mzQC: Reporting and exchange format for mass spectrometry quality control data

**Document Status**

This document presents a specification of the mzQC data format developed by members of the Human Proteome Organisation (HUPO) Proteomics Standards Initiative (PSI) Quality Control (QC) Working Group. Distribution is unlimited.

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# Abstract

The Human Proteome Organisation (HUPO) Proteomics Standards Initiative (PSI) defines community standards for data representation in biological mass spectrometry, including proteomics, metabolomics, and lipidomics, to facilitate data comparison, exchange, and verification. The Quality Control Working Group develops standards and recommendations to describe the quality of mass spectrometry data and related analysis results.

This document defines the mzQC file format to report and exchange quality-related information for a mass spectrometry experiment, associated analysis results, or collections thereof. The mzQC specification defines a simple yet versatile file format with a hierarchical structure to store quality metrics, thereby providing support for general quality control, storage of quality decisions, visualisation efforts, and easy persistence and exchange of all of the above. While the mzQC format provides the technical infrastructure to encode the basis to decide about the quality of mass spectrometry experiments, importantly, the standard itself is not opinionated about what constitutes good quality. The format and its specification are realized in the widespread JavaScript Object Notation (JSON) that can be easily implemented in software to produce or consume mzQC files. The mzQC format is complemented by the Quality Control Controlled Vocabulary (QC CV), which includes formal definitions of relevant quality metrics. The combination of the clear, human-readable syntax of the mzQC format and the rich semantic information associated with quality metrics stored in the QC CV provides powerful mechanisms to interpret, store, and enable reuse of quality control data.

# Contents

[**Abstract**](#_gjdgxs) **1**

[**Contents**](#_qgxwkmf98lxg) **2**

[**Introduction**](#_1fob9te) **5**

[Background (from XML to JSON)](#_7xuwei1n8ssv) 6

[Document Structure](#_tyjcwt) 6

[**Notational Conventions**](#_1t3h5sf) **6**

[**mzQC Use Cases**](#_kf79v0qjyze3) **7**

[Identifying Non-Conforming MS Experiments Using Outlier Detection](#_yj7yp1si2r19) 8

[Longitudinal Monitoring of Instrument Performance](#_3rntll9w5vi) 8

[Producing Audience-Targeted Quality Reports](#_356kqbrpm3oo) 8

[Assisting in Novel Instrument Method Development](#_l929i571rrxc) 9

[Quality Control of Metabolomics Experiments](#_he1ngon57e70) 9

[**Relationship to Controlled Vocabularies**](#_2s8eyo1) **9**

[The Quality Control Controlled Vocabulary (QC CV)](#_lnxbz9) 10

[The PSI Mass Spectrometry Controlled Vocabulary (MS CV)](#_6g8q6y1q4t3q) 13

[Other Controlled Vocabularies](#_35nkun2) 14

[**Relationship to Other Data Standard Specifications**](#_dwie4afjb0ek) **14**

[mzML](#_3wn73562cgog) 15

[mzIdentML](#_jhfu7yv0305m) 15

[mzTab and mzTab-M](#_1ksv4uv) 15

[Universal Spectrum Identifier](#_45xhfmj65m4o) 15

MAGE-TAB [Proteomics](#_c23dgf57q999) 16

[**Format Specification**](#_44sinio) **17**

[Schema Sections](#_7xuwei1n8ssv) 27

[**cvParameter Values For Metrics**](#_xafnqqz6zyd2) **28**

[**Validation of mzQC Files**](#_41mghml) **29**

[Syntactic Validation](#_hnie5gn1tj1) 29

[Semantic Validation](#_11mvp63az60x) 29

[**General Recommendations**](#_bhdmmpbnwtkc) **30**

[Recommended File Extension](#_7xuwei1n8ssv) 30

[Compression](#_7xuwei1n8ssv) 30

[Encoding Non-Computable Numbers](#_7xuwei1n8ssv) 30

[Number of runQualities in mzQC Files](#_7xuwei1n8ssv) 30

[Element Order in cvParameter Derived Objects](#_7xuwei1n8ssv) 31

[Metadata](#_7xuwei1n8ssv) 31

[Development](#_7xuwei1n8ssv) 31

[**Pending Issues**](#_z4yaytwcv0ja) **31**

[Inclusion of Graphics](#_7xuwei1n8ssv) 31

[Referencing](#_7xuwei1n8ssv) 32

[Custom Thresholds and Flagging](#_7xuwei1n8ssv) 32

[**Conclusions**](#_z0a8ish3e59t) **32**

[**Authors**](#_2grqrue) **32**

[Additional Contributors](#_mxv7jfu2tkww) 33

[**Software**](#_vx1227) **33**

[**Intellectual Property Statement**](#_28h4qwu) **34**

[**Copyright Notice**](#_37m2jsg) **34**

[**References**](#_3fwokq0) **35**

[**Appendix**](#_2u6wntf) **36**

[Examples](#_7xuwei1n8ssv) 36

[Companion Documents](#_7xuwei1n8ssv) 36

# Introduction

This document systematically describes how the mzQC file format can be used to store quality-related information from MS-based experiments. It is a specification, not a tutorial. As such, the presentation of technical details is deliberately direct. The role of the text is to describe the mzQC file format and justify design decisions. The document does not discuss how mzQC should be used in practice, consider tool support for data capture or storage, or provide comprehensive examples of mzQC files. Tutorials and example material are available via the Appendix and through the [PSI-QC working group repository](https://github.com/HUPO-PSI/mzQC).

## Background (from XML to JSON)

Unlike most previous PSI standards, which are predominantly XML-based file formats (e.g., mzML [(Martens et al. 2011)](https://sciwheel.com/work/citation?ids=454825&pre=&suf=&sa=0), mzIdentML [(Jones et al. 2012; Vizcaíno et al. 2017)](https://sciwheel.com/work/citation?ids=1239999,6684068&pre=&pre=&suf=&suf=&sa=0,0), mzQuantML [(Walzer et al. 2013)](https://sciwheel.com/work/citation?ids=5390998&pre=&suf=&sa=0)), mzQC is defined using JSON syntax. A disadvantage of XML-based file formats is that XML is quite verbose, adding substantial formatting overhead to the already large data volume of mass spectrometry files. Additionally, producing or consuming an XML file is not natively supported in most programming languages and requires specialized software libraries, whose usage can be complex. Because qcML [(Walzer et al. 2014)](https://sciwheel.com/work/citation?ids=1240267&pre=&suf=&sa=0) was never an official PSI Standard, and the XML format has several disadvantages, the mzQC format is defined using JavaScript Object Notation (JSON) syntax. This file format has been developed to be a lightweight and universal data interchange format. As such, JSON files have a small memory footprint and have a wide built-in support in many programming languages. JSON makes use of two data structures: key–value pairs and ordered lists of values. JSON files are easy to understand, which explains their emergence as a replacement for XML in many systems. For example, it has become the *de facto* encoding language of (web) APIs, where easy-to-understand, lightweight data interchange formats are adopted most frequently. Functionally, JSON can be deployed for the same kind of data interchange purposes as XML. By specifying mzQC using a JSON syntax, we can leverage the broader availability of efficient libraries and simpler implementations while adhering to the proven design principles of PSI formats and maintaining the necessary level of compatibility. This lowers the technical threshold for mzQC adoption and will increase the reach of mzQC through an extended use case basis.

