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# EZScreen<sup>™</sup> Glucose Colorimetric Assay Kit (384 Well)

(Catalog # K950-384; 384 assays; Store at -20°C)

#### I. Introduction:

Glucose (C<sub>6</sub>H<sub>12</sub>O<sub>6</sub>, FW: 180.16) is delivered to the cells from the blood stream as the primary biological fuel to generate the universal energy molecule, ATP. Even though human body has evolved an advanced metabolic pathway to keep the blood glucose level in normal range, failure happens and that leads to develop conditions of persistent high or low blood sugar. Diabetes mellitus is one such most important metabolic disorder where an increased blood-sugar level develops compare to normal blood sugar level. Glucose level in blood is also a key diagnostic parameter for many metabolic disorders and its measurement is very important in both research and drug discovery. BioVision's EZScreen<sup>TM</sup> Glucose Colorimetric Assay Kit, uses a glucose enzyme mix which oxidizes glucose specifically and produces a product which reacts with a chromophore generating a stable signal (OD: 590 nm). This color is directly proportional to the amount of glucose present in the sample. The method is quantitative, rapid, simple, sensitive, and designed to be used in high throughput settings. The kit can detect 0.5 to 5 mM of glucose in various biological samples.

#### II. Applications:

- · Measurement of Glucose in various biological samples
- · Growth Media
- · Analysis of carbohydrate metabolism
- · Analysis of glucose content in foodstuff

## III. Sample Type:

- · Serum, plasma, & other body fluids
- · Growth media
- · Bacteria, yeast cultured samples
- · Milk, juice, etc.

#### IV. Kit Contents:

Components	K950-384	Cap Code	Part Number
EZScreen <sup>™</sup> Glucose Assay Buffer	25 ml	WM	K950-384-1
EZScreen <sup>™</sup> Glucose Probe	0.8 ml	Red	K950-384-2
EZScreen <sup>™</sup> Glucose Enzyme Mix (lyophilized)	1 vial	Green	K950-384-3
EZScreen <sup>™</sup> Glucose Standard (100 mM)	100 µl	Yellow	K950-384-4

## V. User Supplied Reagents and Equipment:

- 384-well clear plate with flat bottom
- · Multi-well spectrophotometer with 384-well plate reading capability

#### VI. Storage Conditions and Reagent Preparation:

Store kit at -20°C, protected from light. Briefly centrifuge small vials prior to opening. Read entire protocol before performing the assay.

- EZScreen<sup>TM</sup> Glucose Assay Buffer: Warm to room temperature prior to use. Store at -20°C or 4°C.
- EZScreen<sup>TM</sup> Glucose Probe: Ready to use as supplied. Warm to room temperature prior to use to thaw contents. Store at -20°C, protect from light and moisture. Use within two months. Aliquot to avoid multiple freeze-thaw cycles.
- EZScreen<sup>™</sup> Glucose Enzyme Mix: Dissolve in 220 µl EZScreen<sup>™</sup> Glucose Assay Buffer. Aliquot & store at -20°C. Keep on ice while
  in use. Use within two months.

#### VII. Glucose Assay Protocol:

1. Sample Preparation: Add 1.0 to 12.5 µl of sample directly to a 384 well clear flat bottom plate. Adjust the volume to 12.5 µl/well with EZScreen<sup>TM</sup> Glucose Assay Buffer. For serum, limit sample volume to 1 µl or dilute serum. Serum in healthy patients contains <6.0 nmol/µl glucose. Adjust the final volume to 12.5 µl with EZScreen<sup>TM</sup> Glucose Assay Buffer.

