Experiment Number: A61460

Test Type: Genetic Toxicology - Micronucleus

Route: Gavage

Species/Strain: Mouse/B6C3F1

G04: In Vivo Micronucleus Summary Data

Test Compound: Glycoluril

Date Report Requested: 09/20/2018 Time Report Requested: 22:24:15

CAS Number: 496-46-8

NTP Study Number: A61460

72 Hours **Study Duration:**

Study Methodology: Slide Scoring

Male Study Result: Negative **G04: In Vivo Micronucleus Summary Data**

Test Compound: Glycoluril

CAS Number: **496-46-8**

Date Report Requested: 09/20/2018
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Route: Gavage

Species/Strain: Mouse/B6C3F1

Test Type: Genetic Toxicology - Micronucleus

Experiment Number: A61460

Tissue: Bone marrow; Sex: Male; Number of Treatments: 3; Time interval between final treatment and cell sampling: 72 h

		MN PCE/1000		% PCE
Dose (mg/kg)	N	Mean ± SEM	p-Value	Mean ± SEM
Vehicle Control ¹	5	1.90 ± 0.62		55.60 ± 4.25
500.0	5	2.40 ± 0.70	0.3034	60.60 ± 5.84
1000.0	5	1.70 ± 0.85	0.5890	62.40 ± 1.94
2000.0	5	1.60 ± 0.29	0.6339	54.70 ± 5.72
Trend p-Value		0.7180		
Positive Control ²	5	25.00 ± 1.56	< 0.001 *	51.60 ± 3.45
Trial Summary: Negative				

G04: In Vivo Micronucleus Summary Data

Test Compound: Glycoluril CAS Number: 496-46-8

Date Report Requested: 09/20/2018
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Route: Gavage

Species/Strain: Mouse/B6C3F1

Experiment Number: A61460

LEGEND

Test Type: Genetic Toxicology - Micronucleus

MN = micronucleated, PCE = polychromatic erythrocyte, NCE = normochromatic erythrocyte

CAS Number = Chemical Abstracts Service registry number

N = Number of subjects

Values given as Mean or Mean ± Standard Error Mean

Results were tabulated as the mean of the pooled results from all animals within a treatment group, plus or minus the standard error of the mean

Pairwise comparison to the concurrent control, dosed groups significant at p = 0.025/number of treatment groups; positive control value is significant at p = 0.05

Cochran-Armitage trend test, significant at p = 0.025

- * Statistically significant pairwise or trend test
- 1: Vehicle Control: Hcl
- 2: 20.0 mg/kg Cyclophosphamide

** END OF REPORT **