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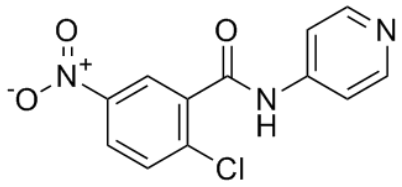
Data Sheet

Global Supplier of Chemical Probes, Inhibitors & Agonists.

Product Name	:	T0070907
Cat. No.	:	PC-43387
CAS No.	:	313516-66-4
Molecular Formula	:	$C_{12}H_8ClN_3O_3$
Molecular Weight	:	277.6632
Target	:	PPAR
Solubility	:	10 mM in DMSO



Biological Activity

T0070907 is a potent, selective, covalent **PPARγ** antagonist with binding IC50 of 1 nM, >800-fold preference for PPARγ over PPARα and PPARδ.
T0070907 specifically blocks PPARγ function in both cell-based reporter gene and adipocyte differentiation assays, with no effect on FXR, LXR, PR.
T0070907 modulates the interaction of PPARγ with cofactor proteins by affecting the conformation of helix 12 of the PPARγ ligand-binding domain (LBD).
T0070907 significantly more efficient at causing cancer cell death than the activators troglitazone and rosiglitazone.

References

Lee G, et al. ***J Biol Chem.*** 2002 May 31;277(22):19649-57.

Schaefer KL, et al. ***Cancer Res.*** 2005 Mar 15;65(6):2251-9.

Masuda T, et al. ***Clin Cancer Res.*** 2005 Jun 1;11(11):4012-21.

Zaytseva YY et al. ***Anticancer Res***. 2011, 31(3), 813-823.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

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