RESOLUTION OIV-OENO 675D-2022

SPECIFIC MONOGRAPHS FOR PROFISETINIDINS/PROROBITENIDINS

THE GENERAL ASSEMBLY,

IN VIEW OF Article 2, paragraph 2 iv of the Agreement of 3 April 2001 establishing the International Organisation of Vine and Wine,

CONSIDERING the work of the "Specifications of Oenological Products" Expert Group,

CONSIDERING the resolution OIV-OENO 624-2022 "Update to the monograph on oenological tannins" which describes a general monograph,

CONSIDERING the need to set up specific monographs for each family of tannins,

DECIDES, at the proposal of Commission II "Oenology", to add the monograph COEI-1-PROFIS to the *International Oenological Codex*:

OENOLOGICAL TANNINS

Specific monograph for oenological tannins containing profisetinidins/prorobitenidins

Profisetinidins/prorobitenidins are a sub-class of condensed (or proanthocyanic) tannins. Tannins from the quebracho and acacia spp. tree are included in this sub-class.

1. Method for the determination of sub-class affiliations

1.1. Characterisation by high-performance liquid chromatography (HPLC)

1.1.1. Principle

This method is designed to verify the presence of characteristic components of condensed tannins of the profisetinidin/prorobitenidin sub-group and measure their total concentration. It is applicable to preparations of oenological tannins that are declared as pure and thus as not containing tannins with small masses from different families or sub-families (or classes).

1.1.2. Reagents, material and apparatus

1.1.2.1. Reagents

(+)-catechin, CAS No. 154-23-4
Ultrafiltered water (resistivity: 18.3 MΩ·cm)
Water (HPLC quality)
Methanol (HPLC quality)
Formic acid (HPLC quality)

1.1.2.2. Materials

100-mL borosilicate-glass flask
Filters with 0.45 μm pore size diameter
Plastic 1-mL syringe

1.1.2.3. Apparatus

Technical balance with precision of 0.01 g
Analytical balance with precision of 0.1 mg
Class-A volumetric glassware
Mass chromatographic system with spectrometry detection composed of:

- Gradient pump for binary or quaternary mix

- Injector fitted with a loop of 10 μL

- Spectrophotometric detector at 280 nm fixe wavelength

- Column Eclipse Plus C-18 (for example): 2.1 x 100 mm, 1.8 μm particle size

- ESI-SIM (Single Ion Monitoring mode via Electro Spray Ionisation) ionisation source

- Mass spectrometer detector: quadrupole time of flight (Q-TOF)

1.1.3. Preparation of samples and standards

<u>Samples</u>: weigh approximately 0.5 g of oenological tannins on the analytical balance and make a note of the weight. Dissolve the oenological tannins in 100 mL of ultrafiltered water in a 100-mL borosilicate-glass flask and mix well.

Preparation of standard solutions: put 10 mg of (+)-catechin in solution into 50 mL of ultrafiltered water, corresponding to a 200 mg/L concentration. Then carry out dilutions in ultrafiltered water to obtain 5, 10, 20, 40, 60, 80 and 100 mg/L concentrations.

Solvent A: water (HPLC quality) containing 0.1% of formic acid.

Solvent B: methanol containing 0.1% of formic acid.

1.1.4. Procedure

The sample solutions and standard solutions are filtered on 0.45 μm (pore size diameter) filters and analysed by chromatography under the following conditions given by way of example:

Injected volume: 10 μL of sample solution or standard solution of (+)-catechin

Detection at 280 nm

Composition of elution gradient: (time, % of solvent A)

0min, 99.0%; 0.5 min, 94.0%; 20 min, 50.0%; 25 min, 0.0%; 32 min, 94.0% and 10 min for equilibrium.

Flow rate: 0.3 mL/min

Quantification and detection of the characteristic components of condensed tannins from the profisetinidin/prorobitenidin sub-group or sub-class according to the ESI-SIM scan and Q-ToF detection (for example).