JSON data structures are built using two structures:

* An (unordered) collection of key–value pairs, generally called an **object**.

An object begins with a left brace { and ends with a right brace }. Each name is followed by a colon and the key–value pairs are separated by a comma.

Syntax: {key\_1: value\_1, key\_2: value\_2}

Example: {"precursor\_charge": 2, "precursor\_mz": 261.54}

* An (ordered) list of values, generally called an **array** or list.

An array begins with a left bracket [ and ends with a right bracket ]. Values are separated by a comma. Value types can be mixed.

Syntax: [value\_3, value\_4, value\_5]

Example: [true, {"subobject": "yes"}, 42]

Values can be a string in double quotes, a number, a boolean value (true or false), null, an object, or an array. These structures can be nested; for example, multidimensional tables can be represented as arrays of arrays. Strings and values are like the respective C or Java structures. [[json.org](https://www.json.org/)] Syntactic validation is available through the usage of [JSON Schema](https://json-schema.org/) for annotation and validation of JSON documents.

### qcML

The predecessor to the mzQC format, the qcML format [(Walzer et al. 2014)](https://sciwheel.com/work/citation?ids=1240267&pre=&suf=&sa=0), is based on a data representation in XML, which constitutes the major difference between the formats. While JSON has a large feature intersection with XML, its simpler structure and widespread language support will facilitate implementation of mzQC producing and consuming software, as exemplified by the various software tools that already support mzQC. The organizational structure of the data is largely preserved between qcML and mzQC, like the hierarchical structure and representation of quality metrics through cvTerm-like objects. An important difference between the qcML format and the mzQC format is that complex data types, such as lists or tables of QC metrics can be represented natively in the more versatile mzQC format while their support in qcML is limited and cumbersome. Additionally, lessons learned from the use of qcML led to a refinement of the actual value representation of quality metrics in the mzQC format. The use of a CV to specify and exchange quality metrics was also adopted and evolved to ensure machine readability and to reflect a broader base of use cases. Finally, as opposed to the mzQC format, the qcML format was not established as an official PSI standard.

### Resources

Users new to JSON can find detailed information via the following resources:

* [JSON website](https://www.json.org/)
* [formal JSON specification](http://www.ecma-international.org/publications/files/ECMA-ST/ECMA-404.pdf)

## **Document Structure**

The remainder of this document is structured as follows. Section 2 describes a variety of conventions for the notation that apply throughout the document. Section 3 lists use cases for the mzQC format. Sections 4 and 5 outline the relationships between mzQC and controlled vocabularies or other file format specifications, respectively. Sections 6 and 7 detail the format specification. Section 8 provides information on validation of mzQC files. Section 9 provides general recommendations on using mzQC, Section 10 lists a few pending issues, and Section 11 contains a brief summary and conclusion. Sections 12, 13, 14, 15, and 16 include the list of authors, software, intellectual property statement, copyright notice, and references, respectively. An appendix introduces tutorial material and companion documents explaining various aspects of the format and its use cases.

# Notational Conventions

The keywords “MUST,” “MUST NOT,” “REQUIRED,” “SHALL,” “SHALL NOT,” “SHOULD,” “SHOULD NOT,” “RECOMMENDED,” “MAY,” and “OPTIONAL” are to be interpreted as described in RFC 2119 [(Bradner 1997)](https://sciwheel.com/work/citation?ids=3528074&pre=&suf=&sa=0).

In this document, we colloquially refer to the data resulting from a single mass spectrometry experiment as a “run”. For example, a run may represent a shotgun LC-MS experiment containing tens of thousands of MS/MS scans. Alternatively, it might originate from a DIA experiment quantifying thousands of peptides, or from a Selected Reaction Monitoring (SRM) experiment spanning hundreds of transitions. The mzQC format is also intended to represent mass spectrometry quality when liquid chromatography is not directly coupled; a run may also represent a set of spectra resulting from a plate of MALDI-TOF data or from a GC-MS experiment. In all of these cases, an mzQC file will typically contain quality metrics for a collection of multiple spectra. The mzQC specification is equally applicable, however, to describe the quality (for example, the identifiability) of a single mass spectrum. The specification also extends to quality representation for “sets” of runs, either as groupings informed by the experimental design, longitudinal data, or differentiated by instrument, batch, site, etc.

# mzQC Use Cases

The mzQC format is intended to be applicable for any type of biological mass spectrometry. As an initiative of the HUPO-PSI, a primary goal is to support proteomics use cases. Additionally, the mzQC format explicitly aims to support QC information coming from alternative applications of biological mass spectrometry, including metabolomics. As such, the following use cases have been deliberately kept general to cover a wide variety of applications.

Importantly, mzQC provides an objective medium to facilitate data comparison, exchange, and verification. The goal of the mzQC format is not to judge the quality of the data that it describes or mandate the inclusion of specific QC metrics for an experiment to be valid.

Qualitative or quantitative interpretation of data encoded in mzQC files will likely be context dependent. For example, an LC-MS/MS experiment might contain some number of MS scans and some number of MS/MS scans; these can be recorded in a run-level mzQC file. These same values for a large number of experiments could be recorded in a summary-level mzQC file. To judge that an individual LC-MS/MS experiment represents an outlier for the overall experiment, however, requires the context of the other experiments. The mzQC file itself does not impose a judgment on an individual run, but it may record the judgment of an external tool that is able to take the experimental context into account.

The mzQC format is intended to:

* Report quality metrics calculated by QC tools (such as Quameter [(Ma et al. 2012)](https://sciwheel.com/work/citation?ids=8920549&pre=&suf=&sa=0), PTX-QC [(Bielow et al. 2016)](https://sciwheel.com/work/citation?ids=3033444&pre=&suf=&sa=0), and QCloud [(Chiva et al. 2018)](https://sciwheel.com/work/citation?ids=5027181&pre=&suf=&sa=0)) coming from MS-based experiments.
* Enable the presentation of quality reports to researchers for assessment of instrument performance.
* Record longitudinal QC metrics to monitor instrument health over time.
* Perform quality control of collections of MS experiments across individual runs, batches, biological or technical conditions, studies, or laboratories.
* Store and archive QC metrics next to their originating raw MS files and derived results in public data repositories or internal laboratory information management systems (LIMS).
* Explore and select datasets within public data repositories based on desired data characteristics.
* Provide QC metrics as input for visualization in reports and dashboards.

Several use cases in which mzQC files can be used are described in more detail below.

## Identifying Non-Conforming MS Experiments Using Outlier Detection

When working with a set of mass spectrometry experiments, researchers frequently want to determine whether the collection of files are of consistent quality. High-level quality metrics for each MS experiment (such as the number of identified spectra, peptides, and proteins; the mass calibration error; the width of chromatographic peaks; etc.) can be generated using QC software and exported to individual mzQC files. Subsequently, these separate mzQC files can be combined into a single mzQC for further unified analysis. An outlier detection algorithm can be used to project the experiments in a single principal components analysis [(Wang et al. 2014)](https://sciwheel.com/work/citation?ids=8920314&pre=&suf=&sa=0). Experiments that lie far away from the main group of experiments may then be interrogated as outliers to investigate potential data issues and sources of performance degradation.

