(FOR RESEARCH USE ONLY. DO NOT USE IT IN CLINICAL DIAGNOSTICS !)

Elabscience® Elab Fluor® Violet 450 Labeling Kit

Catalog No: E-LK-E010C

Product size: 3 Reactions/10 Reactions

This manual must be read attentively and completely before using this product.
If you have any problems, please contact our Technical Service Center for help.

Toll-free: 1-888-852-8623
Tel: 1-832-243-6086
Fax: 1-832-243-6017
Email: techsupport@elabscience.com
Web: www.elabscience.com

Please refer to specific expiry date from label outside of box.
Please kindly provide us with the lot number (on the outside of the box) of the kit for more efficient service.

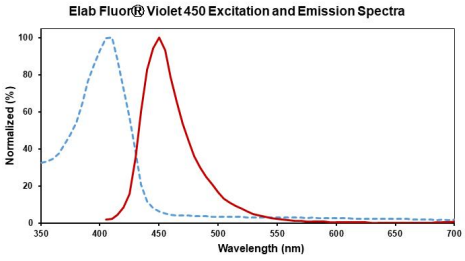
Introduction

Elabscience® Elab Fluor® Violet 450 Labeling Kit provides all the reagents required for labeling, which can label proteins containing primary amino-group (-NH_2) molecules simply and effectively.

Characteristic

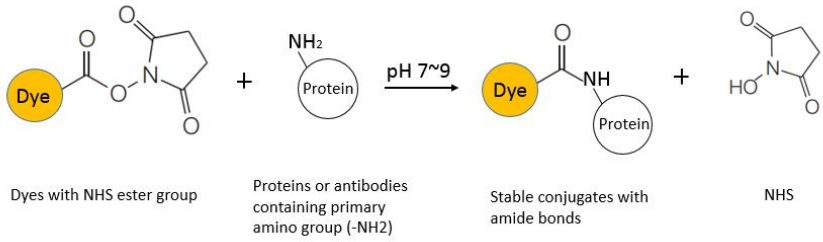
- ✓ **Fast**: The whole process takes only 90 min.
- ✓ **Convenient**: The dye has been activated and can be used directly. Filtration tube desalts without dialysis.
- ✓ **Flexible use**: It can be used for both micro-labeling and large-scale labeling, and can label 0.1-1 mg proteins each time.

Essential Information

Excitation/Absorption maximum (nm)	405
Emission maximum (nm)	444
Molar extinction coefficient ε ($L \cdot mol^{-1} \cdot cm^{-1}$)	35000
Correction coefficient of 280nm(CF280)	0.238
Dye spectrum	

Labeling Principle

Within a certain pH range, Elab Fluor® Violet 450 specifically reacts with primary amino groups (N-terminal and lysine residue side chains) to form a stable amide bond, so as to realize the coupling of Elab Fluor® Violet 450 with protein.

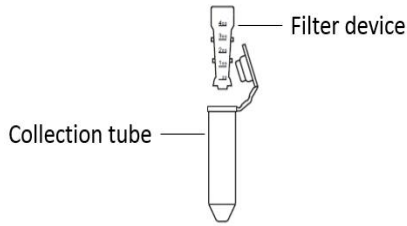


Components

Cat.	Products	3 Reactions	10 Reactions	Storage
E-LK-E10L	Elab Fluor® Violet 450 NHS ester	0.11mg×3	0.11mg×10	-20°C, shading light
E-LK-011	Labeling Buffer II	20 mL	20 mL×2	2~8°C
E-LK-006	DMF	500 μL	500 μL	2~8°C, shading light
E-LK-007	1×PBS(pH7.4)	10 mL	10 mL×2	2~8°C
E-LK-008	1M Tris(pH8.7)	500 μL	500 μL×2	2~8°C
E-LK-001C	50 KD Filtration tube*	3 set**	10 set	RT

*The filtration tube is purchased from Millipore. Please refer to the appendix Ⅲ for usage.

