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: Perfluorooctanoic Acid

CAS Number: 335-67-1

C91070B

Female

See web page for date of PWG Approval

Date Report Requested: 01/17/2019

Time Report Requested: 14:33:59

Lab: Battelle

Test Type: TOX
Route: Oral Gavage

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Lab: Battelle

		Female			
Dose (mg/kg/day)	0	6.25	12.5	25	
(mmol/kg/day)	0	0.0151	0.0302	0.0604	
Plasma Concentration (ng/ml)	BD	491 ± 72 (10)	1153 ± 187 (10)	2960 ± 481 (10)	
Plasma Concentration (uM)	BD	1.2 ± 0.2 (10)	$2.8 \pm 0.5 (10)$	7.1 ± 1.2 (10)	
Normalized Plasma Concentration (uM/mmol/kg)		78.5 ± 11.5 (10)	92.2 ± 15.0 (10)	118.4 ± 19.2 (10)	

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Lab: Battelle

Female				
Dose (mg/kg/day)	50	100		
(mmol/kg/day)	0.121	0.242		
Plasma Concentration (ng/ml)	9326 ± 1821 (10)	23444 ± 3247 (9)		
Plasma Concentration (uM)	22.5 ± 4.4 (10)	56.6 ± 7.8 (9)		
Normalized Plasma Concentration (uM/mmol/kg)	186.5 ± 36.4 (10)	234.4 ± 32.5 (9)		

Test Type: TOX

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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

Values adjusted for molar concentration were calculated by dividing the absolute measurement by the molecular weight of 414.06 g/mol Normalized values were calculated by dividing the absolute measurement by the dose.

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

** END OF REPORT **