## Longitudinal Monitoring of Instrument Performance

QC samples in proteomics can range from a simple peptide mixture to a single protein digest to a complex whole-cell lysate (QC samples in metabolomics are explained in more detail in use case 3.5). Each of these types of samples can be employed in a specific fashion to analyze the system performance. Common QC metrics to assess instrument performance include the sequence coverage of a single protein digest, such as BSA, or the number of identified proteins in a complex mixture. As instrument performance gradually degrades over time, such metrics should be monitored carefully and preventative tuning and calibration needs to be performed regularly to avoid data quality issues. Longitudinal performance tracking is essential to assess whether the instrument is still within its operational parameters and to schedule timely interventions. The QC metrics can be stored in the mzQC format after individual experiments are performed to record the instrument performance at a specific point in time. As such, system suitability test results can be consulted prior to data acquisition, and this information can be used to contextualize the experimental data quality. Additionally, QC metrics for multiple experiments, covering a longer time span, can be compared across multiple mzQC files. QC metrics can be consistently queried from multiple mzQC files corresponding to individual experiments or can be combined in a single mzQC file for easy analysis of instrument performance over time. Additionally, the information stored within the mzQC file(s) can be used to assess the longitudinal data quality across batches within an experiment or for inter-lab quality assessments (given that the sample content originates from the same biological resource and the data was acquired over different timepoints).

## Producing Audience-Targeted Quality Reports

Most MS experiments conducted are intended to be shared with other parties such as a specific collaborator, customer, or a broader audience, either immediately or at a later point. Examples include data acquisition by a core facility or an external collaborator, and data submission to a public repository, which is often required prior to publication in a scientific journal. Downstream use of the data can be facilitated if it is characterized by an accompanying quality report, particularly if the report targets a specific audience or a certain downstream use case. By including the metrics relevant for the envisioned audience(s) in an mzQC file, quality reports can be specifically tailored to the users’ preferences. For example, the report can be as simple as only reporting the quality of sample runs, it can include metrics derived for multiple runs in experimental groups, or it can even be combined with instrument health information from longitudinal monitoring before and during sample run acquisitions.

## Assisting in Novel Instrument Method Development

Novel instrument methods, such as data-independent acquisition [(Gillet et al. 2012)](https://sciwheel.com/work/citation?ids=614309&pre=&suf=&sa=0) and BoxCar acquisition [(Meier et al. 2018)](https://sciwheel.com/work/citation?ids=5221571&pre=&suf=&sa=0), are being developed to expand the robustness, depth, and coverage of MS experiments. For DIA experiments, simple constant-window strategies can mean that some isolation windows are crowded with peptides or fragments, while others have relatively low density. Quality metrics can guide researchers to variable-size windows that are more uniform in peptide density, leading to better subsequent identification through spectral libraries and improved quantitative precision. Novel software tools to characterize DIA experiments and report their metrics via mzQC are currently already in development.

## Quality Control of Metabolomics Experiments

Similar to proteomics, in metabolomics quality control samples can be used to provide a mechanism to judge the quality of the dataset or assay produced and to assess the analytical variance of individual MS runs and instrument variance over time. Several different types of quality samples can be used within an experimental setup. A common QC sample is a pooled sample, for which a small aliquot of each biological sample in the study set is mixed together. Several different pooled QC samples might be used for a study, placed in certain locations or randomly throughout the sample acquisition sequence, depending on the experimental setup. Blank samples are used to assess the carryover and ‘leakage’ from the columns and to determine the limit of detection. Often labs can use a commercially available QC sample, for example, NIST SRM 1950 human serum, or alternatively, create a stock pooled QC sample that differs from the one created using a mixture of the samples within the current experiment to measure the longitudinal variance of the instrument performance across different batches. Such quality metrics can be used for cross-study and cross-lab comparison. Another type of QC for metabolomics are synthetic cocktails that include multiple representatives from different classes of metabolites expected in a study. Relevant QC metrics to monitor include the distribution of the observations specific features, missing values, the distribution of peak intensities, the trend and/or drift of aggregated intensity values, and coefficients of variation of detected features. Overall, the QC sample measurements provide a qualitative and quantitative representation of the entire collection of samples in a study. An mzQC document can capture such quality indicators, to be fed directly into analysis software, for visualization purposes, for pre-analysis data cleaning, or to create a study quality report.

# Relationship to Controlled Vocabularies

The mzQC format describes quality control information based on the definitions formalized in controlled vocabulary terms. The use of CV terms from different vocabularies as control mechanisms and metric definitions are explained in the following.

## The Quality Control Controlled Vocabulary (QC CV)

The PSI-QC controlled vocabulary is intended to provide terms for the definition of quality metrics and related supporting values. The CV has been generated initially with a collection of published and basic metrics. Further care went into the definition of the metrics and the involved values to help interpretation, use, and visualization. The vocabulary builds on established terms and definitions from chemistry, physics, and biology ontologies in references and term relations. The main purpose of the QC CV lies in the definition of metrics related to mass spectrometry quality control. Additionally, another purpose is to provide the underlying definitions and provide specialized metrics for the data structures usable within mzQC files.

The QC CV is encoded as an OBO (Open Biological and Biomedical Ontologies) file. The OBO file format is a biology-oriented language for building ontologies, similar to the Web Ontology Language (OWL). The OBO Foundry (<http://www.obofoundry.org/>) maintains a comprehensive index of ontologies related to the life sciences.

As recommended by the PSI CV guidelines [(Mayer et al. 2013)](https://sciwheel.com/work/citation?ids=1240098&pre=&suf=&sa=0), [qc-cv.obo](https://github.com/HUPO-PSI/mzQC/blob/master/cv/qc-cv.obo) should be dynamically maintained via either the [psidev-qc-dev@lists.sourceforge.net](mailto:psidev-ms-vocab@lists.sourceforge.net) mailing list or (preferably) the QC working group’s GitHub repository (<https://github.com/HUPO-PSI/mzQC/issues>). This allows any user to request new terms in a transparent manner and in agreement with the community involved. Changes and new entries can be requested and discussed within a dedicated issue list, where a template is available to guide new entry definitions. Once a consensus is reached among the community the new terms are added within a few business days.

Because the QC CV is intimately related to the MS CV (see below), we follow the same principles of CV maintenance as the MS CV to avoid incompatibilities, which are likely to occur in case of duplications, redefinitions, and insufficient referencing. Nevertheless, as there are minor inconsistencies between the MS CV and QC CV in how certain information is encoded, it is currently envisioned that both CVs will be developed in parallel.

Online reference: <https://github.com/HUPO-PSI/mzQC/tree/master/cv>

### CV Term Creation for QC Metric Definition

The CV contains entries for metrics that can be recorded in the mzQC files. While the mzQC format allows storing any metric information, the CV makes it possible to interpret the actual values.

New CV terms should be requested via the mzQC GitHub issue tracker (<https://github.com/HUPO-PSI/mzQC/issues>). Upon creating a new issue, the requester has to select the “Request for new CV term” option. This will produce a template that will guide the requester in providing the necessary information to request their new CV term, as detailed below. If additional information or clarifications beyond the initial request are needed, the mzQC working group will work with the requester to finalize their CV term request. When all the necessary information has been provided, a new CV term will be created based on the request and added to the QC CV.

Each metric (and CV entry request) MUST include the following information:

* Name: A (short) string describing your metric.
* Definition: A longer description. This MUST include information about how the metric should be represented in an mzQC file.
* Comment: OPTIONAL details on how the metric should be interpreted (e.g. is a higher value better, can it only be interpreted relative to...).
* Value type: Is the metric type a single value, an n-tuple, a table, or a matrix?
* Unit: OPTIONAL unit of the value, specified using an existing CV term.
* Categorization: A categorization can OPTIONALLY be supplied. Examples are whether the metric depends on spectrum, peptide, protein, or metabolite identifications; or to describe the metric context.

