

Food & Beverage Series

SIMULTANEOUS MEASUREMENT OF GLUCOSE AND SUCROSE IN CEREAL PRODUCTS



Introduction

Dextrose (D-glucose) and sucrose concentrations in complex matrices such as cereal products can be measured directly and quickly using the YSI 2900 Series Biochemistry Analyzer. YSI's unique enzyme technology provides for specific glucose and sucrose measurements. Measurements are virtually unaffected by color, turbidity, density, pH, or the presence of reducing substances.

When a 2900 Series Biochemistry Analyzer is equipped with a glucose and a sucrose membrane, simultaneous measurement of both analytes is possible. Because glucose interferes with sucrose analysis, it is necessary to follow this protocol when analyzing for sucrose in the presence of glucose.

When a sample is injected into the sample chamber, the sucrose diffuses to the sucrose membrane, which contains invertase, mutarotase, and glucose oxidase. The sucrose is hydrolyzed to α -D-glucose and fructose. The mutarotase

allows for the quick equilibrium of glucose between its a and b forms. In the presence of glucose oxidase, the α -D-glucose (glucose) is oxidized to hydrogen peroxide and D-glucono- α -lactone. The hydrogen peroxide is detected amperometrically at the platinum electrode surface. The glucose in the sample diffuses to both the glucose and sucrose membranes, which contain glucose oxidase, and is oxidized as well.

Subtracting the response of the glucose membrane from the response of the sucrose membrane yields the response due to sucrose alone. The glucose response is

taken directly from the glucose membrane. The algorithm in the instrument software calculates the net concentrations. For more information on this system, refer to the Operations Manual.



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I. Materials & Setup

- A. YSI 2900 Series Biochemistry Analyzer equipped with a 2703 Sucrose Membrane, a 2365 Glucose Membrane and 2357 Buffer.
- B. Glucose (2.50 g/L,25.00 g/L) and Sucrose (5.00 g/L, 25.0 g/L) standard solutions.
- C. Buffer Diluent (40 g/L NaH₂PO₄, 10g/L Na₂HPO₄ in reagent water).
- D. Connect the 2900 Series instrument to a suitable power source.
- E. Perform the instrument and membrane daily checks described in the Operations Manual.

- F. Volumetric glassware (Class A recommended).
- G. The following instrument setup is recommended: Sample Size 10 μL

Probe A Parameters		Probe B Parameters		
Chemistry	Glucose	Chemistry	Sucrose	
Unit	g/L	Unit	g/L	
Calibrator	2.50	Calibrator	5.00 g/L	
End Point	30 Sec	End Point	30 Sec	

Autocal Parameters

Temperature	1°C
Time	30 Mir
Sample	5 Sam
Cal Shift	2%

II. Method

- A. Grind sample to a fine powder.
- B. Weigh 1.000 to 5.000 g of powdered sample.
- C. Transfer the sample to a 100 mL volumetric flask using buffer diluent to rinse and dilute. Fill the flask to the mark with buffer and mix. Allow the solution to equilibrate for about twenty minutes before analysis.
- D. Calibrate the 2900 Series instrument with a 2.50 g/L glucose and 5.00 g/L sucrose standard solutions. (at Stations #1 and #2, respectively).
- E. Check the linearity of the membranes at least once a day by injection of glucose (25.00 g/L) and sucrose (25.0 g/L) linearity check solutions. Refer to the Operations Manual for specifications.

- F. Assay the sample prepared in B by aspiration into the 2900 Series instrument.*
- G. Calibrate frequently as described in the Operations Manual.
 - * The linear range of the system is 0 to 25.0 g/L for both glucose and sucrose. The combined concentration of glucose + sucrose cannot exceed 25 g/L. If the sum of the values reported exceeds this, further dilution of the sample is required. If the glucose concentration exceeds the sucrose concentration, accuracy and precision may be compromised due to the software algorithm that subtracts glucose from sucrose. To avoid compromising accuracy refer to Application Note 204LS.

III. Calculations

To calculate % glucose and sucrose, multiply the values reported by the appropriate dilution factor.

Example: A cereal sample (4.336 g) was prepared and assayed as described. The values reported were 0.62 g/L glucose and 9.88 g/L sucrose.

% Glucose: 0.62 g/L x 0.100L /4.336 g	= 0.0143 g glucose/g cereal = 1.43% (w/w)
% Sucrose: 9.88 g/L x 0.100L /4.336 g	= 0.2279 g sucrose/g cereal = 22.8% (w/w)

Several cereal samples were assayed using YSI technology and HPLC. The results are listed below.

	Sucro	ose (%)	Glucc	se (%)	
Sample	YSI	HPLC	YSI	HPLC	Label (%)*
А	35.0	36.0	1.24	0.96	38.8
В	21.4	22.2	0.85	0.50	21.1
С	22.4	23.1	1.52	1.52	24.6
D	17.4	18.3	0.27	0.07	28.0
Е	25.2	25.4	1.61	1.33	28.2
F	29.8	27.9	3.51	4.14	35.2
G	32.2	31.1	3.01	2.42	38.7
Н	32.9	30.4	0.71	0.43	38.7
I	5.40	6.70	1.03	0.70	7.0





Ordering Information

Biochemistry Analyzer

2900

2365	Glucose Membrane Kit
2776	Glucose Standard Solution (2.50 g/L)
2777	Glucose Standard Solution (25.00 g/L)
2703	Sucrose Membrane Kit
2780	Sucrose Standard Solution (5.00 g/L)
2778	Sucrose Standard Solution (25.0 g/L)
2357	Buffer Kit
2363	Potassium Ferrocyanide Test Solution
2392	NaCl Solution (for membrane installation)



YSI Life Sciences develops and manufactures scientific instruments, sensors and systems that serve a variety of scientific and industrial markets worldwide. YSI has a long history in the life sciences and bioanalytical markets, most notably with our introduction of the world's first commercial whole blood glucose analyzer in 1975. Today there are over 10,000 YSI instruments installed around the world, trusted in critical situations to provide the most accurate data in the shortest time.

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