**1 set 50 KD Filtration tube (0.5 mL) consisted of one filter device and two collection tubes.



Storage

The unsealed kit can be stored at 2~8°C for 1 year, and the dissolved Elab Fluor® Violet 450 can be stored at -20°C or -80°C for 1 week.

Materials Not Supplied

1. Pipettor and tips (0.5-10μL, 2-20μL, 20-200μL, 200-1000μL).
2. Ultraviolet spectrophotometer or nanodrop or multifunctional microplate reader.
3. 37°C incubator.
4. Centrifuge (centrifugal force up to 12,000×g).

Calculation of the usage amount of Elab Fluor® Violet 450:

The amount of dye used in each reaction depends on the mass, concentration and molecular weight of the protein to be labeled. For the protein above 100 KD, the recommended molecular ratio of dye and protein using this kit is 18:1.

Example: Label 1 mg protein(concentration about 2 mg/mL), when the molecular ratio of Elab Fluor® Violet 450 and protein (150KD) is 18:1, the molar concentration of Elab Fluor® Violet 450 is 13.06 mM (refer to the preparation of Elab Fluor® Violet 450), the calculation of the amount of Elab Fluor® Violet 450 to be added is below:

1. Calculate the amount of substance required of Elab Fluor® Violet 450:

$$n_{\text{Elab Fluor® Violet 450}} = \text{n}_{\text{protein}} \times 18 = \frac{1\ mg}{150000 mg/mmol} \times 18$$

$$= 0.00012 mmol$$

2. Calculate the required volume of Elab Fluor® Violet 450:

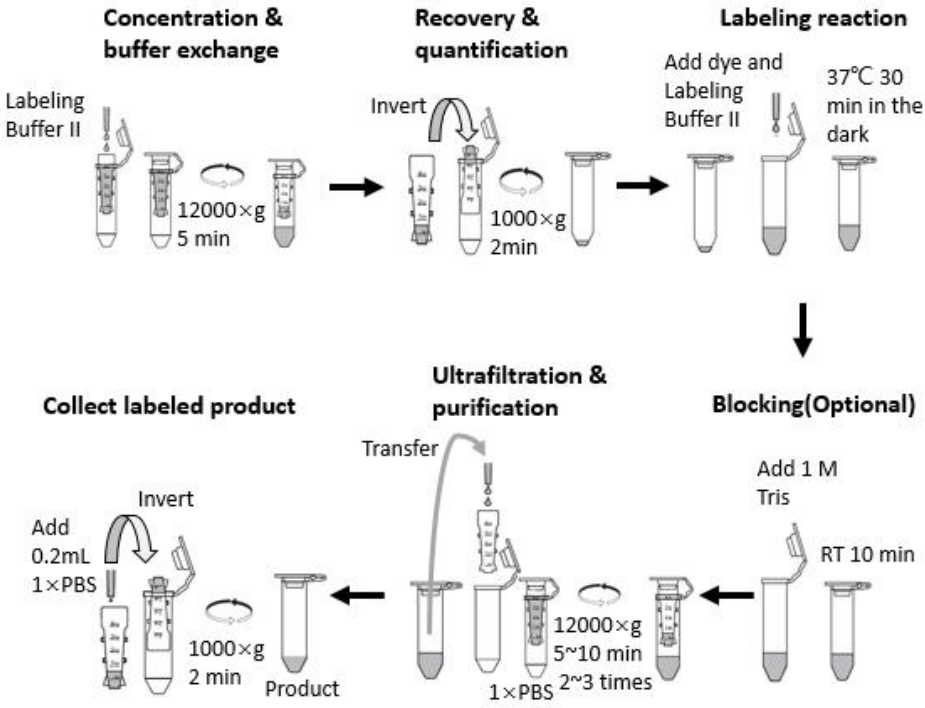
$$V_{\text{Elab Fluor® Violet 450}} = \frac{n_{\text{Elab Fluor® Violet 450}}}{C_{\text{Elab Fluor® Violet 450}}} = \frac{0.00012\text{mmol}}{13.06 mM} = 9.2 \mu L$$