**Example CV term:**

[Term]

id: QC:4000059

name: Number of MS1 spectra

def: "The number of MS1 events in the run." [PSI:QC]

is\_a: QC:4000003 ! single value

is\_a: QC:4000010 ! ID free

is\_a: QC:4000023 ! MS1 metric

comment: A lower number of MS1 spectra acquired during one sample run compared to similar runs can indicate mismatched instrument settings or issues with the instrumentation or issues with sample amounts.

relationship: has\_relation MS:1000579 ! MS1 spectrum

relationship: has\_relation QC:4000013 ! QC metric relation: single run

property\_value: has\_units UO:0000189 ! count unit

synonym: "MS1-Count" EXACT []

**ID**

id: QC:4000059

Each term MUST have a unique ID, specified as QC:XXXXXXX. Metric IDs are immutable and not reusable (e.g. for redefinition), and will be assigned upon inclusion or redefinition.

**Name**

name: Number of MS1 spectra

Each term MUST have a human-readable name. The name SHOULD be informative, SHOULD consist of maximum 100 characters, and SHOULD only consist of alphanumeric 7-bit ASCII characters, spaces, and punctuation marks ([\-\_,\.]).

Both the ID and the name for each term will be given in the mzQC files as well when the term is used.

**Definition**

def: "The number of MS1 events in the run." [PSI:QC]

The definition SHOULD consist of a short explanation of the term and how it should be stored in the mzQC file. The description SHOULD also provide aid in interpreting the values. The definition section SHOULD NOT contain calculation or interpretation details, but rather it should explain the purpose, requirements, and scope of the metric.

**Comment**

comment: A lower number of MS1 spectra acquired during one sample run compared to similar runs can indicate mismatched instrument settings or issues with the instrumentation or issues with sample amounts.

The comment section SHOULD contain calculation and interpretation details, like whether smaller or larger values are desirable. It is also RECOMMENDED to give a short explanation about how the metric works. If the metric calculation is not obvious, the calculation is RECOMMENDED to be briefly described in common terms. For published metrics, it is also RECOMMENDED to refer to the corresponding code.

**Value Type And Unit**

is\_a: QC:4000003 ! single value

property\_value: has\_units UO:0000189 ! count unit

A single value metric with a count as unit (UO:0000189).

is\_a: QC:4000003 ! single value

property\_value: has\_units UO:0000221 ! dalton

property\_value: has\_type STATO:0000237 ! standard deviation

A single value metric with as unit the standard deviation (STATO:0000237) in Dalton (UO:0000221), for example, the standard deviation of the distribution of precursor mass errors of identified spectra.

Each term that reports a value MUST indicate the corresponding value type using an is\_a relation. Different value types are possible: single value, n-tuple, table, or matrix. A value must be associated with a unit, see Metric Categorization. Depending on the value type, different additional categorization is REQUIRED.

* **single value:** Unit specification using has\_units is REQUIRED, type specification using has\_type is RECOMMENDED.
* **n-tuple**: An ordered list/array of length ‘n’. Unit specification using has\_units is REQUIRED, type specification using has\_type is RECOMMENDED. Units and types (optional) MUST be uniform for all values. An n-tuple is represented by a JSON array, which implicitly defines its length ‘n’.
* **table**: A table MUST have one or more columns defined using has\_column and MAY have optional columns defined using has\_optional\_column. A table is represented using a JSON key–value object where key(s) represent the column term names/accessions and the value(s) are JSON arrays of uniform value type and length.
* **table column type definitions**: Unit specification using has\_units is REQUIRED, type specification using has\_type is RECOMMENDED. The term name will be used as the column’s header.
* **matrix**: Unit specification using has\_units is REQUIRED, type specification using has\_type is RECOMMENDED. Units and types (optional) MUST be uniform for all values. A matrix is represented by a JSON array of JSON arrays where the inner arrays MUST be of uniform length, which implicitly defines the matrix dimensions.

Units SHOULD be sourced from the Units of Measurement Ontology ([UO](https://www.ebi.ac.uk/ols/ontologies/uo)), if available, otherwise from the Statistical Methods Ontology ([STATO](http://stato-ontology.org/)) or others as necessary. Protein modifications SHOULD be sourced from [Unimod](http://www.unimod.org/) or [PSI-MOD](https://github.com/HUPO-PSI/psi-mod-CV) where possible.

**Metric Categorization**

is\_a: QC:4000010 ! ID free

is\_a: QC:4000023 ! MS1 metric

relationship: has\_relation MS:1000579 ! MS1 spectrum

relationship: has\_relation QC:4000013 ! QC metric relation: single run

Different types of categorization can be assigned to CV terms. First, it is RECOMMENDED to specify whether a metric requires identification information to be computed (ID based) or not (ID free). Second, additional categories to describe the metric context (from which data the metric is derived, to which element of the instrumental setup the metric pertains, etc.) can be specified as well. It is RECOMMEND to align the categorization of novel metrics to existing terms to facilitate consumption of related metrics.

property\_value: has\_units UO:0000010 ! second

property\_value: has\_column QC:4000117 ! Outliers below

If the metric term has an associated value, its unit MUST be defined using the property\_value tag. “Single”, “n-tuple”, and “matrix” type values MUST be assigned a single, uniform unit type with has\_units. For “table” type values, one or more has\_column\_type/has\_optional\_column\_type specifications MUST be associated with the table. These implicitly define the column units through the has\_units attributes of the corresponding column definitions.

property\_value: has\_type STATO:0000237 ! standard deviation

For full semantic integration, it is RECOMMENDED to specify the value type for automatic processing and interpretation of the value. It is RECOMMENDED to source value types from [STATO](http://stato-ontology.org/).

**Additional Information**

synonym: "MS1-Count" EXACT []

In case of reimplementing, renaming, or redefining a metric, it is RECOMMENDED to also add synonym attributes with either the name or ID of the initial metric. It is not required for the initial metric to be included in any controlled vocabulary, but the name SHOULD be unambiguous and recognizable (e.g. from the source publication). Synonyms can be “RELATED” (the defined metric is similar, but not the same as what is connected with the synonym name), “NARROW” (the metric’s values can be identically interpreted as in the meaning of the synonym metric, however, definition and calculation may somewhat differ), “EXACT” (the defined metric is basically a result of renaming).

## The PSI Mass Spectrometry Controlled Vocabulary (MS CV)

The PSI-MS controlled vocabulary [(Mayer et al. 2013; Mayer et al. 2014)](https://sciwheel.com/work/citation?ids=1240098,5391118&pre=&pre=&suf=&suf=&sa=0,0) has been generated by software vendors and academic groups working in the area of mass spectrometry and proteome informatics and is a rich source of general mass spectrometry terms allowing for the annotation of mzQC files. Some terms describe attributes that must be coupled with a numerical value and optionally a unit for that value. Terms that require a value are denoted by having a value-type reference in the CV itself. Terms that need to be qualified with units are denoted with a has\_units relationship in the CV itself.

