Food & Beverage Series

SIMULTANEOUS MEASUREMENT OF GLUCOSE AND SUCROSE IN BAKED GOODS

Application Note 215LS
YSI Life Sciences

Life Sciences
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**INTRODUCTION**

Dextrose (D-glucose) and sucrose concentrations in complex matrices such as baked goods can be measured directly and quickly using the YSI 2900 Series Biochemistry Analyzer. YSI’s unique enzyme technology provides for rapid 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 α and β forms. In the presence of glucose oxidase, the β-D-glucose (glucose) is oxidized to hydrogen peroxide and D-glucono-δ-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.

**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

<table>
<thead>
<tr>
<th>Probe A Parameters</th>
<th>Probe B Parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chemistry Glucose</td>
<td>Chemistry Sucrose</td>
</tr>
<tr>
<td>Unit g/L</td>
<td>Unit g/L</td>
</tr>
<tr>
<td>Calibrator 2.50</td>
<td>Calibrator 5.00 g/L</td>
</tr>
<tr>
<td>End Point 30 Sec</td>
<td>End Point 30 Sec</td>
</tr>
</tbody>
</table>

**Autocal Parameters**

<table>
<thead>
<tr>
<th>Temperature</th>
<th>Time</th>
<th>Sample</th>
<th>Cal Shift</th>
</tr>
</thead>
<tbody>
<tr>
<td>1°C</td>
<td>30 Min</td>
<td>5 Sam</td>
<td>2%</td>
</tr>
</tbody>
</table>

**II. METHOD**

A. Grind sample to a fine powder.

B. Weigh 1.00 to 5.00 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 2.50 g/L glucose and 5.00 g/L sucrose standard solutions.
E. Check the linearity of the membranes at least once a day by injection of glucose (25.0 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 baked muffin sample (4.654 g) was prepared and assayed as described. The values reported were 0.81 g/L glucose and 8.54 g/L sucrose.

\[
\begin{align*}
\text{% Glucose:} & \quad 0.81 \text{ g/L} \times \frac{0.100 \text{L}}{4.654 \text{ g}} \\
& = 0.0174 \text{ g glucose/g muffin} \\
& = 1.74\% (w/w)
\end{align*}
\]

\[
\begin{align*}
\text{% Sucrose:} & \quad 8.54 \text{ g/L} \times \frac{0.100 \text{L}}{4.654 \text{ g}} \\
& = 0.1834 \text{ g sucrose/g muffin} \\
& = 18.3\% (w/w)
\end{align*}
\]