Standard Operating Procedure for Pipettes

Verification Procedure for Accuracy and Precision

(defined for Users to incorporate in SOPs) In accordance with ISO8655 Standard



Contents

Glossary	3
Introduction	4
Environmental Conditions	5
Technician	6
Pipette Operation	6
Training	6
Pipette Tips	7
Test Equipment	8
Balance	8
Thermometer, Hygrometer and Barometer	8
Weighing Containers	9
Water	9
Verification Procedure	10
Procedure Summary	10
Multichannel Pipettes	10
Test Volumes	11
Estimating the Evaporation Rate (Mass Loss/Cycle)	12
Gravimetric Test	13
Calculations	
Formulae	14
Z-factor	15
Verification Procedure Report	16
Appendix	18
Associated Documents	22
Notes	23

In this document the word "tip" is used in the generic sense, where tip is the disposable part that must be used with the pipette: for Distriman this means "DistriTip", for Microman "capillary-piston" and for Pipetman "Gilson Diamond Tip".

Glossary

adjustment manufacture of an apparatus within

appropriate tolerances, or the supplier's setting of the apparatus, ensuring the metrological performance, as specified in the applicable part of

ISO 8655.

calibration set of operations that establish the

relationship between the dispensed volume and the corresponding nominal or selected volume of the

apparatus.

maximum permissible error upper or lower permitted extreme

value for the deviation of the dispensed volume from the nominal volume or selected volume of a piston-operated volumetric appa-

ratus.

systematic error difference between the dispensed

volume and the nominal volume or selected volume of the piston-oper-

ated volumetric apparatus.

random error scatter of the dispensed volumes

around the mean of the dispensed

volumes.

uncertainty of measurement parameter, associated with the dis-

pensed volume, that characterizes the dispersion of the volumes that could reasonably be attributed to

the dispensed volume.

nominal volume volume specified by the manufac-

turer and used for identification and for indication of the measuring

range.

Introduction

This document describes a verification procedure for the following Gilson pipettes: Pipetman® Ultra (Single and Multichannel), Pipetman® Concept (Single and Multichannel), Pipetman® P, Pipetman® Neo, Pipetman® F, Pipetman® 8X200, Microman® and Distriman®.

The procedure is for verifying pipette performance using gravimetric tests of repeated aspirate and dispense cycles with distilled water (grade 3, ISO 3696), in controlled conditions. The test conditions and methods described herein are fully compatible with ISO 8655 and are often stricter than those specified in the international standard, as are the expected results for maximum permissible errors, which are tabulated in the Appendix.

So, adherence to this procedure assures conformity to Gilson's specifications for accuracy (systematic error) and precision (random error) and to ISO requirements.

The procedure, which for small volumes includes a correction for evaporation loss, evaluates the total system of pipetting: pipette, tip, and operator. Therefore the procedure must be carried out by suitably qualified and trained technicians. In calculating the volumes from balance readings, corrections are made for the temperature and air pressure when the test was made (Z-factor, refer to page 15).

Although the document does not directly concern itself with other tests performed by the user, the method and calculations described herein may be applied in other tests, outside the scope of this document. Users shall establish a regular testing routine at least once a year for their piston pipettes according to: accuracy and precision requirements, frequency of use, number of operators using the pipette, number of operations on each occasion of use and the nature of the liquids being dispensed.

In the case of Pipetman Ultra and Pipetman Concept, the number of cycles can be the basis of your regular testing routine.

Environmental Conditions

The test shall be carried out in a draught-free room with a stable environment.

The test room (laboratory) shall have humidity and temperature control so that the atmospheric conditions of the environment where the procedure will take place and the temperature of the equipment used are stable and homogeneous before and during the procedure.

The use of a chart recorder is recommended.

The temperature of the pipettes being verified and the distilled water (grade 3, ISO 3696) used in the gravimetric test should have stabilized before the procedure commences.

The pipettes, water and test apparatus should have been placed in the test room at least 2 hours before starting the tests.

Ideally, verification takes place under the following conditions:

1) Temperature (t)

ISO 8655 recommends that the gravimetric tests take place where the ambient and water temperature (t) are stable (\pm 0.5 °C) between 15 °C and 30 °C. Gilson recommends a range between 20 °C and 23 °C with a constant temperature (\pm 0.5 °C) between the beginning and the end of gravimetric tests. It is recommended to put water and pipettes at least 2 hours in the calibration room to reach an equilibrium with the room conditions. Z-factor is used to convert mass into volume according to temperature and pressure.

2) Relative humidity (RH)

ISO 8655 states that the RH must be greater than 50%. However, Gilson recommends that a humidity range of between 50% and 75% be maintained throughout the verification procedure. In all cases, the evaporation rate will be evaluated for volumes \leq 50 µL.

3) Barometric pressure

Gilson's tests should take place at 1013 ± 25 hPa. The barometric pressure in the test room shall be recorded to the nearest 0.5 kPa. Z-factor is used to convert mass into volume according to temperature and pressure.

Technician

Pipette Operation

Consistency of pipetting technique contributes significantly to the reproducibility of the results of the Verification Procedure. Inexperienced technicians can cause substantial variations in apparent pipette performance. For meaningful test results, the technicians must be well-trained and qualified.

