

Complex Precision Summary

Instrument	Analyte	- Sample	Days (Tot/Excl)	No. Runs with Outliers	Within Run SD/CV	Total SD/CV
Immulite	✓ CPEP	Seronorm 1	20 / 2	None	0,012 / 5,4	0,012 / 5,8
	✓ CPEP	Seronorm 2	20 / 2	None	0,023 / 4,2	0,026 / 4,6

Required parameters are missing.

Experiment 'passes' (or adequate number of results).

X Experiment 'fails' (or not enough results).

Experiment has outliers.



	Source	Lot Number	Expiration	Analytes	
Reagents	Siemens	313		CPEP	
Calibrators	Siemens	143		CPEP	



Instrument Immulite

Sample Name Seronorm 1

Alternate Precision

Claim Evaluation

User's Concentration: 0,214 Claim Concentration: 0,2

		Standard Deviation						
	df	User's % CV	User's	Claim	Verification Value (95%)	Pass/Fail		
Within Run	36	5,4	0,012	0,01	0,012	Pass		
Between Run		1,6	0,003					
Between Day		1,0	0,002					
Total	70	5,8	0,012	0,02	0,023	Pass		
Medical Read	70	5.8	0.012					

The calculated value passes if it does not exceed the verification value.

20

15

Precision Plot

(Different plotting symbols represent different runs)

2

1

-1

-2

-3

10

Day Number

Upper 95% tole	rance limit for 9	5% of user	estimates
df for user's	within run	total	
experiment	SD	SD	_
10	0,016	0,017	
20	0,015	0,015	
30	0,014	0,015	This table provides
40	0,014	0,015	data for a manufact-
50	0,014	0,014	urer to include in
60	0,013	0,014	published materials
70	0,013	0,014	for users.
80	0,013	0,014	101 43613.
90	0,013	0,014	
100	0,013	0,014	

5

Su	pp	10	tir	ng	Da	at	a	

Analyst	Labonovum
Analysis Date	22 dec 2022 to 19 jan 2023
Days (Tot/Excl)	20 / 2
Runs per Day	2
Reps per Run	2
Critical Value	95%
Units	nmol/l
Verify Mode	Verify Vendor Claim
TEa	
Random Error Budget	
Allow Rand. Err.	
Control	
Reagent	Siemens 313
Calibrators	Siemens 143
Comment	

Accepted by:

Signature Date

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Instrument Immulite Sample Name Seronorm 1

Alternate Precision

Experimental Results

Date	Resul	ts	Date	Resul	ts	Date	Resul	ts
22 dec 2022	0,21 0,21	0,21 0,20	03 jan 2023	0,21 0,23	0,21 0,22	12 jan 2023	0,22 0,21	0,22 0,22
23 dec 2022	0,20 0,20	0,20 0,21	04 jan 2023	0,21 X 0,23	0,38 0,38	13 jan 2023	0,23 0,22	0,22 0,21
27 dec 2022	0,21 0,23	0,21 0,22	05 jan 2023	0,20 0,20	0,20 0,21	16 jan 2023	0,22 0,22	0,21 0,21
28 dec 2022	0,20 0,21	0,21 0,24	06 jan 2023	0,19 0,24	0,21 0,22	17 jan 2023	0,21 0,21	0,21 0,21
29 dec 2022	0,22 X 0,44	0,20 0,21	09 jan 2023	0,20 0,21	0,22 0,22	18 jan 2023	0,21 0,21	0,21 0,20
30 dec 2022	0,22 0,26	0,21 0,19	10 jan 2023	0,22 0,22	0,23 0,22	19 jan 2023	0,21 0,21	0,22 0,20
02 jan 2023	0,21 0,21	0,24 0,20	11 jan 2023	0,22 0,23	0,22 0,24			

"X" indicates an excluded run, "O" indicates an outlier run, and "S" indicates a day that does not have a full complement of results. In all of these cases, the entire day is excluded from the calculations.

