

Drug-endobiotic interaction effect of UGT enzymes inhibition on systemic bile acids levels in rat

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SUPPLEMENTARY MATERIAL

Table S1. Optimized mass spectrometer conditions used in the analysis of ezetimibe and IS

Mass Spectrometer Parameter				Value			
Curtain gas flow rate				35 L/h			
Collision gas flow rate				Medium			
Ion spray voltage				(-)4500 volts			
Ion source temperature				500 °C			
Nebulizer gas				50 psi			
Drying gas				45 psi			
MRM conditions used in the optimized method							
Analyte	Q1 Mass (Da)	Q3 Mass (Da)	Dwell time (msec)	DP (V)	CE (V)	CXP (V)	Rt (min)
Ezetimibe	408.0	271.1	200	-70	-22	-8	2.1
Ezetimibe-Glucuronide	584.5	271.1	200	-84	-35	-13	1.9
Zafirlukast	574.1	462.3	200	-81	-45	-10	1.6
Telmisartan (IS)	513.2	287.2	200	-82	-10	-10	2.1

DP – Declustering potential; EP – Entrance potential; CE– Collision energy; CXP – Collision cell exit potential; Rt – Retention time and IS – Internal standard; V – volts

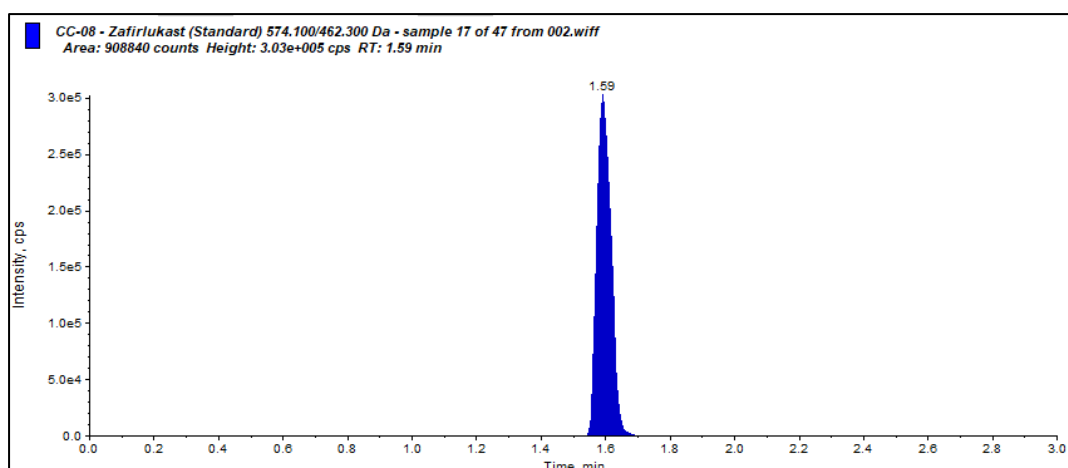


Fig. S1 (a) Chromatogram of zafirlukast from calibration curve standard (900 ng/mL)

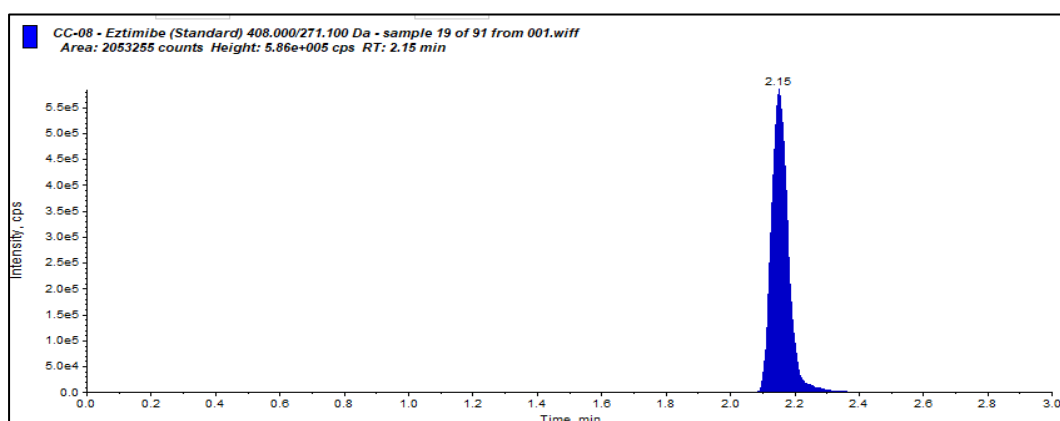


Fig. S1 (b) Chromatogram of ezetimibe from calibration curve standard (550 ng/mL)

