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## Product Data Sheet

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Product Name: Brentuximab vedotin

Cat. No.: GC68296

### Chemical Properties

Cas. No. 914088-09-8

Formula M.Wt

Solubility Storage -80°C, protect from light

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution: ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure

### Background

IC50: 2.5 ng/mL (CD30)<sup>[2]</sup>

Brentuximab vedotin (cAC10-vcMMAE) is an antibody-drug conjugate (**ADC**) comprising an anti-**CD30** antibody and the cytotoxic agent Monomethyl auristatin E (MMAE).

Brentuximab vedotin inhibits **CD30**-positive cells with an **IC<sub>50</sub>** of 2.5 ng/mL.

Brentuximab vedotin can be used for the research of relapsed and refractory Hodgkin lymphoma<sup>[1][2]</sup>.

Brentuximab vedotin (cAC10-vcMMAE) (1 µg/mL; 96 h) shows cytotoxicity to CD30<sup>+</sup> in Karpas 299 cells<sup>[2]</sup>.

Brentuximab vedotin (CAC10-VCMAE) (1 µg/mL; 12, 24 and 48 h) selectively induces growth arrest in G2/M phase then lead to apoptotic cell death<sup>[2]</sup>.

Cell Cytotoxicity Assay<sup>[2]</sup>

Cell Line: Karpas 299 cells

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

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Concentration: 1 µg/mL

Incubation  
Time: 96 h

Result: Showed cytotoxicity to CD30<sup>+</sup> Karpas 299 cells with an IC<sub>50</sub> value of 2.5 ng/mL.

### Cell Cycle Analysis<sup>[2]</sup>

Cell Line: L540 cells

Concentration: 1 µg/mL

Incubation  
Time: 12, 24, and 48 h

Result: Selectively induced growth arrest in G2/M phase to apoptotic cell death.

Brentuximab vedotin (cAC10-vcMMAE) (10-120 mg/kg; i.p. for 3 weeks) the maximum tolerated dose (MTD) is between 30 and 40 mg/kg<sup>[2]</sup>.

Brentuximab vedotin (cAC10-vcMMAE) (0.3, 1 mg/kg; flanks injection; every 4 days for a total of 4 doses 1 mg/kg) induces tumor CD30 regression<sup>[2]</sup>.

Animal Model: SCID mice<sup>[2]</sup>

Dosage: 10 to 120 mg/kg

Administration: Intravenous injection; 10 to 120 mg/kg; for 3 weeks

Result: Showed an maximum tolerated dose between 30 and 40 mg/kg.

Animal Model: SCID mice<sup>[2]</sup>

Dosage: 0.3 and 1 mg/kg

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Administration: Flanks injection; 1 mg/kg every 4 days for a total of 4 doses; 0.3 mg/kg every 4 days for a total of 4 doses

Result: Induced complete and durable tumor regression, but 0.3 mg/kg provided lower therapy than 1 mg/kg dose.

[1]. Shea L, Mehta-Shah N. Brentuximab Vedotin in the Treatment of Peripheral T Cell Lymphoma and Cutaneous T Cell Lymphoma. *Curr Hematol Malig Rep.* 2020 Feb;15(1):9-19.

[2]. Francisco JA, et al. cAC10-vcMMAE, an anti-CD30-monomethyl auristatin E conjugate with potent and selective antitumor activity. *Blood.* 2003 Aug 15;102(4):1458-65.

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