

Experiment Number: 05049 - 03
Test Type: 14-DAY
Route: DOSED WATER
Species/Strain: RATS/HSD

P10: STATISTICAL ANALYSIS OF NON-NEOPLASTIC LESIONS
Ionic liquid Toxicity
CAS Number: IONICLIQUIDS

Date Report Requested: 04/24/2020
Time Report Requested: 12:10:10
First Dose M/F: 03/09/09 / 03/10/09
Lab: MBA

NTP Study Number:	C05049		
Lock Date:	12/04/2009		
Cage Range:	ALL		
Date Range:	ALL		
Reasons For Removal:	ALL		
Removal Date Range:	ALL		
Treatment Groups:	Include 001 Control Male	Include 012 1.0 mg/ml NBuPy	Include 013 3.0 mg/ml NBuPy
	Include 014 6.0 mg/ml NBuPy	Include 015 Control Female	Include 026 1.0 mg/ml NBuPy
	Include 027 3.0 mg/ml NBuPy	Include 028 6.0 mg/ml NBuPy	
Study Gender:	Both		
TDMSE Version:	3.0.2.3_002		
PWG Approval Date:	NONE		

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SUMMARY OF STATISTICALLY SIGNIFICANT ($P \leq .05$) RESULTS IN THE ANALYSIS OF IONIC LIQUID TOXICITY

MALE RATS

Organ

Liver

FEMALE RATS

Organ

Thymus

Morphology

Depletion Glycogen

Morphology

Necrosis

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**STATISTICAL ANALYSIS OF NON-NEOPLASTIC LESIONS IN RATS(HSD)
TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Males			
	Control Male	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Heart				
Cardiomyopathy				
LESION RATES				
OVERALL (a)	8/10 (80%)	0/0 (0%)	0/0 (0%)	4/5 (80%)
POLY-3 RATE (b)	8/10.00	0/0.00	0/0.00	4/5.00
POLY-3 PERCENT (g)	80%	0%	0%	80%
TERMINAL (d)	8/10 (80%)	0/0 (0%)	0/0 (0%)	4/5 (80%)
FIRST INCIDENCE	15 (T)	---	---	15 (T)
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.738
POLY 1.5	(e)	(e)	(e)	P=0.738
POLY 6	(e)	(e)	(e)	P=0.738
COCH-ARM / FISHERS	P=0.634	(e)	(e)	P=0.736N
MAX-ISO-POLY-3	(e)	(e)	(e)	P=1.000

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TERMINAL SACRIFICE AT 3 WEEKS

DOSE	Males			
	Control Male	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Liver				
Depletion Glycogen				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/5 (0%)	1/5 (20%)	4/5 (80%)
POLY-3 RATE (b)	0/10.00	0/5.00	1/5.00	4/5.00
POLY-3 PERCENT (g)	0%	0%	20%	80%
TERMINAL (d)	0/10 (0%)	0/5 (0%)	1/5 (20%)	4/5 (80%)
FIRST INCIDENCE	---	---	15 (T)	15 (T)
STATISTICAL TESTS				
POLY 3	P<0.001**	(e)	P=0.356	P<0.001**
POLY 1.5	P<0.001**	(e)	P=0.356	P<0.001**
POLY 6	P<0.001**	(e)	P=0.356	P<0.001**
COCH-ARM / FISHERS	P<0.001**	(e)	P=0.333	P=0.004**
MAX-ISO-POLY-3	P<0.001**	(e)	P=0.101	P<0.001**

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Males			
	Control Male	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Lung				
Metaplasia Osseous				
<hr/>				
LESION RATES				
OVERALL (a)	1/10 (10%)	0/0 (0%)	0/0 (0%)	0/5 (0%)
POLY-3 RATE (b)	1/10.00	0/0.00	0/0.00	0/5.00
POLY-3 PERCENT (g)	10%	0%	0%	0%
TERMINAL (d)	1/10 (10%)	0/0 (0%)	0/0 (0%)	0/5 (0%)
FIRST INCIDENCE	15 (T)	---	---	---
<hr/>				
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.636N
POLY 1.5	(e)	(e)	(e)	P=0.636N
POLY 6	(e)	(e)	(e)	P=0.636N
COCH-ARM / FISHERS	P=0.427N	(e)	(e)	P=0.667N
MAX-ISO-POLY-3	(e)	(e)	(e)	P=0.274N