You should operate the pipette according to the instructions given in the user's guide of the pipette under test. Attention should be given to maintain a steady rhythm when aspirating and dispensing samples, speed and smoothness when pressing and releasing the push-button, and tip immersion depth. The test cycle time shall be kept to a minimum. It should not exceed 60 s.

Training

The Verification Procedure described in this document must be carried out by a suitably qualified technician. We strongly recommend that the technician succefully completes a suitable Gilson training program. Please contact your local Gilson distributor for details.

Pipette Tips

In accordance with the instructions given in its user's guide, the pipette under test must be clean (refer to the decontamination procedure), correctly assembled (refer to the user's guide), and fitted with a new Gilson tip before starting the Verification Procedure.

Because the quality of the tip used is a significant factor in ensuring that a pipette performs to specifications, tip selection is specially important in verification procedure.

For example, all models of Pipetman are calibrated at the factory using Gilson Diamond tips, which are of the highest quality. Therefore, for Pipetman, you must only use the Gilson Diamond tips when carrying out the gravimetric test to have the best performance and results.

Pipette model	Volume range	Tips	Filter Tips
P2, P2N, U2 P10, P10N, U10 C10, C8x10, C12x10	0.2 μL to 2 μL 1.0 μL to 10 μL 0.5 to 10 μL	D10, DL10	DF10, DFL10
P20, P20N, U20 U8x20, U12x20	2 μL to 20 μL 1 μL to 20 μL	DL10	DF30
P100N P100, U100, C100 C8x100, C12x100	10 μL to 100 μL 20 μL to 100 μL 5 μL to 100 μL 5 μL to 100 μL	D200	DF100
P200 F2 to F200* P200N, U200 P8x200	50 μL to 200 μL 2 μL to 200 μL 20 μL to 200 μL 20 μL to 200 μL	D200	DF200
U8x300, U12x300 C300 C8x300, C12x300	20 μL to 300 μL	D300	DF300
P1000N P1000, U1000 F250 to F1000	100 μL to 1000 μL 200 μL to 1000 μL 250 μL to 1000 μL	D1000	DF1000
C1200 C8x1200, C12x1200	100 μL to 1200 μL	D1200	DF1200
P5000, U5000 F5000 C5000	1 mL to 5 mL 5 mL 0.5 mL to 5 mL	D5000	
P10ml, U10ml, C10ml	1 mL to 10 mL	D10ml	

*: Not valid with filter tips D200.

Test Equipment

To ensure the integrity of the Verification Procedure, all of the measuring instruments: balances, hygrometer and thermometers should be checked regularly.

Balance

Information on suitable balances (some of which have more than one sensitivity range) is available from the International Organization of Legal Metrology (OIML). Appropriate balances, conforming to OIML R76-1, should be used. Balances should be serviced, calibrated and certified by qualified technicians using weights traceable to an internationally recognized authority (OIML).

Nominal Volume (μL)	Display (mg)	Balance Sensitivity
< 100	0.001	10 ⁻⁶ g
100 μL to 1000	0.01	10⁻⁵g
> 1000	0.1	10⁴g

Note: These requirements are more rigorous than those specified in ISO 8655-6, Table 1.

The sensitivity of the balance chosen must be consistent with the accuracy required, which is one tenth of the deviation to be assessed.

For Pipetman (all models) and Microman, select the sensitivity according to the pipette's nominal volume (see table).

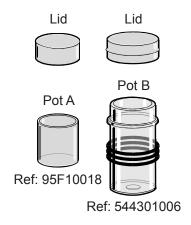
For Distriman, test volumes are specified for each DistriTip model according to specific aliquot volumes (choose the sensitivity accordingly).

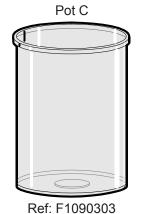
The table on which the balance is placed must be equipped with a marble surface plate that is independent of the perimeter, to avoid transmitting vibrations. For the same reason, the table must not be in contact with a wall. Avoid placing the balance near to a window or near to a door to avoid too long a response time for the balance and irregular evaporation caused by drafts or greenhouse effects.

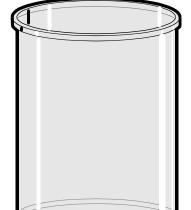
Thermometer, Hygrometer, and Barometer

Use a calibrated thermometer to measure the water temperature at the beginning and at the end of each test series. Use a thermometer with a maximum uncertainty of measurement of 0.2 °C. The hygrometer shall have a standard maximum uncertainty of 10% and the barometer a standard uncertainty of less than or equal to 0.5 kPa.

Test Equipment







Pot D

Ref: F1490343

Weighing Containers

Special containers are used to receive water from the test pipette during weighing. Controlling evaporation during the gravimetric test is essential. To minimize evaporation, Gilson uses custom-designed cylindrical flat-bottomed weighing containers made of nonporous plastic.

A Weighing Kit (ref: F144700) consisting of four sizes of container (Pot A, B, C, and D), filters (ref: F123854) for cleaning them and tweezers (ref: F144706), is available from Gilson. Individual components of the kit are available as spares.

