**Experiment Number:** K99050

Species/Strain: Rat/F344/NCrl

Route: Gavage

## **Toxicokinetics Data Summary**

Compound/Analyte: Ginkgo Biloba Extract/Quercetin

**CAS Number:** 90045-36-6

Request Date: 11/27/2019 Request Time: 2:30:16

Lab: Battelle

### Male

	Treatment Group (mg/kg)			
	30 Gav <sup>a</sup>	100 Gav <sup>b</sup>	300 Gav <sup>a</sup>	
		Plasma		
Cmax_obs (ng/mL)	8.15	13.2	63.4	
Tmax_obs (minute)	120	10.0	480	
Half-life (minute)	2120	570	ND	
AUC_0-T (ng/mL•min)	4780	13500	43400	
AUCinf_pred (ng/mL•min)	12400	17100	ND	

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#### **LEGEND**

**Route:** Gavage

#### MODELING METHOD & BEST FIT MODEL

<sup>a</sup> WinNonlin (Version 7.0, Certara, L.P., Princeton, NJ) non-compartmental library models with no weighting factors, Non-compartmental model, parameter estimates are reported to three significant figures. ND = Not determined. The terminal elimination phase could not be fully characterized for the 300 mg/kg group because there were not enough measurable concentrations of QCT after the Tmax\_obs. In addition, the characterization of the terminal phases for the 30 mg/kg was poor, with an r2 value of 0.0762 and 61.4 percent of the AUCinf pred extrapolated.

<sup>b</sup> WinNonlin (Version 7.0, Certara, L.P., Princeton, NJ) non-compartmental library models with no weighting factors, Non-compartmental model, parameter estimates are reported to three significant figures. ND = Not determined.

#### **ANALYTE**

Quercetin

#### TK PARAMETERS

Cmax = Observed or Predicted Maximum plasma (or tissue) concentration

Tmax = Time at which Cmax predicted or observed occurs

Half\_life = Lambda z Half life, t 1/2, the terminal elimination half-life based on non-compartmental analysis

AUC\_0-T = Area under the plasma concentration versus time curve, AUC, from time ti (initial) to tf (final), AUClast

AUCinf = Area under the plasma concentration versus time curve, AUC, extrapolated to time equals infinity

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#### TK PARAMETERS PROTOCOL

#### TK PARAMETERS

**Route:** Gavage

TK analysis was performed on Ginkgolide A (GLA), Ginkgolide B (GLB), Ginkgolide C (GLC), Ginkgolide J (GLJ), Bilobalide (BLL), Isorhamnetin (ISR), Quercetin (QCT), and Kaempferol (KMF) after a single gavage administration of ginkgo biloba extract (GBE) batch 1 (Lot 020703) in corn oil at doses of 30, 100, and 300 mg/kg. Blood samples were collected prior to dose administration and at 11 time points post dose administration from typically three animals/time point/group. Time points were Pre-dose, 5, 10, 15, 30, 60, 90, 120, 240, 480, 720, and 1440 minutes. The LLOQ was 4 ng/mL for GLA, 1 ng/mL for GLC and GLB, 5 ng/mL for BLL and ISR, 3 ng/mL for QCT, 10 ng/mL for GLJ, and 40 ng/mL for KMF. The LOD is 3 ng/mL for GLA, 0.3 ng/mL for GLC, 2 ng/mL for BLL and ISR, 1 ng/mL for QCT, 0.4 ng/mL for GLB, 5 ng/mL for GLJ, and 40 ng/mL for KMF. Body weight ranges are 111.8-187.5 g, 114.5-187.7 g, and 111.8-186.6 g for 30, 100, and 300 mg/kg dosed male rats, respectively. The plasma samples were hydrolyzed, heated, cooled, then extracted with ethyl acetate and centrifuged. The organic layer for each sample was removed, concentrated to dryness under nitrogen gas and reconstituted in methanol for analysis. The samples were hydrolyzed to convert the flavonol glycosides to corresponding aglycones which are the compounds quantified. A liquid chromatography coupled with tandem mass spectrometric (LC-MS/MS) method was used to quantitate known GBE constituents, terpene trilactones (Ginkgolide A [GLA], Ginkgolide B [GLB], Ginkgolide C [GLC], Ginkgolide J [GLJ], and Bilobalide [BLL]) and aglycones (Isorhamnetin [ISR], Kaempferol [KMF], and Quercetin [QCT]) of flavonol glycosides.