

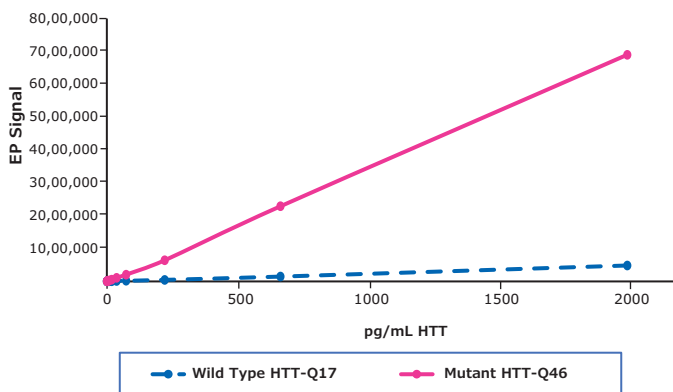
## Sample Testing Services for Measuring Mutant Huntingtin Protein in Cerebrospinal Fluid (CSF) and Brain Homogenates/Lysates

Huntington's Disease (HD) is a progressive brain disorder caused by a defect in the Huntingtin gene. HD is caused by a mutated form of the huntingtin gene, where excessive (more than 36) Glutamine repeats result in formation of an unstable protein. This affects 5-10 in 100,000 people of European descent and is characterized by a decline in cognitive function as well as uncontrolled movement of the body. Mutant huntingtin protein (mHTT) is a key factor in neurodegeneration and accurate analysis in low levels is critical to accelerating research to ultimately find a cure for the disease.

We are happy to offer sample testing services through our SMC™ Custom Assay/Sample Testing (CAST) team to measure mutant Huntingtin protein in your critical samples with our highly validated assay offering:

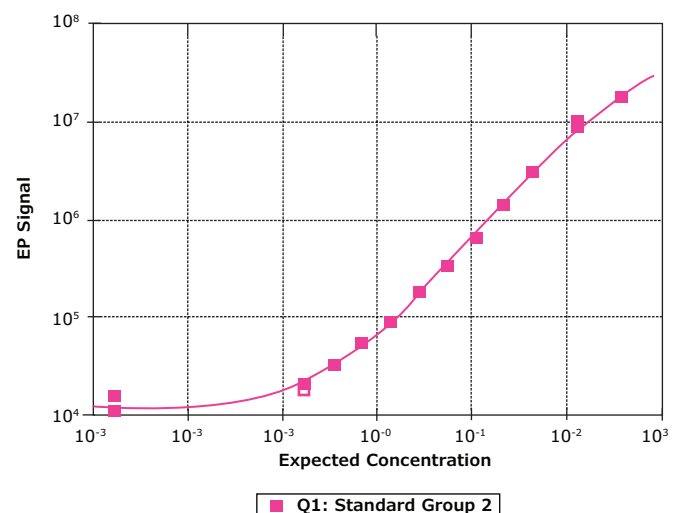
- Measurement down to 0.35 pg/ml for mHTT-Q45 (1-573)
- Measurement in Rat and Human Cerebrospinal Fluid (CSF) and cell lysates.
- Small sample volumes required (10 µL of CSF per well)
- Assay is specific to Mutant Huntingtin (HTT) Protein (>36 poly-glutamine sequence repeats). Assay does not detect Wild type HTT (<23 poly-glutamine sequence repeats).

SMC™ mHTT Assay: Wild Type HTT vs. Mutant HTT



HTT	Description	LLOQ (pg/mL)	MW (g/mol)	M	LLOQ (fM)	Binding Affinity
Q17	(1-3144) full length	74.1	352,178	2.10.E-13	210.40	Very Low
Q46	(1-3144) full length	2.31	355,894	6.49.E-15	6.49	High

EP Signal vs. Expected Concentration



Typical Standard Curve using mHTT-Q45 [pg/mL]

Contact your sales specialist for more information.

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Lit. No. MS\_FL3166EN Ver 1.0 2018 - 18040 11/2018