Experimental Operation

■ Experiment preparation

1. Read the instructions carefully.
2. Bring all reagents to room temperature for 20 min before use (note: the reagent components are temporarily unused are still in the refrigerator).
3. Infiltration of ultrafiltration tube: Add 500 μL of Labeling Buffer II into the dry filter device, stand at room temperature for 10 min, and discard Labeling Buffer II before adding the reagent to be labeled (the filter device should remain moist throughout the labeling process).
4. Preparation of Elab Fluor® Violet 450: Dissolve 0.11 mg of Elab Fluor® Violet 450 NHS ester with 10 μL of DMF, and stand for 10 min until it is fully dissolved. At this time, the concentration of Elab Fluor® Violet 450 is 13.06 mM, and cover the tube for later use.

■ Labeling process



■ **Labeling procedure (This procedure is used to label 1 mg protein)**

1. **Concentration & buffer exchange:** Put the filter device in the collection tube, add 1 mg of protein to be labeled into the filter device, add Labeling Buffer II to the final volume of 0.5 mL, cover the filter tube, centrifuge at 12,000×g for 5 min, and discard the liquid in the collection tube.

   Note:

   a) The maximum volume of the filter device is 0.5 mL.

   b) If the volume of 1 mg protein is greater than 0.5 mL, please add it in several times and concentrate it by centrifugation and ultrafiltration.

   c) If the protein to be labeled contains free amino groups (Tris, amino acids or other interferents, repeat ultrafiltration with Labeling Buffer II to ensure that it is removed fully).

2. **Recovery & quantification:** Invert the filter device into the collection tube, centrifuge at 1000×g for 2 min, collect the protein in the collection tube, take out the filter device, add an appropriate amount of Labeling Buffer II into the collection tube, make sure that the protein concentration is about 2 mg/mL. At the same time, add 0.5 mL Labeling Buffer II into the filter device and put it on a pipe rack for later use.
3. **Labeling reaction:** Immediately add 9.2 μL of 13.06 mM Elab Fluor® Violet 450 to the protein solution, gently blow and mix fully, sealed with a lid, and incubate at 37°C for 30 min in the dark.
4. **(Optional) Blocking:** Add 1 M Tris (pH 8.7) to stop the reaction at the ratio of 10 μL of 1 M Tris (pH 8.7) per 100 μg protein, mix fully and incubate at room temperature for 10 min.
5. **Ultrafiltration & purification:** Add an appropriate amount of 1×PBS into the above reaction solution to the final volume of 0.5 mL, gently mix and transfer the reaction solution to the filter device, make sure that the Labeling Buffer II in the filter device in step 2 should be discard (if the above reaction solution exceeds 0.5 mL, it can be transferred to the

spin-dried filter device for several times after ultrafiltration), and cover the cap after matching with the collection tube, and centrifuge for 5~10 min at the speed of 12,000×g. Discard the liquid in the collection tube, replenish 1×PBS to 500 μL in the filter device, and repeat the centrifugal ultrafiltration operation for 2~3 times until the color of the ultrafiltrate in the collection tube is almost colorless and transparent.

6. **Collect labeled product:** Add 0.2 mL 1×PBS into the filter device and pipet gently. Invert the filter device in another collection tube and centrifuge at 1000×g for 2 min. Collect the solution in the collection tube, which is the protein labeled by Elab Fluor® Violet 450.

■ (Optional) Determining the degree of labeling

1. Use an absorbing light scanning device to set the scanning range of 230 nm~800 nm.
2. Set 1×PBS to as blank control.
3. Take 2 μL Elab Fluor® Violet 450 labeled sample, scan the absorption spectrum (230 nm to 800 nm) and record A280 and A405 data (1 cm optical path).

