Study Number: C06100 Test Type: TOX Route: Oral Gavage Species/Strain: Rat/Harlan Sprague Dawley

C Number:

Study Gender:

PWG Approval Date

PA48: Summary of Tissue Concentration Test Compound: Perfluorohexane sulfonate potassium salt CAS Number: 3871-99-6

C06100

Female

See web page for date of PWG Approval

Date Report Requested: 01/17/2019 Time Report Requested: 14:34:14 Lab: Battelle

Study Number: C06100	PA	48: Summary of Tissue Conc	entration	Date Report Requested: 01/17/2019
Test Type: TOX	Test Con	npound: Perfluorohexane sulfona	ate potassium salt	Time Report Requested: 14:34:14
Route: Oral Gavage		CAS Number: 3871-99-6		Lab: Battelle
Species/Strain: Rat/Harlan Sprague Dawley				
		Female		
Dose (mg/kg/day)	0	3.12	6.25	12.5
(mmol/kg/day)	0	0.0078	0.0156	0.0312
Plasma Concentration (ng/ml)	175 ± 22 (10) **	37030 ± 1651 (10) **	50410 ± 1552 (10) **	63820 ± 3201 (10) **

92.5 ± 4.1 (10) **

11868.6 ± 529.1 (10)

126.0 ± 3.9 (10) **

8065.6 ± 248.3 (10)

159.5 ± 8.0 (10) **

5105.6 ± 256.1 (10)

0.4 ± 0.1 (10) **

Plasma Concentration (uM)

Normalized Plasma Concentration (uM/mmol/kg)

Study Number: C06100 Test Type: TOX Route: Oral Gavage Species/Strain: Rat/Harlan Sprague Dawley

Test Compound: Perfluorohexane sulfonate potassium salt

CAS Number: 3871-99-6

Date Report Requested: 01/17/2019 Time Report Requested: 14:34:14 Lab: Battelle

Female			
Dose (mg/kg/day)	25	50	
(mmol/kg/day)	0.0625	0.125	
Plasma Concentration (ng/ml)	83820 ± 3740 (10) **	95510 ± 3745 (10) **	
Plasma Concentration (uM)	209.5 ± 9.3 (10) **	238.7 ± 9.4 (10) **	
Normalized Plasma Concentration (uM/mmol/kg)	3352.8 ± 149.6 (10)	1910.2 ± 74.9 (10)	

Date Report Requested: 01/17/2019 Time Report Requested: 14:34:14 Lab: Battelle

LEGEND

Data are displayed as mean ± SEM (N) unless otherwise noted.

SD – Study Day

If over 20% of the animals in a group are above the limit of detection, then 1/2 the limit of detection value is substituted for values that are below the limit of detection.

When the control group did not have over 20% of its values above the limit of detection, no mean or standard error were calculated; no statistical analysis was done for the endpoint.

Statistical analysis performed by Jonckheere (trend) and Shirley or Dunn (pairwise) tests (unless otherwise noted).

Statistical significance for the control group indicates a significant trend test

Statistical significance for a treatment group indicates a significant pairwise test compared to the vehicle control group

* Statistically significant at P <= 0.05

** Statistically significant at P <= 0.01

Molar dose concentrations were calculated based on the salt form, using MW 438.2 g/mol, while PFHxS (non-salt) was measured in the plasma and liver with molar concentrations based on MW 400.1 g/mol.

Normalized values were calculated by dividing the absolute measurment by the dose.

BD - Group did not have over 20% of its values above the limit of detection.

** END OF REPORT **