**Example:**

[Term]

id: MS:1001844

name: MS1 feature area

def: "Area of MS1 feature." [PSI:PI]

xref: value-type:xsd\:double "The allowed value-type for this CV term."

is\_a: MS:1002735 ! feature-level quantification datatype

Online reference: <https://github.com/HUPO-PSI/psi-ms-CV>

## Other Controlled Vocabularies

Due to the wide availability of alternative ontologies, these can be used to define terms to represent quality metrics as well. However, the requirements enforced for QC CV terms may not be met in all cases and caution needs to be exercised not to introduce value interpretation ambiguities. We acknowledge the existence of overlaps in the concepts defined in openly available bio-ontologies, and hence only suggest precedence of choice if possible. It is desirable to integrate strictly quality control related concepts in the QC CV, even if relevant terms might already be available elsewhere. Including all relevant terms in the QC CV will help to reduce dependencies on outside projects and ensures that a proper definition is available for all terms included in mzQC files. However, the flexible design of the mzQC format does not preclude the inclusion of third party or custom CVs besides the QC CV. Note that when custom CVs are included the user should make sure that there are no namespace clashes between different CVs. For CVs mentioned in this document and other widely established vocabularies this can be assumed to be the case, but it can be verified with the help of CV lookup services such as [Ontobee](http://www.ontobee.org/).

In general, units SHOULD be sourced from the Units of Measurement Ontology ([UO](https://www.ebi.ac.uk/ols/ontologies/uo)), if available, otherwise from the Statistical Methods Ontology ([STATO](http://stato-ontology.org/)) or others as necessary. Protein modifications SHOULD be sourced from [Unimod](http://www.unimod.org/) or [PSI-MOD](https://github.com/HUPO-PSI/psi-mod-CV) where possible.

# Relationship to Other Data Standard Specifications

The mzQC format describes quality control information from MS-based experiments and is therefore dependent on data in different formats. The mzQC specification tries to minimize the replication of information contained in other PSI formats and uses concepts of and references to related specifications. It is RECOMMENDED that mzQC should be used in conjunction with PSI standards such as mzML, mzTab, or mzIdentML, although it is also possible to use it in conjunction with other formats (for example, peak files in the Mascot Generic Format (MGF)).

## mzML

mzML [(Martens et al. 2011](https://sciwheel.com/work/citation?ids=454825&pre=&suf=&sa=0); <http://www.psidev.info/mzml>) is the PSI standard for capturing mass spectra/peak lists resulting from an MS experiment. As one of the main sources of raw spectral data from which quality metrics may be computed, mzML has a strong relationship to mzQC. In general, mzQC files will refer to mzML files as the input for metric calculation, rather than the converse. USIs are the preferred mechanism to refer to individual spectra in an mzML file. Alternatively, the native spectrum identifier format (MS:1000767; nativeID in the MS CV) can be used.

However, it is also possible to represent mzQC assessments from within an mzML file, since metric entries in mzQC and QC CV terms may be used to annotate spectra (as a <cvParam> element of a <spectrum> object) or even an entire mzML file (via a <cvParam> element of a <run>) to reflect a spectrum filtering strategy. It is RECOMMENDED that in case mzQC elements are used within an mzML file, the corresponding mzQC file is referenced in the <sourceFile> tag of the mzML and registered in the <processingMethod> object. An example of referencing an mzQC file within an mzML file can be found in the mzQC repository (see the Appendix).

## mzIdentML

mzIdentML [(Vizcaíno et al. 2017](https://sciwheel.com/work/citation?ids=6684068&pre=&suf=&sa=0); <http://www.psidev.info/mzidentml>) is the PSI standard for capturing peptide and protein identification data. As with mzML, the connection from mzQC to mzIdentML is strong, as input data to calculate identification-based quality metrics has to be sourced from identification file formats. Additionally, similar to mzML, mzQC information can be referred to in mzIdentML files via <cvParam> elements of <SpectrumIdentificationItem>, <SpectrumIdentificationResult>, and <SpectrumIdentificationList>.

## mzTab and mzTab-M

mzTab [(Griss et al. 2014](https://sciwheel.com/work/citation?ids=1239967&pre=&suf=&sa=0); <https://github.com/HUPO-PSI/mztab>) is the combined “lightweight supplement” PSI standard to report identification and quantification results in a tabular text format. It is easy to parse and contains the essential information required to evaluate results. mzTab bridges the gap between a pure text summary and machine-only readable formats. Its design allows a comprehensive summarization of processed MS results, in addition to referring to more details present in XML-based standard formats. Due to the minimally required comprehensiveness, its content values represent a sufficient base to calculate basic to intermediate quality metrics. mzTab also supports metabolomics use cases via the mzTab-M format [(Hoffmann et al. 2019)](https://sciwheel.com/work/citation?ids=6682900&pre=&suf=&sa=0). mzQC files can be referenced via the external\_study\_uri metadata field in mzTab. Additionally, it might be relevant to store simple QC metrics covered by a CV term from the QC CV directly in an mzTab file. For example, the mass deviation could be reported for identification and quantification data.

## Universal Spectrum Identifier

The Universal Spectrum Identifier [(Deutsch et al. 2020](https://sciwheel.com/work/citation?ids=10755047&pre=&suf=&sa=0); USI) describes a virtual path to locate a spectrum within a public MS dataset available via ProteomeXchange. When storing quality metrics related to individual spectra, USIs can be used to uniquely refer to spectra. In the most likely use case, when reporting quality metrics for multiple spectra simultaneously represented by a tabular QC metric, either an optional USI or nativeID column is RECOMMENDED to refer to individual spectra.

## MAGE-TAB Proteomics

The Proteomics Sample Metadata Project describes the MAGE-TAB standard to encode metadata that captures the relationship between samples and the data generated (<https://github.com/bigbio/proteomics-metadata-standard>). MAGE-TAB Proteomics makes it possible to provide rich metadata information in a consistent and structured format, with the aim to provide a mechanism for including metadata when depositing datasets to public proteomics data repositories.

As raw peak files typically only contain a very limited amount of information about the experimental design, metadata in the MAGE-TAB Proteomics format can be highly valuable to provide details on the study design. This can inform the calculation of relevant QC metrics to assess experimental quality beyond factors that are apparent from the peak files. For example, information on the biological conditions can be used to verify that proper randomization of the samples was performed, or QC metrics can be separately calculated for technical replicates and biological replicates to assess the respective sources of variability.

# Format Specification

|  |
| --- |
| **Figure 1**: Diagrammatic overview of the mzQC schema. |

## Schema Sections

The mzQC format can accommodate multiple metrics, possibly from multiple MS runs. Several dedicated (and named) data structures (arrays) are available for this purpose. These are named as the plural of the elements they provide space for, for example, the controlledVocabularies element contains controlledVocabulary objects. In the following, the schema elements are described in outer-to-inner order, starting with the mzQC root element ([Figure 1](#5gtpcggcwb78)). If elements are allowed in multiple places in this hierarchy, they are listed only once at the first valid occurrence. All objects described are mandatory and of single occurrence, unless stated otherwise. Arrays need to contain at least one element or MUST otherwise be omitted. We use the notational convention to list the acceptable child elements under the “item types” enumeration. The naming convention followed for all schema elements is camelCase (starting lower case) to denote a type. The examples given are sometimes abbreviated for conciseness where putting the example into context would result in verbosity. Symbols and characters used are the same as seen in the legend of [Figure 1](#5gtpcggcwb78).

We recommend consulting the introductory companion documents listed in the Appendix for examples of complete mzQC files and to get more familiar with the schema.

