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# Certificate of Analysis

## www.tocris.com

Print Date: May 4th 2022

### Product Name: Norketamine hydrochloride

Catalog No.: 1970

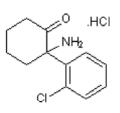
Batch No.: 4

CAS Number:79499-59-5IUPAC Name:2-Amino-2-(2-chlorophenyl)cyclohexanone hydrochloride

## 1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: Batch Molecular Weight: Physical Appearance: Solubility: C<sub>12</sub>H<sub>14</sub>CINO.HCI.H<sub>2</sub>O 278.18 White solid water to 100 mM phosphate buffered saline to 100 mM Desiccate at RT

Storage: Batch Molecular Structure:



### 2. ANALYTICAL DATA

TLC:	R <sub>f</sub> = 0.3 (DCM / MeOH/NH4OH (19:0.9.0.1))			
HPLC:	Shows 99.8% purity			
<sup>1</sup> H NMR:	Consistent with structure			
Mass Spectrum:	Consistent with structure			
Microanalysis:	Carbon Hydrogen Nitrogen			
	Theoretical 51.81 6.16 5.04			

Found 51.76 6.49 5.34

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use



# **Product Information**

Batch No.: 4

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#### Product Name: Norketamine hydrochloride

CAS Number: 79499-59-5

IUPAC Name: 2-Amino-2-(2-chlorophenyl)cyclohexanone hydrochloride

#### **Description:**

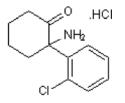
Norketamine hydrochloride is a major metabolite of ketamine that is a potent non-competitive NMDA receptor antagonist ( $K_i$  = 3.6 µM for displacement of [<sup>3</sup>H]-MK 801 in rat brain). Antinociceptive and anesthetic in vivo. R-enantiomer and S-enantiomer also available.

#### **Physical and Chemical Properties:**

Batch Molecular Formula: C<sub>12</sub>H<sub>14</sub>CINO.HCI.H<sub>2</sub>O Batch Molecular Weight: 278.18 Physical Appearance: White solid

#### Minimum Purity: ≥99%

#### **Batch Molecular Structure:**



#### Storage: Desiccate at RT

#### Solubility & Usage Info:

water to 100 mM

phosphate buffered saline to 100 mM

CAUTION - This product is hygroscopic and we recommend that it is desiccated upon arrival. Solutions should be made up as soon as the vial is opened.

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#### Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a 45-60°C water bath).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. Our standard recommendations are:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.

#### Other Information:

#### INFORMATION FOR CUSTOMERS IN THE UK ONLY

This product is a Schedule 2 Home Office controlled substance and customers in the UK are required to hold the relevant licence or be exempt from restrictions in order to purchase and possess this material.

#### Licensing Information:

Sold for research purposes under agreement from Pfizer Inc.

#### **References:**

Shimoyama *et al* (1999) Oral KA is antinociceptive in the rat formalin test: role of the metabolite, norKA. Pain **81** 85. PMID: 10353496. **Ebert** *et al* (1997) NorKA, the main metabolite of KA, is a non-competitive NMDA receptor antagonist in the rat cortex and spinal cord. Eur.J.Pharmacol. **333** 99. PMID: 9311667.

Leung et al (1986) Comparative pharmacology in the rat of KA and its two principal metabolites, norKA and (Z)-6-hydroxynorKA. J.Med.Chem. 29 2396. PMID: 3783598.

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