Table 1: Example chemical formulas and m/z of the different profisetinidins/prorobitenidins (for the quebracho tree)

Compound	Chemical formula	m/z
(+)-catechin	$C_{15}H_{14}O_6$	290.1
(-)-epicatechin	$C_{15}H_{14}O_6$	290.1
Catechin-fisetinidol dimers	$C_{30}H_{26}O_{11}$	562.1
Catechin-fisetinidol trimers	$C_{45}H_{38}O_{16}$	834.2
Catechin-fisetinidol tetramers	$C_{60}H_{50}O_{21}$	1106.3

Table 2: Example chemical formulas and m/z of the different profisetinidins/prorobitenidins (for the acacia tree)

| Compound	Chemical formula	m/z
(+)-catechin	$C_{15}H_{14}O_6$	290.1
(-)-epicatechin	$C_{15}H_{14}O_6$	290.1
(-)-epicatechin-3-O-gallate	$C_{22}H_{18}O_{10}$	442.4
Fisetinidol-gallocatechin dimers	$C_{30}H_{26}O_{12}$	578.1
Robinetinidol-catechin dimers	$C_{30}H_{26}O_{12}$	578.1
Chalcan-flavan dimers (gambiriin)	$C_{30}H_{28}O_{12}$	580.1
Catechine-fisetinidol-robinetinidol or gallocatechin-fisetinidol-fisetinidol trimers	$C_{45}H_{58}O_{17}$	850.2
Robinetinidol-robinetinidol-catechin or gallocatechin-fisetinidol-robinetinidol trimers	$C_{45}H_{38}O_{17}$	866.2
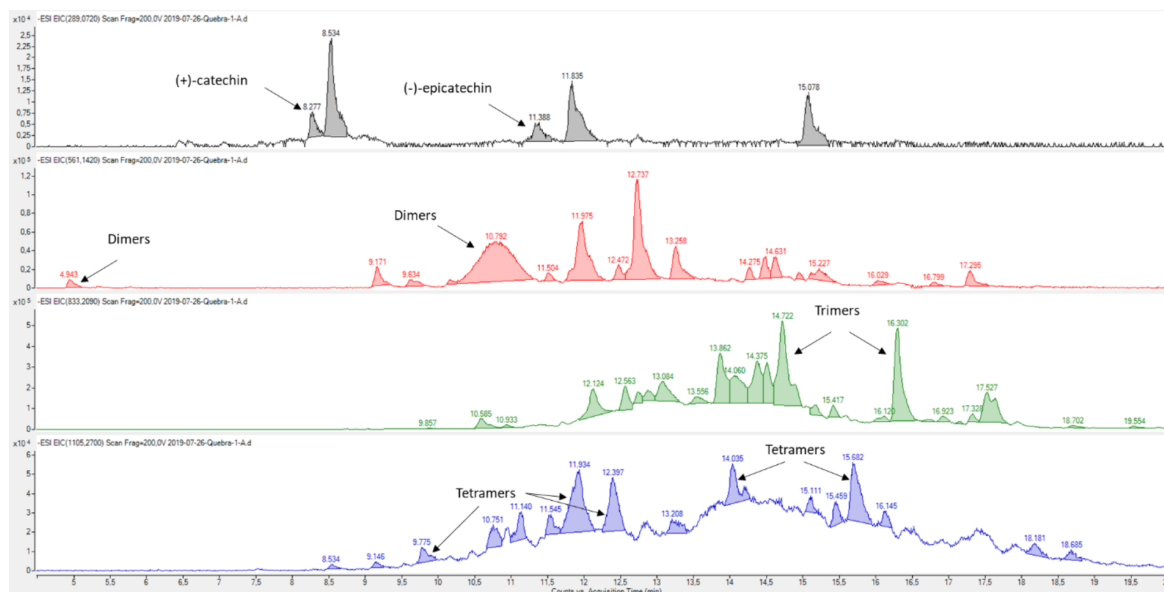
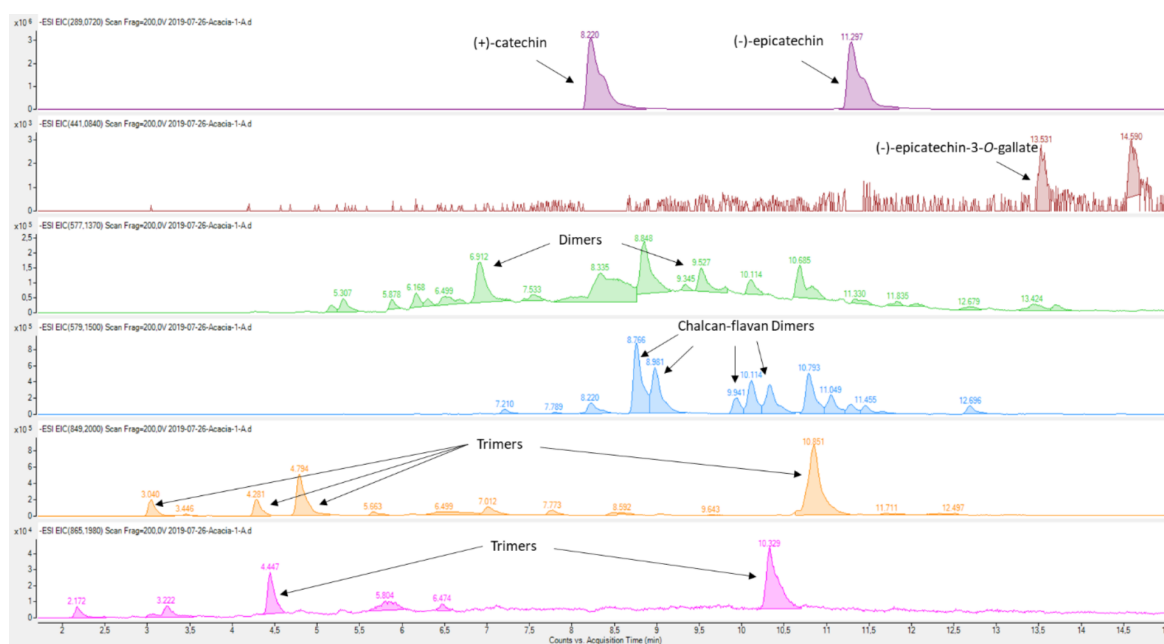*Figure 1: Example ESI-SIM scan of quebracho tree*