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Instrument Immulite

Sample Name Seronorm 2

Alternate Precision

Claim Evaluation

User's Concentration: 0,564 Claim Concentration: 0,5

	Standard Deviation							
	df	User's % CV	User's	Claim	Verification Value (95%)	Pass/Fail		
Within Run	36	4,2	0,023	0,03	0,036	Pass		
Between Run		2,1	0,012					
Between Day		0,0	0,000					
Total	69	4,6	0,026	0,05	0,057	Pass		
Medical Read	69	4.6	0.026					

The calculated value passes if it does not exceed the verification value.

Precision Plot

Day Number

df for user's	within run	total	
experiment	SD	SD	
10	0,032	0,035	_
20	0,029	0,033	
30	0,028	0,031	This table provides
40	0,028	0,031	data for a manufact
50	0,027	0,030	urer to include in
60	0,027	0,030	published materials
70	0,027	0,030	for users.
80	0,027	0,030	ioi users.
90	0,026	0,029	
100	0,026	0,029	

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Analyst	Labonovum
Analysis Date	22 dec 2022 to 19 jan 2023
Days (Tot/Excl)	20 / 2
Runs per Day	2
Reps per Run	2
Critical Value	95%
Units	nmol/l
Verify Mode	Verify Vendor Claim
TEa	
Random Error Budget	
Allow Rand. Err.	
Control	
Reagent	Siemens 313
Calibrators	Siemens 143
Comment	

Accepted by:

Signature Date

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Instrument Immulite Sample Name Seronorm 2

Alternate Precision

Experimental Results

Date	Resul	ts	Date	Resul	ts	Date	Resul	ts
22 dec 2022	0,60 0,56	0,55 0,55	03 jan 2023	0,56 0,61	0,54 0,62	12 jan 2023	0,57 0,57	0,58 0,56
23 dec 2022	0,55 0,58	0,54 0,57	04 jan 2023	0,57 0,59	0,55 0,55	13 jan 2023	0,61 0,57	0,55 0,57
27 dec 2022	0,54 0,55	0,55 0,55	05 jan 2023	0,58 X 0,03	0,76 0,24	16 jan 2023	0,57 0,56	0,54 0,58
28 dec 2022	0,54 0,55	0,56 0,53	-	X 0,45 X 0,57	0,79 0,56	17 jan 2023	0,52 0,57	0,52 0,55
29 dec 2022	0,56 0,54	0,59 0,55	09 jan 2023	0,58 0,55	0,60 0,58	18 jan 2023	0,55 0,54	0,57 0,55
30 dec 2022	0,61 0,57	0,52 0,69	10 jan 2023	0,58 0,57	0,56 0,60	19 jan 2023	0,56 0,56	0,54 0,57
02 jan 2023	0,56 0,55	0,56 0,57	11 jan 2023	0,57 0,54	0,58 0,56			

"X" indicates an excluded run, "O" indicates an outlier run, and "S" indicates a day that does not have a full complement of results. In all of these cases, the entire day is excluded from the calculations.

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Clinical Chemistry -- Labonovum Lab Services

Complex Precision

Report Interpretation Guide

The Complex Precision Module is used in three situations:

- A manufacturer wants to calculate precision statistics using a statistically rugged procedure recognized in the industry, for use in official documents which may then be submitted to regulatory bodies.
- A user wants to determine whether an instrument meets the manufacturer's claim for precision using a statistically valid approach.
- A user wants to determine both within-run and total precision.

Experiment Procedure

- Define the number of replicates per run, runs per day, and number of days for the experiment. CLSI:EP5 recommends 2 replicates per run, 1 or 2 runs per day, for a minimum of 20 days.
- Collect data for a preliminary run of 8-20 results. The preliminary run is used to detect outliers. This step is optional, but strongly recommended. (It is required for EP5 compliance.)
- Collect data for the full duration of the experiment. The number of replicates per run and runs per day must be the same for all days.

Definitions

Precision. Ability to obtain the same result upon repeated measurement of a specimen.

