

## **PRODUCT INFORMATION**

Applications Antibody internalization labeling kit

**Detection method** Cell viability detection with MTT, CCK8, or CTG

**Excitation-Emission** 

**Molecular Weight** The product has a MW of 34 kDa

Formulation & Reconstitution

Lyophilized from sterile PBS, pH 7.4. Normally 5 % 8% trehalose is added as protectants before lyophilization. Please see Certificate of Analysis for specific instructions of reconstitution.

The DiTagTM Eribulin IgG labeling reagents can IgG type be used for human IgG1, IgG2 and IgG4, rabbit

lgG, mouse lgG2a and lgG2b.

Recommended **Dilutions** 

**Background** 

We recommend test antibody to mix with

AME100005 at 2:1 in molar ratio

Description DiTagTM Eribulin IgG labeling reagent

Delivery in Stock

The reagents are supplied in lyophilized form. We

recommend storing the vial(s) at -20°C,

desiccated and protected from light. Once reconstituted, the reagents can be stored at 2-8°C Storage & Shipping

for 1~2 weeks, or with 50% glycerol at -20°C. DiTag™ Eribulin IgG labeling reagents provide an

easy solution for quantifying antibody internalization activities. Leveraging Mal-PEG2-VCP-Eribulin conjugated to an Fc binding protein, these reagents bind to IgG antibodies from the conjugated to the confidence of the confiden

various species, resulting in the formation of an Eribulin-labeled antibody-reagent complex. Upon antibody internalization, the cleavable linker Val-Cit-PABC is enzymatically cleaved by cathepsin B, a protein overexpressed in multiple cancer types. This enzymatic cleavage triggers the release of This enzymatic cleavage triggers the release of

PABC-substituted Eribulin, forming an unstable intermediate that liberates the free drug. Measurement of cell killing or inhibition allows researchers to evaluate the efficiency of antibody internalization into cells. This critical information enhances our understanding of the cellular uptake mechanism of antibodies and aids in

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assessing their efficacy in targeted therapies or diagnostic applications.

Usage Research use only





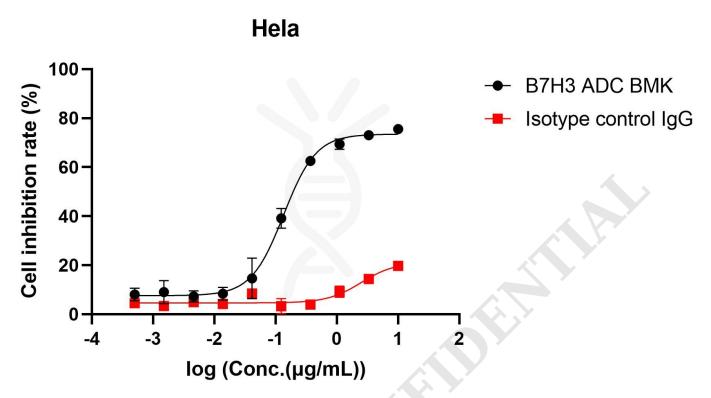


Figure 1. Cell inhibition rate of Hela detected by CCK8 method. The IC50 of B7H3 ADC BMK is 135.1ng/ml, indicating specific internalization.

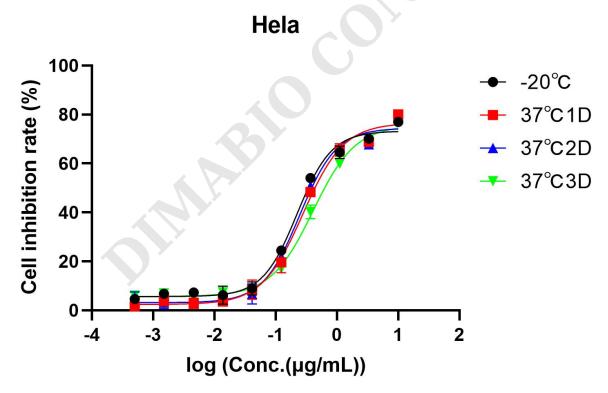
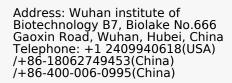


Figure 2. Accelerated stability test of AME100005. After lyophilization, the samples were stored at -20°C (black), 37°C for 1 day (red), 37°C for 2 days (blue), 37°C for 3 days (green), separately. After reconstitution, cell inhibition rate of each samples was detected by CCK8 method. The data indicate that all the samplesexhibit excellent stability.



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