

# Volumetric Measurement in the Laboratory



Learning the basics – how to work with volumetric instruments.



BRAND. For lab. For life.®





# Introduction

Volumetric measurement plays a central role in the laboratory. The user has to determine the degree of accuracy required for each measurement. Based on this, he can choose the appropriate volumetric instrument.

Reliable measurements require the use of precision instruments and their proper handling. To provide a better understanding of volumetric instruments and their operation, this booklet explains the most important terms for their classification and handling, and illustrates them by using BRAND laboratory equipment as examples.

The brochure 'Information on Volumetric Measurement' is designed to give the reader a quick overview of volumetric instruments. It is not intended to replace the safety instructions or operating manuals of the liquid handling instruments described.

Please do not hesitate to contact us if you have any further questions.

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# Volumetric Instruments – an Overview

## Volumetric instruments glass/plastics

The volumetric measurement of liquids is a routine operation in the laboratory. Therefore, volumetric instruments, such as volumetric flasks, bulb pipettes, graduated pipettes, graduated cylinders and burettes are standard equipment. They can be made from glass or plastic. Suppliers offer volumetric instruments in varying qualities. Graduated beakers, beakers, Erlenmeyer flasks, dropping funnels and the like are not volumetric instruments! They are not precisely calibrated, and the scale serves only as an approximate guide. A selection of volumetric instruments from BRAND is illustrated here:



Volumetric flasks



Bulb and graduated pipette



Graduated cylinder



Burette

## Liquid handling instruments

To meet the ever growing demands on volumetric measuring in the lab, such as testing in series, new devices are constantly being developed, e.g., for dispensing, pipetting and titrating. Usually, the instruments designed by different manufacturers for a particular purpose are more or less similar in functional principle. However, major differences are apparent when the focus is on constructional details and design. This selection of liquid handling instruments from BRAND illustrates the range of instruments available:



Bottletop dispenser



Bottletop dispenser



Bottletop burette

Single-channel  
air interface pipettesMultichannel  
air interface pipettesPositive displacement  
pipetteRepetitive pipette,  
manualRepetitive pipette,  
electronic



# Manufacture of Glass Volumetric Instruments

## From raw material to finished precision instrument

### Blanks

Two types of glass are used for the production of blanks:

- soda-lime glass (e.g., AR-GLAS®) for bulb and graduated pipettes
- borosilicate glass (typically Borosilicate glass 3.3) for volumetric flasks, graduated cylinders and burettes.

These glass types meet the stringent laboratory requirements for chemical and physical resilience.



*Blanks for graduated cylinders*

High-quality blanks and strict statistical testing of the required quality characteristics are the basis for producing high-quality volumetric instruments. For example, thermal stress in the glass blanks must be eliminated by a controlled heating and cooling process.

This results in optimum mechanical stability, a requirement for keeping the volume constant despite any subsequent temperature fluctuations. Therefore, BLAUBRAND® and SILBERBRAND volumetric instruments can be heated up to 250 °C in a drying cabinet or sterilizer without any resulting volume changes.

As with all glass instruments, however it should be noted, that uneven heating or sudden

temperature changes produce thermal stresses, which may result in breakage.

Therefore:  
Always place glass instruments into a cold drying cabinet or sterilizer; then heat slowly. At the end of the drying or sterilizing period, allow instruments to cool off slowly inside the switched-off drying cabinet or sterilizer. Never heat volumetric instruments on a hotplate!

### Calibration

Every glass volumetric instrument is individually calibrated at BRAND, i.e. the instrument is accurately filled with a defined quantity of water, and a calibration mark is applied at the lowest point of the meniscus. In the case of graduated instruments, two calibration marks are applied.

Computer-controlled systems ensure maximum precision in a fully automated production line. 'Statistical Process Control' (SPC) guarantees production of volumetric instruments with the smallest deviation from nominal capacity (accuracy) and narrow scatter of individual values (coefficient of variation).

Volumetric instruments are either calibrated 'to contain' ('In') or 'to deliver' ('Ex').

#### Calibration 'to contain' (TC, In):

The contained quantity of liquid corresponds to the capacity printed on the instrument. These instruments include e.g., graduated cylinders, volumetric flasks, and capillary pipettes up to 200 µl.

#### Calibration 'to deliver' (TD, Ex):

The delivered quantity of liquid corresponds to the capacity printed on the instrument. The wetting residue remaining in the instrument has already been taken into account in the calibration. These instruments include e.g., graduated and bulb pipettes, and burettes.

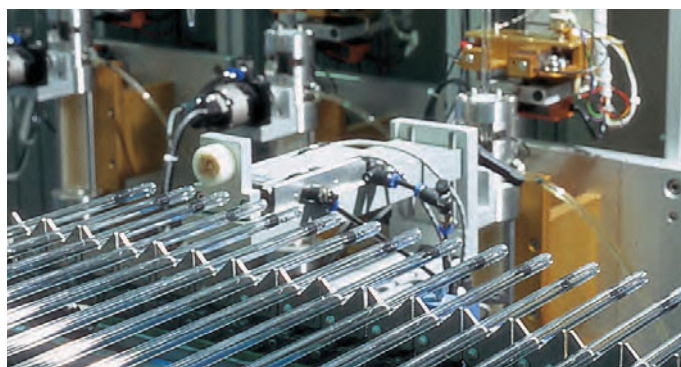
#### Reference temperature

The standard reference temperature, i.e. the temperature at which volumetric instruments will contain or dispense their volumes, is 20 °C.

If the adjustment or calibration is performed at a temperature that deviates from this standard, the corresponding measurement values must be corrected.

#### Note:

Due to the small coefficient of expansion of the glass material, the reference temperature is of minor significance in practical use since the measurement deviations resulting from volume expansion of the measuring instrument are generally smaller than the error limit.



*Calibration of graduated pipettes*



## Silk-screening

Calibration is followed by silk-screen printing of marks and inscriptions. BRAND uses stretchable screen stencils for all graduated pipettes, burettes, graduated cylinders, and mixing cylinders. These stencils can be stretched to match the calibration marks accurately, so that the measuring precision is maintained for all intermediate volumes.

Pipettes are additionally marked with 'color-code' rings at their upper end, which make it easier to identify similar pipette sizes clearly. The ISO 1769 industry standard defines color-coding for different nominal volumes.

## Screening inks

BRAND uses quality inks manufactured especially for glass volumetric instruments:

### Blue enamel:

High contrast, optimum combination of resistance and legibility. Blue enamel is used for BLAUBRAND® volumetric instruments (class A/AS).

### White enamel:

White enamel is used for SILBERBRAND volumetric instruments (class B).

### Amber diffusion stain:

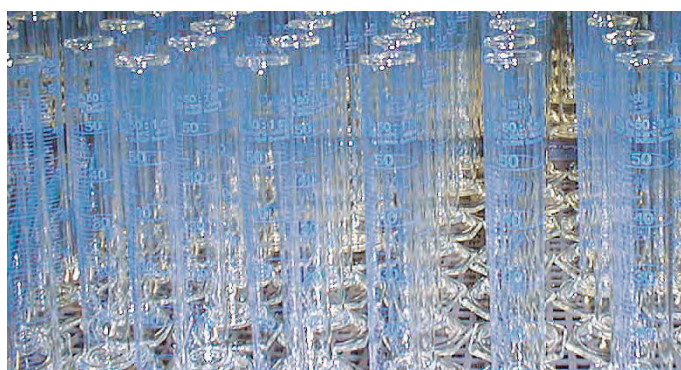
Diffuses into the glass surface and can only be removed by abrasion. It is used for volumetric instruments that are exposed to especially aggressive cleaning conditions. An amber diffusion dye is used for BLAUBRAND® ETERNA volumetric instruments (class A/AS) as well as for SILBERBRAND ETERNA volumetric instruments (class B).



*Automatic silk-screening of volumetric flasks*

## Firing

Firing or annealing the screened ink is the last step on the way from the blank to the finished volumetric instrument. A carefully controlled annealing process, along with the specially produced quality inks, is a prerequisite for durable graduations. This involves a gradual heating and cooling of the volumetric instruments. Depending on the type of glass, temperatures of 400 to 550 °C are reached in this process.



*Silk-screened graduated cylinders before firing*

## Quality assurance

Quality assurance is implemented at BRAND through ongoing testing during production and statistical testing in the final inspection. (For detailed information, see page 39.)

For more information about glass volumetric instruments see the video on our Youtube channel  
**mylabBRAND:**

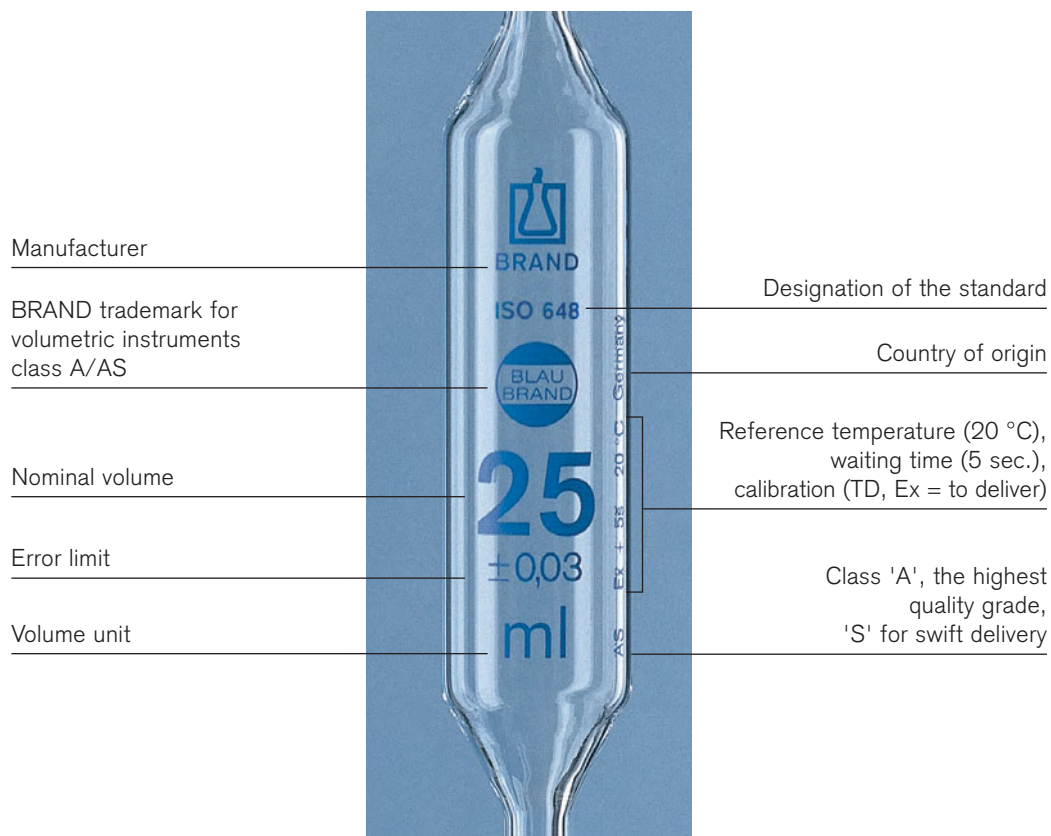


BLAUBRAND® – the right line in volumetric measurement

## Identification of volumetric instruments

Batch 03 with  
year identification DE-M 19

### Example: BLAUBRAND® bulb pipette



The labeling below **must** be printed on every volumetric instrument:

- Nominal capacity
- Unit symbol: ml or cm<sup>3</sup>
- Calibration temperature: 20 °C
- Calibration: Ex or In
- Class: A, AS, AW or B
- Waiting time (if necessary): in the format 'Ex + 5 s'
- Manufacturer's name or logo
- Volumetric flasks and graduated cylinders: glass material

The following information **may** also be added:

- Country of origin
- Error limit
- Manufacturer's trademark (here: BLAUBRAND®)
- Standard, e.g., ISO 648
- Batch number

## Accuracy classification

Volumetric instruments are generally available in two accuracy classes:

### Class A/AS/AW

Volumetric instruments of class A and AS have identical error limits as established by DIN EN ISO. These are generally implemented only in glass volumetric instruments. Exceptions are BRAND plastic volumetric flasks made from PFA and PMP and plastic graduated cylinders made from PMP, which are designed to meet the highest requirements and likewise correspond to class A. Measuring flasks with wide neck are marked with class AW and have a higher error limit compared with class A measuring flasks. For class AS volumetric instruments, calibrated to deliver (TD, Ex), the additional 'S' means swift delivery.

