

Human Tau [pT181] Antibody Bead Kit

INFORMATION SHEET

| Catalog #: | LHB7051 | Description: | Hu Tau [pT18] | 1] | Lot:*: | 436836 | |
|---|--|------------------------------------|--|--------------------------------------|-----------|--------------------------------|--|
| *Note: A letter at the end of the lot number signifies an additional packaging of this same lot. | | | | | | | |
| Intended Use This reagent set comprises the analyte specific components for the measurement of human Tau when phosphorylated at threonine 181 in CSF and tissue culture media. These reagents are also suitable for the measurement of Tau [pT181] in cell lysates. Buffer reagents needed to complete the reaction are sold separately under Catalog # LNB0001 (BioSource's Neuroscience Buffer Reagent Kit). These reagents are intended for use in the Luminex® 100 TM or 200 TM System only. This kit is configured for research use only and is not to be used in diagnostic procedures. | | | | | | | |
| Reagents Provi 1. Antibody F Catalog #: Bead Region: Form: Storage: | Bead Concentrat LM144 69 0.25 mL 10x b | Description: bead concentrate sol | Ms x Hu Tau Beads lution in storage buffer. Conta | Lot: 436854 ins 7.5 mM sodium | | 0.25 mL-100 tests reservative. | |
| Recommended Dilution: Use 25 μL of diluted bead solution per assay. For a single analyte assay, mix 1 part 10x bead concentrate with 9 parts Working Wash Solution. To develop a multiplexed assay that includes other Neuroscience Antibody Bead Kits, mix equal volumes of the different 10x bead concentrates, then dilute the beads with Working Wash Solution to yield a 1x bead mixture. For example, to develop a 3-plex assay, mix equal parts of each of the three bead concentrates and dilute with 7 parts Working Wash Solution. See the Product Insert included in the Neuroscience Buffer Reagent Kit for further information. Note: Beads for Human Total Tau, Tau [pT181], and Tau [pS199] cannot be mixed. Each of these analytes must be measured separately. 2. Detector Antibody Concentrate (10x): | | | | | | | |
| Catalog #: Form: | 0.5 mL of a thymol as a problem Beads and Determined | 10x stock of Detector Antibody. | ctor Antibody Concentrate in tration of antibody is matched | | | | |
| Storage: Store at 2 - 8°C until the expiration date indicated on the kit. Recommended Dilution: Use 50 μL of diluted Detector Antibody solution per assay. For a single analyte assay, mix 1 part 10x Detector Antibody concentrate with 9 parts Detector Antibody Diluent. To develop a multiplexed assay that includes other Neuroscience Antibody Bead Kits, mix equal volumes of the different 10x Detector Antibody Concentrates, then dilute the antibodies with Detector Antibody Diluent to yield a 1x Detector Antibody mixture. For example, to develop a 3-plex assay, mix equal parts of each of the three Detector Antibody Concentrates and dilute with 7 parts Detector Antibody Diluent. See the Product Insert included in the Neuroscience Buffer Reagent Kit for further information. | | | | | | | |
| 3. Standard (2 Catalog #: Form: | SM055 This Tau [pT | 181] standard (lyo | philized recombinant protein | Lot: R090608 n) is designated in | pg/mL. Th | Single use e protein in this | |
| standard has been calibrated with the respective BioSource phosphoELISA TM kit. Detailed information on calibration is provided on the accompanying page. Storage: | | | | | | | |

Concentration of Reconstituted Standard**: Tau [pT181] 30,000 pg/mL

**Important note: The concentration of reconstituted standard is lot-specific. Please verify all concentration values entered in data analysis software.

Reconstitution: When working with CSF samples and cell lysate samples, reconstitute in 0.5 mL Assay Diluent. When working with tissue culture medium samples, reconstitute with 0.5 mL of a mixture of 50% Assay Diluent and 50% tissue culture medium.

Recommendations for Combining with Other Standards: When combining with one other standard, reconstitute with 0.5 mL. Combine equal volumes of the two standards. When combining with two other standards, reconstitute with 0.333 mL, and combine equal volumes of the three standards. Up to four standards may be combined.

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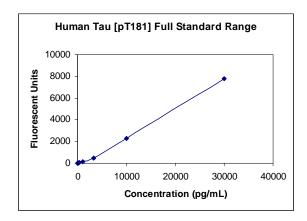
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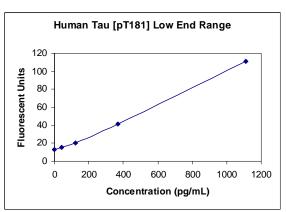
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Recommended Starting Concentration for Standard Curve: Upon reconstitution, the starting concentration of standard is the value cited above. Make serial 1:3 dilutions in Assay Diluent (CSF and cell lysate samples) or mixture of 50% Assay Diluent and 50% tissue culture medium (tissue culture medium samples). Each well designated for standard receives 50 μ L of 1x Detector Antibody followed by 50 μ L standard. Each well designated for sample receives 50 μ L of 1x Detector Antibody followed by 25 μ L assay Diluent and 25 μ L sample. See the Product Insert included in the Neuroscience Buffer Reagent Kit for further information

Performance Characteristics

Analytical Sensitivity: The analytical sensitivity of the Tau [pT181] assay is <50 pg/mL. This was determined by adding two standard deviations to the mean of the fluorescence intensity units obtained when the zero standard was assayed 24 times.





