

**Guidelines on Validation of Non-Regulatory Chemical and Radiochemical Methods**  
**(EPA/600/B-22/001, 2022)**  
**Method Validation Summary Table Template**  
**Version 1.0 (October 2023)**

The Guidelines provide collected information on critical areas of method performance assessment for validation studies. The Method Validation Summary Table from the Guidelines is designed to provide consistency in delivery of summary method validation results in a concise, easy-to-prepare and share format. Below is an example of the Table that you can fill-in with your data. The link to the Guidelines can be found at the end of the Table.

<b>A</b>	<b>Validation Design</b>	<b>Description</b>
1	Number of Laboratories	Add Your Text Here
2	Number of Matrices	Add Your Text Here
3	Types of Matrices Tested (water, soil, sediment, etc.)	Add Your Text Here

Notes on Section A: Optional

<b>B</b>	<b>Method Validation Overview</b>	<b>Description</b>
1	Method Title	Add Your Text Here
2	Organization	Add Your Text Here
3	Date	Add Your Text Here
4	Purpose	Add Your Text Here
5	Qualitative or Quantitative	Add Your Text Here
6	Target Analytes/Parameters	Add Your Text Here

Notes on Section B: Optional

<b>C</b>	<b>Method Development Considerations</b>	<b>Description and/or Results</b>
1	Sample Cost	Add Your Text Here
2	Recommended Sample Holding Times	Add Your Text Here
3	Sample Preservation	Add Your Text Here
4	Waste Generation	Add Your Text Here

Notes on Section C: Optional

<b>D</b>	<b>Method Performance Characteristic</b>	<b>Description and/or Results</b>
1	Bias/Trueness	Add Your Text Here
2	Detection Capability and Quantification Capability	Add Your Text Here

3	Instrument Calibration	Add Your Text Here
4	Measurement Uncertainty	Add Your Text Here
5	Precision	Add Your Text Here
6	Range	Add Your Text Here
7	Ruggedness	Add Your Text Here.
8	Selectivity in the Presence of Interferences	Add Your Text Here

Notes on Section D: Optional

- Add Your Relevant Details for Items in Section D, Optional, See Example Note Below:

“DI X number of validation study samples were tested unspiked or spiked at nominal concentrations of X and XXX ng/L. Each laboratory tested X replicate spiked samples of each matrix at each prepared concentration. Samples were prepared centrally and shipped to laboratories...”

**Related Links:**

1. Guidelines on Validation of Non-Regulatory Chemical and Radiochemical Methods:  
<https://www.epa.gov/system/files/documents/2022-12/GUIDELINES%20ON%20VALIDATION%20NRCRM%20EPA%20600B-22-001.PDF>