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## [Overview presentation]

The appropriate sample-handling procedure for measuring the plasma  $\beta$ -amyloid level using a fully automated immunoassay

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Overview	Objectives
presentation	Plasma $\beta$ -amyloid (A $\beta$ ) assays are a promising tool for Alzheimer's disease diagnosis
	in clinical practice. To obtain reliable results, establishing an appropriate sample-
	handling procedure for each analytical platform is warranted. This study proposes an
	appropriate sample-handling procedure using HISCL analyzer by elucidating the
	individual/combined effects of pre-analytical parameters on plasma A $\beta$ 42/A $\beta$ 40 levels.
	Methods
	We investigated the effects of various pre-analytical parameters, including storage
	times for whole blood, plasma, and freezing conditions, on plasma A $\beta$ 42/A $\beta$ 40 levels,
	and confirmed if these values met the acceptable criteria.
	Results
	Plasma A $\beta$ 42/A $\beta$ 40 levels were acceptable in all conditions. We determined our
	protocol by confirming that plasma A $\beta$ 42/A $\beta$ 40 levels remained acceptable when
	combining pre-analytical parameters.
	Conclusion
	We established an appropriate sample-handling protocol that ensures reliable
	measurement of plasma A $eta$ 42/A $eta$ 40 levels using HISCL analyzer. We believe the A $eta$
	assay, with our protocol, shows promise for aiding AD diagnosis in clinical settings.
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