

A Validated Method for the Quantification of Clopidogrel in Human Plasma at the 2.5 pg/mL Level Using Xevo TQD and ACQUITY UltraPerformance LC

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Abstract

This method allows for the analysis of a pro-drug in human plasma that is highly sensitive and provides carryover less than 20% of the lower limit of quantification.

Introduction

Clopidogrel is a thienopyridine derivative antiplatelet pro-drug used in the prevention of atherosclerotic events. Following oral administration the dosed compound undergoes hepatic metabolism to give rise to the active thiol-metabolite, and the inactive carboxylic acid metabolite. The inactive metabolite accounts for the majority of circulating clopidogrel related material in humans, while the active metabolite and unchanged pro-drug are present at very low levels. The mechanism of action is derived from the binding of the active thiol metabolite to cell receptor P2Y₁₂, irreversibly inhibiting the platelet activation process.¹

Due to the reactivity of the active thiol metabolite, coupled with the low levels of the unchanged pro-drug, most quantitative studies are based on the circulating levels of the inactive metabolite. In this application note, we report the development of a highly sensitive solid phase extraction and LC-MS/MS assay for the analysis of the pro-drug clopidogrel in human plasma with an assay sensitivity of 2.5 pg/mL.

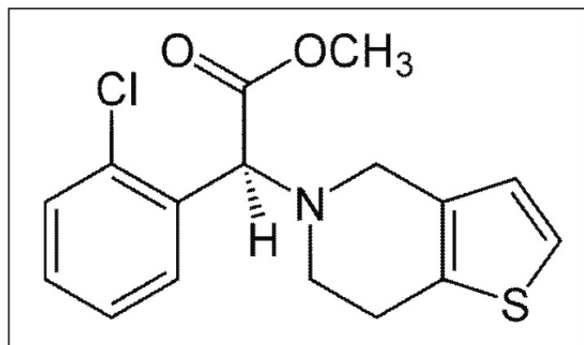


Figure 1. Structure of Clopidogrel.

Experimental

~~The sample was prepared using an Oasis MCX μ Elution Solid Phase Extraction Plate. 350 μ L of plasma sample~~

was mixed with 20 μ L of internal standard solution and 350 μ L of aqueous buffer. The samples were applied to the solid phase extraction plate, which had previously been primed with methanol and aqueous buffer. The sample was washed with an aqueous – methanol solution and eluted with a 2 x 25 μ L of 5% NH_4OH in 60:40 IPA:ACN, then further diluted with 25 μ L of aqueous buffer prior to injection. The analysis was performed on an ACQUITY UPLC H-Class System. A 10 μ L aliquot of the sample was injected onto an ACQUITY UPLC C_{18} 2.1 x 50 mm 1.7 μ m Column. The column was operated under gradient conditions over 3 minutes at a flow rate of 600 μ L/min. The column effluent was monitored using a Xevo TQD Mass Spectrometer operated in multiple reaction monitoring (MRM) positive ion electrospray mode. The transition $322 \Rightarrow 212$ was employed for the clopidogrel and the transition $326 \Rightarrow 216$ was employed for the d4 internal standard.

Results and Discussion

Clopidogrel was eluted with a retention time of 2.10 minutes, as shown in Figure 2. The peak produced by the chromatography system was very symmetrical with a width at the base of 3 seconds. The sharpness of the peak and the symmetrical nature enabled efficient processing and peak integration. The 3 minute analysis time allowed for a total analysis time of five hours for a 96-well SPE plate, allowing two plates to be processed per day. The data displayed in Figure 2 illustrates the injection of an extracted plasma blank injection immediately following the analysis 500 pg/mL standard. We observed from this data that there was no discernable carryover in the blank chromatogram. The extremely low carryover exhibited by the ACQUITY UPLC H-Class System allowed the full sensitivity of the Xevo TQD Mass Spectrometer to be exploited.

