

# Product Sheet



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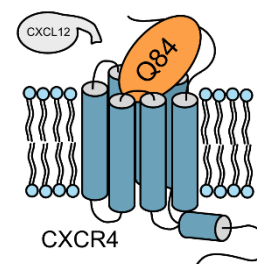
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## CXC chemokine receptor type 4 (CXCR4) / fusin

**Catalogue no.:** Q84c  
**Clone name:** QX4-C2 / VUN400

**Product:** VHH directed against CXCR4  
**Target:** The CXC chemokine receptor type 4 (CXCR4 / fusin, UniProtKB P61073) is a 7-transmembrane spanning class A (rhodopsin-like) G protein-coupled receptor (GPCR). Binding of the chemokine CXCL12/SDF1 $\alpha$  activates heterotrimeric Gai, promoting cytoskeleton rearrangements and migration of e.g. immune cells to sites of inflammation. CXCR4 is important during embryonic development and regulates the homing and retention of hematopoietic stem cells in bone marrow. Upregulation of CXCR4 and CXCL12 contributes to the progression and metastasis of many tumor types. In addition, CXCR4 acts as a co-receptor for entry of HIV-1 and HIV-2 into cells.<sup>1-5</sup>



**Source:** Recombinant monoclonal VHH (Llama glama), purified from *S.cerevisiae* using affinity chromatography. Immunization with CXCR4-containing nanodiscs and cells. Phage-display selection on captured CXCR4-containing lipoparticles with total elution.<sup>5</sup>

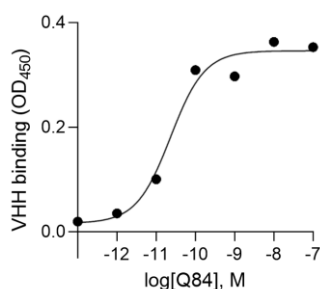
**Specificity:** Q84 and Q85 bind to the extracellular part of human CXCR4 and compete for CXCL12 binding.<sup>5,6</sup>

**Formulation:** 0.2  $\mu$ m filtered solution in PBS. The products are equipped with a C-terminal C-Direct tag with an unpaired cysteine for directional conjugation.

**Mol. Weight:** 15.0 kDa  
**Ext. Coeff. ( $\epsilon$ ):** 20065 M<sup>-1</sup> cm<sup>-1</sup>  
**A<sub>280</sub> at 1g/L:** 1.3

**Storage:** Shipped on blue ice. Store at 4°C or -20°C (aliquots). Addition of 0.02% sodiumazide is optional.

**Applications:** ELISA, IF, antagonism



Binding of Q84 to CXCR4 in immobilized membrane extracts of CXCR4-expressing HEK293T cells.

### References:

- 1 Bleul et al. (1996) Nature 382, 829-833
- 2 Gonzalo et al. (2000) J Immunol 165, 499-508
- 3 Domanska et al. (2004) Eur J Cancer 49, 219-230
- 4 Deng et al. (1996) Nature, 381, 661-666
- 5 Jahnnichen et al. (2010) PNAS, 107, 20565-20570
- 6 van Hout et al. (2018) Biochem Pharmacol, 158, 402-40127