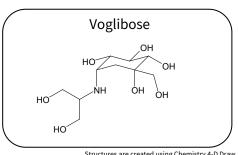
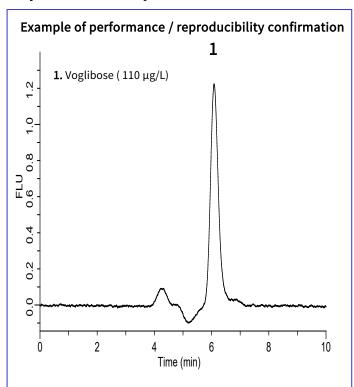
# Analysis of Voglibose Tablet Based on the Japanese Pharmacopoeia, 18th Edition.

Voglibose tablet is listed in the Japanese Pharmacopoeia and the post-column HPLC-fluorescence method is adopted for dissolution and quantitation method. In pharmacopoeia, cooling down is required by the specified pipe arrangement and at the specified temperature, after the reaction in post-column system. This time we used Chromaster (HITACHI), installing two sets of ovens can fulfill the requirement. (K. Suzuki)

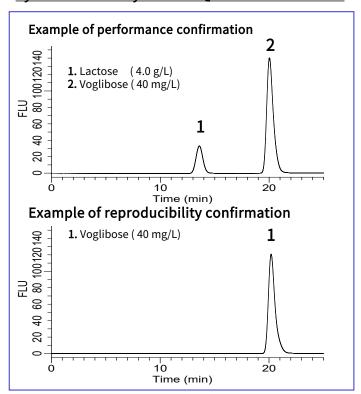


Structures are created using Chemistry 4-D Draw which is provided by ChemInnovation Software, Inc.

## **System Suitability Test for Dissolution**



## System Suitability Test for Quantitation Method



#### System suitability test

- For each 100 μL of standard solution, the peak theoretical plate number and the symmetry factor are more than 2000 plates and less than 1.5, respectively.
- 2. When the analysis is repeated 6 times, the relative standard deviation of peak area is less than 3.0%.

#### Result this time

Theoretical plate number  $: 2,433 \ (\ge 2,000)$ Symmetry factor  $: 1.19 \ (\le 1.5)$ Relative standard deviation  $: 0.99 \ \% \ (\le 3.0)$ 

Table 1: Reproducibility of peak area value

	Peak area value
Average	92130
RSD(%)	0.99

#### System suitability test

- 1. For each  $50\mu L$  of this test solution, lactose and voglibose elute in order and the degree of separation is greater than 4
- 2. When the analysis is repeated 6 times, the relative standard deviation of peak area is less than 2.0%.

#### Result this time

Order of elution :  $\frac{OK}{Resolution}$  :  $\frac{5.7}{(\ge 4)}$  :  $\frac{1.23 \%}{(\le 2.0)}$ 

Table 2: Reproducibility of peak area value

	Peak area value
Average	21149069
RSD(%)	1.23

## **HPLC** condithions

#### Dissolution

#### **HPLC** condition

Column : Inertsil NH2 (5  $\mu$ m, 150  $\times$  4.0 mm I.D.)

Cat. No. :5020-05535 Eluent : A) CH<sub>3</sub>CN

B) 10 mM Na<sub>2</sub>HPO<sub>4</sub> (pH 6.5; 10 mM NaH<sub>2</sub>PO<sub>4</sub>)

A/B = 500/500, v/v (premix)

: 12 mM NaIO<sub>4</sub> + 50 mM Taurine Reaction Solution

Reaction coil : PTFE 0.5 mm ID x 22 m

(20 m (inside of oven) + 2 m(for connection))

Flow rate : Eluent; 0.85 mL/min

Reacthion solution; 0.85 mL/min

Adjusted in order for the retention time to be 6 min.

Column temperature :25°C :100°C Reaction temperature Cooling temperature : 25 °C : PTFE 0.33 mm ID x 2.5 m Cooling coil

(2 m (Inside of oven) + 0.5 m(for connection))

Detection : FL Ex 350 nm, Em 430 nm (PMT Low)

Injection volume : 100 µL

## Quantitation method

#### **HPLC** conditions

Flow rate

: Inertsil NH2 (5  $\mu$ m, 150  $\times$  4.0 mm I.D.) Column

:5020-05535 Cat. No. Eluent : A) CH<sub>3</sub>CN

B) 10 mM Na<sub>2</sub>HPO<sub>4</sub> (pH 6.5; 10 mM NaH<sub>2</sub>PO<sub>4</sub>)

A/B = 600/300 v/v (premix)Reaction solution: 12 mM NaIO<sub>4</sub> + 50 mM Taurine

Reaction coil: PTFE 0.5 mm ID x 22 m

(20 m (inside of oven) + 2 m(for connection)) : eluent; 0.56 mL/min

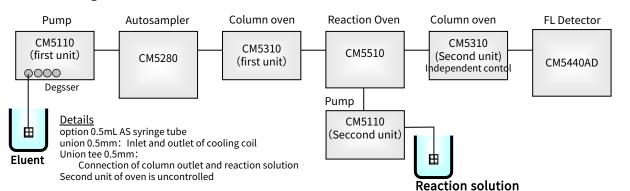
reaction solution; 0.56 mL/min \* Adjusted in order for the

retention time to be approximately 20 min. Column temperature Reaction temperature :100°C : 15 °C Cooling temperature Cooling coil : PTFE 0.33 mm ID x 2.5 m

(2 m (inside of oven) + 0.5 m(for connection)) Detection : FL Ex 350 nm, Em 430 nm (PMT Low)

Injection volume :50 µL

## Flow daigram



## Instrument used (only for reference)

## HPLC System Hitachi HPLC system Chromaster

No.	Name of product	Model number
1	Organizer	
2	Detector (FL)	5440AD
3	Column oven	5310
4	Autosampler	5280
5	Pump	5110
6	Reaction oven	5510



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