

# AMD3100 octahydrochloride, CXCR4 antagonist ab120718

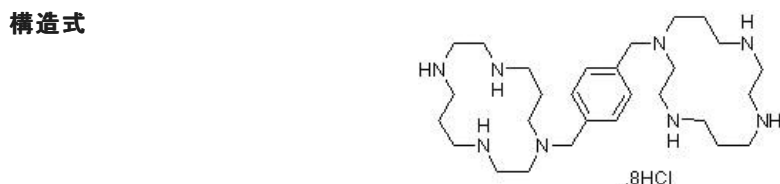
**23 References**    画像数 2

### 製品の概要

製品名	AMD3100 octahydrochloride, CXCR4 antagonist
製品の詳細	Highly selective CXCR4 antagonist
生理活性の詳細	<p>Plerixafor (hydrochloride) is a macrocyclic compound that acts as an irreversible antagonist against the binding of CXCR4 with its ligand, SDF-1 (CXCL12).</p> <p>It suppresses infection by HIV with an IC<sub>50</sub> value of 1-10 ng/ml with selectivity toward CXCR4-tropic virus. Plerixafor mobilizes hematopoietic stem and progenitor cells for transplant better than G-CSF alone. It also increases T-cell trafficking in the blood and spleen as well as the central nervous system. Plerixafor regulates the growth of primary and metastatic breast cancer cells and inhibits dissemination of ovarian carcinoma cells.</p>

精製度 > 99%

CAS 番号 155148-31-5



### 製品の特性

体系名	1,1'-[1,4-Phenylenebis(methylene)]bis-1,4,8,11-tetraazacyclotetradecane octahydrochloride
分子量	794.48
分子式	C <sub>28</sub> H <sub>54</sub> N <sub>8</sub> .8HCl
PubChem 登録番号	65014
保存方法	Store at -20°C. Store under desiccating conditions. The product can be stored for up to 12 months.
溶解性	Soluble in PBS, pH 7.2, at 10 mg/ml.
使用に関する注意	Wherever possible, you should prepare and use solutions on the same day. However, if you need to make up stock solutions in advance, we recommend that you store the solution as aliquots in tightly sealed vials at -20°C. Generally, these will be useable for up to one month. Before use, and

prior to opening the vial we recommend that you allow your product to equilibrate to room temperature for at least 1 hour.

Need more advice on solubility, usage and handling? Please visit our [frequently asked questions \(FAQ\) page](#) for more details.

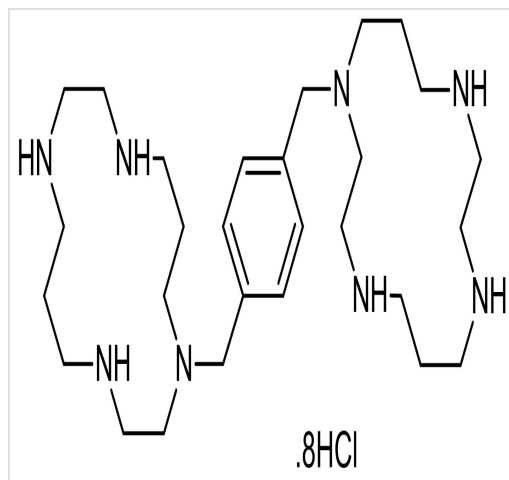
#### SMILES 線形表記

C1CNCCNCCCN(CCNC1)CC2=CC=C(C=C2)CN3CCCNCCNCCCNCC3.Cl.Cl.Cl.Cl.Cl.Cl.Cl.Cl

#### 由来

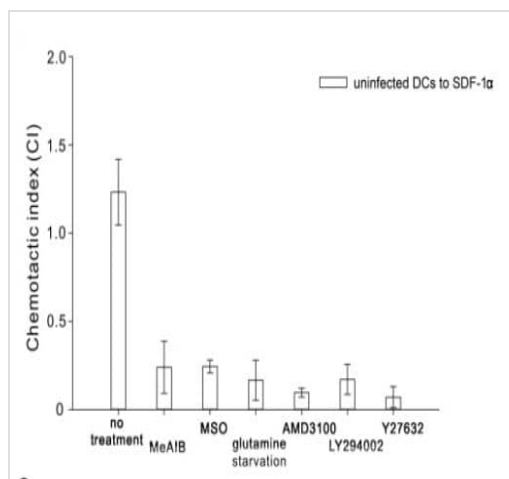
Synthetic

#### 画像



2D chemical structure image of ab120718, AMD3100 octahydrochloride, CXCR4 antagonist

Chemical Structure - AMD3100 octahydrochloride, CXCR4 antagonist (ab120718)



Cellular activation - AMD3100 octahydrochloride, CXCR4 antagonist (ab120718)

Image from Lee IP, et al. Plos One, 9(10), e109803. Fig 3B; doi: 10.1371/journal.pone.0109803

Uninfected control DCs were treated with MeAIB, MSO, inhibitors of CXCR4 (AMD3100), PI3K (LY294002, [ab120243](#)) or Rho kinase (Y27632, [ab120129](#)), or Gln starvation for 2 hours before assessing migration to 100 ng/ml SDF-1  $\alpha$ . Chemotactic index (CI) is defined as the fold increase in the number of migrating DCs to SDF-1  $\alpha$  over the spontaneous migration. One-way ANOVA reveals an effect of pharmacological treatments on the SDF-1  $\alpha$ -induced migration ( $F(6,44) = 6.700$ ,  $P < 0.001$ ). Asterisks indicate  $P < 0.05$  (Dunnett's post hoc).

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES, NOT FOR USE IN HUMANS"

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