



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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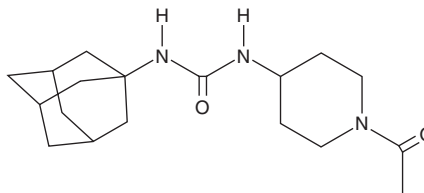
# PRODUCT INFORMATION



**AR-9281**

Item No. 38346

**CAS Registry No.:** 913548-29-5  
**Formal Name:** N-(1-acetyl-4-piperidiny)-N'-tricyclo[3.3.1.1<sup>3,7</sup>]dec-1-yl-urea  
**Synonyms:** APAU, UC1153  
**MF:** C<sub>18</sub>H<sub>29</sub>N<sub>3</sub>O<sub>2</sub>  
**FW:** 319.4  
**Purity:** ≥95%  
**Supplied as:** A solid  
**Storage:** -20°C  
**Stability:** ≥4 years



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

## Laboratory Procedures

AR-9281 is supplied as a solid. A stock solution may be made by dissolving the AR-9281 in the solvent of choice, which should be purged with an inert gas. AR-9281 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of AR-9281 in ethanol is approximately 30 mg/ml and approximately 20 mg/ml in DMSO and DMF.

## Description

AR-9281 is an inhibitor of soluble epoxide hydrolase (sEH; IC<sub>50</sub> = 7 nM for the human enzyme).<sup>1</sup> It reduces LPS-induced increases in plasma levels of dihydroxyeicosatrienoic acids (DiHETs), prostaglandin E<sub>2</sub> (PGE<sub>2</sub>; Item No. 14010), and PGD<sub>2</sub> (Item No. 12010) in rats when administered at a dose of 3 mg/kg.<sup>2</sup> AR-9281 (1 and 3 mg/kg) inhibits LPS-induced allodynia in a rat model of inflammatory pain and increases the paw withdrawal threshold in a rat model of diabetic neuropathic pain induced by streptozotocin (Item No. 13104).

## References

1. Jones, P.D., Tsai, H.J., Do, Z.N., *et al.* Synthesis and SAR of conformationally restricted inhibitors of soluble epoxide hydrolase. *Bioorg. Med. Chem. Lett.* **16(19)**, 5212-5216 (2006).
2. Wagner, K., Inceoglu, B., Dong, H., *et al.* Comparative efficacy of 3 soluble epoxide hydrolase inhibitors in rat neuropathic and inflammatory pain models. *Eur. J. Pharmacol.* **700(1-3)**, 93-101 (2013).

### WARNING

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

### SAFETY DATA

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.

### WARRANTY AND LIMITATION OF REMEDY

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