## PRODUCT INFORMATION



## 4-methoxy-α-Pyrrolidinopentiophenone (hydrochloride)

Item No. 14097

CAS Registry No.: 5537-19-9

Formal Name: 1-(4-methoxyphenyl)-2-(1-pyrrolidinyl)-1-

pentanone, monohydrochloride

Synonyms: 4-MeO-α-PVP,

4-methoxy-α-Pyrrolidinovalerophenone,

4'-methoxy-2-(1-pyrrolidinyl)-Valerophenone

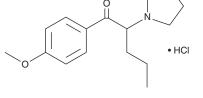
MF:  $C_{16}H_{23}NO_2 \bullet HCI$ 

FW: 297.8 **Purity:** ≥98%

 $\lambda_{\text{max}}$ : 225, 291 nm UV/Vis.: A crystalline solid Supplied as:

-20°C Storage: Stability: ≥5 years

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



## Description

Pyrrolidinophenones are a family of compounds that feature a pyrrolidinyl group affixed to the narcotic cathinone. Many have been identified as components of designer drugs. 4-methoxy-α-Pyrrolidinopentiophenone (4-MeO-α-PVP) (hydrochloride) shares structural features with the stimulant  $\alpha$ -PVP. The mechanism of action is unknown for  $\alpha$ -PVP, but is believed to be similar to that of the designer drug MDPV (Item No. 10624), which acts as a norepinephrine-dopamine reuptake inhibitor. 1.2 This product is intended to be used for forensic and research applications.

## References

- 1. Heron, C., Costentin, J., and Bonnet, J.J. Evidence that pure uptake inhibitors including cocaine interact slowly with the dopamine neuronal carrier. Eur. J. Pharmacol. 264(3), 391-398 (1994).
- 2. Meltzer, P.C., Butler, D., Deschamps, J.R., et al. 1-(4-methylphenyl)-2-pyrrolidin-1-yl-pentan-1-one (pyrovalerone) analogs. A promising class of monoamine uptake inhibitors. J. Med. Chem. 49(4), 1420-1432 (2006).

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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