User's Concentration. Grand mean, computed by adding the results (across all days, replicates, and runs) and dividing the total by the number of results.

Claim Concentration. Concentration at which the manufacturer's SD claims were determined. Laboratorians often think of precision in terms of CV, which is somewhat constant across concentrations. However, the statistical calculations in this module are intended to verify SD. Thus the sample tested should be at approximately the manufacturer's claim concentration. In the report summary, the CV is provided in addition to the SD.

Standard Deviation (SD). SD is the primary measure of Precision (variation of the individual results about the mean). The point of the Complex Precision experiment is either 1) to determine whether the SD meets the manufacturer's claim, or 2) to compute within-run and total SDs to establish such a claim.

SD Components: The experimental results are analyzed by a random-effects Analysis of Variance (ANOVA) procedure to partition the SD into separate components. The two

components usually cited in precision claims are within-run and total SD. The intent of the following definition of the components is to be intuitive, NOT to be mathematically correct

- Within-run SD. Measures the "average" SD computed over replicates that occur within the same run
- Between-run SD. SD computed from the means of the results for each run.
- **Between-day SD.** SD computed from the means of the results for each day.
- Total SD. A composite of within-run, between-run, and between-day SDs. This is not the answer you would get if you computed an ordinary SD on all the data, ignoring replicate number, day number, and run number.

Claim Value (of SD). Two kinds of claims may be verified: Manufacturer's claims and Medical Requirements. A manufacturer will typically provide separate values for within-run and total SD. A medical requirement is for Total SD only.

Verification Value. You can pass the precision test even if your measured SD is greater than the claim, as long as the difference is not statistically significant. The Verification Value is the largest SD that is not significantly different from the claim. It varies with sample size — the larger the sample, the closer the Verification Value is to the claim value.

The precision test passes if the computed SD does not exceed the Verification Value.

Critical Value. The confidence level on which the Verification Value is based. Normally the Critical Value is 95%. This means that the Verification Value is equivalent to a 95% confidence limit -- the observed SD meets the manufacturer's claim if its lower 95% confidence limit does not exceed the claim value.

Coefficient of Variation (CV). SD expressed as a percent of the mean.

Degrees of Freedom (df). df is like an "effective N" for an SD component. As df gets larger the confidence limit around the computed SD narrows, and the verification value gets closer to the claimed value.

Outlier. A result so far from the others as to arouse suspicion that it was generated by a different mechanism.

Outlier Rejection

The program first calculates the SD of the preliminary run. This preliminary SD is multiplied by a user-defined Multiplier (usually 5.5) to compute the Maximum Acceptable Difference between Replicates. Any run whose range exceeds this

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Complex Precision

Report Interpretation Guide

maximum is declared an outlier, and the entire day is excluded from analysis.

Report Labeling

The Report title is "EP5 Precision" if the experiment meets all requirements of the CLSI:EP5 guideline. Specifically:

- Two replicates per run, either one or two runs per day, for at least 20 days.
- Uses a preliminary run of at least 8 results to establish a preliminary SD. Excludes a replicate pair if the difference between the duplicates exceeds 5.5 times this preliminary SD.

Reports that use other numbers of replicates per run, runs per day, or duration are labeled "Alternate Precision".

An option in Preferences/Reports can change the Report Summary sort order to be by sample name rather than by analyte name.

Pass or Fail?

The Complex Precision experiment "passes" as long as neither the within-run or total SD exceeds its verification value. However, the experiment might warrant further review if more than 5% of the runs were rejected as outliers.

Preliminary Report

The word PRELIMINARY printed diagonally across the report indicates that the data is incomplete, and the report is not acceptable as a final report. Some or all of the statistics may be missing. Causes:

- Less than 3 days.
- Less than 6 runs.

References

1. CLSI Document EP5-A. Evaluation of precision performance of quantitative measurement methods; Approved guideline. CLSI, 940 West Valley Road, Suite 1400, Wayne, PA 19087-1898 USA, 1999. (References to this document will be to CLSI:EP5)