

Study Number: C91070B

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

Male

See web page for date of PWG Approval

Date Report Requested: 01/17/2019

Time Report Requested: 14:34:02

Lab: Battelle

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Dose (mg/kg/day)	Male							
	0	0.625		1.25		2.5		
(mmol/kg/day)	0	0.0015		0.003		0.006		
Plasma Concentration (ng/ml)	98 ± 6 (10) **	50690 ± 2207 (10) **	73480 ± 3206 (10) **	95430 ± 4036 (10) **				
Plasma Concentration (uM)	0.2 ± 0.0 (10) **	122.4 ± 5.3 (10) **	177.5 ± 7.7 (10) **	230.5 ± 9.7 (10) **				
Normalized Plasma Concentration (uM/mmol/kg)		81104.0 ± 3531.6 (10)	58784.0 ± 2564.7 (10)	38172.0 ± 1614.4 (10)				
Liver Concentration (ng/g)	BD	54610 ± 2233 (10)	85220 ± 3186 (10)	110740 ± 4467 (10)				
Liver Concentration (uM)	BD	131.9 ± 5.4 (10)	205.8 ± 7.7 (10)	267.4 ± 10.8 (10)				
Normalized Liver Concentration (uM/mmol/kg)		87376.0 ± 3572.0 (10)	68176.0 ± 2548.4 (10)	44296.0 ± 1786.7 (10)				
Liver/Plasma Ratio	BD	1.08 ± 0.02 (10)	1.16 ± 0.02 (10)	1.17 ± 0.03 (10)				

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Dose (mg/kg/day)	Male					
	5			10		
(mmol/kg/day)	0.0121			0.0242		
Plasma Concentration (ng/ml)	110720	± 3891	(10) **	148570	± 15405	(10) **
Plasma Concentration (uM)	267.4	± 9.4	(10) **	358.8	± 37.2	(10) **
Normalized Plasma Concentration (uM/mmol/kg)	22144.0	± 778.2	(10)	14857.0	± 1540.5	(10)
Liver Concentration (ng/g)	109030	± 3557	(10)	124470	± 9251	(10)
Liver Concentration (uM)	263.3	± 8.6	(10)	300.6	± 22.3	(10)
Normalized Liver Concentration (uM/mmol/kg)	21806.0	± 711.4	(10)	12447.0	± 925.1	(10)
Liver/Plasma Ratio	0.99	± 0.03	(10)	0.87	± 0.05	(10)

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LEGEND

Data are displayed as mean \pm 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 $P \leq 0.05$

** Statistically significant at $P \leq 0.01$

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 ****