

5 June 2014 EMA/CHMP/CVMP/QWP/63699/2014 Committee for Human Medicinal Products (CHMP) Committee for Veterinary Medicinal Products (CVMP)

Addendum to EMA/CHMP/CVMP/QWP/17760/2009 Rev 2: Defining the Scope of an NIRS Procedure

The latest revision of the 'guideline on the use of near infrared spectroscopy by the pharmaceutical industry and the data requirements for new submissions and variations' introduces the concept of the NIRS procedure **scope**, to facilitate continuous improvement and life cycle management.

Changes *within* the approved scope of the NIRS procedure are subject to GMP only. Changes *outside* of the approved scope of the NIRS procedure are subject to variation application. The definition of the scope is given in Section 4.1.1 of the guideline and the use of this scope to manage change control is further explained in Section 7 of the guideline.

As the concept of the NIRS procedure scope is new, this addendum has been produced to give a fictitious example of the scope for an NIRS procedure used for release testing for assay and content uniformity of the active substance in a finished product (uncoated tablet) and how changes to this scope would be managed according to the guideline.

Table 1 gives the method scope for the approved NIRS procedure.

Table 2 details those changes that would be considered WITHIN the scope of the procedure; therefore maintained under GMP only.

Table 3 details those changes that would be considered OUTSIDE of the scope of the procedure; therefore subject to variation application. The proposed changes may be the subject of a post approval change management protocol, with consequential downgrade of the variation type.



Table 1: Approved method scope for an assay and content uniformity NIRS procedure

Parameter	Procedure scope
Instrument	Foss XDS Masterlab: 3010-1019
	Foss XDS sampling module: 3010-0954
Software	Foss Vision Version 1
Mode	Transmission
Scan Rate; Number	30/second; number of scans 150
Sample presentation	At-line sample trays customised for the tablet shape (40 tablets)
Concentration range	75-125 %
Spectral pre-processing	- Standard normal variate (SNV)
	- 15 points Savitzky Golay 2 ND derivative on 800-1130 nm
Spectral quality check algorithm	Mahalanobis distance (MD) and residual variance (RV)
Spectral quality check algorithm	MD: max match value +1x standard deviation
threshold	RV: max match value +3x standard deviation
Chemometric algorithm	PLS
PLS model parameter	PLS spectral range: 1030-1140.
	Number of latent variables : 4
Statistical attributes	SEP
	Bias
	Slope
	Intercept
Reference method	Method code-0032 (HPLC method using UV detection)

Table 2: Changes $\underline{\text{within}}$ the scope of the approved NIRS procedure

Please note:

- 'NA' means that change cannot be implemented without prior regulatory approval;
- A change can occur solely or in combination with other changes

Parameter	Change that can be made within the scope	
Instrument	Foss XDS masterlab: 3010-1019 (no change)	
	Foss XDS sampling module: 3010-0954 (no change)	
	Change lamp (to same type, for maintenance)	
	Change detector (to same type, for maintenance)	
	Change spectrophotometer (same vendor) if demonstrated in reproducibility tests	
Software	Changed to Foss Vision Version 2	
Mode	NA	
Scan Rate; Number	Changed to 40/second or 20/second and / or number of scans changed to 100 or 200, if already demonstrated in robustness studies	
Sample presentation	Change of size of at-line sampling trays customised for the tablet shape (60	
	tablets or 30) if already demonstrated in robustness studies	
Concentration range	NA	
Spectral pre-processing	NA	
Spectral quality check algorithm	NA	
Spectral quality check	MD: max match value +1x standard deviation (no change)	
thresholds	RV: max match value +2x standard deviation (changed from 3 to 2)	
Chemometric algorithm	NA	
PLS model parameter	PLS spectral range: 1030-1120 (range tightened)	
	Number of latent variables : 4 (no change)	
	Change to introduce new samples to the calibration model (outliers, OOS) compliant with the scope of the procedure	
Statistical attributes	Met (no change to requirements)	
Reference method	Method code-0032	
	(changes to the reference method within Ph. Eur. tolerances, for which the principle of operation has not changed)	

Table 3: Changes outside of the scope of the approved NIRS procedure

Parameter	Changes considered beyond the scope of the procedure	Type of variation
Instrument	Change of vendor, e.g. to ABB FT NIR	Type 1B variation - BIId2.d
Software	Change of vendor, e.g. to GRAMS 9.0	Type 1B variation - BIId2.d
Mode	Changed to reflectance	Type 1B variation - BIId2.d
Scan Rate; Number	Changed to 40/second or 20/second and / or number of scans changed to 100 or 200, if not demonstrated in robustness studies	Type 1A variation - BIId2.a
Sample presentation	Change to on-line measurement rather than at-line	Type 1B variation - BIId2.d
Spectral pre- processing	Changed to: -10 point Savitzky Golay 2 ND derivative on 800-1130 nm -Multiple Scatter Correction (MSC) with first derivative	Type 1B variation - BIId2.d
Spectral quality check algorithm	Changed from PLS to SIMCA	Type 1B variation - BIId2.d
Spectral quality check threshold	MD: max match value +2x standard deviation (change to widen) RV: max match value +3x standard deviation (no change)	Type 1B variation - BIId2.d
Chemometric algorithm	MLR, PCR	Type 1B variation - BIId2.d
Model parameter	Changes such as: - Spectral range change: 1020-1170 - Number of latent variables: 5 (change from 4)	Type 1B variation - BIId2.d
Statistical attributes	Widened beyond initial approved statistical attributes	Such a change would not generally be submitted in isolation and would be related to another change. The change in statistical attributes would therefore be considered under the variation for the related change.
Reference method	Method reference-0050 (change in detection mode e.g. UV replaced by fluorescence, a change in mode of operation) or Method reference-0032 (Changes outside Ph.Eur. tolerances)	Implications for the NIRS procedure (e.g. its revalidation over time and the need for a change to the reference method stated in the scope of the NIRS procedure) should be considered under the related variation for the change in reference method

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