Class AS volumetric instruments have become quite well established. The risk of clogging is low in pipettes and burettes with a larger tip opening. The delivery behavior of various liquids is compensated by observing the defined waiting time (see 'Delivery and waiting times' on page 11).



#### Class A/AS/AW

- Always stands for the highest-grade accuracy
- 'S' stands for swift delivery (pipettes and burettes)
- Only classes A/AS/AW have DE-M marking
- Graduation: The long graduation marks extend over at least 90% of the tube perimeter or are present as ring markings.

### Class B

Volumetric instruments of class B are available in glass or plastic. Class B instruments generally have twice the error limits of class A/AS.

Examples of BRAND class B plastic volumetric instruments are volumetric flasks and graduated cylinders made of PMP or PP, as well as bulb and graduated pipettes made of PP.

For class B measuring instruments calibrated to deliver (TD, Ex), no waiting time is specified.



#### Class B

- Generally twice the error limits of class A/AS
- Graduation: The long graduation marks extend about 20 - 40% of the tube perimeter.

## The choice of volumetric instruments – glass or plastic?

There is no all-round material to meet every single requirement in the laboratory. The decision to use glass or plastic is guided by the application, product design and after considering the specific properties of the materials and economic aspects. Volumetric instruments of plastic excel by their high resistance to breakage and low weight. PP, PMP and PFA are proven materials.

The accuracy of bulb pipettes, graduated pipettes, volumetric flasks, and graduated cylinders made from PP corresponds to that of class B error limits. PMP and PFA are also used for measuring instruments that correspond to class A error limits, e.g., volumetric flasks (PMP/PFA) and graduated cylinders (PMP). Due to its higher purity, PFA is preferably used in trace analysis applications.



*Graduated cylinder made of PMP, class A*

# Working with Volumetric Instruments

## The liquid meniscus

The term meniscus describes a curvature in the surface of the liquid.

The meniscus may be curving upwards or downwards. The curvature develops as a function of the interplay between the adhesion and cohesion forces.

If the liquid molecules are attracted more strongly by the glass wall (adhesion) than by their own kind (cohesion), the meniscus is curved downwards, or concave; the edge of the liquid surface is slightly raised. This is the case, e.g., with aqueous solutions.

If the diameter of a pipette is narrow enough – as with capillary pipettes – the adhesion force is strong enough to pull up not only the edge but also the entire liquid level (capillary effect).

If the cohesion force of a fluid is stronger than the adhesion force of the glass wall, an upwardly curved (convex) meniscus is formed. This happens with mercury, for example.

## Meniscus setting

Correct meniscus setting is a prerequisite for accurate volumetric measurement.



*Concave meniscus in a graduated pipette.*

In the case of a concave meniscus, the volume has to be read at the lowest point of the liquid level. The lowest point of the meniscus has to touch the upper edge of the graduation mark.



*Convex meniscus in a graduated pipette.*

With the meniscus curved upwards, the volume should be read at the **highest** point of the liquid level (edge). During the reading, the **highest** point must touch the upper edge of the graduation mark.



*Appearance of the meniscus at the Schellbach stripe in a burette.*

The Schellbach stripe is a narrow blue band at the center of a white stripe. Schellbach stripes are imprinted on the back of volumetric instruments to improve readability. The refraction of light causes two arrow points to appear at the meniscus. The reading point is where the two arrows meet.

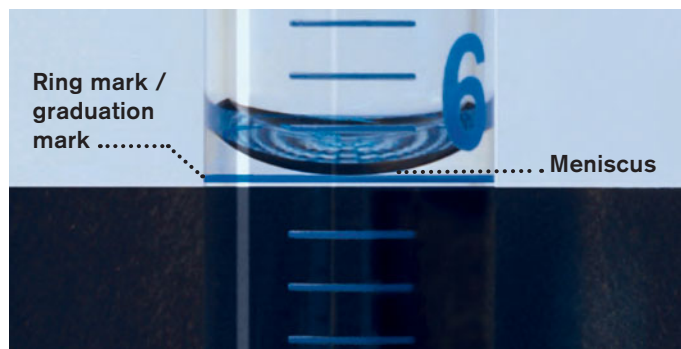
### Important note:

The temperature of the liquid and the environment during use are important. While the expansion of a glass volumetric instrument is negligible, the expansion of the liquid at different temperatures must be taken into account. To minimize volume errors as much as possible, the volumes of all the liquids in contact with one another should be measured at a common (prevailing daily) temperature. Especially in the preparation of standard solutions, for example, the pipetting of the samples and the titration should be done at the same temperature to the extent possible. Significant temperature differences between the measuring instrument and the liquid should likewise be avoided.

## Reading the meniscus

For parallax-free adjustment of the meniscus, the volumetric instrument is held upright and the observer's eye must be at the same height as the meniscus. In this position the ring mark will appear as a line.

The meniscus will appear darker and more easily readable in front of a light background if a piece of dark paper is held behind the instrument immediately beneath the ring mark or graduation mark.



## Delivery and waiting times

In volumetric instruments for liquid delivery (calibrated to deliver, TD, Ex), the volume delivered is always smaller than the volume contained in the measuring instrument. This is caused by the fact that a certain amount of liquid remains as a film on the inner surface of the instrument. The volume of this liquid film depends on the delivery time and should be taken into account when calibrating the measuring instrument.

### Possible volume errors:

The volume delivered from a pipette or burette becomes smaller if the tip is broken off (shorter delivery time), or increases if the tip is not clean and the liquid outflow is impeded (longer delivery time). Likewise, the volume increases if the residual liquid in the tip after pipetting is blown out by mistake. (for the proper handling of pipettes, see page 13.)

### Delivery time

The delivery time is defined as the period of time required for the free fall of the meniscus (discharge of water due to gravity) from the upper volume mark to reach the lower volume mark or the tip. This is connected with the defined waiting time for class AS volumetric instruments.

### Waiting time

The waiting time begins when the meniscus comes to rest at the lower volume mark or in the tip. During the waiting time, residual liquid continues to flow down from the glass wall.

Waiting time for class AS:  
The established waiting time of 5 s for class AS bulb and graduated pipettes is the time after which the meniscus visibly comes to rest in the tip, and before the tip can be removed from the inner surface of the receiving vessel.  
The waiting time of 5 s must be indicated on the pipette by the manufacturer (see page 8).

## Examples of delivery and waiting times for different classes

(25 ml bulb pipette)

### Class AS (DE-M marking)

Delivery time 15 - 20 sec. + Waiting time 5 sec.

### Class A (DE-M marking)

Delivery time 25 - 50 sec. (no waiting time)

### Class B

Delivery time 10 - 50 sec. (no waiting time)



# Working with Volumetric Instruments

## Pipettes, general

Pipettes are volumetric instruments for measuring volumes of liquid and are generally calibrated 'to deliver'. During the manufacturing process, they are individually volumetrically calibrated and provided with one or more calibration marks.

We distinguish between bulb and graduated pipettes (calibrated to deliver, TD, Ex) and disposable micropipettes up to 200 µl (calibrated to contain, TC, In).



Bulb pipette with 1 mark

### Bulb pipettes

- Calibration:  
Class AS: 'Ex + 5 s'  
Class B: 'Ex'
- Generally higher measurement accuracy than graduated pipettes
- Bulb pipette models:  
The most important model is the bulb pipette with 1 mark (total delivery).  
Less common is the model with 2 marks (partial delivery).

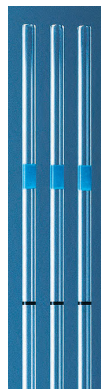
Bulb pipettes are also called single volume pipettes.



Graduated pipette, type 2, nominal volume at the top

### Graduated pipettes

- Calibration:  
Class AS: 'Ex + 5 s'  
Class B: 'Ex'
- Scaling permits the reading of partial volumes
- Types of graduated pipettes:  
Type 2 – nominal volume at top, total delivery also for partial volumes  
Type 1 – nominal volume at bottom, partial delivery for all volumes  
Type 3 – nominal volume at bottom, total delivery only for the nominal volume



### Capillary pipettes e.g., BLAUBRAND® intraMark

- Calibrated to contain (TC, In)
- One ring mark
- Volume limited by one end and the ring mark



### Capillary pipettes e.g., BLAUBRAND® intraEnd

- Calibrated to contain (TC, In)
- No ring mark
- Volume limited by both ends (end-to-end capillaries)

## Handling of pipettes

### Pipettes calibrated 'to deliver' ('TD, Ex')

Proper pipetting with bulb pipettes with 1 mark (here: nominal volume 25 ml) and graduated pipettes type 2, class AS (here: partial volume 3 ml)  
Utility: pipetting aid (see page 18)

#### Filling

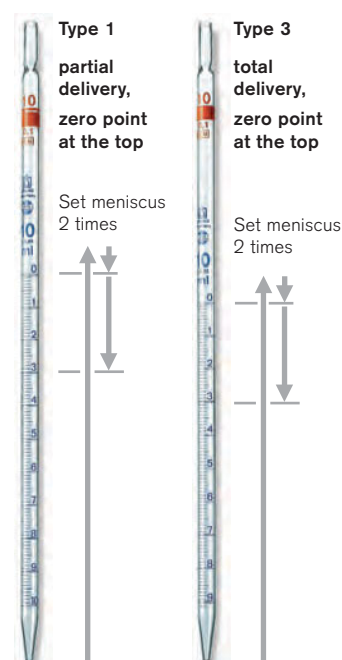
1. Fill the pipette with a pipetting aid to approx. 5 mm above the selected graduation mark.
2. Remove any liquid remaining on the outside of the tip of the pipette with a tissue.
3. Set the meniscus.
4. Wipe off any drop of liquid adhering to the tip.

#### Delivering

5. Hold the pipette vertically. Deliver the liquid with the tip of the pipette in contact with the inner surface of the inclined receiving vessel.
6. When the meniscus comes to a rest in the tip, the waiting time of 5 s begins (class AS only).
7. After the waiting time, draw the pipette tip upwards along the inner wall through a distance of about 10 mm to remove residual liquid.

#### Note:

The residual liquid still left in the tip has already been taken into account during calibration and must not be discharged into the vessel, such as by blowing out.



#### Type 1 and type 3

When using type 1 or type 3 graduated pipettes (zero point at top), the meniscus must

- 1.) first be set at the zero point and then the liquid be run out to slightly above the desired partial volume.
- 2.) After a waiting time of 5 s, the meniscus should be set a second time.



Fill pipette



Wipe off



Set meniscus



Deliver

Working with type 2 graduated pipettes is significantly quicker and simpler. With type 1 and type 3 pipettes there is a risk that with the always necessary second meniscus setting too much liquid is released, and that the sample needs to be prepared again (as also with bulb pipettes that have 2 marks).



## Handling of pipettes

### Pipettes calibrated 'to contain' (TC, In)

Correct pipetting with capillary pipettes

Utility: pipetting aid (see page 18)

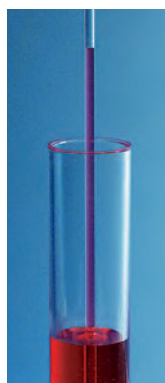
Capillary pipettes are pipettes with a very narrow internal diameter. They are filled either with a pipetting aid, or automatically by capillary action. After emptying, the capillary pipette must be rinsed repeatedly with the diluting medium.

#### Filling

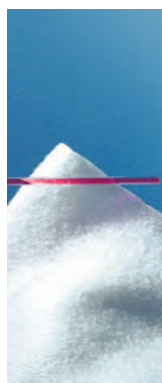
- Aspirate liquid accurately to the desired mark.
- Hold the pipette horizontally and carefully wipe off with a tissue.