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Males			
	Control Male	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Spleen				
Depletion Cellular				
<hr/>				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/0 (0%)	0/5 (0%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/0.00	0/5.00
POLY-3 PERCENT (g)	0%	0%	0%	0%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/0 (0%)	0/5 (0%)
FIRST INCIDENCE	---	---	---	---
<hr/>				
STATISTICAL TESTS				
POLY 3	(n)	(n)	(n)	(n)
POLY 1.5	(n)	(n)	(n)	(n)
POLY 6	(n)	(n)	(n)	(n)
COCH-ARM / FISHERS	(n)	(n)	(n)	(n)
MAX-ISO-POLY-3	(n)	(n)	(n)	(n)

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Males			
	Control Male	1.0 mg/ml NBuPy	3.0 mg/ml NBuPy	6.0 mg/ml NBuPy
Thymus Atrophy				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/5 (0%)	0/5 (0%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/5.00	0/5.00
POLY-3 PERCENT (g)	0%	0%	0%	0%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/5 (0%)	0/5 (0%)
FIRST INCIDENCE	---	---	---	---
STATISTICAL TESTS				
POLY 3	(n)	(n)	(n)	(n)
POLY 1.5	(n)	(n)	(n)	(n)
POLY 6	(n)	(n)	(n)	(n)
COCH-ARM / FISHERS	(n)	(n)	(n)	(n)
MAX-ISO-POLY-3	(n)	(n)	(n)	(n)

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Males			
	Control Male	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Thymus Necrosis				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/5.00	1/5.00
POLY-3 PERCENT (g)	0%	0%	0%	20%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
FIRST INCIDENCE	---	---	---	15 (T)
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.356
POLY 1.5	(e)	(e)	(e)	P=0.356
POLY 6	(e)	(e)	(e)	P=0.356
COCH-ARM / FISHERS	P=0.177	(e)	(e)	P=0.333
MAX-ISO-POLY-3	(e)	(e)	(e)	P=0.101

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Females			
	Control Female	1.0 mg/ml NBuPy	3.0 mg/ml NBuPy	6.0 mg/ml NBuPy
Heart				
Cardiomyopathy				
LESION RATES				
OVERALL (a)	10/10 (100%)	0/0 (0%)	0/0 (0%)	5/5 (100%)
POLY-3 RATE (b)	10/10.00	0/0.00	0/0.00	5/5.00
POLY-3 PERCENT (g)	100%	0%	0%	100%
TERMINAL (d)	10/10 (100%)	0/0 (0%)	0/0 (0%)	5/5 (100%)
FIRST INCIDENCE	15 (T)	---	---	15 (T)
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	(e)
POLY 1.5	(e)	(e)	(e)	(e)
POLY 6	(e)	(e)	(e)	(e)
COCH-ARM / FISHERS	(e)	(e)	(e)	(e)
MAX-ISO-POLY-3	(e)	(e)	(e)	(e)

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DOSE	Females			
	Control Female	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
<hr/>				
Liver				
Depletion Glycogen				
<hr/>				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/5.00	1/5.00
POLY-3 PERCENT (g)	0%	0%	0%	20%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
FIRST INCIDENCE	---	---	---	15 (T)
<hr/>				
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.356
POLY 1.5	(e)	(e)	(e)	P=0.356
POLY 6	(e)	(e)	(e)	P=0.356
COCH-ARM / FISHERS	P=0.177	(e)	(e)	P=0.333
MAX-ISO-POLY-3	(e)	(e)	(e)	P=0.100

