



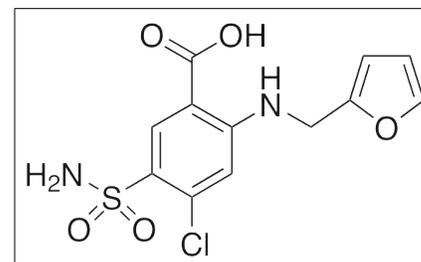
LKT Laboratories, Inc.

## Furosemide

Phone: 888-558-5227  
651-644-8424  
Fax: 888-558-7329  
Email: [getinfo@lktlabs.com](mailto:getinfo@lktlabs.com)  
Web: [lktlabs.com](http://lktlabs.com)

### Product Information

**Product ID** F8270  
**CAS No.** 54-31-9  
**Chemical Name** 5-(Aminosulfonyl)-4-chloro-2-[(2-furanylmethyl)- amino] benzoic acid  
**Synonym** Furseimide, Beronald, Discoid, Durafurid, Eutensin, Fulsix, Furosedon, Impugan, Katlex, Lasix, Macasirrol, Nicorol, Odemase, Rosemide, Urex  
**Formula** C<sub>12</sub>H<sub>11</sub>ClN<sub>2</sub>O<sub>5</sub>S  
**Formula Wt.** 330.74  
**Melting Point** 203-206 °C  
**Purity** ≥97%  
**Solubility** Soluble in acetone, DMF, and methanol. Slightly soluble in water, chloroform and ether.



**Bulk quantities available upon request**

Product ID	Size
F8270	5 g
F8270	10 g
F8270	25 g

**Store Temp** Ambient

**Ship Temp** Ambient

**Description** Furosemide is a loop diuretic that is clinically used to treat congestive heart failure, edema, and hypertension; it inhibits the Na<sup>+</sup>/K<sup>+</sup>/Cl<sup>-</sup> (NKCC) symporter and CFTR Cl<sup>-</sup> channels in the loop of Henle, decreasing reabsorption of Na<sup>+</sup> and K<sup>+</sup> and increasing secretion of Mg<sup>2+</sup> and Ca<sup>2+</sup>. Furosemide exhibits antihypertensive, anxiolytic, and potentially antiepileptic/anticonvulsant activities. Furosemide noncompetitively inhibits GABA-A receptors and decreases conditioned anxiety in animal models of contextual fear conditioning and fear-potentiated startle.

**References** Ju M, Scott-Ward TS, Liu J, et al. Loop diuretics are open-channel blockers of the cystic fibrosis transmembrane conductance regulator with distinct kinetics. *Br J Pharmacol.* 2014 Jan;171(1):265-78. PMID: 24117047.

Krystal AD, Sutherland J, Hochman DW. Loop diuretics have anxiolytic effects in rat models of conditioned anxiety. *PLoS One.* 2012;7(4):e35417. PMID: 22514741.

Somasekharan S, Tanis J, Forbush B. Loop diuretic and ion-binding residues revealed by scanning mutagenesis of transmembrane helix 3 (TM3) of Na-K-Cl cotransporter (NKCC1). *J Biol Chem.* 2012 May 18;287(21):17308-17. PMID: 22437837.

Korpi ER, Kuner T, Seeburg PH, et al. Selective antagonist for the cerebellar granule cell-specific gamma-aminobutyric acid type A receptor. *Mol Pharmacol.* 1995 Feb;47(2):283-9. PMID: 7870036.

**Caution:** This product is intended for laboratory and research use only. It is not for human or drug use.