### mzQC

Root element of an mzQC file. It MUST enclose the listed elements and MUST be the sole root element in an mzQC file.

Type: object

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| version | string | true | Version of the mzQC format. |
| creationDate | date-time | true | Creation date of the mzQC file. |
| contactName | string | false | Name of the operator/creator of this mzQC file. |
| contactAddress | string | false | Contact address (mail/e-mail or phone) . |
| description | string | false | Description and comments about the mzQC file contents. |
| runQualities | array of runQuality | true\* | Description in section 6.1.3. |
| setQualities | array of setQuality | true\* | Description in section 6.1.4. |
| controlledVocabularies | array of controlledVocabulary | true | Description in section 6.1.17. |

\*At least one of runQualities or setQualities MUST be present.

The version string MUST be in the format “major.minor.patch”, each separated by a period. The creation date string MUST be in [ISO8601](https://en.wikipedia.org/wiki/ISO_8601#Combined_date_and_time_representations) format, for example, 2019-10-29T14:40:17.

Example:

"mzQC": {

"version": "1.0.0",

"creationDate": "2019-10-29T14:40:17",

"runQualities": [..],

"controlledVocabularies": [..]

}

### baseQuality

Base element from which both runQuality and setQuality elements are derived. baseQuality is an abstract element; only its derived runQuality and setQuality elements SHALL be used in a valid mzQC document.

Type: object

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| metadata | object | true | Description in section 6.1.8. |
| qualityMetrics | array of qualityMetric | true | Description in section 6.1.15. |

Example: see runQuality or setQuality.

### runQualities

OPTIONAL list of runQuality elements. However, it is REQUIRED that at least one of runQualities or setQualities is present. If specified, the runQualities list MUST contain at least one runQuality element.

Although it is possible to include multiple runQuality elements in the runQualities list, a consideration for using JSONPath is that querying of individual runQuality elements can become less efficient.

Type: array

Item types: runQuality

Min/max: (1, -)

Example:

"runQualities": [..]

### setQualities

OPTIONAL list of setQuality elements. However, it is REQUIRED that at least one of runQualities or setQualities is present. If specified, the setQualities list MUST contain at least one setQuality element.

Although it is possible to include multiple setQuality elements in the setQualities list, a consideration for using JSONPath is that querying of individual setQuality elements can become less efficient.

Type: array

Item types: setQuality

Min/max: (1, -)

Example:

"setQualities": [..]

### runQuality

Element containing metadata and qualityMetrics for a single run.

Type: baseQuality

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| metadata | object | true | Description in section 6.1.8. |
| qualityMetrics | array of qualityMetric | true | Description in section 6.1.15. |

Example:

runQualities: [

{

"metadata": {

"label": "file\_1\_4h",

"inputFiles": [..],

"analysisSoftware": [..]

},

"qualityMetrics": [..]

}

]

### setQuality

Element containing metadata and qualityMetrics for a collection of related runs (“set”). It is REQUIRED that setQuality only contains qualityMetrics which describe properties which apply to the set as a whole (as opposed to a list of values where each value can be attributed to a single run in isolation and would thus rather be stored as runQuality).

Type: baseQuality

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| metadata | object | true | Description in section 6.1.8. |
| qualityMetrics | array of qualityMetric | true | Description in section 6.1.15. |

It is REQUIRED that all qualityMetrics contained in the setQuality are derived from the same set of input files. For example, if qualityMetric *m1* contained in setQuality *s1* is derived from runs *r1*, *r2*, and *r3*, then qualityMetric *m2* contained in *s1* MUST also be derived from runs *r1*, *r2*, and *r3*. If this is not the case, *m2* MUST be contained in a different setQuality *s2*.

Example:

setQualities: [

{

"metadata": {

"label": "groupA",

"inputFiles": [..],

"analysisSoftware": [..]

},

"qualityMetrics": [..]

}

]

### cvParameter

Base element for a term that is defined in a controlled vocabulary, with OPTIONAL value.

Type: object

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| accession | string | true | Accession number identifying the term within its controlled vocabulary (pattern: ^[A-Z]+:[A-Z0-9]+$). |
| name | string | true | Name of the controlled vocabulary term describing the parameter. |
| description | string | false | Definition of the controlled vocabulary term. |
| value | any | false | Value of the parameter. |

The description attribute MAY be omitted, especially in larger documents to reduce their size, since the CV term definition can be easily retrieved from the respective CV via the accession.

Example:

{

"accession": "MS:1003151",

"name": "SHA-256",

"value": "3522254348badf0723d5d2f406cff412ff45111ae31323a306b033147fc0cdcc"

}

### metadata

Metadata describing the runQualityor setQualityto which it belongs.

Type: object

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| label | string | true | Unique name for the run (for runQuality) or set (for setQuality). |
| inputFiles | array of inputFile | true | Description in section 6.1.9. |
| analysisSoftware | array of analysisSoftware | true | Description in section 6.1.13. |
| cvParameters | array of cvParameter | false | Description in section 6.1.14. |

We RECOMMEND that the label is informative, for example so that it can be used to label values in a figure. Relevant information that the label can convey relates to the experimental design, the base name of the input file(s) to which the QC metrics correspond (for a runQuality), or a descriptive label of a grouping (for a setQuality, e.g. “timepoint4h”).

Example:

"metadata": {

"label": "groupA",

"inputFiles": [..],

"analysisSoftware": [..],

"cvParameters": [..]

}

### inputFiles

List of input files from which the QC metrics have been generated. At least one input file MUST be present and it is RECOMMENDED that this inputFile corresponds to a raw or peak file.

Type: array

Item types: inputFile

Min/max: (1, -)

Example:

"inputFiles": [..]

### inputFile

Input file used to generate the QC metrics. We RECOMMEND that only meta information about the files used are stored here.

Type: object

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| name | string | true | The name MUST uniquely match to a location (specified below) listed in the mzQC file. |
| location | string (URI) | true | Unique location representing the complete file path, REQUIRED to be specified as a [URI](https://tools.ietf.org/html/rfc3986). The file URI is RECOMMENDED to be publicly accessible. |
| fileFormat | cvParameter | true | Description in section 6.1.11. |
| fileProperties | array of cvParameter | false | Description in section 6.1.12. |

We RECOMMEND to use a short, yet descriptive, name, such as the base file name (without file extension), which may lend itself as a label on a plot.

Example:

{

"name": "H2-1-1",

"location": "ftp://massive.ucsd.edu/MSV000081205/ccms\_peak/RAWs/H2-1-1.mzML",

"fileFormat": {

"accession": "MS:1000584",

"name": "mzML format"

},

"fileProperties": [

{

"accession": "MS:1000747",

"name": "completion time",

"value": "2012-02-03T11:00:41"

},

{

"accession": "MS:1003151",

"name": "SHA-256",

"value": "3522254348badf0723d5d2f406cff412ff45111ae31323a306b033147fc0cdcc"

}

]

}

### fileFormat

Type of input file.

Type: cvParameter

Example:

"fileFormat": {

"accession": "MS:1000584",

"name": "mzML format"

}

### fileProperties

Detailed information of the input file.

RECOMMENDED properties and their CV accessions are:

* Completion time of the input file (MS:1000747).
* Checksum of the input file (any child of: MS:1000561 ! data file checksum type).