When the volumes to be tested are less than 200 μ L, Gilson uses weighing containers equipped with lids (see below).

Pot A container and lid, both of which must be manipulated using tweezers to avoid handwarming, for volumes up to 20 μ L.

Pot B container and lid for volumes from 20 to 200 μ L. This container is fitted with P 5000 O-rings to avoid hand-warming.

Pot C container (50 mm x 35 mm) for volumes from 200 to 5000 μ L.

Pot D container (70 mm x 50 mm) for volumes greater than 5000 μL.

Water

The liquid used for testing must be distilled or deionized water grade 3 (degassed) conforming to ISO3696 at room temperature. To avoid fluctuations in water temperature, use a large container as a water reservoir (Pot C or Pot D). The reservoir should contain sufficient water for all of the tests.

Procedure Summary

The Verification Procedure certifies both pipette accuracy and precision. Environmental conditions, test equipment, and other qualifications previously described in this document should be implemented to assure the validity of the test results.

After pre-rinsing the tip, record ten individual weighings per selected volume. For variable volume pipettes, three volume settings are selected per pipette model based on the pipette's useful volume range (nominal, approximately 50 % and minimum volume or 10 % of the nominal volume). For fixed volume pipettes (Pipetman F) only the nominal volume is used.

- 1 Set the pipette to its test volume (see table opposite).
- 2 Estimate the evaporation loss (for small volumes).
- 3 Perform the gravimetric test: record the weighings on the Verification Procedure Report.
- 4 Perform the calculations: record the results on the Verification Procedure Report.
- 5 Compare the results with the accuracy and precision specifications given in the user's guide of the test pipette.

Multichannel Pipettes

According to ISO 8655-6, "For the purpose of the test, each channel shall be regarded as a single channel and reported as such". Consequently, for each of the following procedures: fill all channels simultaneously when aspirating the test liquid, then expel only the test liquid aspirated by the channel being tested into the weighing vessel.

Note: Gilson recommends simplifying the process by fitting a tip to the tested channel, only.

Test Volumes	Minimum Volume (µL)	Mid Range Volume (μL)	Nominal Volume (µL)
Pipetman			
P2N P2, U2 P10, U10, P10N P20, U20, P20N P100N P100, U100 P200N P200, U200 P1000N P1000, U1000 P5000, U5000 P10ml, U10ml	0.2 0.5 1 2 10 20 20 50 100 200 1000 1000	1 1 5 10 50 50 100 100 500 500 2500 5000	2 2 10 20 100 100 200 200 200 1000 5000 10000
Microman			
M10 M25 M50 M100 M250 M1000	1 3 20 10 50 100	5 10 - 50 125 500	10 25 50 100 250 1000
Distriman			
DistriTip Micro DistriTip Mini DistriTip Maxi	2 20 200	5 50 500	10 100 1000
Pipetman Multichar	nnel		
8X200	20	100	200
Pipetman Ultra Mul	tichannel		
8x20 12x20 8x300 12x300	2 2 30 30	10 10 150 150	20 20 300 300
Pipetman Concept			
C10 C100 C300 C1200 C5000 C10ml	1 10 30 120 500 1000	5 50 150 600 2500 5000	10 100 300 1200 5000 10000
Pipetman Concept	Multichannel		
C8x10, C12x10 C8x100, C12x100 C8x300, C12x300	1 10 30	5 50 150	10 100 300

Estimating the Evaporation Rate (Mass Loss/Cycle)

Weighing requires special care, for small volumes (< 50 μ L, according to ISO 8655) use tweezers and weighing containers fitted with lids (Pot A). The goal is to minimize, control and quantify evaporation loss during the weighing cycle.

Apart from the design of the weighing vessel, the test cycle time is important. Evaporation is estimated by performing a series of four simulated weighings, repeating the weighing cycle without dispensing to the weighing container. The total difference attributable to evaporation is calculated and divided by 4 to obtain an average. The rate is expressed in mg/cycle (or for one cycle the loss may be expressed in mg).

For example, evaporation rates usually range for Pot A between 0.010 mg to 0.025 mg per weighing cycle. Recalculate the evaporation rate every 4 hours or whenever ambient conditions change (temperature, pressure, and humidity).

- 1 Add water to the weighing container until it is about one-third full.
- 2 Fit the weighing container with its lid and use tweezers to place it on the balance pan.
- 3 Using the pipette, aspirate a sample from the reservoir at the test volume setting.
- 4 Tare the balance and remove the weighing container from the balance pan.
- 5 Use tweezers to remove the lid.
- 6 Dispense the sample into the reservoir or to waste, *not the* weighing container.
- 7 Fit the weighing container with its lid and use tweezers to put it back on the balance pan.
- 8 Record the result e_1 .
- 9 Repeat steps 3 through 8 three times to obtain e_2 , e_3 , and e_4 .
- 10 Calculate the loss/cycle: $e = |e_1 + e_2 + e_3 + e_4|/4$ (mg).
- 11 The evaporation loss/cycle *e* (mg) should be added to the mean mass before calculating the mean volume.

Gravimetric Test

According to ISO 8655-6: "The test shall be carried out in a draught-free room with stable environment."

