

**TABLE OF CONTENTS**

<b>1.0</b>	<b>Overview of GeneBLAzer® Technology</b> .....	<b>1</b>
<b>2.0</b>	<b>Materials Supplied</b> .....	<b>1</b>
<b>3.0</b>	<b>Materials Required, but Not Supplied</b> .....	<b>2</b>
3.1	Optional Equipment and Materials .....	2
<b>4.0</b>	<b>Cell Culture Conditions</b> .....	<b>3</b>
4.1	Media Required .....	3
4.2	Growth Conditions (See section 7.0 for detailed protocol) .....	3
<b>5.0</b>	<b>Assay Procedure</b> .....	<b>4</b>
5.1	Quick Reference Guide (for more detailed protocol information, see Section 5.2).....	4
5.2	Detailed Assay Protocol.....	5
5.3	Detection.....	5
<b>6.0</b>	<b>Data Analysis</b> .....	<b>6</b>
6.1	Background Subtraction .....	6
6.2	Visual Observation of Intracellular Beta-lactamase Activity Using LiveBLAzer™-FRET B/G Substrate (CCF4-AM).....	6
6.3	Representative Data.....	7
<b>7.0</b>	<b>Detailed Cell Handling Procedures</b> .....	<b>7</b>
7.1	Thawing Method.....	7
7.2	Propagation Method.....	7
7.3	Freezing Method .....	7
<b>8.0</b>	<b>References</b> .....	<b>8</b>
<b>9.0</b>	<b>Purchaser notification</b> .....	<b>9</b>

**1.0 OVERVIEW OF GENEBLAZER® TECHNOLOGY**

GeneBLAzer® Beta-lactamase Reporter Technology provides a highly accurate, sensitive, and easy to use method of monitoring cellular responses to drug candidates or other stimuli (1). The core of the GeneBLAzer® Technology is a Fluorescence Resonance Energy Transfer (FRET) substrate that generates a ratiometric reporter response with minimal experimental noise. In addition to the two-color (blue/green) readout of stimulated and unstimulated cells, this ratiometric method reduces the absolute and relative errors that can mask the underlying biological response of interest. Such errors include variations in cell number, transfection efficiency, substrate concentration, excitation path length, fluorescence detectors, and volume changes. The GeneBLAzer® Beta-lactamase Reporter Technology has been proven effective in high-throughput screening (HTS) campaigns for a range of target classes, including G-protein coupled receptors (GPCRs) (2, 3), nuclear receptors (4-6), and kinase signaling pathways (7).

**2.0 MATERIALS SUPPLIED**

<b>Cell Line Name:</b>	NFκB- <i>bla</i> ME-180
<b>Description:</b>	CellSensor™ NFκB- <i>bla</i> ME-180 cells contain a beta-lactamase reporter gene under control of the NFκB response element that is stably integrated into ME-180 cells. NFκB- <i>bla</i> ME-180 cells respond to Tumor Necrosis Factor α (TNF-α).
<b>Product Number:</b>	K1171
<b>Shipping Condition:</b>	Dry Ice
<b>Storage Condition:</b>	Liquid nitrogen. <b>Immediately upon receipt, cells must be stored in liquid nitrogen or thawed for immediate use. Cells stored at -80°C can quickly lose viability.</b>
<b>Quantity:</b>	~2,000,000 (2 x 10 <sup>6</sup> cells/ml)
<b>Application:</b>	This cell line can be used as a parental cell line to detect agonists/antagonists of the NFκB signaling pathway
<b>Growth Properties:</b>	Adherent
<b>Cell Phenotype:</b>	Epithelial
<b>Selection Marker:</b>	Blasticidin (5 µg/ml)
<b>Vector Used</b>	pLenti- <i>bsd</i> /NFκB- <i>bla</i> vector
<b>Mycoplasma Testing:</b>	Negative
<b>BioSafety Level:</b>	2

### 3.0 MATERIALS REQUIRED, BUT NOT SUPPLIED

Media/Reagents	Recommended Source	Part #
LiveBLAzer™ -FRET B/G Loading Kit LiveBLAzer™ -FRET B/G Substrate (CCF4-AM), 5 mg DMSO for Solution A Solution B Solution C	Invitrogen	K1030 Other sizes and Loading Kits are available
Solution D	Invitrogen	K1157
Recovery™ Cell Culture Freezing Medium	Invitrogen	12648-010
DMEM, with GlutaMAX™	Invitrogen	10569-010
DMSO	Fluka	41647
OptiMEM® I Reduced Serum Medium	Invitrogen	11058-021
Fetal bovine serum (FBS), dialyzed, tissue-culture grade <b>(DO NOT SUBSTITUTE!)</b>	Invitrogen	26400-044
Non-essential amino acids (NEAA)	Invitrogen	11140-050
Penicillin/Streptomycin (antibiotic)	Invitrogen	15140-122
Phosphate-buffered saline without calcium and magnesium [PBS(-)]	Invitrogen	14190-136
HEPES (1 M, pH 7.3)	Invitrogen	15630-080
Sodium Pyruvate	Invitrogen	11360-070
Tumor Necrosis Factor Alpha (TNF-α)	R&D Systems	210-TA-010
0.05% Trypsin/EDTA	Invitrogen	25300-054
Blasticidin (antibiotic)	Invitrogen	R210-01
Consumables	Recommended Source	Part #
Black-wall, clear-bottom, 384-well assay plates (with low fluorescence background)	Corning Life Sciences	3712
Compressed air	Various	—
Equipment	Recommended Source	
Fluorescence plate reader with bottom-read capabilities	Various	
Filters, if required for plate reader (see <b>Section 5.3</b> )	Chroma Technology Corp.	