Type: array

Item types: cvParameter

Min/max: (1, -)

Example:

"fileProperties": [

{

"accession": "MS:1000747",

"name": "completion time",

"value": "2012-02-03T11:00:41"

},

{

"accession": "MS:1003151",

"name": "SHA-256",

"value": "3522254348badf0723d5d2f406cff412ff45111ae31323a306b033147fc0cdcc"

}

]

### analysisSoftware

Software tool(s) used to generate the QC metrics.

Type: array

Item types: extension of cvParameter

Min/max: (1, -)

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| accession | string | true | Accession number identifying the term within its controlled vocabulary (pattern: ^[A-Z]+:[A-Z0-9]+$). |
| name | string | true | Name of the controlled vocabulary term describing the software tool. |
| description | string | false | Definition of the controlled vocabulary term. |
| value | any | false | Name of the software tool. |
| version | string | true | Version number of the software tool. |
| uri | string (URI) | true | Publicly accessible URI of the software tool or documentation. |

Example:

"analysisSoftware": [

{

"accession": "MS:1009002",

"name": "QuaMeter IDFree",

"version": "1.1.21070",

"uri": "http://proteowizard.sourceforge.net/"

}

]

### cvParameters

OPTIONAL list of parameters containing additional metadata about its parent runQuality/setQuality. If specified, the cvParameters list MUST contain at least one cvParameter element.

Type: array

Item types: cvParameter

Min/max: (1, -)

Example:

"cvParameters": [..]

### qualityMetrics

The collection of qualityMetrics for a particular runQuality or setQuality.

Type: array

Item types: qualityMetric

Min/max: (1, -)

Example:

"qualityMetrics": [..]

### qualityMetric

Element containing the value and description of a QC metric defined in a controlled vocabulary.

Type: cvParameter

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| accession | string | true | Accession number identifying the term within its controlled vocabulary (pattern: ^[A-Z]+:[A-Z0-9]+$). |
| name | string | true | Name of the controlled vocabulary element describing the metric. |
| description | string | false | Definition of the controlled vocabulary term. |
| value | Single value, n-tuple, table, matrix | false | Value of the metric (see section 7). |
| unit | cvParameter / array of cvParameter | false | One or more controlled vocabulary elements describing the unit of the metric (if applicable — see section 7). |

The description attribute MAY be omitted, especially in larger documents to reduce their size, since the CV term definition can be easily retrieved from the respective CV via the accession. We RECOMMEND to include the description only if added verbosity is expected to help (human) readability. In any case, the description text MUST NOT be altered from the term definition. If a qualityMetric does not take a value, both the value and unit attributes MUST NOT be present. To facilitate independent consumption of an mzQC file without explicitly having to parse all listed CVs, it is REQUIRED to repeat the unit as defined in the CV. For table metrics, the unit MUST be an array with the order of the units matching the order of the table columns (see Section 7).

Example:

{

"accession": "QC:4000059",

"name": "Number of MS1 spectra",

"description": "The number of MS1 events in the run.",

"value": 7787,

"unit": {

"accession": "UO:0000189",

"name": "count unit"

}

}

### controlledVocabularies

Collection of controlled vocabulary elements used to refer to the source of the used CV terms in the qualityMetric objects (and others).

Type: array

Item types: controlledVocabulary

Min/max: (1, -)

Example:

"controlledVocabularies": [..]

All CVs containing terms used in an mzQC document MUST be referenced in the controlledVocabularies element. To allow direct JSONPath queries of mzQC documents, there MUST NOT be any namespace clashes between the used CVs. For CVs mentioned in this document and other widely established vocabularies this can be assumed to be the case, but it can be verified with the help of CV lookup services such as [Ontobee](http://www.ontobee.org/) or [OLS](https://www.ebi.ac.uk/ols/index).

### controlledVocabulary

Element describing a controlled vocabulary used to refer to the source of the used CV terms in qualityMetric objects (and others).

Type: object

Object definition:

| Name | Type | Required | Description |
| --- | --- | --- | --- |
| name | string | true | Full name of the controlled vocabulary. |
| uri | string (URI) | true | Publicly accessible URI of the controlled vocabulary. |
| version | string | false | Version of the controlled vocabulary. |

The uri is RECOMMENDED to be a public URL. In cases where a code repository is referenced, the uri MUST be stable, i.e. refer to either a specific commit (for that version) or a (direct) release link of the file.

Example:

{

"name": "Proteomics Standards Initiative Mass Spectrometry Ontology",

"uri": "[https://raw.githubusercontent.com/HUPO-PSI/psi-ms-CV/  
5e02c611b8392301ce00e8d7bb3a4d63848ef8b3/psi-ms.obo](https://raw.githubusercontent.com/HUPO-PSI/psi-ms-CV/5e02c611b8392301ce00e8d7bb3a4d63848ef8b3/psi-ms.obo)",

"version": "4.1.45"

}

# cvParameter Values For Metrics

Values of qualityMetrics are defined by their corresponding controlled vocabulary term. They can be one of four different types.

Type:string / number / boolean / array / object

Object definition:

| Name | CV Accession | Description | Example |
| --- | --- | --- | --- |
| single value | QC:4000003 | A single string, number, or boolean value. | "target" or  1.0 or  true |
| n-tuple | QC:4000004 | An array of values of the same type (and unit). | [2, 3, 5, 7, 11] |
| table | QC:4000006 | An object with each column a key–value pair of the form column name: [array of values]. All columns MUST have equal length. Values in the table MAY have different types in each column (in contrast to a matrix). The actual structure of the table is defined by the CV term. | {  score: [0, 1, 1],  intensity: [0, 13, 9] } |
| matrix | QC:4000007 | An array of child arrays with values. All child arrays MUST have the same length and represent a row of the matrix, i.e. the matrix elements are indexed as [row, column]. All values in a matrix MUST have the same type. | [[1, 2, 3],  [4, 5, 6],  [7, 8, 9]] |

For most metric types only a single unit is needed (single value, n-tuple, matrix). In contrast, the table metric type can have different units for each individual column. In this case, the unit element MUST be specified as an array whose order matches the table columns. For a concrete example we refer to the companion document on CV term use (Appendix).

# Validation of mzQC Files

Here we give a brief overview of syntactic and semantic validation requirements of mzQC files. Full validation is implemented in the [mzqc-pylib](https://github.com/bigbio/mzqc-pylib) reference implementation. A living document with up-to-date validation information can be found in the [online mzQC GitHub documentation](https://github.com/HUPO-PSI/mzQC/blob/master/doc/validation.md).

## Syntactic Validation

With the help of the mzQC JSON schema, mzQC instances can be readily checked for syntactic schema compliance. There are a [number of validators already implemented for use in various programming languages](https://json-schema.org/implementations.html) that support validation of JSON schema draft-07 from which the mzQC schema is designed.

## Semantic Validation

Due to the advanced design of the mzQC JSON schema, many aspects of a valid mzQC file can already be verified via syntactic validation without the need of explicit semantic validation. There are, however, a few additional semantic rules that need to be followed to create fully compliant mzQC files.

* All used controlledVocabulary elements MUST reference a valid ontology.
* All used CV terms (including the quality metrics) must be present in one of the controlled vocabularies listed in the files controlledVocabularies element, matching in name and accession.
* All qualityMetric elements MUST be unique within their runQuality or setQuality parent.
* The value type of qualityMetric elements MUST match the specification in the CV (i.e. single value, n-tuple, matrix, or table).
* The value of qualityMetric MUST match its unit definition (and type implicitly).
* All columns of table qualityMetric elements MUST have the same length.
* label attributes in the metadata elements MUST be unique within its mzQC file.
* All inputFile elements MUST have a unique location attribute within their runQuality or setQuality element.