- 1 Place distilled or deionized water from the container in the weighing vessel to a depth of at least 3 mm.* (Refit lid for Pot A and B.)
- 2 Record the test conditions (ambient and water temperature, relative humidity, barometric pressure).
- 3 Select the test volume of your variable-volume piston pipette.
- 4 Fit the tip or capillary/piston assembly to the pipette (the manufacturer specifications are valid only when test executed with the manufacturers tips).
- Wet pipette tip five times to reach equilibrium in the dead air volume (not needed for Distriman and Microman), but do not take into account for calculations.

One test cycle should take less than 1 min.

A consistent rhythm during weighing operation should be maintained.

6 Change tip.

- 7 Pre-wet the tip once.
- 8 Pipette the test volume.
- 9 Determine tare mass (reset balance).
- 10 Remove the lid if needed (using the tweezers for pot A)
- 11 Open balance door, retrieve weighing container, deliver sample, refit its lid, if needed, using the tweezers, replace on the balance and close the door.
- 12 After allowing display to stabilize and record the mass.
- 13 Repeat the test cycle until ten measurements have been recorded as a series of masses m_1 to m_{10} .
- 14 For sample below or equal to 50µl, estimate evaporation loss by repeating steps 8 to 10 exactly as a normal sample weighing but without actually adding any sample to the weighing container. Record absolute value (ei) and repeat several (m) times.
- 15 Record the test conditions (ambient temperature, relative humidity, barometric pressure). Check that values are still within recommended limits.

Note *: ISO recommends that the orifice of the tip be immersed to between 2 mm and 3 mm below the surface of the water. However, you should first consult the user's guide for the model of Gilson pipette that you are testing.



Formulae

$$t = (t_1 + t_2)/2$$

$$B = (B_1 + B_2)/2$$
 2 l

$$V_i = Z (m_i + \overline{e})$$

 V_i = individual volumes (µL)

 m_i = individual masses (mg)

 \overline{e} = evaporation loss (mg)

Z = Z-factor (μ L/mg)

1 Calculate the mean temperature (t) of the distilled water (rounded to the nearest 0.5 °C).

2 Use the average barometric pressure (*B*) and mean temperature (*t*) to find the corresponding *Z-factor* from the table.

3 Multiply the weighings (mg), after any required correction for evaporation, by the *Z-factor* to obtain a series of volumes (μ L).

$$\overline{V} = \frac{\sum_{i=1}^{n} V_{i}}{n}$$

 V_i = individual volumes

V = mean volume

n = number of weighings

4 Compute the mean volume from the series of volumes (μL).



e = systematic error

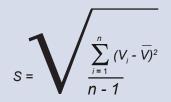
 \overline{V} = mean volume

 V_s = selected volume

 $e_{0} = 100 (\overline{V} - V_{0})/V_{0} \%$

5 Calculate the systematic error, which is the difference between the mean volume of actual measurements and the true value as specified by the volume setting of the pipette (selected volume). For fixed volume pipettes, replace V_s with V_o = nominal volume.

Accuracy may be expressed in μL or as a percentage.



V_i = individual volumes (calculated as above)

V = mean volume

n =number of measurements

s = repeatability standard deviation

nents

$$CV = 100 \times s / \overline{V}$$

between individual weighings. Quantifies the magnitude of scatter due to random error. Also known as Repeatability Standard Deviation (RSD).

6 Calculate the random error, which is the closeness of agreement

As a percentage, also known as coefficient of variation (CV).

Calculations

Z= Conversion factor (μ L/mg) t = Average temperature (°C) B = Air pressure (kPa)

Z-factor

 \boldsymbol{Z} correction factors for distilled water as a function of test temperature and air pressure.

