ALDEFLUOR™ Kit

For the identification, evaluation, and isolation of stem and progenitor cells expressing high levels of ALDH

Catalog #01700 1 Kit



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Product Description

ALDEFLUOR™ is a reagent kit that is used to identify human cells that express high levels of the enzyme aldehyde dehydrogenase (ALDH). The activated ALDEFLUOR™ Reagent, BODIPY-aminoacetaldehyde (BAAA), is a fluorescent non-toxic substrate for ALDH, which freely diffuses into intact and viable cells. In the presence of ALDH, BAAA is converted into BODIPY-aminoacate (BAA), which is retained inside the cells. The amount of fluorescent reaction product is proportional to the ALDH activity in the cells and is measured using a flow cytometer. Viable ALDH-bright (ALDH^{br}) cells can, in principle, be isolated using a cell sorter. Active efflux of the reaction product is inhibited by an efflux inhibitor in the ALDEFLUOR™ Assay Buffer. A specific inhibitor of ALDH, diethylaminobenzaldehyde (DEAB), is used to control for background fluorescence.

ALDEFLUOR™ is optimized for the detection of ALDH^{br} hematopoietic cells in human blood and bone marrow, but it can also be used with non-hematopoietic cells. For a full list of ALDEFLUOR™ products, visit www.stemcell.com.

Product Information

COMPONENT NAME	COMPONENT #	SIZE	STORAGE	SHELF LIFE
Dry ALDEFLUOR™ Reagent*†	01703	50 µg	Store at 2 - 8°C.	Stable until expiry date (EXP) on label.
ALDEFLUOR™ Diethylaminobenzaldehyde (DEAB) Reagent, 1.5 mM in 95% ethanol**	01705	1 mL	Store at 2 - 8°C.	Stable until expiry date (EXP) on label.
Hydrochloric Acid (HCl, 2 N)**†	01704	1.5 mL	Store at 2 - 8°C.	Stable until expiry date (EXP) on label.
Dimethylsulphoxide (DMSO)**†	01706	1.5 mL	Store at 2 - 8°C.	Stable until expiry date (EXP) on label.
ALDEFLUOR™ Assay Buffer**	01702	2 x 55 mL	Store at 2 - 8°C. Do not freeze.	Stable until expiry date (EXP) on label.

^{*}ALDEFLUOR™ Reagent is not cytotoxic. The combination of Dry ALDEFLUOR™ Reagent, DMSO, and HCl shows no cytotoxic or phototoxic effects at concentrations 100-fold above those used in this assay.

Directions for Use

Please read the entire protocol before proceeding.

A) ALDEFLUOR™ ACTIVATION

Dry ALDEFLUOR™ Reagent is provided in a stable, inactive form (BODIPY-aminoacetaldehyde-diethyl acetate, BAAA-DA). For use, Dry ALDEFLUOR™ Reagent is dissolved in DMSO, converted to the fluorescent-activated ALDEFLUOR™ Reagent (BAAA) by treatment with 2 N HCl, and diluted with ALDEFLUOR™ Assay Buffer:

- 1. Assemble all necessary supplies and allow kit reagents to come to room temperature (15 25°C) before use.
- 2. Add 25 µL of DMSO to the vial of Dry ALDEFLUOR™ Reagent, mix well, and let it stand for 1 minute at room temperature.

 NOTE: Dry ALDEFLUOR™ Reagent is an orange-red powder that changes to a bright yellow-green color upon addition of DMSO.
- 3. Add 25 µL of 2 N HCl and mix well. Incubate this mixture for 15 minutes at room temperature. NOTE: Adding 2 N HCl before DMSO will render the product inactive.

^{**}Please refer to the Safety Data Sheet for hazard information.

[†]This component is sold as part of the ALDEFLUOR™ Kit (Catalog #01700) and is not available for individual sale.

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- 4. Add 360 μL of ALDEFLUOR™ Assay Buffer to the vial and mix.
 - NOTE: Upon addition of the ALDEFLUOR™ Assay Buffer, the solution may appear slightly cloudy. This does not affect the assay performance.
- 5. Keep the activated ALDEFLUOR™ Reagent at 2 8°C during use.
- 6. Aliquot the remaining activated ALDEFLUOR™ Reagent and store at -20°C.