Figure 2: Example ESI-SIM scan of acacia tree

1.2.   Conclusion

An oenological tannin is recognised as a profisetinidin/prorobitenidin when:

- its total polyphenol content is higher than 65% (gravimetric method in Annex 1 of the general monograph OIV-OENO 624-2022),

- its molecular procyanidin/prodelphinidin content as characterised by the HPLC method is higher than 20 mg (for quebracho tannins) and 150 mg (for acacia tannins) equivalent of (+)-catechin per gram of oenological tannins.

2.   Methods of measurement of properties and functionalities

The following compliance methods and criteria are only applicable when the property/functionality is claimed on the preparation of tannins.

2.1.   Antioxidant ability

2.1.1.   Principle

Determination of profisetinidins/prorobitenidins' antioxidant ability to contribute to the protection of must and wine from oxidation.

2.1.2.   Products

2.1.2.1.   Antioxidant capacity

DPPH (2,2-diphenyl-1-picrylhydrazyl): MM = 394.32

Trolox (6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid): MM = 250.29

Methanol at 99.9% volume

96-well microplate reader (FLUOstar Omega - BMG Labtech, for example)

2.1.2.2.   Direct oxygen consumption (OCR)

Ethanol at 96% volume, CAS No. 64-17-5

Tartaric acid: MM = 150.09, CAS No. 87-69-4

Iron (III) chloride hexahydrate: MM = 270.30, CAS No. 7705-08-0

Copper (II) sulfate pentahydrate: MM = 249.68, CAS No. 7758-98-7

Clear glass bottles with inserted pills of 0.75-L capacity

NomaSens oximeter, for example

2.1.3. Protocols

2.1.3.1. Antioxidant capacity (DPPH assay)

<u>0.15 g/L oenological tannin solution</u>: dissolve 37.5 mg of oenological tannins in 500 mL of model wine solution (distilled water, 12% vol. of ethanol, 4 g/L of tartaric acid and pH adjusted to 3.5). Dilution of oenological tannins solution could be needed if the measurement absorbance is higher than 1 unit (in this case the dilution should be included in the calculation).

<u>1mM Trolox solution</u>: dissolve 125 mg of Trolox in 500 mL of model wine solution (distilled water, 12% vol. of ethanol, 4 g/L of tartaric acid and pH adjusted to 3.5).

<u>Calibration curve</u>: dissolve in 1, 0.8, 0.6, 0.4, 0.2 and 0.1 mL of 1 mM Trolox solution into 0, 0.2, 0.4, 0.6, 0.8 and 0.9 mL of model wine solution. These quantities correspond to 1, 0.8, 0.6, 0.4, 0.2 and 0.1 mM final concentration of Trolox respectively.

<u>6.10^{-5}M DPPH solution</u>: dissolve 2.36 mg of DPPH in 100 mL of methanol. The solution should be freshly prepared.