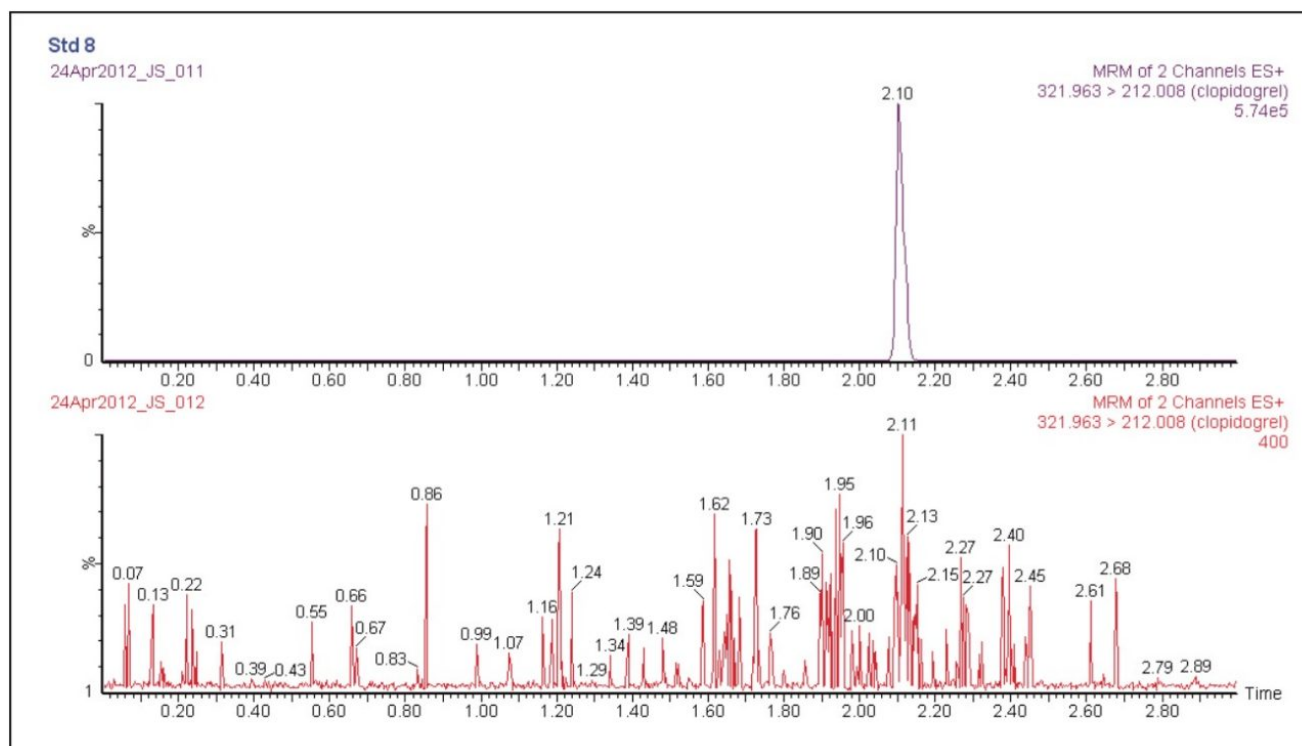


Figure 2. LC-MS/MS Chromatogram of extracted blank and 100 pg/mL clopidogrel standard.

The lower limit of quantification for the assay was determined to be 2.5 pg/mL. The assay was validated with separate 96 well batches on three consecutive days over the range 2.5 to 500.0 pg/mL. A typical calibration obtained for the assay is shown in Figure 3; the correlation coefficient ranged between 0.9990 and 0.9994 using a 1/x weighting linear regression. The intra-day precision and accuracy validation data is shown in Tables 1, 2, and 3. The validation data show that the coefficient of variation ranged from 5.8% to 12.1% for the 2.5 pg/mL LLOQ with a bias between -4.7% and -13.2%. For the high QC (300 pg/mL) the coefficient of variation ranged from 1.3% to 3.0% with a bias between 1.6% and 1.7%. The inter-day precision and accuracy data are displayed in Table 4. The coefficient of variation was determined to be 9.2% for the 2.5 pg/mL LLOQ with a bias of -7.9. For the high QC (300 pg/mL) the coefficient of variation was determined to be 2.3% with a bias of 1.7%.