   Note: In this case, it is necessary to scan the absorption value in the range of 230nm~800nm, rather than measuring the value at A280 and A405. By observing the absorption spectrum curve, some abnormal values can be eliminated, such as the measurement error caused by bubbles in the sample.
4. The DOS and protein concentration can be calculated based on the molar extinction coefficient of Elab Fluor® Violet 450 dye, the A280 correction value, the molar extinction coefficient of the protein, the molecular weight of the protein, etc. The calculation formula is as follows:

   $DOS = (A_{405} \times \varepsilon_{IgG}) / (\varepsilon_{\text{Elab Fluor® Violet 450}} \times (A_{280} - CF_{280} \times A_{405}))$

   $\text{Protein concentration (mg/mL)} = (A_{280} - CF_{280} \times A_{405}) \times 150000 / \varepsilon_{IgG}$

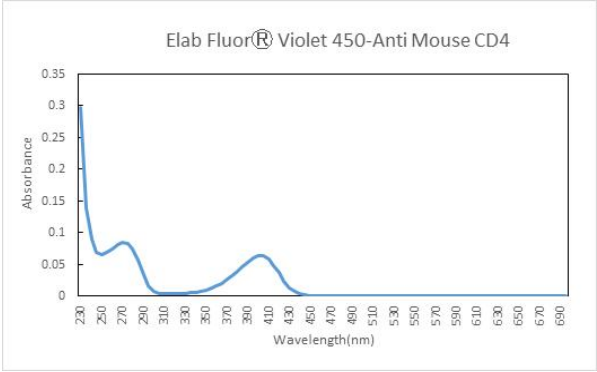
Parameter	Implication	Value

DOS	The average number of fluorescein labeled per protein.	
A_{405}	The OD value of Elab Fluor® Violet 450 dye at 405 nm wavelength with 1 cm optical path.	Refer to measurements
ε_{IgG}	The molar extinction coefficient of IgG ($L \cdot mol^{-1} \cdot cm^{-1}$).	210000
$\varepsilon_{Elab\ Fluor® Violet\ 450}$	The molar extinction coefficient of Elab Fluor® Violet 450 ($L \cdot mol^{-1} \cdot cm^{-1}$).	35000
A_{280}	The OD value of Elab Fluor® Violet 450 labeled protein at 280 nm wavelength with 1 cm optical path.	Refer to measurements
CF_{280}	The correction coefficient of OD value of Elab Fluor® Violet 450 dye at 280nm wavelength	0.238

■ **The storage and use of protein**

Add 0.05~0.2% Proclin 300 or 0.05% sodium azide and stabilizer protein (such as 0.1% BSA) to the labeled protein, the protein can be stored at 2~8°C in the dark for 6 months. Or add the same volume of glycerol, the protein can be stored at -20°C for 6 months.

Typical Results



Notes

1. Please select the appropriate kit according to the molecular weight of the protein to be labeled. The kit provides a 50 KD Filtration tube.
2. Elab Fluor® Violet 450 is susceptible to moisture hydrolysis failure, and should be stored at -20°C or -80°C with the desiccant. In order to prevent water vapor from condensing into the dye, it is necessary to equilibrate the dye to room before the experiment.
3. The kit can also be used to label other proteins containing free amino groups. The specific labeling ratio is determined according to the number of available amino groups in the marker or set different molar ratios for labeling.
4. The optimal molecular ratio of the dye and protein (150KD) recommended for labeling with this kit is 18:1, which can ensure that the dye can label the protein, and DOS is in the normal range, but it cannot ensure the best experimental results. The optimal labeling ratio may vary according to the difference of protein, and the user can optimize according to the actual situation.