#### Delivering

- To empty capillaries blow out the liquid with a pipetting aid, and rinse two to three times with the diluting medium (required due to calibration 'to contain').
- End-to-end capillaries are often placed directly in the dilution solution and washed out by shaking.



*Fill capillary*



*Wipe off*



*Deliver*



*Pipette holder with end-to-end-capillaries*



## Handling of volumetric flasks



Volumetric flasks, class A and B, are calibrated to contain (TC, In), and are mainly used for the preparation of highly accurate dilutions and standard solutions.

Modern analytical methods require small-capacity volumetric flasks. Small-sized conventional volumetric flasks (up to approx. 50 ml) tend to tip over easily due to their high center of gravity and their small base area. Trapezoidal volumetric flasks are much more stable. Their center of gravity is lower, and their base area is more than twice as large compared to the equivalent standard flasks.

How to prepare a standard solution with a volumetric flask:

1. Insert the precisely weighed amount of substance, or rinse in a standard solution concentrate.
2. Fill the flask with distilled water to about half. Swirl the flask to dissolve and mix the contents thoroughly.
3. Top up the flask with distilled water to just below the ring mark.
4. Top up the remaining volume, using a wash bottle or pipette, until the meniscus is exactly at the ring mark. Important: meniscus must be read at eye level. The wall of the flask must not be wetted above the mark.
5. Close the flask and shake upside down to mix contents.

## Handling of graduated and mixing cylinders

### Graduated cylinders:

Graduated cylinders, class A and B, are measuring instruments that are calibrated to contain (TC, In), i.e. they indicate the exact volume taken up.



### Handling:

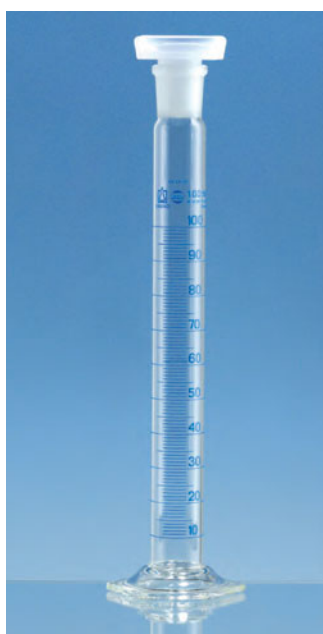
- Fill with liquid.
- Set the meniscus to the required mark (read at eye level!)
- The wall of the cylinder must not be wetted above the mark.
- The indicated volume is the amount of liquid contained.

### Note:

In the laboratory, graduated cylinders are frequently used just like measuring instruments that are calibrated to deliver (TD, Ex). Measurements with water showed that the dispensed volume is reduced by approximately the amount of the error limit of the graduated cylinder due to the residue amount from wetting. Prerequisite: The liquid is slowly dispensed in one portion, and to finish delivery of the liquid the cylinder is held at an incline for a further 30 s.

### Mixing cylinders

The mixing cylinder is calibrated to contain (TC, In), just as the graduated cylinder. They are additionally supplied with a ground glass joint and stopper.



Mixing cylinders can be used in the preparation of standard solutions and dilutions, the same as volumetric flasks.

- After portions of various liquids have been measured out, these can then be mixed by shaking directly in the mixing cylinder.

### Note:

When two liquids are mixed, a volume change may occur.

## Handling of burettes

Burettes are glass volumetric instruments calibrated to deliver (TD, Ex) which are used for titration in volumetric analysis.

### Note on waiting time:

Compared to pipettes, the handling of burettes is different during practical use and calibration. Typically, the volume used in a titration is less than the nominal volume, and the standard solution is added dropwise when close to the color change to avoid overtitration.

### Calibration

Class AS: 'Ex + 30 s'

Class B: 'Ex'

In practice the time required for this dropwise addition is the same or even longer than the established waiting time. As a result, it is not necessary to observe the established waiting time of 30 s with class AS burettes during practical use.

Types of burettes:



*Burette with lateral stopcock*

*Automatic burette, Pellet pattern*

*Automatic burette, Dr. Schilling pattern*

*Compact burette*

### Handling

1. Rinse the burette with the standard solution to be used, and align burette tube vertically.
2. Fill the burette slightly above the zero mark. To prime the stopcock, drain the burette no further than the nominal capacity.
3. Refill the burette with titrant free of air bubbles to approx. 5 mm above the zero mark.
4. Drain liquid to set the zero point accurately. Important: Meniscus must be read at eye level (parallax-free level). Automatic burettes: Fill to approx. 5 mm above the zero mark. This is adjusted automatically after air release.
5. Wipe off any drops adhering to the discharge tip.
6. Open the stopcock and slowly add titrant to the sample (containing the indicator). The discharge tip must not touch the wall of the vessel. Keep swirling the sample vessel lightly while adding titrant, or place it on a magnetic stirrer.
7. Read the discharged volume at eye level.
8. Any drops remaining on the tip of the stopcock should be wiped against the vessel wall and rinsed down. It is part of the titrated volume.

### Notes on titration:

In addition to burettes, volumetric flasks, bulb pipettes and Erlenmeyer flasks are required for titrating. For easiest recognition of the color change, place the sample vessel on a white surface. Make sure that the solution is completely homogenous. There

must be no cloudiness, flocculation or precipitation present. The titration is finished, when the color change is reached. Before each new titration, reset the zero point and start the titration from there.

## Handling of density bottles

Density bottles are mainly used to determine the density of liquids of moderate viscosity. They are not volumetric instruments, however, they are calibrated 'to contain' as in the case of volumetric flasks.



*Density bottle with stopper*

### Handling

1. Determine the weight of the dry and empty density bottle.
2. Fill the density bottle with liquid, avoiding bubbles. The ground neck should be covered to about 1/3.
3. In a thermostatic bath, adjust the temperature of the bottle and contents to 20 °C.
4. Align the stopper respectively, the thermometer of the density bottle according to the marking, and insert carefully. The capillary tube fills up and the displaced liquid comes out.
5. Carefully dry the outer surfaces of the stopper (respectively, the side capillary) and the density bottle with tissue.

#### ATTENTION:

Be careful not to remove any liquid from the capillary. The sample liquid must be exactly level with the upper end of the capillary.

6. Determine the weight of the filled density bottle.

Calculate the density from the mass (weight) and the volume of the liquid at the reference temperature of 20 °C. The volume is engraved on the bottle. The equation is:

$$\text{Density } (\rho) = \text{Mass } (m) / \text{Volume } (V)$$

Take the buoyancy of the air into account for weighing.

#### Note:

Calibrated density bottles always carry a unique identification number on all component parts.

Use only parts with the same number together.

## Working with pipetting aids

Pipetting aids are indispensable for working with pipettes.

### Types of pipetting aids:

- motorized pipetting aids
- manual pipetting aids

Pipetting using the mouth or with a tube and mouthpiece is not allowed. A pipetting aid should always be used for this. This significantly decreases the risk of infection or injury.

### Motorized pipetting aids

Motorized pipetting aids are ideal for pipetting longer series (e.g., for cell cultures).

#### e.g., **accu-jet® pro** from BRAND

The variable motor speed control and a special valve system enable sensitive operation with pipettes from 0.1 to 200 ml.

An integrated check valve in connection with a membrane filter effectively protects against penetrating liquids. To protect against corrosion, an active pressure compensation diverts vapors outside.

### Handling

Pipetting is controlled through two large function buttons:

#### ▲ Filling

To fill the pipette, press the upper button. Intake rate is continuously variable by trigger pressure.

#### ▼ Delivery

Discharge rate is continuously variable by trigger pressure.

Choose:

**Free delivery**

or

**Power delivery**

### Dispensing liquid: free delivery or power delivery?

The choice of delivery modes is determined by the application. In the analytical laboratory, the **'free delivery'** mode is often preferred in order to obtain the required volumetric accuracy. To achieve the accuracy indicated on the pipettes, it is necessary to allow the liquid to run off freely, and to observe

the delivery and waiting times. In microbiology, however, volumetric accuracy is less significant. Here, the uniform and rapid delivery of nutrient solutions, etc. is of primary importance. Therefore, the **'power delivery'** mode is usually preferred in this field.



## Manual pipetting aids

Manual pipetting aids are used when pipetting a short series, primarily in chemical laboratories.

### e.g., macro pipette controller from BRAND

The macro is compatible with the full range of bulb and graduated pipettes from 0.1 to 200 ml. The special valve system allows easy

meniscus adjustment. A hydrophobic membrane filter protects the system against penetrating liquid.

## Handling



### Create negative pressure

Squeeze bellow.



### Filling

Move pipetting lever upward. The farther up the lever is pressed, the faster the pipette will fill.



### Adjust meniscus / dispensing by 'free delivery'

Press pipetting lever slightly downward. The meniscus falls – release the lever, the meniscus stops.

To drain the pipette, press the lever all the way down. To comply with class A accuracy, do not blow out the residual liquid!

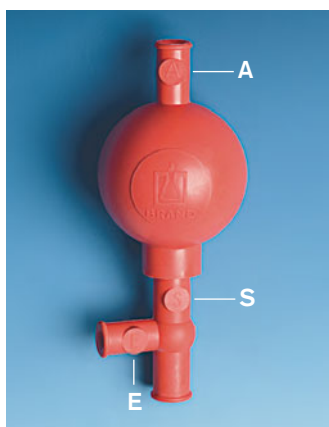


### Blow out

When pipetting viscous media with 'free delivery', the pipette tip frequently does not empty completely. In these cases, blow out any remaining residues by pressing on the rubber bulb of the macro pipette controller.

## Pipette filler

The classic standard pipetting aid for bulb and graduated pipettes.



### Handling

1. Insert top of pipette
2. Press 'A' and squeeze the ball (create negative pressure)
3. Press 'S' to aspirate liquid just above the desired mark
4. Press 'E' to discharge liquid to the desired mark, or to drain the pipette

### Blow out

To blow out viscous media, the side opening must be closed and the small ball compressed.

### Attention!

The pipette filler should not be stored in the primed state, so that no liquid will be taken up!



## Manual pipetting aids for small-volume pipettes up to 1 ml

Special pipetting aids have been developed for these pipettes. They are used in the medical field with capillary pipettes, blood diluting pipettes and blood sugar pipettes up to a max. of 1 ml.

e.g., micro pipette controller from BRAND

### Handling



### Filling / Delivery

Turn the thumb wheel to take in and deliver liquid. Pipettes calibrated to contain (TC, In) should be washed multiple times by filling and emptying with dilution solution.

### Dispensing by 'free delivery'

To deliver liquid from pipettes calibrated to deliver (TD, Ex), press the air release button (observing the waiting time if necessary).

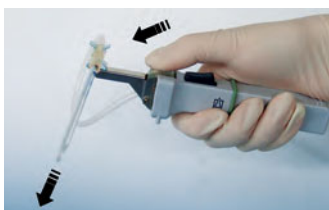
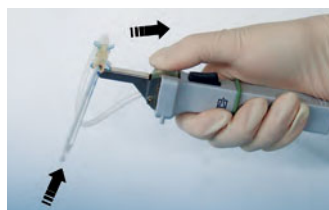
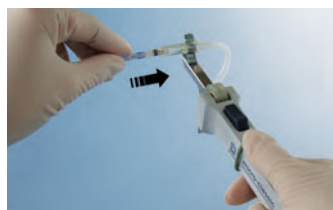
### Eject

The large ejector key allows ejection of used pipettes without hand contact.

e.g., micro-classic pipette controller from BRAND

Due to its angled design, it is especially suitable for work under a microscope in IVF laboratories and in medical laboratories.

### Handling



### Attaching the pipette

Always attach the short end of the pipette, i.e. hold at the color code of the pipette and carefully slide it into the adaptor.

### Filling

Turn the thumb wheel backward until liquid reaches the desired mark.

### Delivery

#### Pipettes calibrated 'to contain':

Turn the thumb wheel forward until liquid is delivered. Rinse the pipette at least three times with the dilution solution.

#### Pipettes calibrated 'to deliver':

For 'free delivery' press the air release button (observe waiting time if required).



