

Amyloid Fluorescent Staining Kit

Cat. No. CSR-SYN02

Updated on March 18th, 2020

X RIKEN (Japan)'s technology is used in Amyloid Fluorescent Staining Kit

[I] Kit Components

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Storage: 4°C

• component: This kit can be used up to 100 tests when used as described in below [III]

Component	Size	Quantity	Note
Amyloid fluorescent staining reagent	100 μL	1	Keep in a cool dark place once unpacked.
Nuclear staining reagent	100 μL	1	Use appropriate protective equipment (such as gloves and glasses) when handling. Keep in a cool dark place once unpacked.
Fluorescence enhancer	5 g	1	Use appropriate protective equipment (such as gloves and glasses) when handling. Keep at room temperature.

Required but not provided:

- 50% ethanol
- PBS(-)
- Purified water
- 10% neutral buffered formalin

[II] Preparation of working solutions

- Amyloid fluorescent staining solution: Dilute 200 fold with 50% ethanol (Just before use)
- Nuclear staining solution: Dilute 500 fold with PBS(-) (Just before use)
- Fluorescence enhancer: Into 10 mL purified water, dilute 1g of Fluorescence enhancer, vortex thoroughly (Please prepare necessary amount). Since fluorescence enhancer will not dissolve completely & precipitate, use the supernatant of saturated solution.

(III) How to use the kit

- Below example is for using α -Synuclein Aggregation Assay Kit (Cosmo Bio, cat. no. CSR-SYN01-COS) for 24 well plate size.
- 1. Assay as described in α -Synuclein Aggregation Assay Kit (Cosmo Bio, cat. no. CSR-SYN01-COS) manual.
- 2. Remove culture medium, add 0.5 mL 10% Neutral buffered formalin to each well, leave it at least overnight at room temperature to fix cells.
- 3. Remove formalin solution, add diluted 0.2 mL Amyloid structure fluorescent staining solution to each well, incubate at room temperature for 30 min. with protection from light (still standing).
- 4. Remove Amyloid structure fluorescent staining solution, add 0.3 mL Fluorescence enhancer solution to each well, incubate at room temperature for 5 min. with protection from light (still standing).



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- 5. Remove Fluorescence enhancer solution, add 0.5 mL 50% Ethanol to each well, wash once with PBS(-).
- 6. Add 0.2 mL Nuclear staining solution to each well, incubate 10 min. at room temperature with protection from light (still standing).
- 7. Remove Nuclear staining solution, wash once with PBS(-), followed by Fluorescence microscope observation. If high background, repeat wash with PBS(-).

Fluorescence property of each solution is follow.

Nuclear staining solution: λ ex = 352 nm, λ em = 461 nm

Amyloid structure fluorescent staining solution: λ ex = 390 nm, λ em = 511 nm

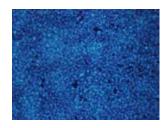
(IV) Detection example of α – Synuclein aggregation complex

 α -Synuclein Aggregation Assay Kit (Cosmo Bio, cat. no. CSR-SYN01) was assayed using SH-SY5Y cell line, followed by Amyloid Fluorescent staining with the same kit.

Negative control vector (pCMV-NC)



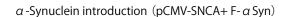
Phase difference



Nuclear staining

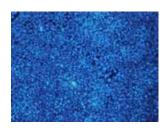


Synuclein aggregate

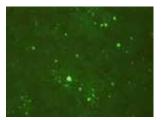




Phase difference



Nuclear staining



Synuclein aggregate



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[V] Related products

Description	Cat. No.	Quantity
lpha -Synuclein Aggregation Assay Kit	CSR-SYN01	1 kit (300 test)
lpha -Synuclein Fibrils	CSR-SYN03	0.1 MG
Human $lpha$ -Synuclein, Recombinant, E.coli	CSR-SYN04	0.1 MG / 1 MG
Mouse α -Synuclein Fibrils	CSR-SYN05	1 MG
Mouse $lpha$ -Synuclein, Recombinant, E.coli	CSR-SYN06	0.1 MG / 1 MG



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