# Thyroxine sulfate

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Cat. No.:	HY-101406	
CAS No.:	77074-49-8	
Molecular Formula:	C <sub>15</sub> H <sub>11</sub> I₄NO <sub>7</sub> S	ļ
Molecular Weight:	856.93	
Target:	Thyroid Hormone Receptor; Drug Metabolite; Endogenous Metabolite	
Pathway:	Vitamin D Related/Nuclear Receptor; Metabolic Enzyme/Protease	1 0
Storage:	-20°C, sealed storage, away from moisture	
	* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

## SOLVENT & SOLUBILITY

In Vitro	DMSO : 140 mg/mL (163.37 mM; Need ultrasonic)					
	Preparing Stock Solutions	Mass Solvent Concentration	1 mg	5 mg	10 mg	
		1 mM	1.1670 mL	5.8348 mL	11.6696 mL	
		5 mM	0.2334 mL	1.1670 mL	2.3339 mL	
		10 mM	0.1167 mL	0.5835 mL	1.1670 mL	
	Please refer to the sol	ubility information to select the app	propriate solvent.			
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 5.75 mg/mL (6.71 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 5.75 mg/mL (6.71 mM); Suspended solution; Need ultrasonic					
	3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.58 mg/mL (3.01 mM); Clear solution					

BIOLOGICAL ACTIVITY				
Description	Thyroxine sulfate is a thyroid hormone metabolite.			
IC <sub>50</sub> & Target	Human Endogenous Metabolite			
In Vitro	Thyroxine sulfate (T4S) is a normal component of human serum and amniotic fluid, and it is mostly derived from thyroxine peripherally and accumulates when type I 5-monodeiodinating activity is low in fetuses or inhibited by drugs, such as ipodate <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

**Product** Data Sheet

#### In Vivo

Significant amounts of thyroxine sulfate (T4S) in fetal sheep serum, meconium, bile, and amniotic and allantoic fluids are observed. T4S concentration in amniotic fluid from women at 18-19 weeks of gestation (25.5 ng/dL) is higher than that at 14-15 weeks of gestation (14.3 ng/dL). A significant rise in serum T4S is detected in hyperthyroid patients 1 day after ingestion of 1 g of ipodate<sup>[1]</sup>. Thyroxine undergoes significant sulfation in rats, and biliary excretion of T4S is enhanced if its type I deiodination is inhibited<sup>[2]</sup>. Serum T4S levels are clearly elevated compared with healthy references, and the decreased deiodination by liver D1 during critical illness appears to play a role in this increase in serum T4S levels<sup>[3]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### **CUSTOMER VALIDATION**

• Toxicology. 2022 Dec 23;153411.

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### REFERENCES

[1]. Wu SY, et al. Identification of thyroxine-sulfate (T4S) in human serum and amniotic fluid by a novel T4S radioimmunoassay. Thyroid. 1992 Summer;2(2):101-5.

[2]. Rutgers M, et al. Effects of propylthiouracil on the biliary clearance of thyroxine (T4) in rats: decreased excretion of 3,5,3'-triiodothyronine glucuronide and increased excretion of 3,3',5'-triiodothyronine glucuronide and T4 sulfate. Endocrinology. 1989 Oct;125(4):2175-86.

[3]. Peeters RP, et al. Increased thyroxine sulfate levels in critically ill patients as a result of a decreased hepatic type Ideiodinase activity. J Clin Endocrinol Metab. 2005 Dec;90(12):6460-5.

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