# Working with Liquid Handling Instruments

The ever increasing demands on the quality of analytical results, and the growing number of samples to be processed result in a need for volumetric instruments which help to handle the routine work in sample preparation as efficiently as possible.

Manufacturers of laboratory equipment have responded to this need by developing specialized liquid handling instruments. These appliances represent an improvement over the traditional volumetric instruments made of glass or plastic, and allow efficient work with a superior degree of precision and ease of operation.

The liquid handling instruments from most manufacturers share a similar operating principle, however the design details and materials used differ somewhat from one manufacturer to the next.

On the following pages, we will explain the functional principles and application of some of the most common liquid handling instruments, using appliances made from BRAND as examples.



**Bottletop dispenser**  
Dispensette®



**Bottletop dispenser**  
seripettor®



**Bottletop burette**  
Titrette®



**Single-channel  
air interface pipette**  
Transferpette® S  
(manual)



**Multichannel  
air interface pipette**  
Transferpette® S-8/-12  
(manual)



**Single-channel  
air interface pipette**  
Transferpette® electronic



**Multichannel  
air interface pipette**  
Transferpette®-8/-12  
electronic



**Positive displacement  
pipette**  
Transferpettor



**Repetitive pipette**  
HandyStep®  
(manual)



**Repetitive pipette**  
HandyStep® touch  
(electronic)

## Dispensing with bottletop dispensers

### Definition of 'dispensing':

The term 'dispensing' is understood to mean the discharge of defined quantities.

For easy, rapid and precise dispensing of reagents, bottletop dispensers are widely employed to replace poured transfers into graduated cylinders. They can be mounted directly on commercial laboratory bottles, either directly or by means of adapters. It is no longer necessary to transfer or decant chemicals. Serial dispensing in particular is greatly facilitated.



### Functional principle of bottletop dispensers

By an upward movement of the piston, the preset amount of liquid is aspirated from the reagent bottle into the dispenser cylinder. By the subsequent downward movement of the

piston, the liquid is released through a valve system and the discharge tube.

There is no need to set a meniscus or to observe a waiting time.

We distinguish between bottletop dispensers with **floating** pistons and with **wiping-seal** pistons.



### Bottletop dispensers with floating pistons

This system requires no piston seal and is therefore very durable and maintenance-friendly. The piston fits into the dispensing cylinder without contact. Piston and cylinder are separated by a gap only a few thousandths of a millimeter wide and filled with liquid. This film of liquid acts as a lubricant which makes the piston glide very smoothly.

e.g., Dispensette® from BRAND



### Range of application

For dispensing aggressive reagents, e.g., concentrated acids such as  $\text{H}_3\text{PO}_4$ ,  $\text{H}_2\text{SO}_4$ , bases like NaOH, KOH, saline solutions, as well as many organic solvents: Dispensette® S

For dispensing organic solvents, such as chlorinated and fluorinated hydrocarbons (e.g., trichlorotrifluoroethane and dichloromethane), or acids such as concentrated HCl and  $\text{HNO}_3$ , as well as for trifluoroacetic acid (TFA), tetrahydrofuran (THF) and peroxides: Dispensette® S Organic

For dispensing HF we recommend the use of the Dispensette® S Trace Analysis with a platinum-iridium valve spring.

### Materials

Depending on the requirements, parts that come into contact with liquid are made from various specially resistant materials, e.g., ceramic, platinum-iridium, tantalum, ETFE, PFA.

### Safety always comes first!

When choosing a bottletop dispenser, the safety features of the instrument should be kept in mind. For example, does it reduce the risk of injury due to glass breakage? How does it avoid accidental splashing when the instrument is primed? How is contact with the medium minimized when the dispensing tube is closed?

Likewise, the suitability of the dispenser for the medium to be dispensed should be checked by the user. Information about this can generally be found in the operating manual in the chapter 'Function and limitations of use'. When in doubt, contact the manufacturer directly. Information on maintenance and monitoring of measuring instruments are also found in the operating manual.

### Monitoring of measuring instruments / calibration

In regard to the monitoring of measuring instruments according to ISO and GLP guidelines, the accuracy of volumetric instruments should be checked regularly and recalibrated if necessary (see page 33).

### Bottletop dispensers with wiping-seal piston

In addition to the 'floating piston' operating principle, instruments with a 'wiping-seal' piston are also used. It is frequently reported that these systems require higher operating forces and that the frictional wear can cause defective seals.

e.g., **seripettor®** from BRAND

The design of the system allows a replacement of the complete dispensing cartridge. The somewhat higher operating forces during filling are minimized by a spring with automatic lifting action. The present example reflects a competitively-priced dispenser for simple dispensing tasks in the volume range of 0.2 to 25 ml.

### Range of application and materials

The range of application includes the daily routine dispensing of bases, acids in low concentration, biological buffers, cell culture media, biological detergents and polar solvents.

The **seripettor® pro** dispenser is suitable for dispensing acids such as concentrated HCl, polar solvents (e.g., acetone) and UV-sensitive media. In contrast to the **seripettor®** dispenser, valves made from chemically resistant materials are used in this instrument.



## Titration with bottle-top burettes

### Definition of 'titration':

Titration is a volumetric method used for the quantitative analysis of a dissolved substance.

### How to titrate?

Using a bulb pipette, a defined portion of a sample (liquid with an unknown fraction of dissolved material, e.g., acetic acid) is placed in an Erlenmeyer flask.

After dilution with water, 3 drops of an indicator solution are added. Then, with continuous swirling, a suitable titrant of known concentration (e.g., 0.1 M NaOH) is added from a burette until a color change in the indicator signals the endpoint of the titration.

Using the chemical equation and the volume of titrants used, the amount of substance dissolved in the sample can be calculated.



### Functional principle of bottle-top burettes

Bottle-top burettes are mounted directly upon the reservoir bottle. By the upward movement of the piston, liquid is aspirated from the reagent bottle into the burette cylinder.

By the subsequent downward movement of the piston, the liquid is released slowly, and added to the sample through the discharge tube until the titration is finished, e.g., by change of color.

### Reading the volume

The discharged volume can be read directly from the display of the bottle-top burette. There are no meniscus reading errors.

### e.g., Titrette® from BRAND in 10 ml, 25 ml, and 50 ml sizes



The piston moves when the hand wheels are turned, and this takes up or discharges the liquid. The electronics of the instrument automatically recognize the direction of rotation, whether filling or titration is taking place.

The liquid can be taken in quickly, and can then be delivered exactly, very slowly, drop by drop. A recirculation valve makes it possible to run the liquid back into the bottle during priming. Thus, air bubbles can be removed without loss of medium. The instrument can readily be disassembled in the laboratory for cleaning and maintenance.

### Range of application

It can be used in many applications for aqueous and non-aqueous solutions (e.g., alcoholic KOH) up to 1 M.

### Materials

Parts that come into contact with liquid are made from various specially resistant materials, e.g., borosilicate glass, PTFE, platinum-iridium,  $\text{Al}_2\text{O}_3$  ceramic.

For more information about titrating with bottle-top burettes see the video on our Youtube channel

mylabBRAND:



Titrette® – Titration with the digital bottle-top burette

## Pipetting with air interface pipettes

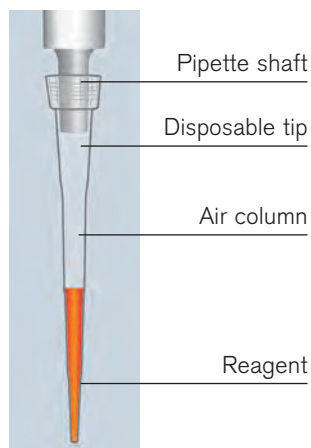
### Definition of 'pipetting':

Pipetting is the accurate one-time uptake and delivery of liquids.

An air interface pipette is used for pipetting aqueous liquids in the microliter to milliliter range. It operates by the air interface principle.

### Functional principle

The up and down movement of the piston inside the pipette shaft creates a negative or positive pressure of the air column. As a result, liquid is either aspirated into the tip or expelled from it. The air column (air interface) keeps the liquid separated from the piston.



### Benefits

There is no wetting of the instrument; the liquid only enters the tip. Tips are used only once, which eliminates any carry-over. This is particularly important for applications where sterile conditions are required, or no carry-over is allowed.

### Calibration

In regard to the monitoring of measuring instruments according to ISO and GLP guidelines, the accuracy of volumetric instruments should be calibrated regularly (i.e. checked) and adjusted if necessary (see page 33).

## Manual single-channel pipettes

e.g., Transferpette® S from BRAND

In the routine lab and in research, precision and functionality are the standards to expect from air interface piston-operated pipettes today.

### Operation



#### Aspirate reagent

1. Press pipetting button to the first stop. Hold the pipette vertically and immerse the tip into the liquid.

Volume range	Immersion depth	Waiting time
0.1 µl - 1 µl	1 - 2 mm	1 s
> 1 µl - 100 µl	2 - 3 mm	1 s
> 100 µl - 1000 µl	2 - 4 mm	1 s
> 1000 µl	3 - 6 mm	3 s

2. Allow the pipetting button to slowly slide back so that the liquid will be aspirated.



#### Discharge reagent

1. Place the pipette tip against the wall of the vessel. Hold the pipette at an angle of 30-45° relative to the container wall and press the pipetting button slowly to the first stop and hold it down.
2. The blow-out stroke empties the tip completely: press the pipetting button down to the second stop and wipe the pipette tip on the vessel wall.



#### Eject tip

Press the tip ejector.





## Manual multichannel pipettes

These pipettes also function by the air interface principle. They allow 8 or 12 pipetting operations to be carried out simultaneously.

Microtiter technology requires pipetting into microtiter plates of 8 x 12 cavities (96-well plates) with standardized spacing. This technology allows e.g., the detection of minute quantities of proteins. This method can only be employed efficiently with multichannel pipettes. Multichannel pipettes are ideal for the efficient transfer of samples, for serial dilutions and for washing of microtiter plates.

### Areas of application

- Clinical diagnostics
- Food analysis
- Immunology
- Biochemistry
- Cell culture

### Analytical techniques

- Immunofluorescence (IF)
- Radio immunoassay (RIA)
- Enzyme immunoassay (EIA, ELISA)
- Cell culture dilution



e.g., Transferpette® S-8/-12 from BRAND

### Operation



#### Aspirate reagent

1. Press pipetting button to the first stop. Hold the pipette vertically and immerse the tips into the liquid.

Volume range	Immersion depth	Waiting time
0.1 µl - 1 µl	1 - 2 mm	1 s
> 1 µl - 100 µl	2 - 3 mm	1 s
> 100 µl - 1000 µl	2 - 4 mm	1 s
> 1000 µl	3 - 6 mm	3 s

2. Allow the pipetting button to slowly slide back so that the liquid will be aspirated.



#### Discharge reagent

1. Place the pipette tips against the wall of the vessel. Hold the pipette at an angle of 30-45° relative to the container wall and press the pipetting button slowly to the first stop and hold it down.
2. The blow-out stroke empties the tips completely: press the pipetting button down to the second stop and wipe the pipette tips on the vessel wall.



#### Eject tips

Press the tip ejector.

Ergonomics and Strain

Intensive, repeated operations done on mechanical instruments without proper ergonomic design result in prolonged stress that can lead to a number of muscular problems, known collectively as RSI (repetitive strain injuries). At particular risk are the muscles in the neck area, shoulders, arms and thumbs. Thus, laboratory work is often accompanied by the appearance of, inter alia, tendonitis and carpal tunnel syndrome. Especially with microliter pipettes, the need for fatigue-free operation is paramount.

Electronic single- and multichannel pipettes

Operating principle

Pressing the pipetting button starts the aspiration or discharge mechanisms (incl. blow-out). The pipette's piston is moved by a motor, aspiration and discharge are controlled by a microprocessor. Various pipetting programs can be selected with the control keys.

Advantages of electronic pipettes

The combination of motor-controlled pipettes and ergonomic design enables stress-free, fatigue-free operation. It also reduces the demands on the thumb while carrying out lengthy series which would otherwise increase the risk of RSI syndrome!

