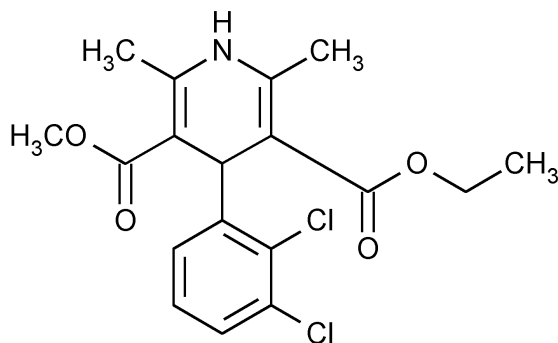




U.S. Pharmacopeia  
The Standard of Quality<sup>SM</sup>

# USP Certificate

## Felodipine LOT H01178



### Molecular Formula

**C<sub>18</sub>H<sub>19</sub>Cl<sub>2</sub>NO<sub>4</sub>**

### Molecular Weight

**384.26**

### CAS Number

**72509-76-3**

### LABEL TEXT



**USP**

### REFERENCE STANDARD

**FELODIPINE 300 mg**

Warning! Harmful if swallowed.  
May cause damage to organs (cardiovascular system).



Do not dry. For quantitative applications, use a value of  
0.998 mg of felodipine per mg of material on the as is basis.  
Keep container tightly closed. Protect from light.

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

CAT No. 1269389

Material mfd in Sweden

Intentionally over-labeled for GHS compliance

For use with specified USP compound at  
certified purity only. See SDS prior  
to use at [www.usp.org](http://www.usp.org).

Wash thoroughly after handling. If swallowed, Call a poison center/doctor if  
you feel unwell. Rinse mouth. If exposed or concerned: Call a poison  
center/doctor. 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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