

Certificate of Analysis (Ver.1.0)

Enniatins mix

1. General information

This document is designed and the certified value(s) and uncertainty(ies) are determined in accordance with ISO Guide 31[1].

2. Description of the Reference Material (RM)

Name:	Enniatins mix	
Catalog number:	STD#3219	
CAS number:	Enniatin A:2503-13-1;	Enniatin A ₁ :4530-21-6;
	Enniatin B:917-13-5;	Enniatin B ₁ :19914-20-6
Formula:	Enniatin A:C ₃₆ H ₆₃ N ₃ O ₉ ;	Enniatin A ₁ :C ₃₅ H ₆₁ N ₃ O ₉ ;
	Enniatin B:C ₃₃ H ₅₇ N ₃ O ₉ ;	Enniatin B ₁ :C ₃₄ H ₅₉ N ₃ O ₉
Formula weight:	Enniatin A: 681.90;	Enniatin A ₁ :667.87;
	Enniatin B:639.82;	Enniatin B ₁ :653.85
Lot #:	2B00D27	
Starting material :	Enniatin A,lot#J200110P,Pribolab Pte. Ltd. ; Enniatin A ₁ ,lot#J200110P,Pribolab Pte. Ltd. ; Enniatin B,lot#J200110P,Pribolab Pte. Ltd. ; Enniatin B ₁ ,lot#J200110P,Pribolab Pte. Ltd.	
Amount:	0.012mg dried down, 10µg/mL after reconstitution with 1.2 mL solvent	
Result concentration:	Enniatin A:10.23±0.12µg/mL ;Enniatin A ₁ :10.07±0.12µg/mL ; Enniatin B:10.38±0.12µg/mL ;Enniatin B ₁ :10.39±0.12µg/mL	
Production date:	26/Apr/2022	
Expiry Date:	25/Oct/2023	
Name of the supplier:	Pribolab Pte.Ltd.	

2.1 Intended use of the RM

- for laboratory use only
- internal standard[2]

2.2 Reconstitution instruction

The standard that you have received may appear at first glance, as an empty vial. The target compound (s) is (are) in a film at the bottom of the vial. Do not open the vial until you are ready to reconstitute

To reconstitute this RM use the following procedure:

1. Add 1.2mL of acetonitrile with a graduated syringe or a volumetric pipette.
2. Cap vial tightly.
3. Mix vigorously on a vortex mixer and repeat for several times over a longer period or sonicate at room temperature.
4. Always keep vial tightly capped.
5. Store the reconstituted standard at -20°C in a dark environment for max. 8 weeks.
6. Store immediately at -20°C after usage to avoid degradation.

Note: Thorough mixing and sufficient mixing time is required to ensure complete reconstitution of the dried down standard!

2.3 Instruction for the correct use of the RM

The compound should be stored at -20°C in a dark place. Before usage of the RM, the compound should be allowed to warm to temperature (20±3°C). The recommended minimum sub-sample amount for all kinds of application is 100 µL. The expiry date of this RM is based on the current knowledge and holds only for proper storage conditions in the originally closed flasks/packages.

2.4 Hazardous situation

The normal laboratory safety precautions should be observed when working with this RM. Further details for the handling of this RM are available as safety data sheet.

3. Certified values and their uncertainties

Enniatins mix					
Compound	Mass concentration ^a		Compound	Mass concentration ^a	
	Certified value ^b	Uncertainty ^c		Certified value ^b	Uncertainty ^c
Enniatin A	10.23 µg/mL	±0.12 µg/mL	Enniatin A ₁	10.07 µg/mL	±0.16 µg/mL
Enniatin B	10.38 µg/mL	±0.14 µg/mL	Enniatin B ₁	10.39 µg/mL	±0.18 µg/mL

3.1 Calculation of uncertainty

The uncertainty of the calibrant solution was calculated on the basis of preparation [4].

Uncertainty components	Description	Standard uncertainty(u)	
Purity (P) of solid Enniatin A Enniatin A ₁ Enniatin B Enniatin B ₁	P ₁ =99.1±0.9% P ₂ =98.6±1.4% P ₃ =98.9±1.1% P ₄ =98.4±1.6%	u(P₁)=0.5% u(P₂)=0.8% u(P₃)=0.6% u(P₄)=0.9%	a
Weighing procedure: Weighted sample: m _{Enniatin A} =2.065mg m _{Enniatin A1} = 2.043mg m _{Enniatin B} = 2.100mg m _{Enniatin B1} = 2.112mg	U _(m) =0.0000008g+1.30*10 ⁻⁵ *m _{Toxin} u _(m) =U _(m) /2	u_(m)=0.0004mg	b
Dilution procedure Volumetric flask:V _f =200mL	calibration: 200±0.15mL repeatability: 0.03mL volume expansion solvent	u(cal)=0.06mL u(rep)=0.03mL u(Vol.exp.)=0.47mL u(v)=0.5mL	c d e f

a Maximum tolerance of purity (rectangular distribution) was divided by $\sqrt{3}$

b Calculation of this u-value is based upon the uncertainty formula for the weighed amount as given in the calibration report from annual balance calibration

c A triangular distribution (division by $\sqrt{6}$) was chosen for the calculation of u(cal)

d Based on a series of ten fill and weigh experiments on a typical 200mL flask; the value was used directly as a standard deviation

e Based on the density of 0.7857 g/cm³ at temperature T=20°C and a maximum temperature variation of ±3°C,of volume expansion, relative volume expansion coefficient of acetonitrile is 1370*10⁻⁶/°C[5],volume expansion term(rectangular distribution)was divided by $\sqrt{3}$

f The three contributions are combined to give the u(v)= $\sqrt{u(\text{cal})^2 + u(\text{rep})^2 + u(\text{Vol.exp})^2}$