An additional advantage is the execution of pipetting programs such as the gel electrophoresis mode (with precise display of delivered volume) and a dispensing mode, which are not possible with manual pipettes.

e.g., Transferpette® electronic single- and multichannel pipette from BRAND

Program keys .....  
Pipetting button .....  
Tip ejection key .....



With electronic single- and multichannel pipettes, handy design, balanced weight distribution, intuitive software and clear, readily understood technical documentation should be standard.

Operation

Aspirate reagent

Immerse the tip in the reagent and press the pipetting button once – the set volume will be aspirated.

Volume range	Immersion depth	Waiting time
0.1 µl - 1 µl	1 - 2 mm	1 s
> 1 µl - 100 µl	2 - 3 mm	1 s
> 100 µl - 1000 µl	2 - 4 mm	1 s
> 1000 µl	3 - 6 mm	3 s

Discharge reagent

Place the pipette tip against the wall of the vessel. Hold the pipette at an angle of 30-45° relative to the container wall and press the pipetting button again – the liquid will be discharged. The blow-out takes place automatically! During this process, wipe the tip on the vessel wall.

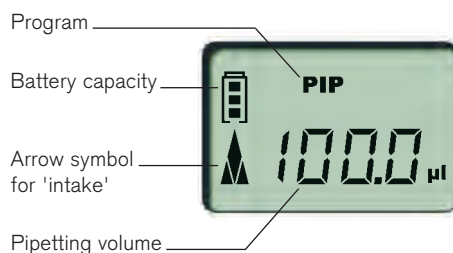
Eject tips

Press the tip ejector.



## Pipetting programs of the microliter pipette Transferpette® electronic

### The display



### The pipetting programs



#### Pipetting

The standard program. The set volume is aspirated by the pipette, and then discharged.



#### Mixing of Samples

Program for mixing of liquids. The sample is repeatedly aspirated and discharged, and the number of mixing cycles is displayed.



#### Reverse Pipetting

Program specially designed for the pipetting of liquids with a high viscosity, high vapor pressure or foamy media. For a set volume, the blow-out volume is aspirated additionally. This volume remains in the tip after delivery to prevent undefined running out, splashing, or the formation of foam or bubbles.



#### Pipetting with Electrophoresis

Program for loading of electrophoresis gels. The required sample volume is aspirated. During discharge, the volume being dispensed is tracked continuously, allowing the user to stop discharge to avoid over-filling sample wells. The pipette records the exact volume dispensed to ensure accuracy of sample mass calculations. GEL mode may also be used for microtitrations.



#### Dispensing

A program for the dispensing of liquids in a series of equal aliquots. A volume that has been aspirated is dispensed in steps.

### Additional functions

Depending on the quality and design, electronic pipettes can offer further instrument-specific functions in addition to the pipetting programs.

The Transferpette® electronic, for example, offers a program to simplify and speed up the calibration of the instrument as well as a battery refresh function.

#### What does 'reverse' mean?

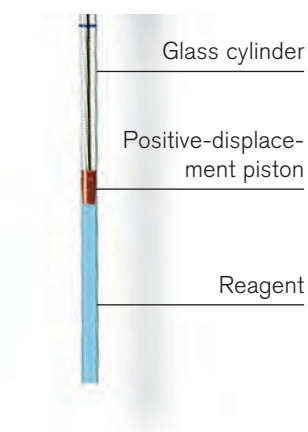
A reversal of the stops used to measure a volume.

Works with mechanical pipettes as follows:

To aspirate the reagent, press the pipetting button down to the **second stop** and let it slide all the way back. Then press down just to the **first stop** to discharge the set volume.

## Pipetting with positive-displacement pipettes

Positive-displacement pipettes are the ideal alternative for difficult applications where air interface pipettes reach their physical limits. They are well suited even for media of very high or low viscosity, high vapor pressure, or a tendency to foam.



### Functional principle

In contrast with air interface pipettes, the positive-displacement piston is in direct contact with the pipetted liquid. The piston wipes the walls of the tips/capillaries completely clean – literally to the last drop which can be observed leaving the tip orifice. This principle always provides reproducible results, regardless of the

physical properties of the liquid. There is no need to discard tips or capillaries after each pipetting operation, since the minimal residual wetting is negligible for most applications. If carry-over is a concern, such as with infectious or radioactive media, an air interface pipette with disposable tips should be preferred.

### Benefits

Highest accuracy and speedy operation. The tips or capillaries are reusable. No need for reading a meniscus for pipetting.

e.g., Transferpettor from BRAND



### Range of application



Highly viscous media, such as highly concentrated protein solutions, oils, resins and fats.



Media with high vapor pressure such as alcohols, ether and hydrocarbons.



Media which tend to foam, such as tenside solutions/surfactants.

### Operation (similar to air interface pipettes)

#### Volume setting

Select desired volume by turning the volume setting knob.

#### Aspirate reagent

Press piston to the stop. Immerse tip into reagent and slowly let the piston return to aspirate reagent.

#### Discharge reagent

Place the capillary/tip against the vessel wall and press the pipetting button down a second time to the stop. Positive-displacement pipettes have no blow-out!

# Dispensing with repetitive pipettes

The distribution of liquids is one of the most important and common activities in medical, pharmaceutical and biological laboratories. The most common techniques are pipetting and dispensing. Dispensing refers to the repeated discharge of identical quantities of liquid. The dispensers described in this chapter eliminate the need for repeated intake after each step – a major time-saving compared to pipetting. Since dispensing is such a common technique, the ergonomic design of the devices plays an essential role.

Dispensing tasks in the laboratory are rarely handled by fully automatic systems requiring no manual intervention. Generally, repetitive pipettes are used for these routine jobs.

- Types of repetitive pipettes:**
- Manual repetitive pipettes
  - Motorized electronic repetitive pipettes



## Functional principle

With manual repetitive pipettes, the volume delivered in each step results from the length of the stroke, defined by the number of steps on a toothed rack, and the size of the tip. Therefore, only a limited number of defined dispensing steps is available.

No intermediate volumes can be selected. A main advantage of these devices is their robustness; their drawback is their often fatiguing operation. Repetitive pipettes work on the proven positive-displacement principle. Therefore, even difficult media with high vapor

pressure, high viscosity or a tendency to foam pose no problem to the repetitive pipettes. According to volume range, the repetitive pipette can be fitted with PD-Tips of different sizes.

## Manual repetitive pipette e.g., HandyStep® from BRAND



The repetitive pipette simplifies serial pipetting by taking up a medium once and then delivering it step by step. With one filling, up to 49 steps between 2 µl and 5 ml can be dispensed, depending on the size of the PD-Tip. The volumes and number of steps are a combined result of the setting of the volume selector key (1-5) and the size of the tip used. Positive-displacement tips (PD-Tips) from BRAND are available in 10 different sizes, sterile or non-sterile. Compatible tips of other manufacturers may also be used.

## Available combinations with positive-displacement tips (PD-Tips) of different sizes from BRAND

Steps and volume ranges

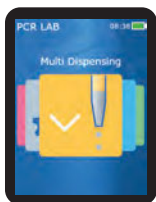
Setting	Tip size (ml)										Steps
	0.1	0.5	1	1.25	2.5	5	10	12.5	25	50	
1	2	10	20	25	50	100	200	250	500	1000	49
1.5	3	15	30	37.5	75	150	300	375	750	1500	32
2	4	20	40	50	100	200	400	500	1000	2000	24
2.5	5	25	50	62.5	125	250	500	625	1250	2500	19
3	6	30	60	75	150	300	600	750	1500	3000	15
3.5	7	35	70	87.5	175	350	700	875	1750	3500	13
4	8	40	80	100	200	400	800	1000	2000	4000	11
4.5	9	45	90	112.5	225	450	900	1125	2250	4500	10
5	10	50	100	125	250	500	1000	1250	2500	5000	9
Volume (µl)											

## Ergonomics and design

Repetitive working with hand-operated appliances can lead to a variety of muscular ailments which may particularly affect the neck, shoulders, arms and thumb. Fatigue-free operation is therefore a crucial requirement for repetitive pipettes, since they are used almost exclusively for serial dispensing. Ergonomic design is indispensable for the stress-free performance of extended pipetting operations in the same working position.

## Motorized electronic repetitive pipettes

### Functional principle



Intake and discharge are controlled by a single key. The piston inside the tips is driven by a motor, with a microprocessor controlling the volume and number of steps.

The ergonomic design results in fatigue-free operation. The piston wipes the walls of the tips completely clean, providing precisely reproducible results without the influences of an air interface.



The positive-displacement tips allow the dispensing of media of high density, high vapor pressure or volatility, or with a tendency to foam.



### e.g., HandyStep® touch from BRAND

Thanks to the patented encoding on the piston head, BRAND Dispenser Tips feature automatic size detection. After insertion, the detected tip size is shown on the display. This prevents errors and both the volume to be dispensed, and the desired work program can now be easily set via the touch display. All the important information is visible at a

glance. Operating the instrument using the STEP button located on the back allows fatigue-free (ergonomic) working. When a new PD-Tip of the same size is inserted, all instrument settings are maintained.

In contrast to manual repetitive pipettes which only permit a limited number of volume

settings, the electronic repetitive pipettes allow the continuous selection of intermediate volumes, such as 1.01 ml.

Automatic tip ejection rounds out the workflow and allows the contamination-free ejection of the tip.



### The HandyStep® touch is available in two versions:

The HandyStep® touch with its three modes covers the requirements of standard applications. The HandyStep® touch S additionally features the modes Sequential Dispensing, Multi-Aspiration and Titration for more comprehensive applications.

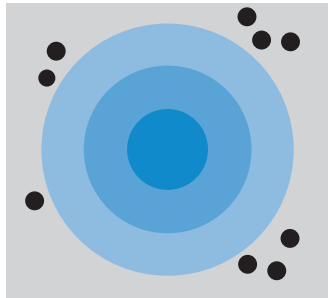
Mode	Function	HandyStep® touch	HandyStep® touch S
Multi Dispensing (MULTI-DISP)	Successive dispensing of a set volume	✓	✓
Auto Dispensing (AUTO-DISP)	Automatic dispensing by the instrument	✓	✓
Pipetting (PIP)	Pipetting a defined volume	✓	✓
Sequential Dispensing (SEQ-DISP)	Successive dispensing of different volumes	—	✓
Multi-Aspiration (MULTI-ASP)	Aspirating different volumes	—	✓
Titration	Manual or successive delivery of liquids for titration	—	✓
Favorites	Saving favorites for quick access	✓	✓
Settings	Individual customization of the device (e.g. brightness or device name)	✓	✓

# Precision Defined

What do 'Error Limit, Accuracy, Coefficient of Variation and Precision' mean in volumetric measuring?

## An illustration of Precision and Accuracy

The dart board simulates the volume range around the centered specified value, the black dots simulate the different measured values of a specified volume.



### Poor accuracy:

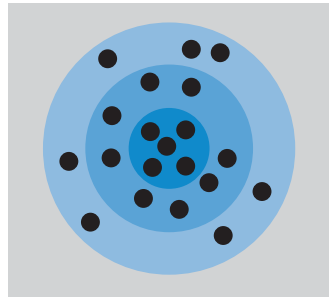
Hits far off center.

### Poor reproducibility:

Hits widely scattered.

### Result:

These volumetric instruments are of inferior quality.



### Good accuracy:

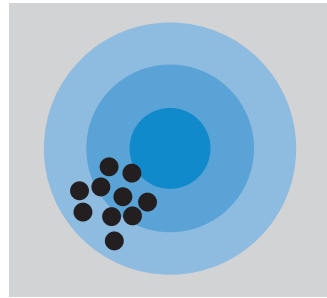
On average, hits are evenly distributed around center.

### Poor reproducibility:

No gross errors, but hits widely scattered.

### Result:

All deviations are 'equally probable'. Instruments exceeding the permissible limit should be removed from service.