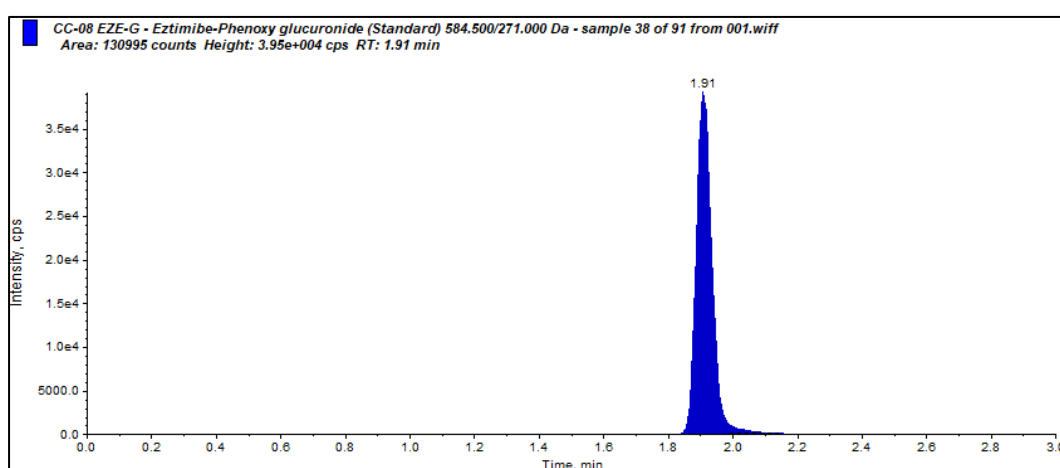


Fig. S1 (c) Chromatogram of ezetimibe- β -D-glucuronide from calibration curve standard (1000 ng/mL)

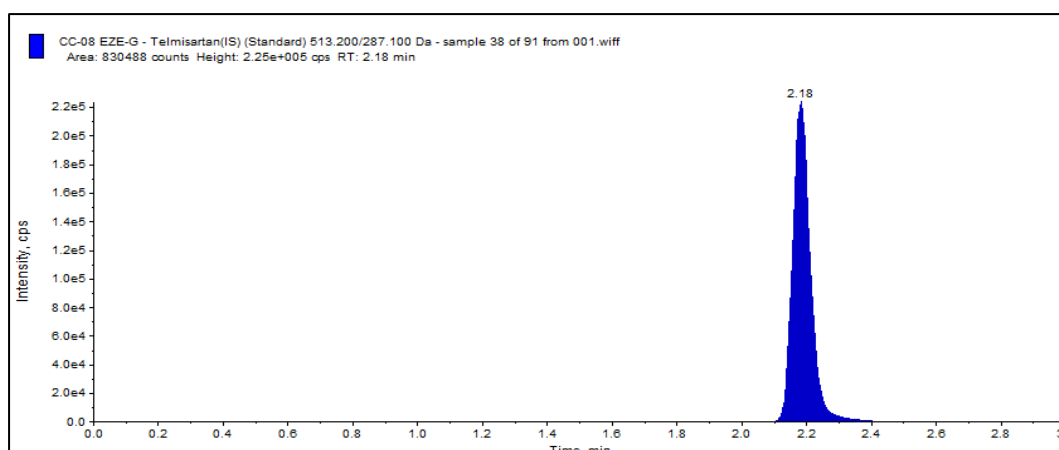


Fig. S1 (d) Chromatogram of telmisartan (IS) at a concentration of 100 ng/mL

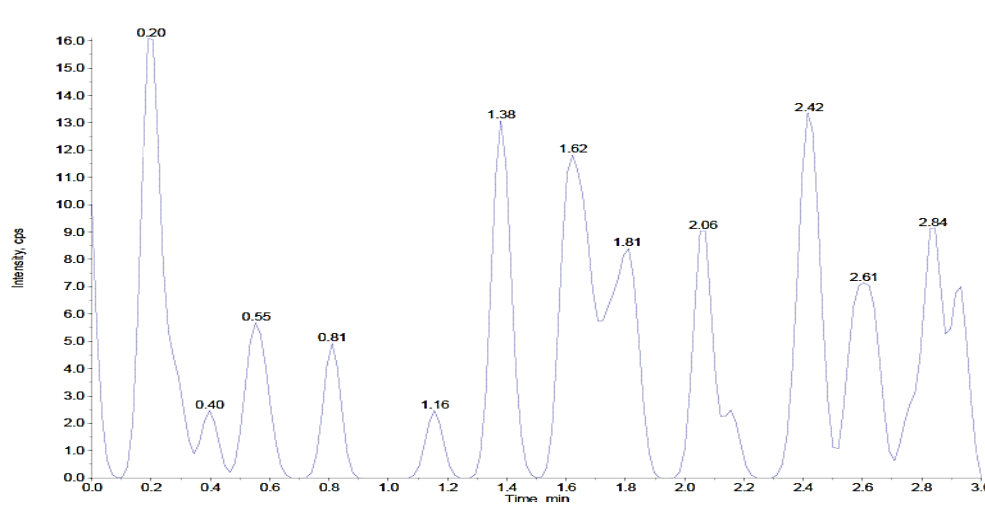


Fig. S1 (e) Chromatogram of blank sample