### Notes:

- a. For unknown samples, we suggest performing a pilot experiment & testing different sample dilutions with the EZScreen<sup>™</sup> Assay Buffer to ensure the readings are within the Standard Curve range. Though this kit has been optimized using serum from a healthy donor by adding sample directly to the well, pilot experiments are strongly encouraged to be carried out if samples are suspected to have abnormal Glucose concentrations.
- b. For sample having background, prepare parallel background well(s) containing same amount of sample as in the test well.
- c. Endogenous enzyme activity may cause loss of glucose. Samples should be deproteinized using a 10 kDa Spin Column (Cat. # 1997)
- d. Instrument reader settings need to be adjusted according to the chosen 384-well plate. (The right dimension of the used 384-well plate may be available in the manual provided by the plate-manufacturer).
- 2. Glucose Standard Curve: Dilute the Glucose Standard to 0.5 mM by adding 5 μl of the EZScreen<sup>TM</sup> Glucose Standard to 995 μl of EZScreen<sup>TM</sup> Glucose Assay Buffer, mix well. Add 0, 2, 4, 6, 8, 10 μl into a series of wells on a 384 well plate. Adjust volume to 12.5 μl/well with Glucose Assay Buffer to generate 0, 1, 2, 3, 4, 5 nmol/well of Glucose Standard..
- 3. Glucose Reaction Mix: Mix enough reagents for the number of assays to be performed. For each well, prepare 12.5 µl Mix containing:





	Reaction Mix	*Background Control Mix
EZScreen <sup>™</sup> Glucose Assay Buffer	10.0 µl	10.5 µl
EZScreen <sup>™</sup> Glucose Probe	2.0 µl	2.0 µl
EZScreen <sup>™</sup> Glucose Enzyme Mix	0.5 μΙ	

Mix and add 12.5  $\mu$ I of the Reaction Mix to each well containing the Standard, and test samples. Mix well.

- \* For samples having background, add 12.5 µl of the background control mix to sample background control well(s).
- 4. Measurement: Incubate the reaction for 30 min. at 37°C, protect from light. Measure absorbance (OD 590 nm) in a microplate reader.
- **5. Calculation:** Subtract 0 Standard reading from all readings which will be the corrected absorbance readings. If the sample background control reading is significant then subtract the sample background control reading from sample reading. Plot the Glucose Standard Curve. Apply the corrected absorbance of the sample to the Glucose Standard Curve to get B nmol of Glucose in the sample well.

## Sample Glucose concentration (C) = B/V X D nmol/µl or mM

Where: **B** = is the amount of Glucose in the sample well from Standard Curve

V = sample volume added into the reaction well ( $\mu$ I).

**D** = Dilution Factor

Glucose molecular weight: 180.2 g/mol

 $1 \text{ mM} \equiv 18.08 \text{ mg/dl}$ 

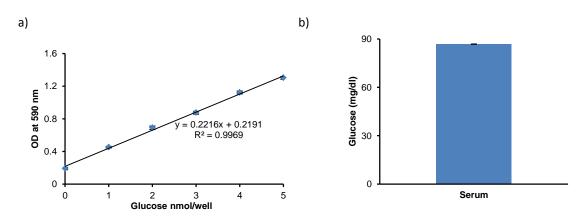


Figure: (a) Glucose Standard Curve; (b) Quantitation of Glucose in human serum. Serum samples were deproteinized using a 10 kDa Spin Column (10000xg, 10 min, 4°C). Undiluted serum filtrate (1 μl) samples were added to the wells directly. Assays were performed according to the kit protocol. Calculated concentration: 86.78 ± 0.1 mg/dl.

## **VIII. RELATED PRODUCTS:**

Glucose Colorimetric/Fluorometric Assay Kit (K606)
Glucose Colorimetric Assay Kit II (K686)
Glucose-6-Phosphate Dehydrogenase Assay Kit (K657)
PicoProbe™ Glucose Fluorometric Assay Kit (K688)
Glucose uptake Fluorometric Assay Kit (K666)
Glucose Dehydrogenase Activity Colorimetric Assay Kit (K786)

2-NBDG Glucose Uptake Assay Kit (Cell-Based) (K682) Glucose-6-Phosphate Dehydrogenase Inhibitor Screening Kit (Colorimetric) (K757)

Glucose Oxidase Activity Colorimetric/Fluorometric Assay Kit (K788) Glucose and Sucrose Colorimetric/Fluorometric Assay Kit (K616) Maltose and Glucose Colorimetric/Fluorometric Assay Kit (K618) PicoProbe™ Glucose-6-Phosphate Fluorometric Assay Kit (K687) Glucose uptake Colorimetric Assay Kit (K676) 10 kDa Spin Column (1997)

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