### Poor accuracy:

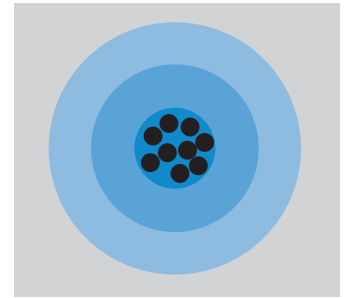
Although all hits are close together, the center (true volume) is still missed.

### Good reproducibility:

All hits are close together.

### Result:

Improperly controlled production, with systematic variation. Instruments exceeding the permissible limit should be removed from service.



### Good accuracy:

All hits are near the center, i.e., the specified value.

### Good reproducibility:

All hits are close together.

### Result:

The volumetric instruments have minute systematic errors, narrow scatter; the permissible limit is not exhausted. These instruments should remain in service.

To describe accuracy, the term "Error limit" is used for glass volumetric devices, while for liquid handling devices the statistical terms "Accuracy [%]" and "Coefficient of Variation [%]" have become established.

### 1 Error limit

The term "Error limit" (EL) in the corresponding standards defines the maximum permissible deviation from the specified value.

### 2 Accuracy (A)

Accuracy (A) indicates the closeness of measured mean volume to the specified value, i.e., systematic measurement variation. Accuracy is defined as the difference between the measured mean volume ( $\bar{V}$ ) and the specified value ( $V_{\text{spec.}}$ ), related to the specified value in percent.

### 3 Coefficient of Variation (CV)

Coefficient of variation (CV) indicates the closeness of values of repeated measurements, i.e., random measurement variation. Coefficient of variation is defined as standard deviation in percent, related to the mean volume.

### 4 Partial volumes

Generally A and CV are related to the nominal volume ( $V_{\text{nominal}}$ ). These values are in % and have to be converted for partial volumes ( $V_{\text{part.}}$ ). In contrast, there is no conversion for partial volumes, if A and CV are stated in volume units (e.g., ml).

### 5 Error limit of A and CV

A good estimate for the error limit (EL) of the instrument, e.g., for the nominal volume ( $V_{\text{nominal}}$ ), can be calculated using the values for accuracy and coefficient of variation.

### 6 Precision (reproducibility)

It describes the closeness in volume units between the different values in a set of measurements.

$$\begin{array}{lllll}
 \text{1} & \text{2} & \text{3} & \text{4} & \text{5} \\
 \text{EL} \geq |V_{\text{measured}} - V_{\text{spec.}}| & A [\%] = \frac{\bar{V} - V_{\text{spec.}}}{V_{\text{spec.}}} \cdot 100 & CV [\%] = \frac{s \cdot 100}{\bar{V}} & A_{\text{part.}} [\%] = \frac{V_{\text{nominal}}}{V_{\text{part.}}} \cdot A_{\text{nominal}} \% & \text{EL} \geq \frac{|A\%| + 2CV\%}{100\%} \cdot V_{\text{nominal}} \\
 & & & (\text{analog } CV_{\text{part.}} [\%]) &
 \end{array}$$

# Monitoring of Measuring Instruments

## Which devices have to be monitored?

All measuring instruments which are used to confirm warranted product qualities are subject to monitoring.

Analytical laboratories have to verify and document the accuracy of all measuring devices used in order to achieve reliable results. This especially applies to laboratories which operate according to GLP guidelines, are accredited to DIN EN ISO/IEC 17025, or are certified to DIN EN ISO 9001. All these standards and guidelines require the availability of written instructions describing the monitoring procedure in detail.

The error limits or accuracy and coefficient of variation have also to be defined, and there must be instructions on how to proceed if the acceptable limits are exceeded.

### Timing and frequency of monitoring

The instrument's accuracy and measuring uncertainty must be known and documented before its admission for use. In addition, the instrument has to be tested again at defined intervals (see DIN EN ISO 10012).

#### Reason:

The performance of measuring instruments may be affected by e.g., the use of aggressive chemicals, and the manner and frequency of cleaning. Since the required accuracy largely depends on the circumstances of each application, the user has to determine the intervals for routine testing.

Typical intervals for liquid handling instruments and plastic volumetric instruments are every 3-12 months, for glass volumetric instruments, every 1-3 years.

### Testing procedures

Volumetric instruments are tested gravimetrically. Testing for liquid handling instruments is performed according to ISO 8655; for glass volumetric instruments, ISO 4787 is applied.

Many influences must be considered when carrying this out. Therefore, BRAND provides Standard Operating Procedures (SOP), including detailed testing instructions, for every type of volumetric instrument. The testing procedure is outlined step by step. To make it even easier, BRAND offers software which can perform all calculations, store them in a database, and print out a detailed test report.

### Time required for testing

The monitoring of measuring devices should not become the main occupation in the laboratory; it should be limited to a reasonable extent. The demand is for simple procedures which are quick and inexpensive to follow.

The combination of testing instructions (SOP), specially developed EASYCAL™ calibration software together with volumetric instruments delivered with individual or batch certificates are best suited to minimize the time needed for this procedure.

### Monitoring of DE-M marking instruments

Volumetric instruments which are DE-M marked to "MessEV" are also subject to the monitoring procedure.

There is no clear directive whether such instruments have to undergo initial testing or not. The user is responsible for answering this question. However, to be on the safe side, the initial test of a representative random sample is recommended. In addition, such a test will document the initial state in relation to subsequent tests.

Another option would be the purchase of volumetric instruments with a certificate from the manufacturer.

## Terms used in the monitoring of measuring devices

### Calibration

Calibration in the strictest sense consists of determining the actual volume delivered.

The calibration procedure should be quick and simple, eliminating potential sources of error. Therefore, BRAND provides detailed testing instructions for every type of volumetric instrument, free of charge.

### Adjustment

Adjustment consists of correcting the deviation of the measured value from the nominal value.

Depending on the manufacturer, the adjustment of liquid handling instruments is generally accomplished by turning an adjustment screw. After setting, a new calibration check is required. This procedure must be repeated until the volume is within the error limits.



## Procedure for volumetric testing

### e.g., Microliter pipette Transferpette® adjustable volume, 20-200 µl

We recommend a calibration of the Transferpette®, as described below, once every 3-12 months. Depending on frequency of use and pipetted media, shorter testing intervals may be defined by the user.

#### Preparation:

1. Verify instrument type and nominal capacity.
2. Read serial number.
3. If instrument is soiled, disassemble and clean if necessary (see operating manual).
4. Check for damage (housing, shaft tip, ejector, piston, seal). Obtain spare parts as required.
5. Place the Transferpette® pipette (unpacked) including accessories into the testing room for at least 2 hours to adjust to room temperature.

#### Functional test:

1. Mount a new pipette tip.
2. Pre-rinse the tip once with testing liquid (distilled/deionized water).
3. Hold the filled pipette vertically and observe for approx. 10 seconds whether a drop is forming at the tip. Make sure the tip is not being heated, e.g., by the sun. Discharge the liquid. In case of smaller volumes (approx. < 50 µl) no drop will form even if there is a leak, due to surface tension.

A tip for finding leaks in small volume pipettes:  
Discharge a small drop from the filled tip so that a small air cushion (air bubble) is present below the liquid. If, during observation, the air cushion falls, there is a leak.

#### Gravimetric test:

1. Determine temperature of the liquid for testing.
2. Mount a new pipette tip.
3. Condition the instrument: aspirate and release testing liquid five times. This will improve the accuracy of the test.
4. Mount a new pipette tip and pre-rinse it once.
5. Place the weighing vessel on the balance and tare.
6. Remove weighing vessel from the balance.
7. Discharge testing liquid into weighing vessel, pressing the pipetting key to the second stop to empty tip completely.
8. Place weighing vessel on the balance. Read and note value.
9. Retare the balance.
10. Repeat steps 2. to 9. ten times. Enter values obtained at 100%, 50% and 10% of nominal volume into the test record.

#### Gravimetric testing values at 21.5 °C (Z = 1.0032)

Tested volume (µl):	200.0000
Specified value (mg):	199.3620
x <sub>1</sub>	200.2000
x <sub>2</sub>	199.6000
x <sub>3</sub>	199.4900
x <sub>4</sub>	199.7000
x <sub>5</sub>	199.7000
x <sub>6</sub>	199.2900
x <sub>7</sub>	199.3500
x <sub>8</sub>	199.4100
x <sub>9</sub>	199.2000
x <sub>10</sub>	199.1900

Values obtained by gravimetric testing only indicate the mass (weight) of the pipetted volume. To obtain the actual pipetted volume, the values have to be multiplied by a correction factor, taking into account the temperature (see below). For all BRAND liquid handling instruments detailed testing instructions are available for download at [www.brand.de](http://www.brand.de).

#### 1. Calculation of the mean volume:

A mean volume ( $\bar{x}$ ) of the weighing values is calculated by dividing the sum of the weighings by the number of weighings made. This mean mass is then multiplied by a correction factor (Z, units of µl/mg) to give the mean volume ( $\bar{V}$ ) delivered. The factor Z combines density of water at the testing temperature and effects of atmospheric pressure. For a typical temperature of 21.5 °C and air pressure of 1013 mbar (hPa), Z = 1.0032 µl/mg.

$$\bar{V} = \bar{x} \cdot Z$$

$$\bar{V} = \frac{x_1 + x_2 + x_3 + \dots + x_{10}}{n} \cdot Z$$

$$\bar{V} = \frac{200.2 + 199.6 + 199.49 + \dots + 199.19}{10} \cdot 1.0032$$

$$\bar{V} = 199.513 \cdot 1.0032$$

$$\bar{V} = 200.1514$$

#### Extract from the table 'Factor Z for liquid handling Instruments'

Temperature °C	Factor z ml/g
18	1.00245
18.5	1.00255
19	1.00264
19.5	1.00274
20	1.00284
20.5	1.00294
21	1.00305
21.5	1.00316
22	1.00327
22.5	1.00338
23	1.00350
23.5	1.00362
24	1.00374
24.5	1.00386
25	1.00399
25.5	1.00412
26	1.00425

\* table refers to 1013 hPa

#### 2. Calculation of accuracy:

$$A [\%] = \frac{\bar{V} - V_{\text{spec.}}}{V_{\text{spec.}}} \cdot 100$$

$$A [\%] = \frac{200.1514 - 200}{200} \cdot 100$$

$$A [\%] = 0.076$$



### 3. Calculation of the standard deviation, necessary for the determination of coefficient of variation

$$s = Z \cdot \sqrt{\frac{(x_1 - \bar{x})^2 + (x_2 - \bar{x})^2 + (x_3 - \bar{x})^2 + \dots + (x_{10} - \bar{x})^2}{n - 1}}$$

$$s = 1.0032 \cdot \sqrt{\frac{(200.2 - 199.51)^2 + (199.6 - 199.51)^2 + (199.49 - 199.51)^2 + \dots + (199.19 - 199.51)^2}{9}}$$

$$s = 1.0032 \cdot \sqrt{\frac{0.8393}{9}}$$

$$s = 0.306$$

### 4. Calculation of the coefficient of variation:

$$CV [\%] = \frac{s \cdot 100}{\bar{V}}$$

$$CV [\%] = \frac{0.306 \cdot 100}{200.1514}$$

$$CV [\%] = 0.153$$

### The result for the calculated example is:

Results of the gravimetric testing:

<b>Tested volume (µl):</b>	200.0000
<b>Mean volume (µl):</b>	200.1514
<b>A [%]</b>	0.076
<b>CV [%]</b>	0.153
<b>A [%] specified*</b>	0.600
<b>CV [%] specified*</b>	0.200

\* Error limits in operating manual.

⇒ This pipette meets specifications.

If the calculated values for Accuracy (A [%]) and Coefficient of Variation (CV [%]) are less than or equal to the error limits, the instrument is calibrated to operate within specifications.

#### Note:

For checking partial volumes, the values  $A_{\text{nominal}} [\%]$  and  $CV_{\text{nominal}} [\%]$  which are related to the nominal volume  $V_{\text{nominal}}$  must be converted.