t (°C)	B (kPa)	80	85	90	95 Ζ (μL/mg)	100	101.3	105
15.0		1.0017	1.0018	1.0019	1.0019	1.0020	1.0020	1.0020
15.5		1.0018	1.0019	1.0019	1.0020	1.0020	1.0021	1.0021
16.0		1.0019	1.0020	1.0020	1.0021	1.0021	1.0021	1.0022
16.5		1.0020	1.0020	1.0021	1.0021	1.0022	1.0022	1.0022
17.0		1.0021	1.0021	1.0022	1.0022	1.0023	1.0023	1.0023
17.5		1.0022	1.0022	1.0023	1.0023	1.0024	1.0024	1.0024
18.0		1.0022	1.0023	1.0023	1.0024	1.0025	1.0025	1.0025
18.5		1.0023	1.0024	1.0024	1.0025	1.0025	1.0026	1.0026
19.0		1.0024	1.0025	1.0025	1.0026	1.0026	1.0027	1.0027
19.5		1.0025	1.0026	1.0026	1.0027	1.0027	1.0028	1.0028
20.0		1.0026	1.0027	1.0027	1.0028	1.0028	1.0029	1.0029
20.5		1.0027	1.0028	1.0028	1.0029	1.0029	1.0030	1.0030
21.0		1.0028	1.0029	1.0029	1.0030	1.0031	1.0031	1.0031
21.5		1.0030	1.0030	1.0031	1.0031	1.0032	1.0032	1.0032
22.0		1.0031	1.0031	1.0032	1.0032	1.0033	1.0033	1.0033
22.5		1.0032	1.0032	1.0033	1.0033	1.0034	1.0034	1.0034
23.0		1.0033	1.0033	1.0034	1.0034	1.0035	1.0035	1.0036
23.5		1.0034	1.0035	1.0035	1.0036	1.0036	1.0036	1.0037
24.0		1.0035	1.0036	1.0036	1.0037	1.0037	1.0038	1.0038
24.5		1.0037	1.0037	1.0038	1.0038	1.0039	1.0039	1.0039
25.0		1.0038	1.0038	1.0039	1.0039	1.0040	1.0040	1.0040
25.5		1.0039	1.0040	1.0040	1.0041	1.0041	1.0041	1.0042
26.0		1.0040	1.0041	1.0041	1.0042	1.0042	1.0043	1.0043
26.5		1.0042	1.0042	1.0043	1.0043	1.0044	1.0044	1.0044
27.0		1.0043	1.0044	1.0044	1.0045	1.0045	1.0045	1.0046
27.5		1.0045	1.0045	1.0046	1.0046	1.0047	1.0047	1.0047
28.0		1.0046	1.0046	1.0047	1.0047	1.0048	1.0048	1.0048
28.5		1.0047	1.0048	1.0048	1.0049	1.0049	1.0050	1.0050
29.0		1.0049	1.0049	1.0050	1.0050	1.0051	1.0051	1.0051
29.5		1.0050	1.0051	1.0051	1.0052	1.0052	1.0052	1.0053
30.0		1.0052	1.0052	1.0053	1.0053	1.0054	1.0054	1.0054

Sample of a Verification Procedure Report

Pipetting	g System	Informa	ation								
	Pipette										
	Serial num					Manufac	turer:				
	Number of										
	Tips										
	Tip model:	:		Manu	facturer:		E	Batch nu	ımber:		
Environr	nental Fa	actors									
	Temperatu Hygrometi Z-factor:	ry (%):				Tempera	ture wa	ter (°C):	:		
General	Informat	ion									
	Decontam Adjustmer	ination (Yent (Yes/No)									
Statistics	s Summa	ry									
		Mean Volume (µL)		ults	ematic err Target μL %	Status		ults		get	Status
tion	$egin{aligned} V_{min} \ V_{int} \ V_{nom} \end{aligned}$	$V_{mean,min}$ $V_{mean,int}$ $V_{mean,nom}$	 	 	 	 	 	 	 	 	
riation Variat or	or										
SD: Standard Deviation CV: Coefficient of Variation Es: Systematic Error	Selecte Volum (µL)	ed Mean e Volume (µL)	Res		ematic err Target µL %	or Status	Res	ults	ndom er Tarç SD (µL)	get	Status
SD: Sta CV: Co Es: Sys	1 V _{min1} 1 V _{int1} 1 V _{nom1}	$egin{array}{c} V_{\it mean,min1} \ V_{\it mean,int1} \ V_{\it mean,nom1} \end{array}$		 	 	 	 	 	 	 	
	V _{min} V _{int} V _{nom}	$V_{mean,min}$ $V_{mean,int}$ $V_{mean,nom}$	 	 	 	 	 	 	 	 	

 V_{minN}

 V_{intN}

Ν

 $V_{{\it mean}, {\it minN}}$

V_{mean,intN} _ _ V

Sample of a Verification Procedure Report

Calibration Details

	ected ume	Minimum Volume (value μL)	Mid Range Volume (value μL)	Nominal Volume (value μL)
1		V_{a1}	V _{b1}	V _{c1}
2				
3				
4				
 10				
		V _{a10}	V _{b10}	V _{c10}
or				
Ch. #	Selected Volume	Minimum Volume (value μL)	Mid Range Volume (<mark>value</mark> μL)	Nominal Volume (<mark>value</mark> μL)
1	1	V _{a1,1}	V _{b1,1}	V _{c1,1}
	2			
	3			
	4			
	10	V _{a1,10}	$V_{b1,10}$	V _{c1,10}
i	1	$V_{ai,1}$	V _{bi,1}	V _{ci,1}
	2			
	3			
	4			
	10	V _{ai,10}	V _{bi,10}	V _{ci,10}
N	1	V _{aN,1}	V _{bN,1}	V _{cN,1}
	2			
	3			
	4			
	10	V _{aN,10}	$V_{bN,10}$	V _{cN,10}





Pipetman Ultra



Only for Pipetman Neo.

Except for Pipetman Neo.

Here are comparative tables for maximum permissible errors between ISO 8655 and Gilson. ISO 8655 maximum permissible errors are very wide, so as to have a conformity-basis for all pipettes. At Gilson our knowledge and kno-how allows us to be more stringent, which means the best pipette-performance.

