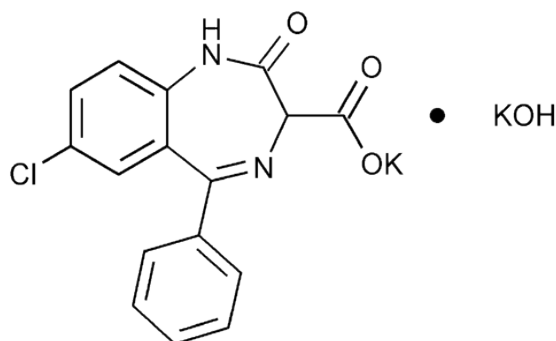




U.S. Pharmacopeia  
The Standard of Quality<sup>SM</sup>

# USP Certificate

## Clorazepate Dipotassium LOT G1F159



### Molecular Formula

$C_{16}H_{11}ClK_2N_2O_4$

### Molecular Weight

408.92

### CAS Number

57109-90-7

### LABEL TEXT



Lot No.: G1F159



### REFERENCE STANDARD

CLORAZEPATE DIPOTASSIUM CIV 125 mg

NDC #00216-0307-04



Warning! Harmful if swallowed. May cause drowsiness or dizziness.  
Suspected of damaging fertility or the unborn child.



Do not dry. For quantitative applications, use a value of 0.994 mg of  
clorazepate dipotassium per mg of material on the as is basis.  
Store under inert gas. Keep container tightly closed. Protect from light.

USP, 12601 Twinbrook Pkwy, Rockville, MD, +1-301-881-0666

CAT No. 1140509

Material mfd in United States

Intentionally over-labeled for GHS compliance

For use with specified USP compendial  
tests. Not for use as a drug. See SDS prior  
to use at [www.usp.org/cds](http://www.usp.org/cds).

Obtain special instructions before use. Do not handle until all safety  
precautions have been read and understood. Use only outdoors or in a  
well-ventilated area. Wash thoroughly after handling. Wear protective  
gloves/protective clothing/eye protection/face protection. If swallowed: Call a  
poison center/doctor if you feel unwell. Rinse mouth. If inhaled: Remove  
person to fresh air and keep comfortable for breathing. Call a poison  
center/doctor if you feel unwell. If exposed or concerned: Get medical  
advice/attention. Store in a well-ventilated place. Keep container tightly  
closed. Store locked up. Dispose of contents/container in accordance with  
local/regional/national/international regulations.

USP certifies that the USP Reference Standards Committee, in accordance with their rules and procedures, determined that this USP Reference Standard lot is suitable to assess compliance with the monograph standards for which it is specified. The critical characteristics of this lot are usually determined independently in three or more laboratories, including USP, government, academic, and industrial collaborators.

*Jeri L. Toth*

QA Director

### **Calculation Value**

Unless otherwise stated on the Reference Standard label, a value of 100.0% should be used in USP or NF compendial applications for which the use of this Reference Standard is intended. Please refer to the specific Reference Standard label for further information.

### **Expiration**

Current lots are identified in the Official USP Reference Standards catalog. In some cases, the previous lot may still be considered official. If so, it is identified in the column marked "Previous Lot/Valid Use Date." Ordinarily, the previous lot is carried in official status for about one year after the current lot enters distribution.

It is the responsibility of each user to determine that this lot is current when used. To ensure up-to-date information, USP publishes the Official USP Reference Standards Catalog, which contains official lot designations. This information is also available on the USP web site, at [www.usp.org](http://www.usp.org), as well as in the bimonthly subscription publication, *Pharmacopeial Forum*.

### **Instructions for Use**

Follow the instructions in the appropriate USP or NF Monographs and General Requirements for Tests and Assays of the current *USP–NF*. In the event that instructions on the label of this lot differ from those found in the current *USP–NF*, those on the label supersede any instructions listed in Chapter <11>.

### **Non-Monograph Use**

The suitability of this Reference Standard for use in non-compendial applications is solely the responsibility of the user.

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