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The Requirements

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For Single-Unit Containers (see Figure 2)— The average volume of liquid obtained from the 10 containers is not less

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than 100%, and the volume of each of the 10 containers lies within the range of 95% to 110% of the volume declared in the labeling. If A, the average volume is less than 100% of that declared in the labeling, but the volume of no container is outside the range of 95% to 110% ...

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VOLUME. The following tests are designed to provide assurance that oral liquids will, when transferred from the original container, deliver the volume of dosage form that is declared on the label of the article.

usp31nf26s1_c698,

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VOLUME (698): Meets

the requirements for
Oral Suspension

packaged in multiple-
unit containers LIMIT

OF 4-AMINOPHENOL A.

N/1øthnnnl fnrmir and
wafer (7 S' 2 '42 S h

Official Monographs /

Acetaminophen 1569

sonicate for 5 min, and

dilute with Mobile

phase to vol- ume.

Pass a portion of this

solution through a filter

of

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meets the

requirements for oral
suspension packaged
in multiple unit

containers official
monographs

acetaminophen 1569'

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**Usp 37 Monograph -
Maharashtra**

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VOLUME - 2012-10-01

Monograph Title

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VOLUME Errata

Identifier 0f3f3944-736

8-941c-4e43-45627374

a185 Figure 1, right

branch, left box:

Change Volume of 1

more containers is less

than 95% LV to:

Volume of 1 or more

containers is less than

95% LV Section

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2012-10-01 - USP-NF

applications for new

packaging or other

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changes that may affect the fill volume.

36 . 37 In general, ...

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Official Text.

Allowable Excess

Volume and Labeled

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transdermal system.

ters Aerosols, Nasal

Sprays, Metered-Dose

Inhalers, and Dry

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Troche (not preferred;
see Lozenge): A solid
dosage form Powder
Inhalers [601],
Deliverable Volume
[698], Density of

<1160>

**PHARMACEUTICAL
CALCULATIONS IN
PRESCRIPTION
COMPOUNDING**

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chapters Aerosols,
Nasal Sprays, Metered-
Dose Inhalers, and Dry
Powder Inhalers 601,

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699, Osmolality and
Osmolarity 785, pH
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ompounding—Nonsteril
e Preparations 795,
Pharmaceutical
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Gravity 841, Cleaning
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Medicine Dropper 1101
...

General Chapters:

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<1160>

**PHARMACEUTICAL
CALCULATIONS IN ...**

Should you have any questions about this General Chapter, please contact Desmond Hunt (301-816-8341 or dgh@usp.org). For any questions about the PDG and its processes, please see the Pharmacopeial Harmonization Group or contact Richard Lew at (240-221-2060 or

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rll@usp.org).

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**Extractable Volume |
USP**

USP c698 <698>
Deliverable Volume.
Data Sheet by United
States Pharmacopeia,
2009. View all product
details

**USP c698 -
Techstreet**

Figure 1, right branch,
left box: Change
Volume of 1 more
containers is less than

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95% LV to: Volume of 1
or more containers is
less than 95% LV

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VOLUME -

2012-10-01 | USP-NF

English term or phrase:

Deliverable Volume: To

meet the requirements

of the USP (755)

Minimum Fill and (698)

Deliverable Volume

tests, target fill levels

greater than 100%

must be

established. This article

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proposes a criterion for establishing an appropriate target fill level such that a sample will have a 95% probability of passing these USP tests

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English to Spanish |
Medical ...**

childhood dreams to
winning olympic gold,
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volume 698 meets the
requirements,
underwater ocean

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sea life super fun
coloring books for kids,
unit 2 macroeconomics
lesson 3 denton,
twinkle twinkle little
star, trivia questions
and answers for teens,
universe 10th edition,
un

Manual Yamaha Yz 250f

- General Chapter
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Volume and General
Chapter <755>

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Chapter <1087>

Apparent Intrinsic
Dissolution Testing
Procedures for Rotating
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<1216> Tablet

Friability 2. Improving
the Visibility and
Efficacy of
Pharmacopeial Forum
(PF) and Stimuli

**2. Improving the
Visibility and ... -
U.S. Pharmacopeia**

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For Multiple-Unit Containers (see Figure 1)— The average volume of liquid obtained from the 10 containers is NLT 100%, and the volume of no container is less than 95% of the volume declared in the labeling. If A, the average volume is less than 100% of that declared in the labeling, but the volume of no container is less than 95% of the

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labeled amount, or if B,
the average volume is
NLT 100% ...

□□□□-□□□□□□□□

USP develops public
standards. USP is
typically silent on if,
when, or how
frequently to test. If
tested - must pass - for
its entire shelf life.

USP, through its
informational general
chapters, can speak
broadly to standards
development. -

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Through PDG this can be harmonized - Help develop broad, globally-acceptable standards or best ...

Sample Sizes in Uniformity Measurements - The Role of USP

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PHARMACOPEIA 1NF 33
THE NATIONAL FORMULARY Volume
4/a By authority of the
United States

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Pharmacopeial
Convention Prepared
by the Council of
Experts and its Expert
Committees Official
from May 1, 2015 The
designation on the
cover of this
publication, "USP NF
2015," is for ease of
identification only.

**2015 USP 38 THE
UNITED STATES
PHARMACOPEIA**

Assay— Dilute an
accurately measured

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volume of Oral Solution
with water to obtain a
solution containing
about 50 mg of
potassium iodide per
mL. To 10.0 mL of this
solution, in a 150-mL
beaker, add about 40
mL of water, 25 mL of
alcohol, and 1.0 mL of
1 N nitric acid. Titrate
with 0.1 N silver nitrate
VS, determining the
endpoint
potentiometrically,
using silver-calomel
electrodes and a salt ...

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