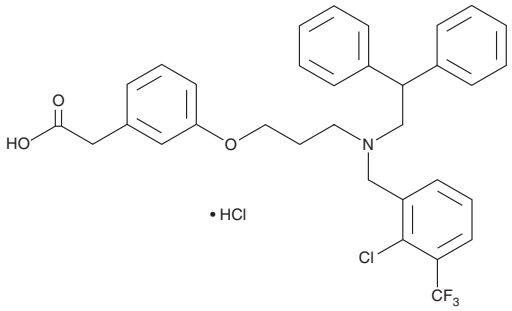
# PRODUCT INFORMATION

## GW 3965 (hydrochloride)

*Item No. 10054*

**CAS Registry No.:** 405911-17-3
**Formal Name:** 3-[3-[[[2-chloro-3-(trifluoromethyl) phenyl]methyl](2,2-diphenylethyl) amino]propoxy]-benzeneacetic acid, monohydrochloride
**MF:** $C_{33}H_{31}ClF_3NO_3 \bullet HCl$
**FW:** 618.5
**Purity:** ≥98%
**Stability:** ≥2 years at -20°C
**Supplied as:** A crystalline solid
**UV/Vis.:** $\lambda_{max}$: 204, 272 nm




### Laboratory Procedures

For long term storage, we suggest that GW 3965 (hydrochloride) be stored as supplied at -20°C. It should be stable for at least two years.

GW 3965 (hydrochloride) is supplied as a crystalline solid. A stock solution may be made by dissolving the GW 3965 (hydrochloride) in the solvent of choice. GW 3965 (hydrochloride) is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of GW 3965 (hydrochloride) in ethanol is approximately 2 mg/ml and approximately 20 mg/ml in DMSO and DMF.

GW 3965 (hydrochloride) is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, GW 3965 (hydrochloride) should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. GW 3965 (hydrochloride) has a solubility of approximately 0.2 mg/ml in a 1:4 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

### Description

The liver X receptors, LXRα and LXRβ, are nuclear receptors that act as ligand-dependent transcription factors.[1] They modulate cholesterol, fatty acids, and glucose homeostasis. GW 3965 is an orally-active agonist of LXRα and LXRβ, activating the human isoforms with $EC_{50}$ values of 190 and 30 nM, respectively.[2] It alters LXR-regulated gene expression in mice and rats, affecting pathways related to glucose and lipid metabolism.[2-4] GW 3965 also affects inflammation and pressor responses through LXRα and LXRβ.[5-7]

### References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5. Joseph, S.B., Castrillo, A., Lafitte, B.A., *et al. Nat. Med.* **9(2)**, 213-219 (2003).
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