2.1.3.2. Direct oxygen consumption (OCR)

<u>1 g/L oenological tannin solution</u>: dissolve 0.75 g of oenological tannins in 750 mL of model wine solution.

<u>Model wine solution</u>: dissolve 4 g of tartaric acid, 2.25 mg of Iron (III) chloride hexahydrate and 0.225 mg of Copper (II) sulfate pentahydrate in 90 mL of ethanol and 660 mL of distilled water. The pH should be adjusted at 3.5.

2.1.4. Tests

2.1.4.1. Antioxidant capacity

First a blank containing solely the DPPH reagent (RB) is measured at 515 nm by placing 190 µL of DPPH solution (1.3.1) in all the wells of the plate. Then, add 10 µL of oenological tannin solution (samples), distilled water (blank) or Trolox curve solution (standards) into the wells and measure (MS) at 515 nm after 30 min.

See Figure 2 for an example of how to fill the plate.

The formula to be applied for the calculation of the antioxidant capacity is as follows:

$$1)\, RB - MS = x$$

$$2) \; antioxidant \; capacity \; (mg \; éq. \, Trolox \; per \; g \; of \; tannins) = \frac{250{,}29 \; (mg)}{0{,}15 \; (g)} \times \frac{x-b}{a}$$

where "a" and "b" correspond respectively to the slope and the constant of the Trolox calibration curve: Absorbance = f ([Trolox]) ◻ Absorbance = ax + b

In all cases, profisetinidins/prorobitenidins should demonstrate an antioxidant capacity, and more specifically they should have more than 450 ± 50 mg equivalent Trolox per gram of tannins (commercial extract).

	1	2	3	4	5	6	7	8	9	10	11	12
A	T 0.1	T 0.1	T 0.2	T 0.2	T 0.4	T 0.4	T 0.6	T 0.6	T 0.8	T 0.8	T 1	T 1
B	T 0.1	T 0.1	T 0.2	T 0.2	T 0.4	T 0.4	T 0.6	T 0.6	T 0.8	T 0.8	T 1	T 1
C	Blank	OT 1	OT 2	OT 3	OT 4	OT 5	OT 6	OT 7	OT 8	OT 9	OT 10	OT 11
D	Blank	OT 1	OT 2	OT 3	OT 4	OT 5	OT 6	OT 7	OT 8	OT 9	OT 10	OT 11
E	Blank	OT 1	OT 2	OT 3	OT 4	OT 5	OT 6	OT 7	OT 8	OT 9	OT 10	OT 11
F	Blank	OT 1	OT 2	OT 3	OT 4	OT 5	OT 6	OT 7	OT 8	OT 9	OT 10	OT 11
G	Blank	OT 1	OT 2	OT 3	OT 4	OT 5	OT 6	OT 7	OT 8	OT 9	OT 10	OT 11
H	Blank	OT 1	OT 2	OT 3	OT 4	OT 5	OT 6	OT 7	OT 8	OT 9	OT 10	OT 11

T = Trolox   OT = Oenological Tannins

Figure 3: Example 96-well plate

2.1.4.2. Direct oxygen consumption (OCR)

First the model wine solution is saturated with oxygen at 8 mg/L by bubbling with air for 10 min at 20-25 °C. Then, add the oenological tannins to the model wine solution in the bottles filled to 0.75 L. Seal the bottles hermetically and shake to fully homogenise.

1. Measure the oxygen consumed every two days starting 1h after the filled of the bottles.

2. To determine the oxygen consumption rate, follow the pathway as shown in Figure 2:

   - represent the oxygen consumption versus the time,

   - then represent the inverse of the oxygen consumed versus the inverse of the time,

   - the oxygen consumption rate corresponds to the inverse of the slope coefficient: OCR t_0 mg of O2 consumed per day and per g of tannins = 1/A, A being the slope