Model (Reference)	Vol ι (μ		Ma: Gils Systematic error (μL)		rmissible Er ISO 8 Systematic error (µL)	3655
P2 (F144801) P2N (F144561) U2 (F21021)	Min Max.	0.2 0.5 2	± 0.024 ± 0.025 ± 0.030	≤ 0.012 ≤ 0.012 ≤ 0.014	± 0.08 ± 0.08 ± 0.08	≤ 0.04 ≤ 0.04 ≤ 0.04
P10 (F144802) P10N (F144562) U10 (F21022)	Min. Max	1 5 10	± 0.025 ± 0.075 ± 0.100	≤ 0.012 ≤ 0.030 ≤ 0.040	± 0.12 ± 0.12 ± 0.12	≤ 0.08 ≤ 0.08 ≤ 0.08
P20 (F123600) P20N (F144563) U20 (F21023)	Min.	2 5 10 20	± 0.10 ± 0.10 ± 0.10 ± 0.20	≤ 0.03 ≤ 0.04 ≤ 0.05 ≤ 0.06	± 0.20 ± 0.20 ± 0.20 ± 0.20	≤ 0.10 ≤ 0.10 ≤ 0.10 ≤ 0.10
P100 (F123615) P100N (F144564) U100 (F21024)	Min. Max.	10 20 50 100	± 0.35 ± 0.35 ± 0.40 ± 0.80	≤ 0.10 ≤ 0.10 ≤ 0.12 ≤ 0.15	± 0.80 ± 0.80 ± 0.80 ± 0.80	 ≤ 0.30 ≤ 0.30 ≤ 0.30 ≤ 0.30
P200 (F123601) P200N (F144565) U200 (F21025)	Min. Max.	20 50 100 200	± 0.50 ± 0.50 ± 0.80 ± 1.60	≤ 0.20 ≤ 0.20 ≤ 0.25 ≤ 0.30	± 1.60 ± 1.60 ± 1.60 ± 1.60	≤ 0.60 ≤ 0.60 ≤ 0.60 ≤ 0.60
P1000 (F123602) P1000N (F144566 U1000 (F21026)		100 200 500 1000	±3 ±3 ±4 ±8	≤ 0.6 ≤ 0.6 ≤ 1.0 ≤ 1.5	±8 ±8 ±8 ±8	≤ 3.0≤ 3.0≤ 3.0≤ 3.0
P5000 (F123603) and U5000 (F21027)		2000	± 12 ± 12 ± 30	≤ 3 ≤ 5 ≤ 8	± 40 ± 40 ± 40	≤ 15 ≤ 15 ≤ 15
P10ml (F161201) and U10ml (F21028)		1 mL 2 mL 5 mL 10 mL	± 30 ± 30 ± 40 ± 60	≤ 6 ≤ 6 ≤ 10 ≤ 16	± 60 ± 60 ± 60 ± 60	≤ 30 ≤ 30 ≤ 30 ≤ 30

Systematic error: expressed as the deviation of the mean of a tenfold measurement from the nominal or selected volume (see ISO 8655-6).

Random error: expressed as the repeatability standard deviation of a tenfold measurement (see ISO 8655-6).

Pipetman Concept



Model (Reference)	Vol ι (μ		Ma Gils Systematic error (μL)		rmissible Er ISO 8 Systematic error (µL)	3655
C10 (F31012)	Min.	0.5 1 5 10	± 0.040 ± 0.025 ± 0.060 ± 0.080	≤ 0.013 ≤ 0.012 ≤ 0.020 ≤ 0.025	± 0.120 ± 0.120 ± 0.120 ± 0.120	≤ 0.080 ≤ 0.080 ≤ 0.080 ≤ 0.080
C100 (F31013)	Min.	5 10 50 100	± 0.35 ± 0.30 ± 0.38 ± 0.4	≤ 0.10 ≤ 0.10 ≤ 0.12 ≤ 0.15	± 0.8 ± 0.8 ± 0.8 ± 0.8	≤ 0.30 ≤ 0.30 ≤ 0.30 ≤ 0.30
C300 (F31014)	Min.	20 30 150 300	± 0.80 ± 0.70 ± 0.90 ± 1.05	≤ 0.16 ≤ 0.20 ≤ 0.23 ≤ 0.30	± 4.00 ± 4.00 ± 4.00 ± 4.00	≤ 1.50 ≤ 1.50 ≤ 1.50 ≤ 1.50
C1200 (F31015)	Min.	100 120 600 1200	± 2.5 ± 2.4 ± 3.6 ± 6.0	≤ 0.4 ≤ 0.4 ≤ 0.8 ≤ 1.2	± 16.0 ± 16.0 ± 16.0 ± 16.0	≤ 6.0 ≤ 6.0 ≤ 6.0 ≤ 6.0
C5000 (F31016)		500 2500 5000	± 10 ± 15 ± 25	≤ 2 ≤ 4 ≤ 7	± 40 ± 40 ± 40	≤ 15 ≤ 15 ≤ 15
C10ml (F31017)		1000 5000 0000	± 25 ± 30 ± 50	≤ 4 ≤ 8 ≤ 12	± 60 ± 60 ± 60	≤ 30 ≤ 30 ≤ 30