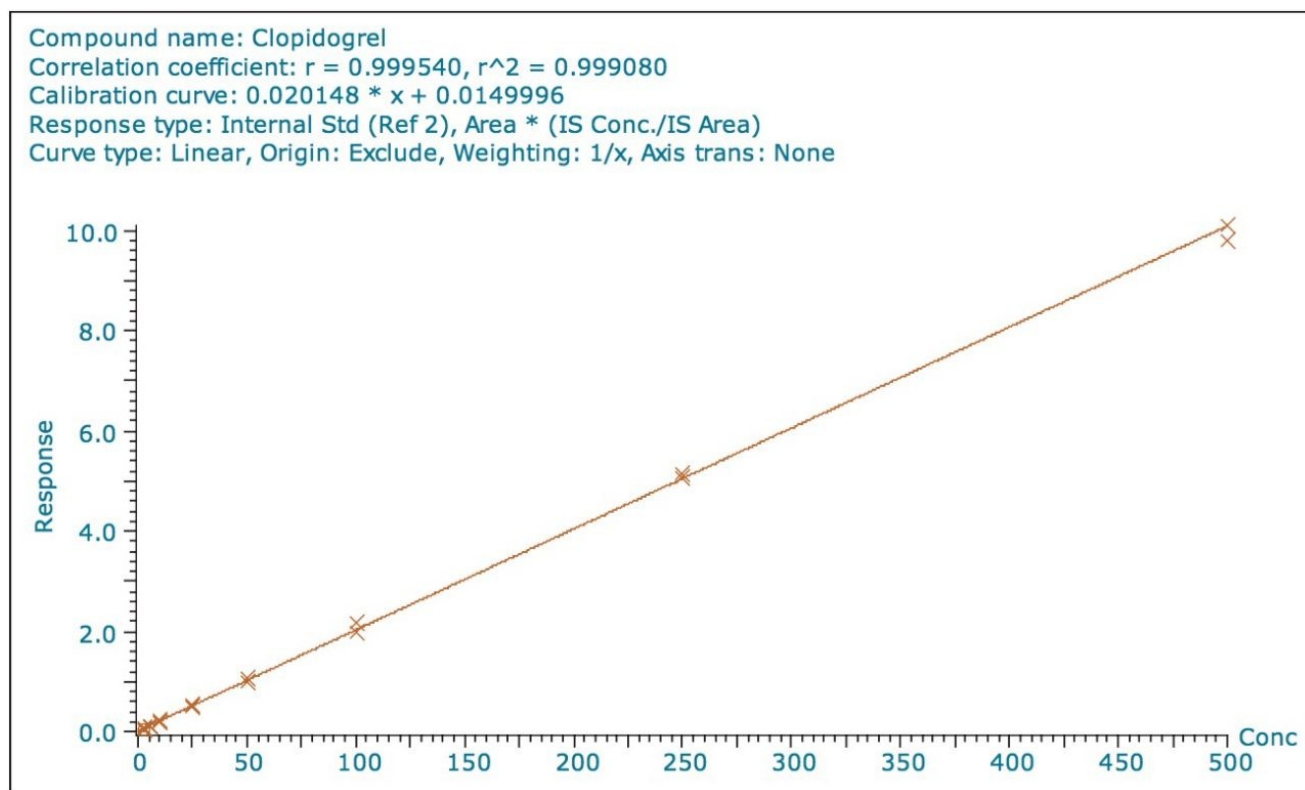


Figure 3. Representative calibration line for the LC-MS/MS quantification of clopidogrel in plasma.