coefficient

In all cases, profisetinidins/prorobitenidins should demonstrate an ability to consume the oxygen directly, and more specifically they should be able to consume at least 0.10 ± 0.05 mg of O2 per litre, per day and per gram of tannins (commercial extract).
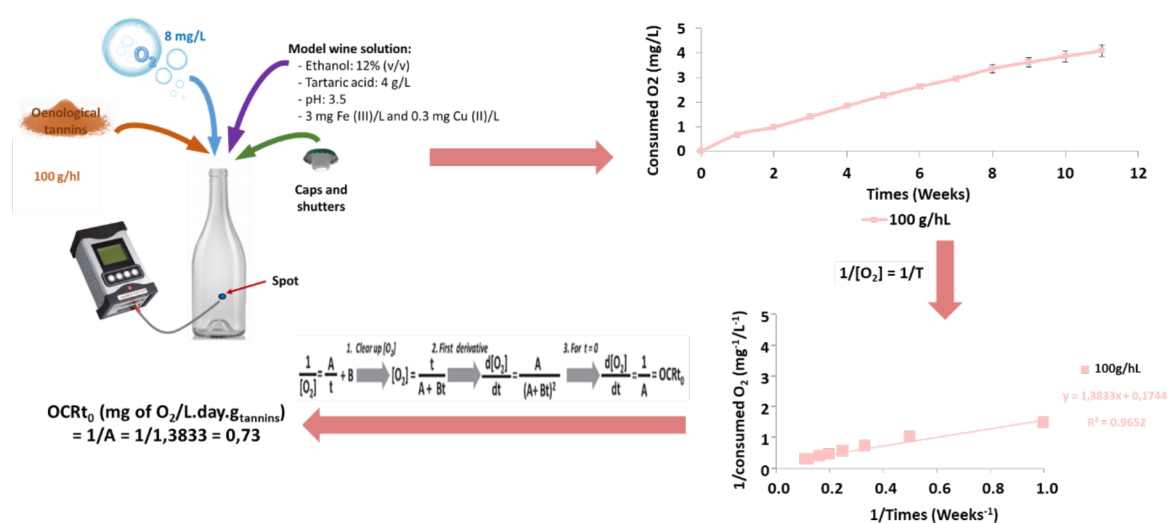

Figure 4: Pathway to determine oxygen consumption rate

2.2. Antioxidasic ability

2.2.1. Principle

Determination of profisetinidins/prorobitenidins' antioxidasic ability to contribute to antioxidasic protection in terms of the laccase activity of compounds in must and wine.

2.2.2. Products

Ethanol at 96% volume, CAS No. 64-17-5

Tartaric acid: MM = 150.09, CAS No. 87-69-4

Sodium acetate: MM = 82.03, CAS No. 6131-90-4

Syringaldazine (4-hydroxy-3,5-dimethoxybenzaldehyde azine): MM = 360.36, CAS No. 14414-32-5

Polyvinylpolypyrrolidone: PVPP, CAS No. 25249-54-1

Must botrytised with laccase activity

Distilled water (HPLC quality)

2.2.3. Protocols

<u>2 g/L oenological tannin solution:</u> dissolve 200 mg of oenological tannins in 100 mL of model wine solution (distilled water, 12% vol. of ethanol, 4 g/L of tartaric acid and pH adjusted to 3.5).

<u>Buffer solution (8.2 g/L):</u> dissolve 410 mg of sodium acetate in 50 mL of distilled water.

<u>Syringaldazine solution (0.06 g/L):</u> dissolve 30 mg of syringaldazine in 500 mL of ethanol.

2.2.4. Tests

1. Add 4 mL of botrytised must to 1 mL of oenological tannin solution in tube, which will correspond to sample modality.

2. Add 4 mL of botrytised must to 1 mL of model wine solution in tube, which will correspond to control modality.

3. After 4 minutes (precisely), add 0.8 g of PVPP in both tube (sample and control modalities), stirred and centrifuged for 10 minutes at 8,500 rpm.

4. Recover 1 mL of the supernatant (for both sample and control modalities), into 1.4 mL of buffer solution and 0.6 mL of syringaldazine solution. Put the mixture into a plastic spectrophotometer cuvette (10 mm path length).

5. Measure the absorbance at 530 nm every minute for 5 minutes (including time measurements at 0 minutes).

6. Then determine the laccase activity and the residual laccase activity by using the following equations and Figure 3:

$$Laccase\ activity = 46{,}15 \times \Delta A\ \mu mol\ L^{-1} min^{-1} = 46{,}15 \times \Delta A\ UL$$