Related Products

Cat.No	Product
E-LK-R002	BSA Removal Kit

Troubleshooting

Symptoms	Causes	Comments
Low DOS value	The initial concentration of protein is inaccurate.	Quantify the protein according to the standard operating procedure, and refer to the instructions of the instrument manufacturer, or use other protein quantification methods such as BCA method.
	Initial proteins contain a large number of interfering components for labeling, such as glycine, imidazole, Tris, sodium azide, thiomersal, proclin, etc.	Use dialysis, desalination or multiple ultrafiltration methods to fully remove the interfering components. In special cases, it may be necessary to quantify the interference components contained in the sample.
	The initial protein contains carrier proteins such as BSA, gelatin.	Use affinity purification or other chromatographic methods to remove the interfering components, or use the BSA removal kit, and then accurately quantify the protein concentration before labeling.
	The value measured is not accurate.	Quantify the measured values according to the standard operation methods.
	During labeling, DMF in the reaction system is excess, which interferes with labeling reaction.	Minimize the amount of DMF to dissolve dyes.
The proteins are not labeled with dye at all.	Improper operation, such as incomplete mixing of dye and protein, excessive ammonium ion or amino component in protein, or other improper operation.	Set a positive control.

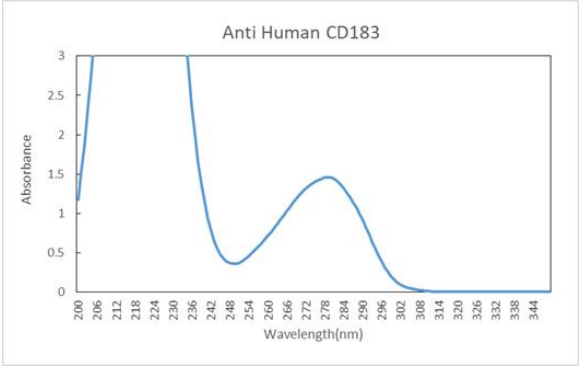
The proteins are not labeled with dye at all.	Improper preservation of dye.	Before labeling, the dye should not be mixed with water. When taking the dye, it should be kept at room temperature for about 5~10 min before unsealing.
	Improper use of ultrafiltration tube.	After ultrafiltration, there is less liquid in the filter device. Do not use the pipette to directly absorb the sample in the filter device. The sample should be centrifuged in an inverted state.
	Due to the difference of centrifuges, the rotation speed is too high during ultrafiltration.	Centrifuge speed is 12,000×g, not 12,000rpm
	Leakage of ultrafiltration tube.	Overload of filter to cause leakage.
Excessive DOS value (such as greater than 10)	The actual concentration of protein is low, such as protein precipitation, uneven concentration, resulting in excessive modification of dye.	Accurately measure the concentration according to the standard operation steps of the instrument and equipment after protein mixed fully.
	Excessive lysine residues of protein.	Reduce the amount of dye added or set different molar ratios for labeling.
	Inappropriate spectral measurements.	Refer to the instrument detection sensitivity, it is generally necessary to ensure that the theoretical concentration of protein is above 0.5 mg/mL after labeling. Measure the range of absorption spectrum, not the maximum absorption value, Determine the accuracy of the measured value based on the absorption spectrum curve, especially the baseline level of absorption spectrum curve.

Low recovery of protein	Protein aggregated and precipitated during the labeling process.	Add 1M Tris to terminate the reaction in time.
	Excessive protein concentration during ultrafiltration.	Do not load too much protein in filter device, such as more than 1 mg of protein.
	The protein cannot be completely dissolved in the labeling buffer.	Choose other labeling kits.