	QC LLOQ 2.5 pg/mL	QC Low 7.5 pg/mL	QC Mid 75.0 pg/mL	QC High 300.0 pg/mL
	2.28	7.51	80.7	315
	2.12	6.60	79.5	307
	2.30	7.41	80.6	302
	2.41	7.27	84.2	301
	1.74	7.50	81.6	293
	*	7.19	77.6	312
Mean	2.17	7.25	80.7	305
St Dev	0.262	0.341	2.18	7.98
%CV	12.1	4.7	2.7	2.6
%Bias	-13.2	-3.4	7.6	1.6

Table 1. Intra-day QC Accuracy/Precision Statistics: Day 1.

*QC LLOQ Rep 6 was a blocked SPE well, no eluent injected.

	QC LLOQ 2.5 pg/mL	QC Low 7.5 pg/mL	QC Mid 75.0 pg/mL	QC High 300.0 pg/mL
	2.32	7.03	73.8	289
	2.13	7.42	76.5	317
	2.40	7.32	71.4	303
	2.72	6.54	74.1	310
	2.5	8.17	75.9	303
	2.22	7.36	82.7	309
Mean	2.38	7.31	75.7	305
St Dev	0.211	0.534	3.86	9.29
%CV	8.8	7.3	5.1	3.0
%Bias	-4.7	-2.6	1.0	1.7

Table 2. Intra-day QC Accuracy/Precision Statistics: Day 2.

	QC LLOQ 2.5 pg/mL	QC Low 7.5 pg/mL	QC Mid 75.0 pg/mL	QC High 300.0 pg/mL
	2.28	7.36	73.0	304
	2.23	8.04	77.3	305
	2.36	7.55	77.2	299
	2.17	6.98	76.7	306
	2.55	7.03	78.7	311
	2.40	7.51	77.5	303
Mean	2.33	7.41	77.6	305
St Dev	0.136	0.389	0.69	4.09
%CV	5.8	5.3	0.9	1.3
%Bias	-6.7	-1.2	3.4	1.6

Table 3. Intra-day QC Accuracy/Precision Statistics: Day 3.

	QC LLOQ 2.5 pg/mL	QC Low 7.5 pg/mL	QC Mid 75.0 pg/mL	QC High 300.0 pg/mL
	2.28	7.51	80.7	315
	2.12	6.60	79.6	307
	2.30	7.41	80.6	302
	2.41	7.27	84.2	301
	1.74	7.50	81.6	293
	*	7.19	77.6	312
	2.32	7.03	73.8	289
	2.13	7.42	76.5	317
	2.40	7.32	71.4	303
	2.72	6.54	74.1	310
	2.50	8.17	75.9	303
	2.22	7.36	82.7	309
	2.28	7.36	78.0	304
	2.23	8.04	77.3	305
	2.36	7.55	77.2	299
	2.17	6.98	76.7	306
	2.55	7.03	78.7	311
	2.40	7.51	77.5	303
Mean	2.30	7.32	78.0	305
St Dev	0.212	0.409	3.22	7.00
%CV	9.2	5.6	4.1	2.3
%Bias	-7.9	-2.4	4.0	1.7

Table

4. Inter-day QC Accuracy/Precision Statistics.

*QC LLOQ Rep 6 was a blocked SPE well, no eluent injected.

Conclusion

- A high sensitivity method has been developed for the analysis of clopidogrel in human plasma.
- The assay showed excellent intra- and inter-day precision and accuracy in a three day validation study.
- The level of quantification was determined to be 2.5 pg/mL with a %CV and bias both considerably below the +/- 20% criteria.
- The carryover was determined to be significantly less than 20% of the LLOQ in an extracted blank following the injection of a high concentration standard.

References

1. J M Pereillo, M Maftouh, A Andrieu, M F Uzabiaga, O Fedeli, P Savi, M Pascal, J M Herbert, J P Maffrand, C Picard. Drug Metabolism and Disposition. 30: 1288-1295, 2002.

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720004401, June 2012



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