#### 3.1 Optional Equipment and Materials

- Epifluorescence- or fluorescence-equipped microscope with appropriate filters
- Microplate centrifuge

## 4.0 CELL CULTURE CONDITIONS

### 4.1 Media Required

Component	Growth Medium	Assay Medium	Freezing Medium
DMEM	90%	—	—
Opti-MEM®	—	99.5%	—
Dialyzed FBS	10%	0.5%	—
NEAA	0.1 mM	0.1 mM	—
Sodium Pyruvate	—	1 mM	—
HEPES (pH 7.3)	25 mM	—	—
Penicillin (antibiotic)	100 U/ml	100 U/ml	—
Streptomycin (antibiotic)	100 µg/ml	100 µg/ml	—
Blasticidin (antibiotic)	5 µg/ml	—	—
Recovery™ Cell Culture Freezing Medium	—	—	100%

**Note:** Unless otherwise stated, have all media and solutions at least at room temperature (we recommend 37°C for optimal performance) before adding them to the cells.

### 4.2 Growth Conditions (See section 7.0 for detailed protocol)

1. Thaw cells in Growth Medium without Blasticidin and culture them in Growth Medium with Blasticidin. Pass or feed cells at least twice a week and maintain them in a humidified 37°C/5% CO<sub>2</sub> incubator. Maintain cells between 10% and 70% confluency. Do not allow cells to reach confluence.

**Note:** We recommend passing cells for three passages after thawing before using them in the beta-lactamase assay.

2. Freeze cells at 2 x 10<sup>6</sup> cells/ml in Freezing Medium.
3. For detailed growth and maintenance directions, see **Section 7.0**.

## 5.0 ASSAY PROCEDURE

The following instructions outline the recommended procedure for monitoring the TNF- $\alpha$ -induced NF $\kappa$ B signaling pathway using LiveBLAzer™-FRET B/G Substrate as the readout. If you use alternative substrates (*e.g.*, ToxBLAzer™ DualScreen, or LyticBLAzer™ Loading kits), follow the loading protocol provided with the product.

### 5.1 Quick Reference Guide (for more detailed protocol information, see Section 5.2)

Plate layouts and experimental outlines will vary; in screening mode, we recommend using at least three wells for each condition: Unstimulated, Stimulated, and Cell-free.

**Note:** Some solvents may affect assay performance. Assess the effect of a test compound solvent before screening. The cell stimulation described below is carried out in the presence of 0.5% DMSO to simulate the effect that a test compound solvent might have on the assay. If you use other solvents and/or solvent concentrations, change the following assay conditions and optimize appropriately.

	Unstimulated Wells	Stimulated Wells	Cell-Free Wells
Step 1 <b>Plate Cells</b>	32 $\mu$ l cells suspended in Assay Medium (12,000 cells/well)	32 $\mu$ l cells suspended in Assay Medium (12,000 cells/well)	32 $\mu$ l Assay Medium (no cells)
Step 2 <b>Incubate Cells</b>	Incubate the plate at in the humidified 37°C/5% CO <sub>2</sub> incubator for 16-20 hours		
Step 3 <b>Add DMSO</b>	Prepare a stock of Assay Medium with 5% DMSO. Add 4 $\mu$ l per well		
Step 4 <b>Add Agonist</b>	4 $\mu$ l Assay Medium	4 $\mu$ l 10X TNF- $\alpha$ in Assay Medium	4 $\mu$ l Assay Medium
Step 5 <b>Incubate Cells</b>	Incubate the plate at in the humidified 37°C/5% CO <sub>2</sub> incubator for 5 hours		
Step 6 <b>Prepare 6X Substrate Mixture</b>	6 $\mu$ l 1 mM LiveBLAzer™-FRET B/G Substrate (CCF4-AM) + 60 $\mu$ l Solution B and mix. Add 874 $\mu$ l Solution C and 60 $\mu$ l Solution D and mix		
Step 7 <b>Load Substrate Mixture</b>	8 $\mu$ l per well		
Step 8 <b>Incubate Substrate + cells</b>	2.5 hours at room temperature in the dark		
Step 9 <b>Detect Activity</b>	See Section 5.3		
Step 10 <b>Analyze data</b>	See Section 6.0		

## 5.2 Detailed Assay Protocol

### 5.2.1 Precautions

1. Work on a dust-free, clean surface. Always handle the 384-well, black-wall, clear-bottom assay plate by the sides; do not touch the clear bottom of the assay plate.
2. If pipetting manually, you may need to centrifuge the plate briefly at room temperature (30 seconds at 14 x g) after additions to ensure all assay components are on the bottom of the wells.

