

Description

SPEAR UltraDetect™ pTau 217 (Tau phosphorylated at threonine 217) was analytically verified for use with human EDTA plasma specimens.**

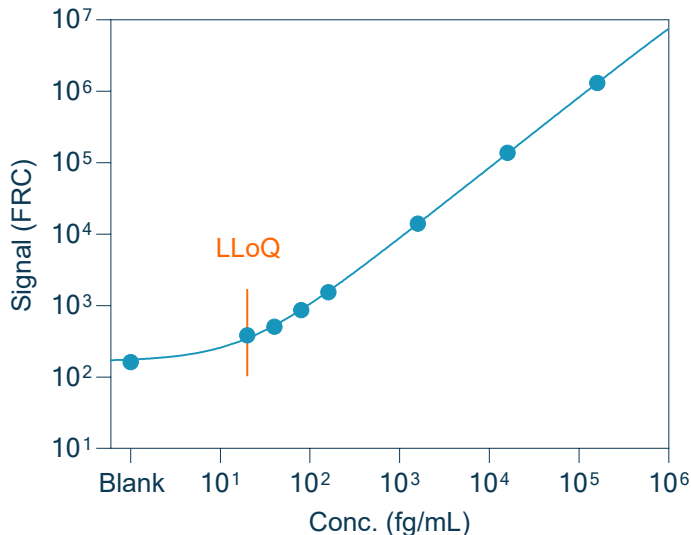
Assay Sensitivity and Range

The lower limit of detection (LLoD) was calculated as 2.5x standard deviations added to the mean blank. The lower limit of quantification (LLoQ) was calculated as lowest concentration with mean CV under 20% and 80-120% recovery. Results were generated from a minimum of 6 independent runs.

Specification	Result
Analytical LLoD (range)	4.92 fg/mL (2.98-7.28)
Functional LLoD	19.7 fg/mL
Analytical LLoQ	20.0 fg/mL
Functional LLoQ	80.0 fg/mL
Minimum required dilution (MRD)	4x (EDTA plasma)
Functional Assay Range	80.0-640000 fg/mL

Calibration Curve

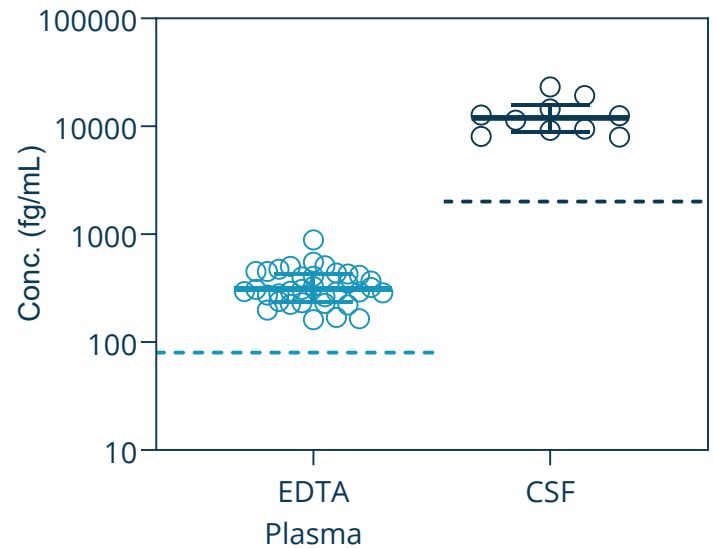
Representative calibration curve fitted with 4PL 1/y² weighting. Analytical LLoQ is indicated in orange.



**All data shown were generated in 96-well runs with F.A.S.T. workflow and qPCR amplification on QuantStudio instruments.

Endogenous Levels

Human EDTA plasma and CSF* from apparently healthy donors were tested. Plasma samples were diluted 4x and CSF 100x. Medians with interquartile ranges are shown for each specimen type with functional LLoQ indicated as dashed horizontal line.



Sample Type (donors)	Mean Conc. (range)	Above LLoD %	Above LLoQ %
EDTA plasma (35 donors)	344 fg/mL (162-885)	100%	100%
CSF* (10 donors)	12838 fg/mL (7964-23184)	100%	100%

*Endogenous levels for CSF were tested for reference only. Full verification of assay performance is recommended before using this matrix in studies.

Benchmark

EDTA Plasma samples (n=100) with concentrations spanning 1.7 logs were measured on SPEAR UltraDetect pTau 217 and an established pTau 217 assay, demonstrating a linear correlation with R² of 0.83 and 2.1x improved stratification of amyloid β PET+ vs. PET- donors by the SPEAR Assay.

Precision

Unspiked EDTA Plasma or EDTA Plasma spiked with CSF, representing six different analyte concentrations, were run in duplicates per run across a minimum of 6 runs to assess intra- and inter-assay precision.

Sample Type	Conc. fg/mL	Mean Intra-assay CV (range)	Inter-assay CV
EDTA Plasma	185	10% (4.4-17%)	6.1%
	1813	2.1% (0.02-4.6%)	6.7%
	3201	3.1% (1.1-7.2%)	4.5%
	6318	1.7% (0.1-3.1%)	3.3%
	6832	1.5% (0.3-2.1%)	2.9%
	14091	1.5% (0.3-5.0%)	3.1%

Spike Recovery

To assess the selectivity of the assay, EDTA Plasma samples from apparently healthy donors (n = 3) were spiked with CSF at low, medium, and high concentrations (relative to endogenous levels and assay dynamic range) and diluted at MRD. Percent recovery was calculated as the endogenous-subtracted spiked sample concentration divided by the measured concentration of the spike material.

Sample Type (donors)	Spike Conc.	Mean Recovery (range)	Recovery
EDTA Plasma (3 donors)	Low	98% (92-102%)	106%
	Med	107% (101-117%)	
	High	115% (102-126%)	

Dilution Linearity

To assess the accuracy of the assay, EDTA Plasma from apparently healthy donors (n = 3) was spiked with CSF and serially diluted 2x to 32x above assay MRD. Percent recovery was calculated as the sample concentration at higher dilution divided by the sample concentration at MRD.

Sample Type (donors)	Dilution	Mean Recovery (range)	Recovery
EDTA Plasma (3 donors)	2x	91% (88-95%)	87%
	4x	89% (85-92%)	
	8x	90% (85-93%)	
	16x	82% (77-87%)	
	32x	84% (83-84%)	

Specificity & Interference

Similar proteins were evaluated for cross-reactivity at 25x the top calibrator concentration and potential interference at 2x endogenous levels. No significant cross-reactivity or interference was observed for:

- Total Tau
- Tau phosphorylated at threonine 181
- Tau phosphorylated at threonine 231

Potential interference was evaluated at low, medium, and high biologically relevant levels for hemoglobin (Hb), triglycerides (TG), conjugated bilirubin (CB), and unconjugated bilirubin (UB). Table below shows the highest concentration at which no significant interference was observed with percent difference from vehicle reported.

Potential Interferent & Conc.		EDTA Plasma % diff.
Hb	High conc. 400 mg/dL	-14.9%
TG	High conc. 500 mg/dL	6.6%
CB	High conc. 2.5 mg/dL	4.3%
UB	High conc. 2.5 mg/dL	4.1%