**FluoProbes®**

**FT-CG2370**

**FSB & other amyloid markers**

**Product Information**

*beta amyloid plaques markers*

<table>
<thead>
<tr>
<th>Product name</th>
<th>cat.number</th>
<th>MW (g·mol⁻¹)</th>
<th>λₑₓc/λₑₓm. max. (nm)</th>
<th>Soluble in</th>
</tr>
</thead>
<tbody>
<tr>
<td>FBS 1% solution</td>
<td>FP-CG2370, 100 µl</td>
<td>420.39</td>
<td>390/511 nm</td>
<td>1% DMSO solution</td>
</tr>
<tr>
<td>BSB</td>
<td>FP-BS6470, 5mg</td>
<td>481.3</td>
<td>340/520 nm</td>
<td>DMSO</td>
</tr>
<tr>
<td>BTA-1</td>
<td>FP-BS6480, 10 mg</td>
<td>240.3</td>
<td>350/NA nm</td>
<td>DMSO</td>
</tr>
<tr>
<td>BTA-2</td>
<td>FP-BS6490, 100 mg</td>
<td>268.4</td>
<td>356/437 nm</td>
<td>DMSO</td>
</tr>
<tr>
<td>Chrysamine G</td>
<td>FP-BS6500, 10 mg</td>
<td>482.5</td>
<td>386/NA nm</td>
<td>DMSO</td>
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<tr>
<td>Half Chrysamine G</td>
<td>FP-BS6520, 10 mg</td>
<td>242.2</td>
<td>342/NA nm</td>
<td>DMSO</td>
</tr>
<tr>
<td>Congo Red</td>
<td>N12511, 1g</td>
<td>696.7</td>
<td>497/NA nm</td>
<td>Water</td>
</tr>
<tr>
<td>ThioFlavin T</td>
<td>FP-BS6530, 1g</td>
<td>318.9</td>
<td>412/482 nm</td>
<td>DMSO</td>
</tr>
</tbody>
</table>

**Storage:**

0-5 ºC, protect from light; Powders: dessicated.

**Introduction**

The formation of amyloid plaques and neurofibrillary tangles are thought to contribute to the degradation of the neurons (nerve cells) in the brain and the subsequent symptoms of Alzheimer’s disease. One of the distinctive features of Alzheimer’s disease is the accumulation of amyloid plaques between nerve cells (neurons) in the brain. These plaques consist primarily of the β-amyloid protein; however, other proteins, such as apoE, are also present.

In recent years there have been extensive studies to investigate the mechanism of Alzheimer’s disease and the roles of β-amyloid in the cause of Alzheimer’s disease. Interchim offers the comprehensive list of Amyloid markers, starting with the great FSB reagent (as well as (labeled) β-amyloid peptides, please inquire).

**FSB**

**Name :**

FSB, 1% DMSO solution

[1-Fluoro-2,5-bis(3-carboxy-4-hydroxystyryl)benzene]

**Catalog Number :**

FP-CG2370 100 µl

**Structure :**

C₁₃H₁₇FO₆ ; MW= 420.39

**Absorption / Emission :**

λₑₓc/λₑₓm = 390/511 nm

FSB, a bis-styrylbenzene analog with fluoride, stains beta amyloid plaques and emits strong fluorescence. The fluorescence intensity of FSB-Beta amyloid complex is twice as high as that of the BSB complex. FSB is also used for MRI detection of plaques due to fluoride in the structure. Higuchi, et al. demonstrated that this compound can be used to detect brain plaques of living mice.
Guidelines for use

1. Permeabilize 30 µm frozen sections of cortical regions
2. Block with 0.5% Triton X-100 and 5% goat serum in phosphate-buffered saline (PBS) for 20 minutes
3. Stain with primary antibodies
4. Wash with PBS/0.1% Triton X
5. Incubate the sections with FluoProbes conjugated secondary antibodies
6. Counterstain with a derivative of 10 µmol/L Congo red (E,E)-1-fluoro-2,5-bis(3-hydroxycarbonyl-4-hydroxy) styrylbenzene (FSB), which specifically binds to the β-sheet conformation of Aβ plaques with an excitation/emission wavelength of 390/511 nm.
7. Mount the sections on slides with Fluoromount G
8. Capture the confocal images of Aβ plaque regions using a confocal microscope with excitation at 488 (for FSB) laser.

Other protocols can be found in the literature.

Related products

- Fluoromount G mounting medium, FP-483331
- Triton X-100, 15851B
- PBS 10X, N14010 or PBS 20X, N13760
- Normal Goat serum, sterile, UP379030
- FluoProbes® 547H Goat anti-Mouse IgG, FP-CB1020
- FluoProbes® 547H Goat anti-Rabbit IgG, FP-CB1050
- FluoProbes® 647H Goat anti-Mouse IgG, FP-CB1040
- FluoProbes® 647H Goat anti-Rabbit, FP-CB1060

References


Other Amyloid markers

| Name: BSB [(trans,trans)-1-Bromo-2,5-bis-(3-hydroxycarbonyl-4-hydroxy)styrylbenzene] | Catalog Number: FP-BS6470 5 mg | MW: 481.3; Soluble in DMSO | \( \lambda_{ex} / \lambda_{em} = 340/520 \text{ nm} \) |

BSB, derived from the structure of Congo Red, is shown to bind to a wide range of amyloid inclusions in situ. More importantly it is also used to label brain amyloids in live animals. BSB recognizes amyloid lesions, and has distinctive properties which allow the quantitative monitoring of the formation of amyloid fibrils assembled from the Ab peptide, a-synuclein and tau. It is a cell-permeable fluorescent probe that specifically binds to and labels intracellular b-amyloid aggregates both in vitro (Ki = 400 nM) and in vivo. It is also used as an antemortem diagnostic tool for animal models of alzheimer's disease.