Representative Standard Curve

Species Reactivity: The Human Tau [pT181] Antibody Bead Kit is specific for human Tau when phosphorylated at threonine 181. It does not cross-react with rat or mouse. Other species were not tested.

Specificity: Buffered solutions of a panel of substances with concentrations ranging between 1 and 35 ng/mL were assayed with the BioSource Human Tau [pT181] Antibody Bead Kit. The following substances were tested and found to have no cross-reactivity: Hu Aβ40, Aβ42, IL-1 α , IL-1 β , IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-10, IFN- α , IFN- γ , GM-CSF, TNF- α , IL-12, IL-13, IL-15, IL-17, MCP-1, MIP-1 α , MIP-1 β , Eotaxin, RANTES, IP-10, MIG, EGF, FGF basic, G-CSF, HGF, VEGF, DR5, IL-1RA, IL-2R, IL-4R, sTNF-RI, sTNF-RII, GRO- α , MCP-2, and MCP-3. This kit does not cross-react with non-phosphorylated Tau.

Precision:

| | Intra-assay | Inter-assay |
|--------------|-------------|-------------|
| | (n=16) | (n=32) |
| Mean (pg/mL) | 12081.10 | 12151.30 |
| SD | 195.42 | 217.36 |
| %CV | 1.62 | 1.79 |

Linearity of Dilution: CSF and tissue culture medium (DMEM containing 10% FBS) were spiked with human Tau [pT181] and serially diluted in Assay Diluent or 50/50 Assay Diluent/tissue culture medium over the range of the assay. Linear regression analysis of samples versus the expected concentration yielded a correlation coefficient of 0.99 for CSF and a correlation coefficient of 0.99 for tissue culture medium.

Recovery:

CSF averaged 82.9%. Tissue culture medium containing 10% FBS averaged 97.5%.

Correlation to ELISA:

A correlation coefficient of 0.99 was calculated when values for natural Tau [pT181], obtained with the Hu Tau [pT181] Antibody Bead Kit, were compared with the BioSource ELISA for Hu Tau [pT181]. With natural sample, Hu Tau [pT181] Antibody Bead Kit (pg/mL) x 0.73 = Hu Tau [pT181] ELISA (pg/mL). Fifty picograms measured with this Hu Tau [pT181] Antibody Bead Kit are equal to 1 Unit, as determined by the BioSource Hu Tau [pT181] ELISA kit. Correlation of results obtained with the Hu Tau [pT181] Antibody Bead Kit to one's own system should be determined to arrive at an appropriate multiplication factor.

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Cell Lysate Sample Preparation:

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In addition to CSF and tissue culture medium samples, this kit has been used to detect Hu Tau [pT181] from lysates made from SH-SY5Y cells, prepared in Cell Lysis Buffer (BioSource Cat. # FNN0011: 10 mM Tris, pH 7.4, 100 mM NaCl, 1 mM EDTA, 1 mM EGTA, 1 mM NaF, 20 mM Na $_4$ P $_2$ O $_7$, 2 mM Na $_3$ VO $_4$, 1% Triton X-100, 10% glycerol, 0.1% SDS, 0.5% deoxycholate, 1 mM PMSF [added fresh from 0.1 M stock in DMSO], and protease inhibitor cocktail [added fresh from stock made using Sigma Cat. # P-2714]) and diluted at least ten-fold in Assay Diluent. To produce a lysate, incubate cells with cell lysis buffer (1-2 x 10^8 cells/mL is recommended) on ice for 30 minutes, vortexing at 10 minute intervals, then clarify the lysate by centrifugation at 13,000 RPM for 10 minutes. Cell lysates may be stored at -80° C for up to three months with one freeze/thaw cycle. Optimization of cell stimulation and cell lysis procedures may be required for each specific application.

By purchasing this Kit, which contains fluorescently labeled microsphere beads authorized by Luminex® Corporation ("Luminex®"), you, the customer, acquire the right under Luminex's patent rights to use this Kit or any portion of this Kit, including without limitation the microsphere beads contained herein, only with Luminex's laser based fluorescent analytical test instrumentation marketed under the name Luminex® 100^{TM} or 200^{TM} System. This product is covered by one or more of the following U.S. patents: 6,046,807

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