

Male					
Treatment Groups (mg/kg)					
	8.4 ^a	8.4 ^b	16.6 ^b	49.8 ^b	8.4 IV ^b
Plasma					
C _{max} (ug/mL)		12.3	24.8	73.1	36.9
T _{max} (minute)		15	15	15	
t _{1/2} (Beta) (minute)		166	122	57.9	208
k ₀₁ (min ⁻¹)	0.0730 ± 0.010				
k ₁₂ (min ⁻¹)	0.0854 ± 0.0078				
Cl (mL/min/kg)					8.5
Cl _{1(F)} (mL/min/kg)		9.1	7.4	9.5	
V ₁ (L/kg)	0.333 ± 0.039				
MRT (minute)		205	193	197	180
AUC _{inf} (ug/mL*min)		927	2231	5237	987
F (percent)	1.11 ± 0.097	0.94	1.14	0.89	

Experiment Number: S0541
Route: Gavage, IV
Species/Strain: Mouse/B6C3F1

Toxicokinetics Data Summary
Test Compound: Gemfibrozil
CAS Number: 25812-30-0

Date Report Requested: 11/09/2016
Time Report Requested: 14:02:59
Lab: Research Triangle Institute

LEGEND

Data are displayed as mean \pm SEM

MODELING METHOD & BEST FIT MODEL

^a Compartmental modeling techniques with established models or models written to simultaneously solve iv and oral data sets (SimuSolv, Version 3.0, The Dow Chemical Company, Midland, MI); 2-compartment model employing a delay term in order to simulate the effect of enterohepatic recirculation

^b Models 200 and 201, PCNONLIN software, SCI Software, Lexington, KY; Non-compartmental analysis

ANALYTE

Gemfibrozil

TK PARAMETERS

C_{\max} = Observed or Predicted Maximum plasma (or tissue) concentration

T_{\max} = Time at which C_{\max} predicted or observed occurs

$t_{1/2(\text{beta})}$ = Half-life for the beta phase

k_{01} = Absorption rate constant, k_a

k_{12} = Distribution rate constant from first to second compartment etc.

Cl = Clearance, includes total clearance

$Cl_{1(F)}$ = Apparent clearance of the central compartment, also $Cl_{(F)}$ for gavage groups in non-compartmental model

V_1 = Volume of distribution of the central compartment, includes V_d and V_{volume} of distribution, V_z apparent volume of distribution NCA, V_{app} apparent volume of distribution for intravenous studies

MRT = Mean residence time

AUC_{inf} = Area under the plasma concentration versus time curve, AUC, extrapolated to time equals infinity

F = Bioavailability, absolute bioavailability

**** END OF REPORT ****