

Produktinformation



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Diagnostik & molekulare Diagnostik



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PRODUCT INFORMATION



Indoxyl Sulfate-d₅

Item No. 34511

CAS Registry No.: 1644451-34-2

Formal Name: 1H-indol-3-yl-2,4,5,6,7-d₅ sulfate, potassium salt

MF: $C_8HD_5NO_4S \bullet K$

FW: 256.3

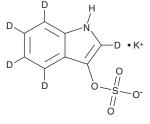
Chemical Purity: ≥95% (Indoxyl sulfate)

Deuterium

Incorporation: \geq 99% deuterated forms (d₁-d₅); \leq 1% d₀

Supplied as: A solid Storage: -20°C Stability: ≥2 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Laboratory Procedures

Indoxyl sulfate-d_s is intended for use as an internal standard for the quantification of indoxyl sulfate (Item No. 16926) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

Indoxyl sulfate- d_5 is supplied as a solid. A stock solution may be made by dissolving the indoxyl sulfate- d_5 in the solvent of choice, which should be purged with an inert gas. Indoxyl sulfate-d₅ is slightly soluble in DMSO and methanol.

Description

Indoxyl sulfate is a uremic toxin and a metabolite of tryptophan (Item No. 29600).¹ It is formed via sulfation of indole, an intermediate generated from tryptophan by intestinal bacteria, by the sulfotransferase (SULT) isoform 1A1 variant 2 (SULT1A1*2) in the liver. 1,2 Indoxyl sulfate activates the aryl hydrocarbon receptor (AhR) in HepG2 40/6 hepatoma cells (EC $_{50}$ = 12.1 nM in a reporter assay).³ It also inhibits the organic anion transporter (OAT) isoforms OAT1 and OAT3 (K_i s = 34.2 and 74.4 μ M, respectively for the rat transporters) in S2 proximal tubule cells. Indoxyl sulfate (0.2 and 1 mM) increases superoxide anion and nitric oxide levels in isolated human mononuclear blood cells.⁵ It increases serum creatinine and blood urea nitrogen (BUN) levels in the 5/6 nephrectomized rat model of chronic renal failure when administered at a dose of 50 mg/kg.⁴

References

- 1. Niwa, T. Uremic toxicity of indoxyl sulfate. Nagoya J. Med. Sci. 72(1-2), 1-11 (2010).
- 2. Banoglu, E. and King, R.S. Sulfation of indoxyl by human and rat aryl (phenol) sulfotransferases to form indoxyl sulfate. Eur. J. Drug Metab. Pharmacokinet. 27(2), 135-140 (2002).
- Schroeder, J.C., Dinatale, B.C., Murray, I.A., et al. The uremic toxin 3-indoxyl sulfate is a potent endogenous agonist for the human aryl hydrocarbon receptor. Biochemistry 49(2), 393-400 (2010).
- Enomoto, A., Takeda, M., Tojo, A., et al. Role of organic anion transporters in the tubular transport of indoxyl sulfate and the induction of its nephrotoxicity. J. Am. Soc. Nephrol. 13(7), 1711-1720 (2002).
- Pieniazek, A., Gwozdzinski, L., Hikisz, P., et al. Indoxyl sulfate generates free radicals, decreases antioxidant defense, and leads to damage to mononuclear blood cells. Chem. Res. Toxicol. 31(9), 869-875 (2018).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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