
Product Data Sheet

Product Name: ZK 93423 hydrochloride

Cat. No.: GC13922

Chemical Properties

Cas. No. 1216574-52-5

Chemical Name ethyl 6-(benzyloxy)-4-(methoxymethyl)-9H-pyrido[3,4-b]indole-3-carboxylate

SMILES CCOC(C1=NC=C2C(C3=C(N2)C=CC(OCC4=CC=CC=C4)=C3)=C1COC)=OFormula $C_{23}H_{22}N_2O_4 \cdot HCl$ M.Wt 426.9

Solubility <11.71mg/ml in DMSO Storage Desiccate at RT

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: 1 nM

ZK-93423 is an anxiolytic drug from the β -Carboline family, which is a nonbenzodiazepine GABAA receptor agonist. The GABAA receptor is an ligand-gated ion channel and ionotropic receptor.

In vitro: The full agonist ZK-93423 is not subtype selective and stimulates $\alpha 1$, $\alpha 2$, $\alpha 3$, and $\alpha 5$ -subunit containing GABAA receptors equally [$\alpha 1\beta 32$ ($K_i = 4.1$ nM), $\alpha 2\beta 32$ ($K_i = 4.2$ nM), $\alpha 3\beta 32$ ($K_i = 6.0$ nM), $\alpha 5\beta 32$ ($K_i = 4.5$ nM), $\alpha 6\beta 32$ ($K_i > 1000$ nM)] [1]. ZK-93423 has also been used as a base to develop new and improved beta-carboline derivatives and help map the binding site of the GABAA receptor. In rats trained to discriminate PTZ from saline, the PTZ cue was antagonized by ZK 93423, which indicated ZK 93423 may exhibit anxiolytic quality [2].

In vivo: ZK 93423 showd to have anxiolytic properties on animal models of anxiety,

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

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA

Product Data Sheet

including social interaction, Vogel, Geller Serfster, 4-plate test and elevated plus-maze. The anxiogenic and anxiolytic actions are mediated by the BDZ binding sites [3].

Clinical trial: So far, no clinical study has been conducted.

References:

[1] Cox ED, Diaz-Arauzo H, Huang Q, Reddy MS, Ma C, Harris B, McKernan R, Skolnick P, Cook JM. Synthesis and evaluation of analogues of the partial agonist 6-(propyloxy)-4-(methoxymethyl)-beta-carboline-3-carboxylic acid ethyl ester (6-PBC) and the full agonist 6-(benzyloxy)-4-(methoxymethyl)-beta-carboline-3-carboxylic acid ethyl ester (Zk 93423) at wild type and recombinant GABAA receptors. *J Med Chem.* 1998 Jul 2;41(14):2537-52.

[2] Stephens DN, Shearman GT, Kehr W. Discriminative stimulus properties of beta-carbolines characterized as agonists and inverse agonists at central benzodiazepine receptors. *Psychopharmacology (Berl).* 1984;83(3):233-9.

[3] File SE, Baldwin HA. Effects of beta-carbolines in animal models of anxiety. *Brain Res Bull.* 1987 Sep;19(3):293-9.

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

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA