

AMIS Product: Best Practice Manual



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1. Receipt of material

- 1.1 For proper handling of our material, please refer to AMIS MSDS
- 1.2 Ensure the material is properly sealed i.e. the sample is not open or tampered with
- 1.3 Ensure that the correct AMIS CRM and quantity is received as per your order
- 1.4 Ensure the samples are stored in a cool dry area, in such a way that it does not compromise the integrity of the CRM

2. Preparation and correct use of material

- 2.1 The sample should be agitated for at least 15 seconds before opening the CRM
- 2.2 Ensure the weighing equipment and the area around the weighing balance is clean and free of contamination. The material should be weighed according to best weighing practices
- 2.3 AMIS certified values are reported on a dry-basis, the user laboratory is required to dry a portion CRM material in air at 105°C in a drying oven to constant mass to determine the moisture content.

3. Moisture factor calculation

Example: Moisture content of the CRM determined by the laboratory = 0.500%

- 3.1 The concentration of the CRM determined by the laboratory is 12.62% (dry basis). Calculating the moisture correction factor using the above values:

$$\text{Moisture correction factor} = 100 - 0.500 / 100 = 0.995$$

- 3.2 Multiplying the factor of 0.995 by determined analyte concentration on an air-dry basis:

$$0.995 \times 12.62\% = 12.56\%$$

4. Recommendations

4.1 Calibrations

- 4.1.1 Ensure all calibration standards and unknown samples are matrixed matched
- 4.1.2 Ensure all calibration standards are treated the same as unknown samples
- 4.1.3 The reported uncertified concentrations are to be used as indicative values only

4.2 Preparing Quality Control charts using AMIS CRMs

- 4.2.1 Over a period of time, analyse a minimum of 10 replicates
- 4.2.2 Evaluation of data:
 - 4.2.2.1 Remove outliers
 - 4.2.2.2 Determine the mean of replicates without the outliers and validate the obtained mean versus the certified value as follows:

The concentration of the CRM determined by the laboratory = 4.59%

CRM Certified Value	Expanded Uncertainty (U)	Coverage Factor (k)	Mean (n=9)	n	Standard Deviation (s)
4.62%	0.08%	2.25	4.59	9	0.01015

The standard uncertainty (u) is found by dividing the expanded uncertainty by the coverage factor:

$$u = \frac{0.08}{2.25} = 0.0356 \%$$

Using the observed mean for the replicate data ($n=9$) obtained for the CRM and substituting into

$$t_{calc} = \frac{|\bar{x} - \mu|}{\sqrt{0.0356^2 + \frac{0.01015^2}{9}}} = \frac{|4.59 - 4.62|}{\sqrt{0.00126 + 0.00001145}} = 0.84$$

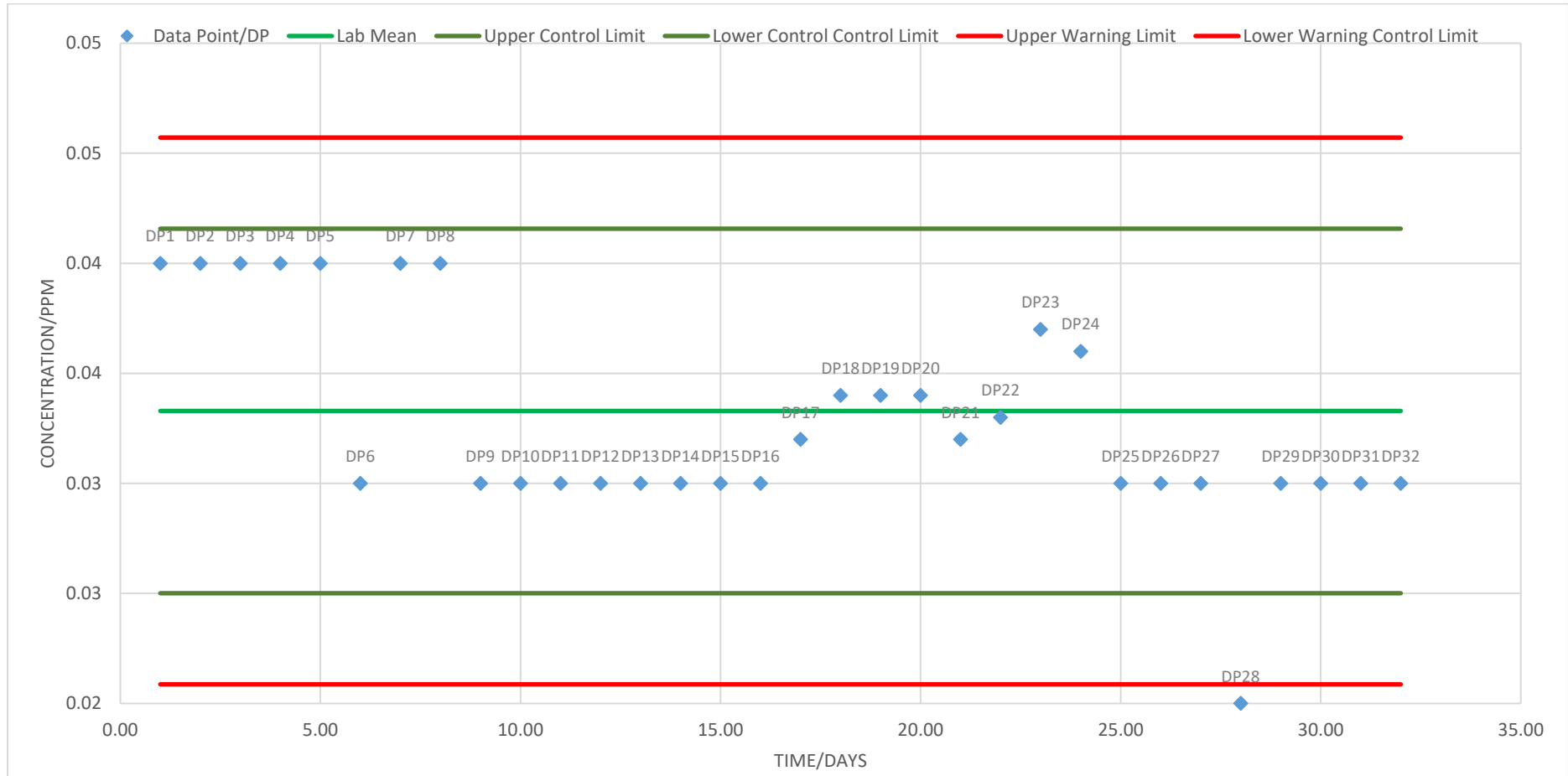
Therefore, $t_{calc} = 0.84$ and $t_{crit}(5\%, 8) = 2.31$ (df is 8, therefore, $t_{crit} = 2.31$) which is > 0.84 .

$t_{crit} > t_{calc}$, hence the laboratory mean is validated

NB: Should $t_{calc} > t_{crit}$, it indicates the method is not accurate and the laboratory should investigate factors affecting accuracy.

- 4.2.3 After the validation of the mean, the standard deviation should be calculated

4.2.4 Plot a graph that has an upper and lower control limit and upper and lower warning controlling limit as per below



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