Declaration

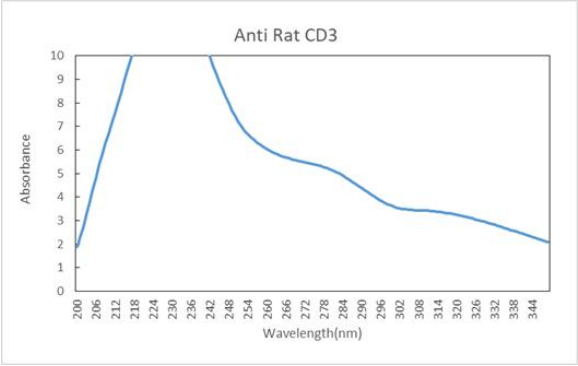
1. This kit is for research use only.
2. Please take safety precautions and follow the procedures of laboratory reagent operation.
3. Although this labeling kit has verified that many different proteins have been labeled, it is still possible that the protein binding affinity is impaired or even lost during the dye-labeled protein process. Although this phenomenon is rare, the reason for this phenomenon is likely that the protein has one or more key lysine amino acid residues directly at the antigen binding site, and its binding function is impaired during the labeling process. In addition, some proteins have key lysine residues (not necessarily at the binding site), which are absolutely critical to maintaining the stability/solubility of the protein. Once modified, the protein is completely unavailable due to precipitation. In this rare case, this labeling kit is not fully responsible. We believe that our customers should be aware that the use of dyes in this kit may in some cases impair the biological function of the modified protein.
4. Although this kit can also be used to label proteins, it should be recognized that changes in various properties of different proteins are quite different from proteins, such as protein solubility in different buffers, pH stability, temperature stability, protein purity, accessibility of labeling sites, and so on. Therefore this labeling kit does not guarantee the quality of labeling proteins other than IgG and IgM proteins.

Appendix Ⅰ: Normal absorbance curve of protein concentration (for reference only)



Description: 1 mg/mL Mouse anti human CD183, the protein type was Mouse IgG1, PBS (pH 7.2, no preservatives), measured by the Nano 100 spectrophotometer, the concentration curve was normal, and A280=1.454, in line with the labeled concentration.

Appendix Ⅱ: Abnormal absorbance curve of protein concentration (for reference only)



Description: 0.5 mg/mL Mouse anti rat CD3, protein type Mouse IgG3, containing protein stabilizer and sodium azide (≤ 0.09%), measured by Nano-100 spectrophotometer, the concentration curve was abnormal, A280=5.195, which does not meet the labeled concentration.

Appendix Ⅲ: Protein retention and concentrate recovery (from Millipore product manual)

(Cite from the User Guide of Millipore: https://www.emdmillipore.com/US/en/product/Amicon-Ultra-0.5-Centrifugal-Filter-Unit,MM_NF-UFC500324#documentation)

For most applications, molecular weight is a convenient parameter to use in assessing retention characteristics. Merck Millipore Ltd. (Millipore) recommends using a membrane with a NMWL at least two times smaller than the molecular weight of the protein solute that one intends to concentrate. Refer to the table below.

Marker/Concentration	Molecular Weight	Device NMWL	% Retention	Spin Time (min)
α -Chymotrypsinogen (1 mg/mL)	25,000	3K	＞95	30
Cytochrome C (0.25 mg/mL)	12,400		＞95	30
Vitamin B-12 (0.2 mg/mL)	1,350		＞42	30
α -Chymotrypsinogen (1 mg/mL)	25,000	10K	＞95	15
Cytochrome C (0.25 mg/mL)	12,400		＞95	15
Vitamin B-12 (0.2 mg/mL)	1,350		＞23	15
BSA (1 mg/mL)	67,000	30K	＞95	10
Ovalbumin (1 mg/mL)	45,000		＞95	10
Cytochrome C (0.25 mg/mL)	12,400		＜35	10
BSA (1 mg/mL)	67,000	50K	＞95	10
Ovalbumin (1 mg/mL)	45,000		~40	10
Cytochrome C (0.25 mg/mL)	12,400		＜20	10
Thyroglobulin (0.5 mg/mL)	677,000	100K	＞95	10
IgG (1 mg/mL)	156,000		＞95	10
Ovalbumin (1 mg/mL)	45,000		＜30	10

Spin Conditions: 40° fixed angle rotor, 14,000×g, room temperature, 500 µL starting volume, n=12.