Calculation of the combined uncertainty u_c and the expanded standard uncertainty U

$$C_{\text{Toxin}} = \frac{10 \times m_{\text{ws}} \times P_1}{V_f} = \frac{10 \times 2.065 \times 99.1}{200} = 10.23 \text{ mg / L}$$

$$\frac{u_c(C_{\text{Toxin}})}{C_{\text{Toxin}}} = \sqrt{\left[\frac{u(P_1)}{P_1}\right]^2 + \left[\frac{u(m)}{m_{\text{ws}}}\right]^2 + \left[\frac{u(V)}{V_f}\right]^2} = \sqrt{\left[\frac{0.5}{99.1}\right]^2 + \left[\frac{0.0004}{2.065}\right]^2 + \left[\frac{0.5}{200}\right]^2} = 0.006$$

$$u_c(C_{\text{Toxin}}) = C_{\text{Toxin}} \times 0.006 = 10.23 \times 0.006 = 0.06 \text{ mg / L}$$

calculation of expanded standard uncertainty U using a coverage factor k=2

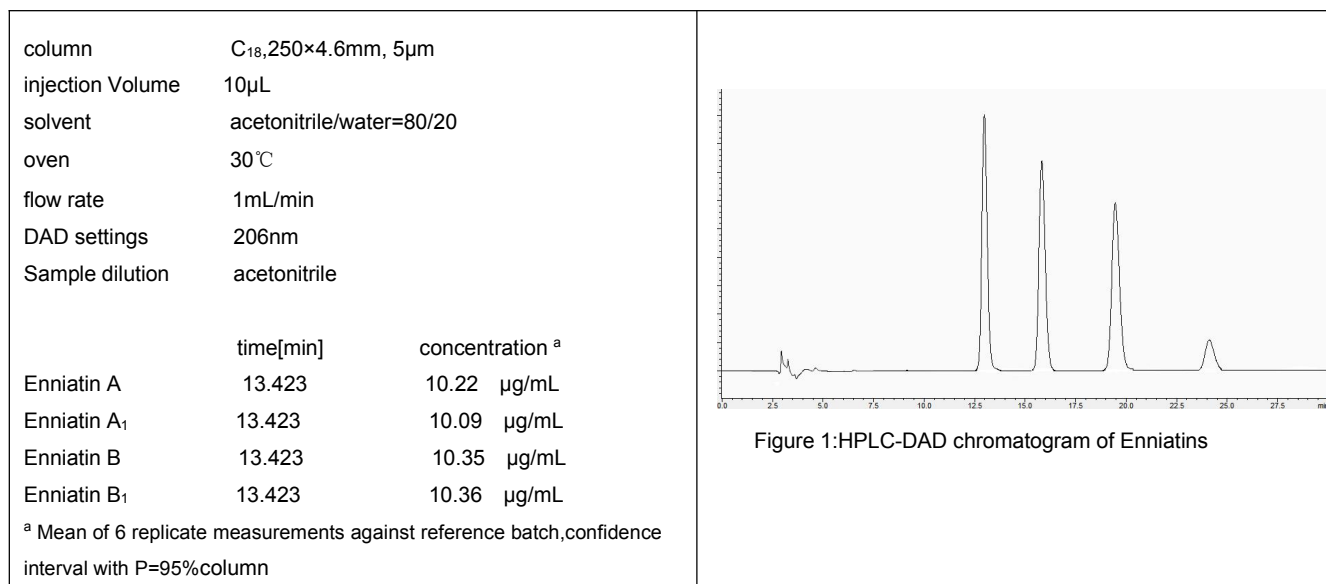
$$U(C_{\text{Toxin}}) = u_c(C_{\text{Toxin}}) \times 2 = 0.06 \times 2 = 0.12 \mu\text{g / mL}$$

4.Discussion of traceability

This calibrant is certified on the basis of gravimetric preparation [5]. Thus the certified value(mass concentration of Enniatins are based on the weighed amount of the starting material and is therefore traceable to the stated purity of the solid raw material.High purity material represents a practical realization of concentration units,through conversion of mass to molar quality.

5. Confirmation of certified value by HPLC-DAD

The certified concentration of Enniatins of the gravimetric prepared solution was confirmed by HPLC-DAD against an independently prepared reference batch of Enniatins.



6. Further information

The purchaser must determine the suitability of this product for its particular use. Pribolab makes no warranty of any kind, express or implied, other than its products meet all quality control standards set by Pribolab. We do not guarantee that the product can be used for a special application.

Inspected by

Zifne Cao
 Quality System Specialist

References:

- [1]ISO Guide 31:2015 - 1-18, "Reference materials – contents of certificates, labels and accompanying documentation"
- [2]G. Häubl, F. Berthiller, R. Krska, R. Schuhmacher, "Suitability of a fully ¹³C isotope labelled internal standard for the determination of the mycotoxin deoxynivalenol by LC-MS/MS without clean-up", Anal. Bioanal. Chem. 384 (3), (2006), 692-696
- [3] International Organization for Standardization (ISO), (2008), "Guide to the expression of uncertainty in measurement", (GUM 1995 with minor corrections) 1st Ed. Geneva, Switzerland
- [4] R.D. Josephs, R. Krska, S. MacDonald, P. Wilson, H. Pettersson, J. AOAC Int. 86, 50-60, (2003), "Preparation of a Calibrant as Certified Reference Material for Determination of the Fusarium Mycotoxin Zearalenone"
- [5] E.W. Flick, (1998), "Industrial Solvents Handbook", 5th Ed., Noyes Data Corp. Westwood NJ