B) CELL SAMPLE PREPARATION

- Prepare fresh or previously frozen cell samples according to standard procedures for the cell type.
- 2. If using blood samples where the red blood cell (RBC) to leukocyte ratio (RBC:WBC) of the specimen is > 2:1, lyse the erythrocytes with Ammonium Chloride Solution (Catalog #07800).
- 3. After RBC lysis, centrifuge the sample for 5 minutes at 250 x g. Remove the supernatant and suspend cells in 1 mL of ALDEFLUOR™ Assay Buffer.
- 4. Perform a cell count.
- If using hematopoietic cells (e.g. peripheral blood, apheresis product, bone marrow, or cord blood), adjust the sample to a concentration of 1 x 10^6 cells/mL with the ALDEFLUOR™ Assay Buffer.

NOTE: For other cell types, different cell concentrations may be more appropriate. For optimization of ALDEFLUOR™ staining conditions for non-hematopoietic cells, refer to the Technical Bulletin: ALDEFLUOR™ Assay Optimization (Document #29902), available at www.stemcell.com.

C) ALDEFLUOR™ ASSAY

- 1. Label one "test" and one "control" tube for each sample to be tested. Place 1 mL of the adjusted cell suspension (section B) into each "test" sample tube.
- 2. Add 5 µL of ALDEFLUOR™ DEAB Reagent to the "control" tube. Recap control tube and DEAB vial immediately.
 - NOTE: ALDEFLUOR™ DEAB is provided in 95% ethanol. Recap immediately to prevent evaporation.
- Add 5 µL of the activated ALDEFLUOR™ Reagent per milliliter of sample to the first sample "test" tube. Mix and immediately transfer 0.5 mL of the mixture to the DEAB "control" tube.
 - NOTE: The ALDH enzymatic reaction begins immediately upon addition of the activated substrate to the cell suspension. It is imperative that an aliquot of the ALDEFLUORTM-reacted cells be added to the DEAB "control" tube without delay.
- 4. Add control and substrate solutions as described in steps 2 and 3 above for each sample to be tested.
- 5. Incubate "test" and "control" samples for 30 to 60 minutes (max) at 37°C. Do not exceed 60 minutes.
 - NOTE: Optimal incubation times may vary between cell types. For suggestions on optimization of ALDEFLUOR™ staining conditions for non-hematopoietic cells, cultured cells, and cell lines, refer to the Technical Bulletin: ALDEFLUOR™ Assay Optimization (Document #29902), available at www.stemcell.com.
- 6. Following incubation, centrifuge all tubes for 5 minutes at 250 x g and remove supernatant. Resuspend cell pellets in 0.5 mL of ALDEFLUOR™ Assay Buffer and store the cells on ice or at 2 8°C.
 - NOTE: If immunophenotyping is to be performed, add and incubate the antibodies after step 6. To prevent efflux of the ALDEFLUOR™ product, it is important that the antibody incubation is performed in ALDEFLUOR™ Assay Buffer. Whenever possible, keep the cells chilled (2 8°C or on ice) to slow down the product efflux.
- 7. OPTIONAL: Perform a viability cell count. If the sample contains fewer than 90% viable cells, it is recommended to stain cells with a DNA dye such as propidium iodide or 7-actinoaminomycin-D in order to stain dead and late apoptotic cells.

D) FLOW CYTOMETER SET-UP AND DATA ACQUISITION

Refer to the Technical Bulletin: The Basic FACS on ALDEFLUOR™ (Document #28000), available at www.stemcell.com.

Notes and Tips

- Fresh or previously frozen samples can be analyzed for ALDH^{br} cells. However, the ALDEFLUOR™ kit will only detect ALDH activity in cells that are viable and have intact cell membranes.
- Removal of erythrocytes from the sample is required. Erythrocytes may be removed by lysis using reagents that do not contain detergents or fixatives (e.g. Ammonium Chloride Solution, Catalog #07800). They may also be removed by density centrifugation.
- When frozen aliquots of the activated ALDEFLUOR™ Reagent are thawed, a small precipitate (pellet) may be observed. Before use, mix the thawed reagent to resuspend the precipitate. This precipitate does not affect assay performance.
- The cell lines A549 (lung carcinoma), SKBR3 (breast cancer), and K562 (CML) express ALDH activity and can be used as positive controls for the ALDEFLUOR™ assay. In addition, commercially available bone marrow mononuclear cells can also be used as positive controls.

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Identification of rare ALDH^{br} cells in heterogeneous cell samples can be improved by removing mature hematopoietic cells on the basis
of lineage antigen expression and enriching for ALDH^{br} cells by using RosetteSep[™] Human Progenitor Cell Basic Pre-Enrichment Kit,
RosetteSep[™] Human Hematopoietic Progenitor Cell Enrichment Cocktail, or EasySep[™] Human CD34 Positive Selection Kit II prior to
performing the ALDEFLUOR[™] assay.

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