

Product Data Sheet

Product Name: Chlorotrianisene

Cat. No.: GC10012

Chemical Properties

Cas. No. 569-57-3

Chemical Name 4,4',4''-(2-chloroethene-1,1,2-triyl)tris(methoxybenzene)

SMILES COC1=CC=C(/C(C2=CC=C(OC)C=C2)=C(Cl)/C3=CC=C(OC)C=C3)C=C1

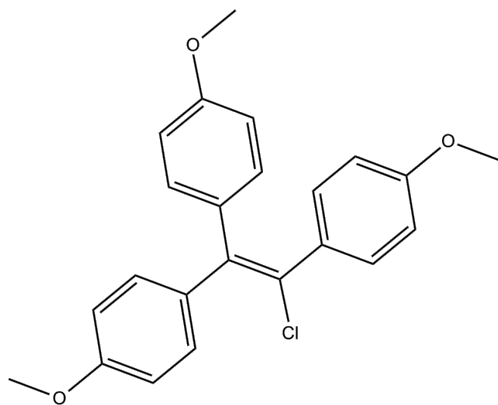
Formula $C_{23}H_{21}ClO_3$ M.Wt 380.86

Solubility $\geq 16.1\text{mg/mL}$ in DMSO Storage Store at -20°C

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

Description:

ED50: 2.0 μM

Glandular kallikrein, a trypsin-like serine protease, has been identified as a major estrogen-induced protein in the rat anterior pituitary. This induction appears to be mediated by increased gene expression since glandular kallikrein mRNA is also estrogen-induced. Chlorotrianisene is classified

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

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as an estrogen agonist.

In vitro: The ED50 of chlorotrianisene were lowered by 3.9-fold. Chlorotrianisene inhibited the growth of both P388 and P388/ADR cells in a concentration-dependent manner [1].

In vivo: Chlorotrianisene markedly varied in its ability to elicit agonist responses on the three measures. On uterine weight and anterior pituitary prolactin, chlorotrianisene behaved as partial agonists. Chlorotrianisene also exhibited strikingly less agonist activity on glandular kallikrein than on PRL or the uterus. Further, the small agonist activity of chlorotrianisene on glandular kallikrein induction differed from that on the uterus or prolactin [2].

Clinical trial: Chlorotrianisene is a estrogen which is highly effective in the suppression of postpartum lactation when given orally in dosage of 48 mg/day for seven days. Recurrence of symptoms and appearance of withdrawal bleeding are virtually eliminated, probably because of the storage of chlorotrianisene in the body fat and its gradual release after cessation of therapy [3].

Reference:

- [1] Ramu A, Glaubiger D, Fuks Z. Reversal of acquired resistance to doxorubicin in P388 murine leukemia cells by tamoxifen and other triparanol analogues. Cancer Res. 1984 Oct;44(10):4392-5.
- [2] Powers CA, Hatala MA, Pagano PJ. Differential responses of pituitary kallikrein and prolactin to tamoxifen and chlorotrianisene. Mol Cell Endocrinol. 1989 Sep;66(1):93-100.
- [3] NULSEN RO, CARMON WB, HENDRICK HO. TACE (chlorotrianisene), a new estrogen for inhibition of lactation. Am J Obstet Gynecol. 1953 May;65(5):1048-51.

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