For a partial volume of 20 µl this means:

$$A_{20 \mu\text{l}} [\%] = \frac{V_{\text{nominal}}}{V_{20 \mu\text{l}}} \cdot A_{\text{nominal}} [\%]$$

$$A_{20 \mu\text{l}} [\%] = \frac{200 \mu\text{l}}{20 \mu\text{l}} \cdot 0.5\%$$

$$A_{20 \mu\text{l}} [\%] = 5\%$$

The calculation of  $CV_{\text{part}}$  is analog.

### What to do if the instrument exceeds the error limits:

1. Review the operating manual to ensure that the instrument was operated properly.
2. Follow the troubleshooting guide in the operating manual for assistance.
3. Recalibrate the instrument in accordance with the operating manual.

If despite these steps the instrument still does not meet the specifications, remove from service and contact the manufacturer for support.

## Calibration software

The monitoring of measuring devices to GLP and DIN EN ISO 9001 is not exactly straightforward. Complex equations easily lead to calculating errors, and the documentation of results can be tiresome. To facilitate this tedious task, some manufacturers have developed special calibration software.

### e.g., EASYCAL™ calibration software from BRAND

EASYCAL™ performs all calculations and generates the complete documentation, automatically. All you need is an analytical balance, a PC, a printer (optional) and EASYCAL™ software.

- Suitable for instruments from all manufacturers
- Specifications of many instruments preloaded
- Testing according to ISO 4787, ISO 8655, etc.

A demo version of the software can be downloaded from the Internet ([www.brand.de](http://www.brand.de)) or can be requested on CD-ROM without charge.

### Monitoring made easy

EASYCAL™ calibration software facilitates the monitoring of measuring devices to GLP and DIN EN ISO 9001, both for liquid handling instruments and for volumetric instruments of glass and plastic. The software is user-friendly and intuitive to use. After determining the type of instrument to be tested, all necessary data are entered step by step on the 'Entry of measuring results' screen.

Two options are available for entering the weighing results:

manual entry, or direct import from the balance via cable, followed by automatic evaluation. After defining the error limits, EASYCAL™ performs all calculations automatically. By one push of a button, a comprehensive test record can be printed out. All results are stored in a database. The test history keeps track of all tested instruments, facilitating the monitoring over time. The test intervals determined in relation to the testing instructions (SOPs) can be defined individually.

## Calibration service for liquid handling instruments

Calibration can be a time-consuming procedure. For this reason, BRAND offers a full calibration service including instrument adjustment and, if required, repair.

Test record printout

# IVD Directive



## Implications and consequences

### IVD Directive 98/79/EC and Regulation (EU) 2017/746

On December 7th, 1998, the EU directive for 'In-vitro-Diagnostic Medical Devices' (IVD Directive) was published in the Official Journal of the European Union and became effective.

The EU regulation 2017/746 for 'In-vitro-Diagnostic Medical Devices' (IVD Regulation) became effective on May 26th, 2017. The transition period ends on May 26th, 2022. Until then the IVD Directive remains valid.

### How to define In-Vitro-Diagnostic Medical Devices (IVD)?

An 'In-Vitro-Diagnostic Medical Device' is any medical device used in-vitro for the examination of specimens, including blood and tissue donations, derived from the human body. IVD can be a reagent, calibrator, control material, kit, instrument, apparatus, equipment, system, or specimen receptacles, intended by the manufacturer to be specifically used for in-vitro diagnostic examination. IVD are mainly used to provide information

- concerning a physiological or pathological state
- concerning a congenital abnormality
- to monitor therapeutic measures.

### What is a Medical Device?

The definition of a 'Medical Device' includes any instrument, apparatus, appliance, material or other article, including the software necessary for its proper application, intended by the manufacturer to be used for human beings for the purpose of:

- diagnosis, prevention, monitoring, treatment or alleviation of disease, injury or handicap
- investigation, replacement or modification of the anatomy or of a physiological process
- control of conception.

Excluded are pharmacological or immunological means, which are regulated by appropriate drug laws.

### CE Marking

The CE mark is the official marking required by the European Community. It shows the user, that this product fulfills all essential safety and environmental requirements as defined in the so-called European Directives. The manufacturer marks the instrument and produces a declaration of conformity describing the instruments' fulfillment with the guidelines and technical requirements.

# Certificates and what they mean

## Standards, requirements, certificates, and explanations

Especially in the pharmaceutical realm, regulations and their compliance assume an immense role. In order not to get lost in the jungle of certificates, the overview on the following page provides you with the most important properties of the most common certificates and statements.

As a rule, one distinguishes between quality certificates (batch and individual certificates), declarations of conformity, and calibration certificates:



① Quality certificate as batch certificate



## Overview of the most important certificates and their characteristics

	① Quality certificate	② Declaration of conformity	③ DAkkS Calibration Certificate
<b>Description</b>	Document that certifies the accuracy of an instrument. Quality certificates are available as individual certificates (with serial number) or batch certificates (with batch number).	Confirmation that the instrument meets the requirements of the German Weights and Measures Act and the ordinances based on it.	Internationally recognized certification of the traceability of the measurement results to national standards.
<b>Issuer</b>	Manufacturer	Manufacturer	Accredited laboratory
<b>Area of application</b>	<ul style="list-style-type: none"> <li>■ To minimize the effort required for initial inspection</li> <li>■ For assuring the demanded traceability of results</li> </ul>	For all instruments in legally regulated areas (e.g. pharmacies, diagnostic laboratories, public offices)	<ul style="list-style-type: none"> <li>■ For especially high-profile calibrations</li> <li>■ For calibration in accredited laboratories</li> </ul>
<b>Reference standard</b>	Selectable: Either manufacturer specifications or industrial standards	Approval for legally regulated areas in accordance with the German Weights and Measures Act and the ordinances based on it.  Marking: <span style="border: 1px solid black; padding: 0 2px;">DE-M</span> 19	ISO/IEC 17 025
<b>Content</b>	<p>All BRAND quality certificates include a declaration of conformity.</p> <p>General:</p> <ul style="list-style-type: none"> <li>■ Manufacturer</li> <li>■ Product</li> <li>■ Batch/serial number</li> <li>■ Single/mean value</li> <li>■ Error of measurement/standard deviation</li> </ul> <p>Additional information on BRAND certificates</p> <ul style="list-style-type: none"> <li>■ All test equipment used</li> <li>■ Measurement method</li> <li>■ Comparative values (volume/error limit)</li> <li>■ QC representative/Date</li> </ul> <p>These specifications are provided either as a mean value of the total batch (batch certificate) or as a value of the individual measuring instrument (individual certificate)</p>	<p>All BRAND declarations of conformity include a quality certificate.</p> <ul style="list-style-type: none"> <li>■ Product designation including ID number</li> <li>■ Date of issue</li> <li>■ Manufacturer</li> <li>■ Laws and technical rules used as a basis</li> <li>■ Signature of a company representative</li> </ul>	<ul style="list-style-type: none"> <li>■ Name of the accreditation body</li> <li>■ Name and address of the issuing laboratory plus accreditation number</li> <li>■ Unique sequential number of the calibration certificate and customer</li> <li>■ Applied methods and conditions (e.g. ambient conditions) under which the calibrations or measurements were executed</li> <li>■ Designation of the calibrated or measured object</li> <li>■ Date of the measurement and date of issue</li> <li>■ Measurement results and measurement uncertainties</li> <li>■ Name, function, and signature of authorized representatives</li> </ul>
<b>Available for</b>	<ul style="list-style-type: none"> <li>■ BLAUBRAND® volumetric instruments and pycnometers</li> <li>■ Class A plastic volumetric instruments</li> <li>■ The following liquid handling instruments: Dispensette®, Titrette®, HandyStep®, Transferpette®, Transferpettor</li> </ul> <p>These instruments are supplied with a batch or individual certificate as standard</p>	<ul style="list-style-type: none"> <li>■ BLAUBRAND® volumetric instruments</li> <li>■ Class A plastic volumetric instruments</li> <li>■ Liquid handling instruments</li> </ul>	<ul style="list-style-type: none"> <li>■ BLAUBRAND® volumetric instruments</li> <li>■ BLAUBRAND® pycnometers</li> <li>■ ASTM centrifuge tubes</li> <li>■ Class A plastic volumetric instruments</li> <li>■ Liquid handling instruments</li> </ul>



# ISO and USP

Many measuring instruments are subject to certain regulations, e.g. laws, guidelines or industry standards. Among others, these regulations define the standards of use, the use of certain measuring instruments for certain applications or specify the design and/or the error limits. Therefore, these regulations are the basis for reliable and reproducible results. For manufacturers of pharmaceutical products for the European market, the EP (European Pharmacopoeia) prescribes the use of glass volumetric instruments in accordance with the current DIN EN ISO standards, which are frequently only designated as ISO (International Organization for Standardization).

Manufacturers of pharmaceutical products for the American market are accredited and audited by the FDA (United States Food and Drug Administration) under the rules of the USP (United States Pharmacopoeia). Both, the USP and the EP are applicable worldwide.

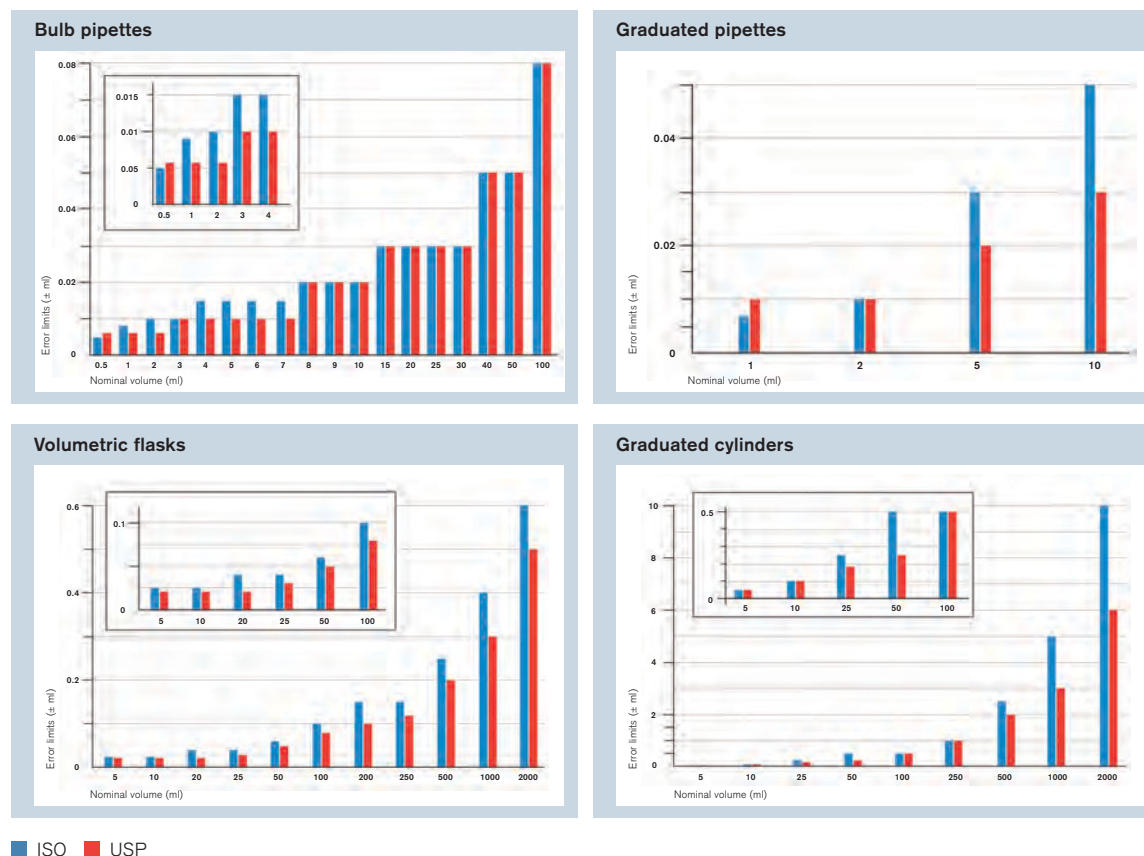
Their respective requirements, however, are not identical. Therefore, measuring instruments such as a 100 ml volumetric flask may exist in two variants (ISO and USP) with different error limits.