References:
BTA-1 is an uncharged derivative of thioflavin-T that has high affinity for Ab fibrils and shows very good brain entry and clearance. The Kd of [3H]BTA-1 for binding to AD brain is very similar to the Kd for binding to synthetic Ab fibrils. BTA-1 does not appear to bind significantly to common neuroreceptors or transporter sites. BTA-1 exhibits high affinity for amyloid deposits (Ki = 11 nM for Ab(1-40)). It crosses the blood brain barrier and displays up to 50-fold higher affinity than ThT. It selectively stains cerebral plaques and cerebrovascular amyloid deposits in the brains of PS1/APP transgenic mice.

Reference:

BTA-2 is an uncharged derivative of thioflavin-T that exhibits high affinity for amyloid deposits (Ki = 143 nM for Aβ(1-40)) and can cross the blood brain barrier. It displays up to 6-fold higher affinity than ThT and stains both plaques and neurofibrillary tangles in post mortem Alzheimer disease brain.

References:

Chrysamine G (CG) is a carboxylic acid analog of Congo red, a histologic dye which stains amyloid. CG binds to the b-amyloid protein of Alzheimer’s disease (AD) in vitro and partitions into the brain of normal mice. The binding of CG is correlated with numbers of senile plaques and neurofibrillary tangles. CG displays both high (Kd = 200 nM; Bmax = 1.13 moles per mole of Ab40) and low (Kd = 38.77 mM; Bmax = 23.10 moles per mole of Ab40) affinity binding sites for b-amyloid (Ab) fibrils. It can cross the blood-brain barrier and serve as an useful probe for detecting senile plaques (Ab aggregate). In addition, CG can be used to stain cerebrovascular amyloid in tissue sections.

References:

Half Chrysamine G (hCG) has a lower affinity for Ab compared with that of CG. Both CG and hCG are equally efficacious in reducing Ab-induced neuronal death at a concentration of 0.1-1 mM, indicating that the mechanism of action for CG is not due to its chelating activity, but rather due to its anti-oxidant activity.

Reference:
FluoProbes®

FT-CG2370

<table>
<thead>
<tr>
<th>Name</th>
<th>Congo Red (CR)</th>
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</thead>
<tbody>
<tr>
<td>Catalog Number</td>
<td>&quot;UltraPure Grade&quot; N12511 1 g</td>
</tr>
<tr>
<td></td>
<td>&quot;Solution&quot; FP-AQ3370 100 tests</td>
</tr>
<tr>
<td>MW</td>
<td>696.67 ; Soluble in water</td>
</tr>
<tr>
<td>( \lambda_{\text{exc}} ), ( \lambda_{\text{em}} )</td>
<td>610/N/A, C_{22}H_{17}FO_{6}</td>
</tr>
</tbody>
</table>

Early diagnosis and classification of amyloid deposition and differentiation from other glomerular fibrillar deposits rely on routine Congo red (CR) histochemistry. CR binding, monitored by characteristic yellow-green birefringence under crossed polarization has been used as a diagnostic test for the presence of amyloid in tissue sections for several decades. This assay is also widely used for the characterization of in vitro amyloid fibrils. CR is sandwiched between two protein molecules causing protein oligomerization. Congo red fluorescence (CRF) is an alternative method based on examination of the CR-stained section by ultraviolet (UV) light. CRF is simple to perform and more pronounced, therefore easier to evaluate than CR in bright light. Congo red, when combined with immunohistochemistry, is still visible under UV whereas CR is masked in bright light. Although not widely used, the CRF method for detecting amyloid is simple to use with a high specificity and sensitivity, and may be applied successfully to frozen sections.

References:

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<td>FP-BS6530, 1g</td>
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<td>( \lambda_{\text{exc}} ), ( \lambda_{\text{em}} )</td>
<td>412/482nm</td>
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The benzothiazole dye thioflavin T (ThT) is a classic amyloid stain for senile plaques containing bA4 peptide in Alzheimer's disease brain. ThT also binds rapidly and specifically to the anti-parallel b-sheet fibrils formed from synthetic b-amyloid (1-40), but does not bind to monomer or oligomeric intermediates. The fibrillar b-sheet-bound dye species undergoes a characteristic 120 nm red shift of its excitation spectrum that may be selectively excited at 450 nm, resulting in a fluorescence signal at 482 nm. ThT is a useful probe for the aggregated fibrillar state of b-amyloid (1-40) fibrils as the amyloid-specific fluorescence reports only fibrillar species. The binding of ThT does not interfere with the aggregation of this peptide into amyloid fibrils. The putative conformational changes detected by the ThT fluorescence suggest that small pharmacologic ligands can perturb and possibly dissociate Ab amyloid fibrils.

References:

Catalog size quantities and prices may be found at [http://www.interchim.com](http://www.interchim.com). Please inquire for higher quantities (availability, shipment conditions).

For any information, please ask : FluoProbes® / Interchim; Hotline : +33(0)4 70 03 73 06

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