Additionally, more detailed checks might be performed to assess the validity of specific metric values on a case-by-case basis (e.g., percentages should sum to 100%).

# General Recommendations

## **Recommended File Extension**

The RECOMMENDED extension for an mzQC file is “.mzqc”.

## **Compression**

We have attempted to remove redundancy from the mzQC format wherever possible during its design. As a result typical mzQC files are not expected to grow overly large, however, due to the diverse nature of quality metric values and because much of the specification contains long form names to maintain human readability, there is ample opportunity to improve storage efficiency through compression. Compression of mzQC files comes at little overhead as text-based formats are inherently ideal targets for compression and common compression libraries are freely available on all computing platforms. Therefore, if a low memory footprint is desired, we RECOMMEND making use of a free compression algorithm, such as gzip.

## **Encoding Non-Computable Numbers**

Although the JSON format does not explicitly support encoding non-computable numbers, such as NaN or infinity, most parsing libraries unofficially support such values. To represent these values in mzQC files, we RECOMMEND to use their JavaScript representation: NaN, Infinity, and -Infinity (note the lack of quotation marks to avoid interpreting these values as strings). This encoding maximimizes compatibility with JSON serialization and deserialization libraries in various programming languages.

## **Number of runQualities**/**setQualities in mzQC Files**

The mzQC specification allows storing multiple runQuality or setQuality elements in a single mzQC file. The main purpose hereof is to group QC information from multiple runs for the subsequent calculation of setQuality metrics (or similarly, higher level setQuality metrics in a hierarchical experimental set-up). However, to facilitate mzQC file processing and storage, we RECOMMEND to restrict individual mzQC files to single runQuality/setQuality elements when possible. An example of the benefits of such a policy can be found in the companion document on the core facility use case.

## **Element Order in cvParameter Derived Objects**

JSON does not enforce a specific order of elements within an object. However, for streaming of large mzQC files, it is RECOMMENDED that the controlledVocabularies element occurs before any runQualities/setQualities element(s). Additionally, for improved readability, we RECOMMEND that in cvParameter derived objects, if possible, “human-readable” elements, like name and value, are listed first.

## **Metadata**

Depending on the use case to which mzQC is applied, there can be supplemental data that is not strictly metadata but is still considered relevant. We RECOMMEND that only meta information about the input files that were used to compute the QC metrics is stored in the metadata section of an mzQC file. If a piece of data does not fit into any metadata section, for example, such as the instrument type, it can be considered a QC metric to a lesser degree and should be stored in the respective qualityMetrics section.

## **Development**

For ongoing development, documentation, and to report issues with the format specification please see the working group’s GitHub page (<https://github.com/HUPO-PSI/mzQC/>).

# Pending Issues

## Inclusion of Graphics

It can be desirable to include graphical metric plots when mzQC files are used to create quality reports or for archival purposes, as metric values can be much more intuitive to grasp when visualised. However, producing such plots is highly system dependent and often requires additional software of specific versions to be installed. To circumvent these limitations, metric plots can be directly stored in an mzQC file as a custom qualityMetric element. We RECOMMEND the metric value to be the base64 encoded string of the plot image file and we RECOMMEND to use a custom metric with a descriptive and unambiguous name that makes it clear from which QC metric(s) the plot is derived and that it is an encoded image.

Further work is needed to make this system work reliably between different methods of mzQC use, for which the respective standardization details are deferred to a later release of the mzQC standard. This includes considerations on the image formats encoded and controlled vocabulary additions for graphical metrics. Until then we RECOMMEND to only use this system in custom applications at your own risk.

## Referencing

Due to the (few) limitations of the JSON format, intra- and inter-file referencing of qualityMetric elements is left to custom mechanisms. The specification of such a mechanism requires considerable effort to benefit only a few applications and is hence deferred to a later release of the mzQC standard. We RECOMMEND using a concept partially outlined in section 9.4 to only store a single runQuality element per mzQC file and replicate qualityMetric elements in-context if necessary. An example where this would be the case is when a setQuality uses multiple QC metrics from multiple MS runs. For good practice, the base runQuality elements should be replicated in an mzQC file that reports such a setQuality, making the references implicit.

## Custom Thresholds and Flagging

In the current version, specifying thresholds for QC metrics or flagging metrics, runs, or sets of runs as “quality unfulfilled” is not directly supported. Once this is supported via the appropriate CV terms, an mzQC file may be enriched in a second pass with thresholds and flags together with a rationale as to why the “quality unfulfilled” flag was set. Note that the mzQC format does not prescribe any thresholds denoting acceptable quality itself, this is left to the generating and interpreting software.

# Conclusions

This document contains the specifications for using the mzQC format to represent quality control metrics in the context of biological mass spectrometry. This specification constitutes a proposal for a standard from the Proteomics Standards Initiative. These artefacts are currently undergoing the PSI document process, which will result in a standard officially sanctioned by PSI.

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In addition to the authors, the following people contributed to format development, reviewed the specification document, or tested mzQC:

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# Software

The following are the initial software implementations supporting the mzQC format.

Core libraries:

* [mzqc-pylib](https://github.com/bigbio/mzqc-pylib): Python library to process and validate mzQC files. It includes a Python object model, (de-)serialisation functions, syntactic validation, and semantic validation of mzQC files.

Metrics generating software:

* [OpenMS](https://github.com/OpenMS/OpenMS): Open-source software C++ library for LC/MS data management and analyses.
* [PTXQC](https://github.com/cbielow/PTXQC): An R package for creation of QC reports from MaxQuant results and OpenMS mzTab files.
* [QCCalculator](https://github.com/bigbio/qccalculator): Python tool for base QC metric calculation from mzML, mzIdentML, and MaxQuant input files.
* [Yamato / SwaMe / Prognosticator](https://github.com/PaulBrack/Yamato): SWATH-MS QC metrics generation tools.

An up-to-date list of software tools that support the mzQC format can be found on the [mzQC GitHub website](https://github.com/HUPO-PSI/mzQC/blob/master/doc/software.md).

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# Appendix

## **Examples**

* [individual runs](https://github.com/HUPO-PSI/mzQC/blob/master/doc/single-run.mzQC.md)
* [set of runs](https://github.com/HUPO-PSI/mzQC/blob/master/doc/set-of-runs.mzQC.md)
* [QC2 Sample](https://github.com/HUPO-PSI/mzQC/blob/master/doc/QC2-sample-example.mzQC.md)
* [metabolomics batch correction](https://github.com/HUPO-PSI/mzQC/blob/master/doc/metabo-batches.mzQC.md)
* [mzML with mzQC metrics](https://github.com/HUPO-PSI/mzQC/blob/master/doc/examples/mzml-mzqc-example.md)

## **Companion Documents**

* [mzQC: a “common currency” for quality control of biological mass spectrometry](https://github.com/HUPO-PSI/mzQC/blob/master/doc/mzQC_common_currency.md)
* [mzQC with multiple experiments](https://github.com/HUPO-PSI/mzQC/blob/master/doc/mzQC_multiple_experiments.md)
* [mzQC format for analytical chemists](https://github.com/HUPO-PSI/mzQC/blob/master/doc/mzQC_for_analytical_chemists.md)