USP Certificate of Performance

## Different error limits for different measuring instruments

With many glass volumetric instruments (bulb pipettes, graduated pipettes, volumetric flasks, and graduated cylinders), the error limits are different between ISO and USP. The American and European specifications for the error limit are only identical for burettes. Subject the regulatory environment suitable measuring instruments must be used in order to obtain standard-conforming results.



# Quality Management

Quality management is indispensable. Ideally, it should already begin at project stage, and should accompany a product's design, development and manufacturing process. This guarantees the greatest possible security in working with laboratory equipment, and the reliability of analytical results.

## Quality management at BRAND

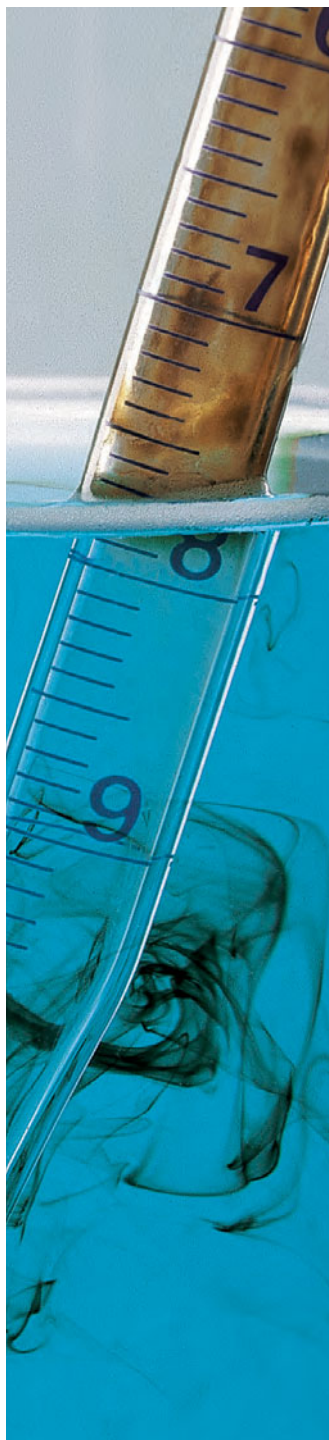
### Quality management is briefly described for liquid handling instruments and BLAUBRAND® volumetric instruments

Quality management at BRAND begins at product conception and continues through the design process and production. Routine checks throughout the entire manufacturing process result in volumetric instruments with the smallest possible deviation from the true volume (accuracy) and narrow scatter of individual values (coefficient of variation). The final step of this Statistical Process Control is random finished product sampling according to DIN ISO 3951.

The quality management system applied at BRAND and certified to DIN EN ISO 9001 is a combination of process monitoring and random checks. The accepted quality level (AQL) is at the very least 0.4., i.e., the limiting values are met with a statistical certainty of at least 99.6 %.

All measuring instruments used in quality control are regularly checked and are referenced to the national standards of PTB (The German Federal Institute of Physics and Metrology). Quality management according to DIN EN ISO 9001 is the basis for issuing of calibration certificates (e.g., our certificates of performance).

All test results are documented and filed for 7 years. If the batch or serial number is known, each specific test result on the date of production can be traced. As BRAND manufactures DE-M marking volumetric instruments, the quality of products is automatically supervised by the 'Eichamt', the German State Office of Weights and Measures. The requirements for monitoring of measuring instruments, traceability to national standards, and staff qualification are fully met.



# Cleaning of Laboratory Equipment

## Manual and machine cleaning

Glass and plastic labware can be cleaned manually, in an immersion bath, or in a laboratory washing machine.

Labware should be cleaned immediately after use – at low temperatures, with brief soaking times, and low alkaline detergents. Labware which has come into contact with infectious substances should first be cleaned and afterwards, if necessary, autoclaved. This is the only way to prevent baking of the substance, and subsequent damage to the labware by any adhering chemical residues.

### Note:

Carefully disinfect labware before cleaning when there is a risk of injury during cleaning procedure.

### Wiping and scrubbing method

The generally accepted wiping and scrubbing method with a cloth or sponge soaked in cleaning solution is the most popular cleaning method. Labware must never be treated with abrasive scouring agents or pads which might damage the surface.

### Immersion method

For the immersion method, labware is soaked in the cleaning solution for 20 to 30 minutes at room temperature, then rinsed with tap water, and finally with distilled water. Only for stubborn residues should the soaking time be extended and the temperature increased.

### Ultrasonic bath

Both glass and plastic labware may be cleaned in an ultrasonic bath. However, direct contact with the sonic membranes must be avoided.

### Machine cleaning

Machine cleaning with a laboratory washing machine is more gentle to labware than cleaning in an immersion bath. The labware is only exposed to the cleaning solution for the relatively short flushing periods when sprayed by the jet or ejector nozzles.

- Lightweight objects will not be tossed and damaged by the jet if they are secured in washing nets.
- Labware is protected against scratching when the wire baskets in the washing machine are plastic coated.

### Glass labware

With glass labware, prolonged immersion times in alkaline media above 70 °C should be avoided. Such treatment, particularly with volumetric instruments, might cause volume changes through glass corrosion, and destruction of graduations.

### Plastic labware

Plastic items generally have smooth, non-wetting surfaces and can usually be cleaned effortlessly under low alkalinity conditions. Polystyrene or polycarbonate labware, e.g., centrifuge tubes, must only be cleaned manually with neutral detergents. Prolonged exposure even to low alkaline detergents will impair their strength. The chemical resistance of these plastics should be verified in each case.

### Cleaning in trace analysis

To minimize metallic traces, laboratory equipment is placed into 1N HCl or 1N HNO<sub>3</sub> at room temperature for not more than 6 hours. (Glass laboratory equipment is often boiled for 1 hour in 1N HNO<sub>3</sub>.) It is then rinsed with distilled water. To minimize organic contamination, laboratory equipment can first be cleaned with alkalis, or a solvent such as alcohol.

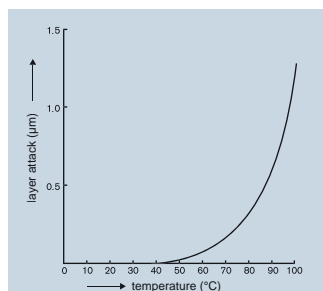
## Gentle cleaning

For gentle treatment of labware, clean immediately after use – at low temperatures, with brief soaking times, and at low alkalinity. Glass volumetric instruments should not be exposed to prolonged immersion times in alkaline media above 70 °C, as such treatment causes volume changes through glass corrosion, and destruction of graduations.

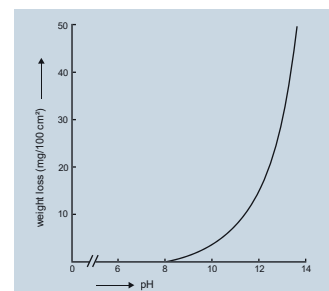
### Information

At 70 °C, a 1N sodium hydroxide solution will corrode a layer of approx. 0.14 µm off the surface of Boro 3.3 (borosilicate glass 3.3) within 1 hour. However, at 100 °C, a layer of

1.4 µm, or tenfold more, will be removed. Therefore, cleaning temperatures above 70 °C should be avoided and low alkaline cleaning agents are preferable.



Alkali attack on Boro 3.3 in relation to temperature, calculated from weight loss. c (NaOH) = 1 mol/l. Exposure time: 1 hour.



Alkali attack on Boro 3.3 in relation to pH value, at 100 °C. Exposure time: 3 hours.

(Graphs are from the brochure 'Technische Gläser' by SCHOTT AG, Mainz.)

## Disinfection and sterilization

### Disinfection

Laboratory instruments that have come into contact with infectious material or genetically modified organisms must be disinfected prior to reuse/disposal; i.e., they must be brought to a condition in which they no longer pose a risk of infection.

Therefore laboratory instruments can be treated with disinfecting detergents for example. If necessary and suitable, the items may subsequently be sterilized (autoclaved).

### Steam sterilization

Steam sterilization (autoclaving) is defined as the destruction or irreversible inactivation of all reproducible microorganisms under exposure to saturated steam at 121 °C (2 bar) according to DIN EN 285.

For correct sterilization procedure, including biological security, please contact your sterilization officer.

### The following points must be observed:

- Efficient steam sterilization is assured only if the steam is saturated and has unrestricted access to all contaminated areas.
- To prevent pressure build-up, containers or vessels must always be open.
- Contaminated reusable labware must be cleaned thoroughly before steam sterilization. Otherwise, residue will bake on during sterilization and microorganisms may not be effectively destroyed if they are protected by the residue. Furthermore, any adhering chemical residues may damage the surfaces due to the high temperatures.
- Not all plastics are resistant to steam sterilization. Polycarbonate, e.g., will lose its strength. Polycarbonate centrifuge tubes cannot be steam sterilized.
- During sterilization (autoclaving), plastic equipment in particular should not be mechanically stressed (e.g., do not stack). Thus, to avoid shape deformation, beakers, flasks, and graduated cylinders should be autoclaved in an upright position.

### Thermal resistance

All reusable BLAUBRAND® and SILBERBRAND volumetric instruments can be heated up to 250 °C in a drying cabinet or a sterilizer, without any subsequent volume deviations. However, as with all glass instruments, irregular heating or sudden temperature changes produce thermal stresses which may result in breakage. Therefore:

- Always place glass instruments into a cold drying cabinet or sterilizer; then heat slowly.
- At the end of the drying or sterilizing period, allow instruments to cool off slowly inside the switched-off oven.
- Do not heat up volumetric instruments on a hot plate.
- Pay attention to the maximum operating temperatures of plastic instruments.

# Safety Information

## Handling of hazardous substances

The handling of hazardous chemicals, infectious, toxic or radioactive substances and genetically modified organisms, calls for a high degree of responsibility on the part of everyone involved, to ensure personal and environmental protection.

The relevant regulations must be scrupulously observed including laboratory, professional association, environmental, radiation, waste disposal and generally accepted technical standards and guidelines (e.g., DIN or ISO).

### Important information on safety

- Before use, laboratory instruments must be examined by the user for suitability and functionality.
- Always hold pipettes near the suction end, and carefully insert the pipette into the adapter of the pipette controller until it is securely and firmly seated. Do not use force. Broken glass can cause injury!
- In the course of repeated use, laboratory instruments should be examined for eventual damage, especially instruments subjected to pressure or vacuum (e.g., desiccators, filter flasks, etc.).
- The hazards of working with defective labware should never be under-estimated (e.g., cuts, burns, risk of infection). If a professional repair is not practical, properly dispose of such items.
- Do not heat volumetric instruments, measuring cylinders and flasks on hot plates.
- Exothermic reactions such as diluting sulfuric acid or dissolving solid alkaline hydroxides must always be carried out while stirring and cooling the reagents, and in suitable vessels such as Erlenmeyer flasks – never in graduated cylinders or volumetric flasks!
- Glass instruments must never be exposed to sudden temperature changes. When taking them out of a drying cabinet while hot, never place on a cold or wet lab bench.
- For compressive loads, only glass instruments intended for this purpose may be used. For example, filtering flasks and desiccators may be evacuated only after confirming that they are in perfect condition.
- Instruments to be repaired must be cleansed of all residues and be sterilized, as necessary. Radioactively contaminated items must be decontaminated as prescribed by ISO. Volumetric glass instruments (e.g., volumetric flasks, graduated cylinders, etc.) should not be repaired when damaged. Exposure to heat may result in residual stress within the glass (greatly increasing the probability of breakage), or an uncontrolled cooling process may lead to permanent volume alterations.
- It is not permissible to simply truncate defective graduated cylinders. This shortens the distance between the upper graduation mark and the spout, as defined by DIN, resulting in an increased danger of chemicals being spilled and the operational safety is no longer guaranteed.
- Waste must be disposed of according to local laws and regulations. This applies also to disposable articles. It must not pose a hazard to human beings or the environment.
- Laboratory equipment must be disposed of according to the materials they are made of, and in a clean state, in accordance with the regulations in force. Please note that laboratory glassware is not recyclable.





## Trademark Index

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