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USP Chapter 41: Accuracy According to the current USP Chapter 41, the "Accuracy" part of the test describes the quality of the weight to be used. One measurement is taken with a single test

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weight, which is required to have a mass between 5% and 100% of the balance's capacity. The deviation of the measured value should be within 0.10% of the test weight value and the measurement uncertainty of the test weight shall not be more than one-third of 0.10%.

USP General Chapter
<41> -

Scaleman.com

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The New USP Chapter 41 Area of Application.

When it was revised and released in December 2013, the title was shortened to “Balances,” which...

Repeatability.

According to the new USP Chapter 41, “Repeatability” defines the starting point of a balance’s operating...

Accuracy. In order to test a ...

USP Chapter 41

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**Regulations |
Weighing with
Analytical ...**

USP Chapter <41>
weighing requirements
are mandatory in a
Pharmaceutical Quality
Control (QC)
laboratory, where
weighing is a
fundamental step in
almost every workflow.
Typically, weighing of a
sample or standard is
the first step in the
analytical procedure,
followed by dilution

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and subsequent analysis by techniques such as HPLC or qNMR.

USP Chapter 41

Weighing

Requirements for

Balances

41 WEIGHTS AND BALANCES. ... class 4 requirements are met by USP XXI class P.) 2. A weight class is chosen so that the tolerance of the weights used does not exceed 0.1% of the

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amount weighed.

Generally, class 2 may be used for quantities greater than 20 mg, class 3 for quantities of greater than 50 mg, and class 4 for quantities of greater ...

General Chapters: **<41> WEIGHTS AND** **BALANCES**

USP General Chapter 41 "Balances" is mandatory and states the requirements for balances used for

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materials that must be accurately weighed. Weighing should be performed using a balance that is calibrated over the operating range and meets the requirements defined for repeatability and accuracy.

USP Chapters 41 and 1251 on Weighing - Mettler Toledo

This chapter states the requirements for

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balances used for Repeatability is satisfactory if two times the standard deviation of the weighed value, divided by the •desired materials that must be accurately weighed (see General Notices, 8.20). Unless otherwise specified, when substances smallest net weight (i.e., smallest net weight that the users

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**BALANCES (IRA
1-Jul-2014) - USP-NF**

52 [41] Weights and Balances / Apparatus
USP 35 [41] WEIGHTS AND BALANCES tently during or subsequent to the manufacturing process. In the case of sterile articles packaged in multiple-dose con-tainers, antimicrobial preservatives are added to inhibit the growth of microorganisms that

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may be introduced
from re-

**<41> WEIGHTS AND
BALANCES -
DrugFuture**

I-2 Acety-Alumi

Combined Index to USP
41 and NF 36

Acetyltriethyl citrate,

5183 Povidone-iodine

topical, 3392 Alkaline N-

Acetyltyrosine, 4418

Terbutaline sulfate

inhalation, 3986 borate

buffer, 5676 N-Acetyl-L-

tyrosine ethyl ester,

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5665 Thimerosal
topical, 4056 cupric
citrate TS, 5750 Acid
Tolnaftate topical,
4135 cupric citrate TS
2, 5750 acrylic, 5665
Triamcinolone
acetonide topical, 4186
...

**Combined Index to
USP 41 and NF 36,
Volumes 1-5**

One Updated General
Announcement (posted
30-Jul-2020) Three
New Notices of Intent

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to Revise (posted
31-Jul-2020)

Cumulative List

Updated (posted

31-Jul-2020) USP-NF

Components. USP-NF is

a combination of two

compendia, the United

States Pharmacopeia

(USP) and the National

Formulary (NF).

Monographs for drug

substances, dosage ...

USP-NF | USP-NF

The United States

Pharmacopeia (USP)

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was created nearly 200 years ago, dedicated to instilling trust where it matters most: in the medicines, supplements and foods people rely on for their health. The quality standards we develop help manufacturers deliver on their promises of safe products, while building confidence among healthcare ...

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USP Harmonization
Status for General
Chapters (as of
26-Apr-2019) PDG#
Method Name CP PDG
Current Official
harmonization Sign-off
Status Stage 4 Web*
Posting Date(s) G01:
Analytical Sieving: Rev
1 <786> USP: S6,
Rev.1 (08-May-2007)
Link to posting and
signoff history: G02:
Bulk Density and
Tapped Density <616>
EP: S6, Rev.3

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(06-Nov-2013) Stage ...

**Harmonization
Status for General
Chapters | USP**

USP General Chapter
<41> - Scaleman.com
USP Chapter <41>
weighing requirements
are mandatory in a
Pharmaceutical Quality
Control (QC)
laboratory, where
weighing is a
fundamental step in
almost every workflow.

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**Usp General Chapter
41 - mamacz.cz**

2.10. Official Text ble
general chapters, and
General Notices.

Unless specifically
Official text is text
contained in USP and
NF, including
exempted elsewhere in
a compendium, the
identity, monographs,
general chapters, and
these General Notices.
strength, quality, and
purity of an article are
determined by

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GENERAL NOTICES AND REQUIREMENTS - USP-NF

The conductivity of the ubiquitous chloride ion (at the theoretical endpoint concentration of 0.47 ppm when it was a required attribute test in USP XXII and earlier revisions) and the ammonium ion at the limit of 0.3 ppm represents a major portion of the allowed

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water impurity level. A balancing quantity of cations, such as sodium ion, is ...

General Chapters:

<645> WATER CONDUCTIVITY

In addition, the tolerance does not correspond to the value of 0.1%, specified under Weights and Balances 41, for weighing material accurately. Rather, the tolerance is

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purposefully tight to reveal possible drift or calibration errors; this tolerance is readily achievable with modern electronic balances.]

General Chapters:

<1251> WEIGHING ON AN ANALYTICAL BALANCE

The modified USP Chapter 41 standard states, 'Repeatability is assessed by weighing one test weight NLT 10

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times. Repeatability is satisfactory if two times the standard deviation of the weighed value, divided by the nominal value of the weight used, does not exceed 0.10%.

Meeting new USP Chapter 41 requirements

General chapters referenced in HMC monographs may include proposed and official USP-NF general

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chapters. When a general chapter is referenced in a monograph, acceptance criteria may be presented after a colon. The following lists (and links to) the USP-NF general chapters that support HMC monographs.

General Chapters | Herbal Medicines Compendium

USP 31 Microbiological
Tests / ... General

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Considerations 7.2, or Soybean-Casein Digest Broth. If necessary, adjust to a pH of 6 to 8. Further dilutions, where necessary, are prepared with the ability of the test to detect microorganisms in the presence of same diluent.

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