$$\%\ of\ residual\ activity = (laccase\ activity\ _{sample}/laccase\ activity_{control}) \times 100$$
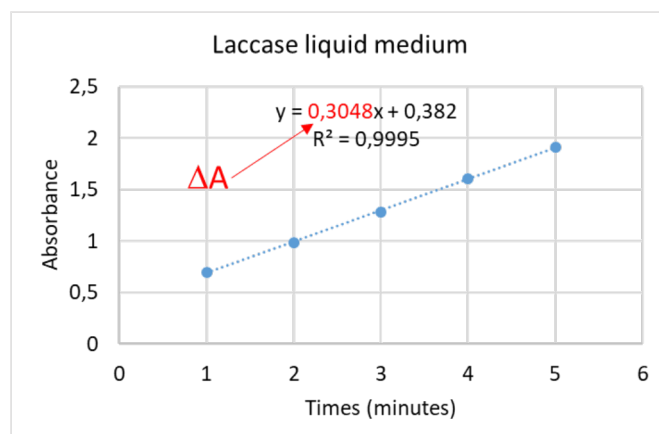
Figure 5: Example determination of ΔA.

In all cases, profisetinidins/prorobitenidins should demonstrate an antioxidasic ability, and more specifically they should be able to reduce the residual laccase activity by at least 40%. This value is valuable for must and wine containing less than 5 UL (units of laccases).

2.3. Colour stabilisation

2.3.1. Principle

Determination of profisetinidins/prorobitenidins' colour stabilisation properties to promote the expression, stabilisation and preservation of colour in red must and wine.

2.3.2. Products

Ethanol at 96% volume, CAS No. 64-17-5

Tartaric acid: MM = 150.09, CAS No. 87-69-4

Malvidin-3-O-glucoside: MM = 528.87, CAS No. 18470-06-9

2.3.3. Protocols

0.8 g/L oenological tannin solution: dissolve 80 mg of oenological tannins in 100 mL of model wine solution (distilled water, 12% vol. of ethanol, 4 g/L of tartaric acid and pH adjusted to 3.5).

0.1 g/L malvidin-3-O-glucoside solution: dissolve 10 mg of malvidin-3-O-glucoside in 100 mL of model wine solution (distilled water, 12% vol. of ethanol, 4 g/L of tartaric acid and pH adjusted to 3.5).

2.3.4.   Tests

1. Place 0.75 mL of oenological tannin solution and 0.75 mL of model wine solution in one 2-mL stoppered conical tube – hereinafter a "tube" – and keep it in the dark at room temperature. This tube will be called "T_0".

2. Place 0.75 mL of malvidin-3-O-glucoside solution and 0.75 mL of model wine solution in one tube and keep it in the dark at room temperature. This tube will be called "M".

3. Place 0.75 mL of oenological tannin solution and 0.75 mL of malvidin solution in one tube and keep it in the dark at room temperature. This tube will be called "T_M".

4. After 7 days, measure the absorbance at 450, 520, 570 and 630 nm of the three tubes (T_M, T_0 and M).

5. Subtract the absorbance values of T_0 to T_M to obtain the absorbance avoiding the interferences due to the "natural" colour of the oenological tannin.

$$A(T_M) - A(T_0) = A(T)$$

6. Then, determine the CIELAB coordinates (L*, a* and b*) corresponding to tannin solution with malvidin-3-O-glucoside (T) and malvidin-3-O-glucoside solution (M) with the free MSCV software (https://www.unirioja.es/color/descargas.shtml) or equivalent.

The formulas to be applied for the calculation of the copigmentation index are as follows:

$$1)\, \Delta E_{ab}.TS = \sqrt{(L^*_T - L^*_W) + (a*_T - a*_W) + (b*_T - b*_W)}$$

$$2)\, \Delta E_{ab}.CS = \sqrt{(L^*_M - L^*_W) + (a*_M - a*_W) + (b*_M - b*_W)}$$

$$3)\, Copigmentation\ Index\ (\%) = 100 \times \frac{\Delta E_{ab}.TS - \Delta E_{ab}.CS}{\Delta E_{ab}.CS}$$

ΔE_{ab}..**TS**: total colour difference between the solution of malvidin-3-O-glucoside containing commercial tannins (T) and a pure white colour solution (W).

ΔE_{ab}..**CS**: total colour difference between the solution of malvidin-3-O-glucoside (M) and a pure white colour solution (W).

The CIELAB coordinates of a pure white colour solution are L* = 100.00, a* = 0.00 and b* = 0.00.

In all cases, profisetinidins/prorobitenidins should demonstrate an ability to stabilise the colour, and more specifically their copigmentation index should read as higher than 3.0 ± 0.5% after 7 days.

<u>Note:</u> Alternative methods of determination can be used in place of any of the methods described, on the condition that these have been internally validated.

3.   Bibliography

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