Pipetman F



Model (Reference)	Volume (µL)	Ma Gils Systematic error (μL)		rmissible Er ISO 8 Systematic error (µL)	8655
F2 (F123770)	2	± 0.08	≤ 0.03	± 0.08	≤ 0.04
F5 (F123771)	5	± 0.10	≤ 0.04	± 0.125	≤ 0.075
F10 (F123772)	10	± 0.10	≤ 0.05	± 0.12	≤ 0.08
F20 (F123604)	20	± 0.20	≤ 0.06	± 0.20	≤ 0.10
F25 (F123775)	25	± 0.25	≤ 0.07	± 0.50	≤ 0.20
F50 (F123778)	50	± 0.40	≤ 0.15	± 0.50	≤ 0.20
F100 (F123784)	100	± 0.80	≤ 0.25	± 0.80	≤ 0.30
F200 (F123605)	200	± 1.60	≤ 0.30	± 1.60	≤ 0.60
F250 (F123787)	250	± 3.00	≤ 0.75	± 4.00	≤ 1.50
F300 (F123788)	300	± 3.50	≤ 0.75	± 4.00	≤ 1.50
F400 (F123789)	400	± 3.60	≤ 0.80	± 4.00	≤ 1.50
F500 (F123790)	500	± 4.00	≤ 1.00	± 4.00	≤ 1.50
F1000 (F123606)	1000	± 8.00	≤ 1.30	± 8.00	≤ 3.00
F5000 (F123607)	5000	± 30.00	≤ 8.00	± 40.00	≤ 15.00



Model (Reference)	Volu (μΙ		Ma Gils Systematic error (μL)	on	rmissible Er ISO 8 Systematic error (µL)	3655 Random
M10 (F148501)	Min. Max	1 5 10		≤ 0.03 ≤ 0.03 ≤ 0.06	± 0.20 ± 0.20 ± 0.20	≤ 0.10 ≤ 0.10 ≤ 0.10
M25 (F148502)	Min. Max.	3 10 25	± 0.27	≤ 0.08 ≤ 0.08 ≤ 0.10	± 0.70 ± 0.70 ± 0.70	≤ 0.30 ≤ 0.30 ≤ 0.30
M50 (F148503)	Min. Max.	20 50		≤ 0.20 ≤ 0.30	± 0.70 ± 0.70	≤ 0.30 ≤ 0.30
M100 (F148504)	Min. Max.	10 50 100		≤ 0.20 ≤ 0.30 ≤ 0.40	± 1.50 ± 1.50 ± 1.50	≤ 0.60 ≤ 0.60 ≤ 0.60
M250 (F148505)	Min. Max.	50 100 250	± 1.70	≤ 0.30 ≤ 0.30 ≤ 0.50	± 6.00 ± 6.00 ± 6.00	≤ 2.00 ≤ 2.00 ≤ 2.00
M1000 (F148506)	Min. Max.	100 500 1000	± 3.00 ± 5.00 ± 8.00	≤ 1.60 ≤ 2.50 ≤ 4.00	± 12.00 ± 12.00 ± 12.00	≤ 4.00 ≤ 4.00 ≤ 4.00



DistriTips Model (Reference)	Volume (µL)		Maxi Gilso Systematic error (µL)	n Random	missible Er ISO δ Systematic error (μL)	8655 Random
125 μL Micro (F164100) Micro ST (F164130	Min.	5	± 0.100 ± 0.125 ± 0.200	≤ 0.080 ≤ 0.075 ≤ 0.100	± 0.20 ± 0.20 ± 0.20	≤ 0.10 ≤ 0.10 ≤ 0.10
1250 μL Mini (F164110) Mini ST (F164140)	Min. Max.	20 50 100	± 0.80 ± 1.00 ± 1.00	≤ 0.20 ≤ 0.40 ≤ 0.60	± 1.50 ± 1.50 ± 1.50	≤ 0.60 ≤ 0.60 ≤ 0.60
12.5 mL Maxi (F164120) Maxi ST (F164150)		200 500 1000	± 6.00 ± 7.50 ± 10.00	≤ 1.00 ≤ 1.50 ≤ 2.50	± 12.00 ± 12.00 ± 12.00	≤ 4.00 ≤ 4.00 ≤ 4.00

ST means Sterilized.



Model (Reference)	Volur (µL)		Max Gils Systematic error (μL)	on Random	missible Errors ISO 8655 Systematic Randor error (µL) error (µl						
8x20 (F21040) and 12x20 (F21041)	Min.	1 2 10 20	± 0.10 ± 0.10 ± 0.20 ± 0.40	≤ 0.08 ≤ 0.08 ≤ 0.10 ≤ 0.20	± 0.40 ± 0.40 ± 0.40 ± 0.40	≤ 0.20 ≤ 0.20 ≤ 0.20 ≤ 0.20					
8x300 (F21042) and 12x300 (F21043)	Min.	20 30 150 300	± 1.00 ± 1.00 ± 1.50 ± 3.00	≤ 0.35 ≤ 0.35 ≤ 0.60 ≤ 1.00	±8 ±8 ±8	≤ 3.00 ≤ 3.00 ≤ 3.00 ≤ 3.00					



