	4, v=17	
	4.6	0.448	0.000		1.87	k= !	5, v=17	
	11	0.430	2.050	*	1.88	k=	6, v=17	

s = 0.013

Note: df used for table values are approximate when v > 20.

dry weight

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ANOVA TABLE

 SOURCE
 DF
 SS
 MS
 F

 Between
 6
 4066.492
 677.749
 1.489

 Within (Error)
 17
 7736.922
 455.113

 Total
 23
 11803.413

Critical F value = 2.70 (0.05,6,17)

Since F < Critical F FAIL TO REJECT Ho:All groups equal

dry weight

File: 1104d Transform: NO TRANSFORMATION

BONFERRONI T-TEST - TABLE 1 OF 2 Ho:Control<Treatment TRANSFORMED MEAN CALCULATED IN GROUP IDENTIFICATION MEAN ORIGINAL UNITS T STAT SIG GRPS 1&2 POOLED 117.350 117.350 0.081 97.100 97.100 1.342 2 0.22 135.000 135.000 -1.170 3 0.62 136.367 Δ 136.367 -1.261 1.6 123*.*400 123.400 5 -0.401 -1.243 4.6 136.100 136.100 6 0.408 7 11 111.200 111.200

Bonferroni T table value = 2.65 (1 Tailed Value, P=0.05, df=17,6)

dry weight

File: 1104d Transform: NO TRANSFORMATION

BONFERRONI T-TEST - TABLE 2 OF 2 Ho:Control<Treatment

NUM OF Minimum Sig Diff % of DIFFERENCE
GROUP IDENTIFICATION REPS (IN ORIG. UNITS) CONTROL FROM CONTROL

1 GRPS 1&2 POOLED 6
2 0.081 3 40.051 34.1 20.250
3 0.22 3 40.051 34.1 -17.650
4 0.62 3 40.051 34.1 -19.017

3 0.22 3 40.051 34.1 -17.650 4 0.62 3 40.051 34.1 -19.017 5 1.6 3 40.051 34.1 -6.050 6 4.6 3 40.051 34.1 -18.750 7 11 3 40.051 34.1 6.150

dry weight

File: 1104d Transform: NO TRANSFORMATION

WILLIAMS TEST (Isotonic regression model) TABLE 1 OF 2

1 GRPS 1&2 POOLED 6 117.350 123.238 2 0.081 3 97.100 97.100 123.238 3 0.22 3 135.000 135.000 123.238 4 0.62 3 136.367 136.367 123.238 5 1.6 3 123.400 123.400 123.238 6 4.6 3 136.100 136.100 123.238 7 11 3 111.200 111.200 111.200	GROU	JP IDENTIFICATION	ORIGINA N M			SOTONIZED MEAN
	2 3 4 5 6	0.081 3 0.22 3 0.62 3 1.6 3 4.6 3	97.100 135.000 136.367 123.400	97.100 135.000 136.367 123.400	123.238 123.238 123.238 123.238	

dry weight

File: 1104d Transform: NO TRANSFORMATION

WILLIAMS TEST (Isotonic regression model) TABLE 2 OF 2

						•
IS	OTONIZED	CALC	. SIG 1	TABLE	DEGREES	OF
IDENTIFICAT	ION ME	EAN 1	WILLIAMS	P=.05	WILLIAMS	FREEDOM
•••••						
GRPS 1&2 P	OOLED 1	23.238				
0.081	123.238	0.390	1.7	74 k	= 1, V=17	
0.22	123.238	0.390	1.8	2 k=	= 2, V=17	
0.62	123.238	0.390	1.8	5 k=	= 3, V=17	
1.6	123.238	0.390	1.87	7 k=	4, v=17	
4.6	123.238	0.390	1.87	7 k≃	5, V=17	
11	111.200	0.408	1.88		6, v=17	

S = 21.333

Note: df used for table values are approximate when v > 20.

EC,

!!!Failure #3: Data not suitable for probit model fit.

Criterion is 3 or more distinct isotone means.