

Product Data Sheet

Proteins

Flubida-2

Cat. No.: HY-D1688 Molecular Formula: $C_{86}H_{96}N_8O_{20}S_2$

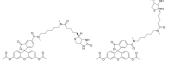
Molecular Weight: 812.93

Target: Fluorescent Dye

Pathway: Others

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.



BIOLOGICAL ACTIVITY

| Description | Flubida-2 is a cell permeable dye which can be hydrolyzed to Fubi-2 by endoesterases in cells (after hydrolysis, Ex=492 nm, Em=517 nm). Flubida-2 can be used to detect pH at a specific site in a cell organelle by directing the probe to where avidin fusion proteins are located ^[1] . |
|-------------|--|
| In Vitro | Guidelines (Following is our recommended protocol. This protocol only provides a guideline, and should be modified according to your specific needs) ^[1] . 1. Dissolve Flubida-2 in DMSO (approximately 2 mM). 2. Mix the stock solution 1: 1 with 20% (w/v in DMSO) Pluronic F-127 (Molecular Probes), and dilute to the desired final concentration (2-4 μM) with serum-free DMEM. 3. 30 to 48 h post-transfection with either AV-KDEL or ST-AV DNA, HeLa cells are rinsed once with serum-free DMEM and loaded with 2-4 μM Flubida-2 for 3-5 h (or overnight for 10-15 h). 4. Chase the labeled cells with normal growth medium for at least 2 h, to allow excess dye-biotin to exit from the cytosol. 5. The strong avidin-biotin interaction ensures stable, specific avidin-Flubi- binding that resisted washing. Biotin starvation of the cells is not necessary before Flubida- loading, as staining was bright and stable. MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

[1]. M M Wu, et al. Studying organelle physiology with fusion protein-targeted avidin and fluorescent biotin conjugates. Methods Enzymol. 2000;327:546-64.

Caution: Product has not been fully validated for medical applications. For research use only.

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