Model (Reference)	Volume (µL)	Ma: Gils Systematic error (µL)	on Random	rmissible Er ISO Systematic error (µL)	8655 Random
8x200 (F161004)	Min. 20 50 100 Max. 200	± 0.50 ± 1.00	≤ 0.25 ≤ 0.25 ≤ 0.50 ≤ 1.00	± 3.20 ± 3.20 ± 3.20 ± 3.20	≤ 1.20 ≤ 1.20 ≤ 1.20 ≤ 1.20

Pipetman Concept Multichannel



Model (Reference)	Volur (µL)		Max Gils Systematic error (μL)	on Random	rmissible Er ISO 8 Systematic error (µL)	3655 Random
C8x10 (F31032) and C12x10 (F31042)	Min. Max.	1 5 10	± 0.04 ± 0.08 ± 0.10	≤ 0.02 ≤ 0.04 ≤ 0.06	± 0.24 ± 0.24 ± 0.24	≤ 0.16 ≤ 0.16 ≤ 0.16
C8x100 (F31033) and 12x100 (F31043)	Min. Max.	10 50 100	± 0.25 ± 0.50 ± 0.80	≤ 0.14 ≤ 0.20 ≤ 0.25	± 1.60 ± 1.60 ± 1.60	≤ 0.60 ≤ 0.60 ≤ 0.60
C8x300 (F31034) and C12x300 (F31044)	Min.	30 150 300	± 1.00 ± 1.50 ± 2.40	≤ 0.18 ≤ 0.38 ≤ 0.60	± 8.00 ± 8.00 ± 8.00	≤ 3.00 ≤ 3.00 ≤ 3.00

Associated Documents

Documents	Gilson Reference
Pipetman Ultra User's Guide	LT801441
Pipetman P User's Guide	LT801117
Pipetman Neo Addendum	LT801511
Pipetman F User's Guide	LT801118
Pipetman 8X200 User's Guide	LT801236
Pipetman Ultra Multichannel User's Guide	LT801462
Pipetman Concept User's Guide	LT801489
Microman User's Guide	LT801502
Distriman User's Guide	LT801285
Decontamination Procedure	LT802288



-	-	_	-	 	 -	-	-	-	-	-	-	 	-	-	-	 	-	-		 _	-	-	-	_	 	 	_	-
-	-	_	_	 	 -	-	-	-	_	_	_	 		_	-	 	-	-		 _	-	_	_	_	 	 	_	-
-	-	_	_	 	 -	-	-	-	-	_	_	 		-	-	 	-	-		 _	-	_	-	_	 	 	_	-
-	-	_	_	 	 -	-	-	-	-	_	_	 		_	-	 	-	-		 _	-	-	-	_	 	 	_	-
-	-	_	-	 	 -	-	-	-	-	_	-	 		_	-	 	-	-	_	 _	-	-	_	_	 	 	_	-
-	-	_	-	 	 -	-	-	-	_	_	_	 		_	-	 	-	-	_	 _	-	-	_	_	 	 	_	-
-	-	_	_	 	 -	-	-	-	-	-	_	 		-	-	 	-	-		 -	-	-	_	_	 	 	-	-
-	-	-	-	 	 _	-	-	-	-	-	-	 		-	-	 	_	-	-	 -	-	-	-	_	 	 	_	-
-	-	_	_	 	 -	-	-	-	_	_	_	 		_	-	 	-	-	-	 -	-	_	_	_	 	 	_	-
_	-	_	_	 	 _	-	-	_	_	-	-	 		_	-	 	_	-		 _	_	_	_	_	 	 	_	-
_	_	_	_	 	 _	_	_	_	_	_	_	 	_	_	_	 	_	_		_	_	_	_	_	 	 	_	_
_	_	_	_	 	 _	_	_	_	_	_	_	 		_	_	 	_	_	_	 _	_	_	_	_	 	 	_	_
-	-	_	_	 	 _	_	-	-	_	_	_	 		_	-	 	_	-	_	_	_	_	_	_	 	 _	_	_
_	_	_	_	 	 _	_	_	_	_	_	_	 			_	 	_	_		 _	_	_			 	 	_	_
	_	_	_	 	 _				_	_	_	 			-	 	_			 _	-				 	 	_	_
- -				 	 							 		. –	- -	 	_			 -					 	 		
						_	_	_	_	_	_	 			_	 	_	_		· _	_	_	_	_		 		_ _ _ _
				 	 _	_	_	_	_	_	_	 		- -	_	 	_	_		 _	_	_	_		 	 		
		_	_	 	 _			_	_		_	 			_	 		_		 	_	_			 	 		
		_	_	 	 _			_	_		_	 			_	 		_		 	_	_			 	 		
		_	_	 	 _			_	_		_	 			_	 		_		 	_	_			 	 		

World Wide Web: www.gilson.com &: sales@gilson.com, service@gilson.com, training@gilson.com

World Headquarters

Gilson, Inc.

3000 Parmenter Street, P.O. Box 620027, Middleton, WI 53562-0027, USA Telephone: (1) 800-445-7661 or (1) 608-836-1551 • Fax: (1) 608-831-4451

Gilson SAS

19 avenue des Entrepreneurs, B.P. 145 95400 Villiers-le-Bel, France Telephone: (33) 1-34-29-50-00 • Fax: (33) 1-34-29-50-20