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Females			
	Control Female	1.0 mg/ml NBuPy	3.0 mg/ml NBuPy	6.0 mg/ml NBuPy
Lung				
Metaplasia Osseous				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/0 (0%)	0/5 (0%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/0.00	0/5.00
POLY-3 PERCENT (g)	0%	0%	0%	0%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/0 (0%)	0/5 (0%)
FIRST INCIDENCE	---	---	---	---
STATISTICAL TESTS				
POLY 3	(n)	(n)	(n)	(n)
POLY 1.5	(n)	(n)	(n)	(n)
POLY 6	(n)	(n)	(n)	(n)
COCH-ARM / FISHERS	(n)	(n)	(n)	(n)
MAX-ISO-POLY-3	(n)	(n)	(n)	(n)

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Females			
	Control Female	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Spleen				
Depletion Cellular				
<hr/>				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/5.00	1/5.00
POLY-3 PERCENT (g)	0%	0%	0%	20%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
FIRST INCIDENCE	---	---	---	15 (T)
<hr/>				
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.356
POLY 1.5	(e)	(e)	(e)	P=0.356
POLY 6	(e)	(e)	(e)	P=0.356
COCH-ARM / FISHERS	P=0.177	(e)	(e)	P=0.333
MAX-ISO-POLY-3	(e)	(e)	(e)	P=0.100

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DOSE	Females			
	Control Female	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
<hr/>				
Thymus Atrophy				
<hr/>				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/5.00	1/5.00
POLY-3 PERCENT (g)	0%	0%	0%	20%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/5 (0%)	1/5 (20%)
FIRST INCIDENCE	---	---	---	15 (T)
<hr/>				
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.356
POLY 1.5	(e)	(e)	(e)	P=0.356
POLY 6	(e)	(e)	(e)	P=0.356
COCH-ARM / FISHERS	P=0.177	(e)	(e)	P=0.333
MAX-ISO-POLY-3	(e)	(e)	(e)	P=0.100

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TERMINAL SACRIFICE AT 3 WEEKS**

DOSE	Females			
	Control Female	1.0 mg/ml NBUpy	3.0 mg/ml NBUpy	6.0 mg/ml NBUpy
Thymus Necrosis				
LESION RATES				
OVERALL (a)	0/10 (0%)	0/0 (0%)	0/5 (0%)	3/5 (60%)
POLY-3 RATE (b)	0/10.00	0/0.00	0/5.00	3/5.00
POLY-3 PERCENT (g)	0%	0%	0%	60%
TERMINAL (d)	0/10 (0%)	0/0 (0%)	0/5 (0%)	3/5 (60%)
FIRST INCIDENCE	---	---	---	15 (T)
STATISTICAL TESTS				
POLY 3	(e)	(e)	(e)	P=0.003**
POLY 1.5	(e)	(e)	(e)	P=0.003**
POLY 6	(e)	(e)	(e)	P=0.003**
COCH-ARM / FISHERS	P=0.007**	(e)	(e)	P=0.022*
MAX-ISO-POLY-3	(e)	(e)	(e)	P<0.001**

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LEGEND

- (a) Number of tumor-bearing animals/number of animals examined at site.
 - (b) Number of tumor-bearing animals/Poly-3 number
 - (d) Observed incidence at terminal kill.
 - (e) Value of statistic cannot be computed.
 - (f) Beneath the control incidence are the P-values associated with the trend test. Beneath the dosed group incidence are the P-values corresponding to pairwise comparisons between the controls and that dosed group.
 - (g) Poly-3 adjusted lifetime tumor incidence.
 - (n) No statistics are calculated if all dose groups have fewer than two tumors.
 - (I) Interim sacrifice
 - (T) Terminal sacrifice
 - # Tumor rates based on numbers of animals necropsied.
 - * To the right of any statistical result, indicates significance at ($P \leq 0.05$).
 - ** To the right of any statistical result, indicates significance at ($P \leq 0.01$).
 - N Indicates a negative trend for all tests
- The Cochran-Armitage and Fishers exact tests compare directly the overall incidence rates.

*** END OF REPORT ***