### 5.2.2 Plate Cells

1. Harvest cells from culture in Growth Medium and resuspend in Assay Medium at a density of  $3.75 \times 10^5$  cells/ml.
2. Add 32  $\mu$ l per well of Assay Medium to the Cell-Free control wells. Add 32  $\mu$ l per well of the cell suspension to the Unstimulated and Stimulated wells.
3. After plating, incubate the plates in a humidified 37°C/5% CO<sub>2</sub> incubator for 16-20 hours.

### 5.2.3 Prepare Stock Solutions

1. Prepare Assay Medium with 5% DMSO.
2. Prepare a 10X stock of TNF- $\alpha$  (at EC<sub>80</sub>) in Assay Medium. We recommend running a dose response curve to determine the EC<sub>80</sub> for your stimulation solution.

### 5.2.4 Stimulate Cells

1. Add 4  $\mu$ l Assay Medium with 5% DMSO to all wells.
2. Add 4  $\mu$ l 10X TNF- $\alpha$  to Stimulated wells and 4  $\mu$ l of Assay Medium to Unstimulated and Cell-Free wells.
3. Incubate the assay plate in a humidified 37°C/5% CO<sub>2</sub> incubator for 5 hours.

### 5.2.5 Substrate Loading and Incubation

This protocol is designed for loading cells with LiveBLAzer™-FRET B/G Substrate (CCF4-AM) or CCF2-AM. If alternative substrates are used please follow the loading protocol provided with the substrate.

Preparation of 6X LiveBLAzer™-FRET B/G Substrate (CCF4-AM) or CCF2-AM Mixture and cell loading should be done in the absence of direct strong lighting. Turn off the light in the hood. Handle the plate gently and do not touch the bottom.

1. Prepare Solution A: 1 mM LiveBLAzer™-FRET B/G Substrate (CCF4-AM, MW = 1096) stock solution in dry DMSO. Store the aliquots of the stock solution at -20°C until use.
2. Prepare 6X LiveBLAzer™-FRET B/G (CCF4-AM) Substrate Mixture:  
Add 6  $\mu$ l of Solution A to 60  $\mu$ l of Solution B and vortex.  
Add 874  $\mu$ l Solution C and 60  $\mu$ l of Solution D to the combined solutions from above step and vortex.
3. Remove assay plate from the humidified 37°C/5% CO<sub>2</sub> incubator.
4. Add 8  $\mu$ l of 6X Substrate Mixture from Step 2 to each well.
5. Cover the plate to protect it from light and evaporation.
6. Incubate at room temperature for 2.5 hours.

## 5.3 Detection

All measurements are made at room temperature from the bottom of the wells, preferably in 384-well, black-wall, clear-bottom assay plates with low fluorescence background. Before reading the plate, remove dust from the bottom with compressed air.

**Note:** If you do not have access to a fluorescence plate reader with bottom-read capabilities, contact our Technical Support for options of other beta-lactamase substrates that can be read with top-reading instruments.

### 5.3.1 Instrumentation, Filters, and Plates

Fluorescence plate reader with bottom reading capabilities

Recommended filters for plate reader:

Excitation filter: 409/20 nm

Emission filter: 460/40 nm

Emission filter: 530/30 nm

**5.3.2 Reading an Assay Plate**

1. Set the fluorescence plate reader to bottom-read mode.
2. Allow the lamp in the fluorescence plate reader to warm up for at least 10 minutes before making measurements.
3. Use the following filter selections:

	Scan 1	Scan 2
<b>Purpose:</b>	Measure fluorescence in the blue channel	Measure fluorescence in the green channel
<b>Excitation filter:</b>	409/20 nm	409/20 nm
<b>Emission filter:</b>	460/40 nm	530/30 nm

**6.0 DATA ANALYSIS****6.1 Background Subtraction**

We recommend that you subtract the background for both emission channels (460 nm and 530 nm).

1. Use the assay plate layout to identify the location of the Cell-Free wells. These control wells are used for background subtraction.
2. Determine the average emission from the Cell-Free wells at both 460 nm (Average Blue Background) and 530 nm (Average Green Background).
3. Subtract the Average Blue Background (data collected at 460 nm) from all of the blue emission data.
4. Subtract the Average Green background (data collected at 530 nm) from all of the green emission data.
5. Calculate the Blue/Green Emission Ratio for each well, by dividing the background-subtracted blue emission values by the background-subtracted green emission values.

**6.2 Visual Observation of Intracellular Beta-lactamase Activity Using LiveBLAzer™ -FRET B/G Substrate (CCF4-AM)**

**Note:** Microscopic visualization of cells will cause photobleaching. Always read the assay plate in the fluorescence plate reader before performing microscopic visualization.

An inverted microscope equipped for epifluorescence and either a xenon or mercury excitation lamp is typically required to view the LiveBLAzer™-FRET B/G Substrate (CCF4-AM) signal in cells. To visually inspect the cells, you will need a long-pass filter passing blue and green fluorescence light so that your eye can visually identify whether the cells are fluorescing green or blue.

Recommended filter sets for observing beta-lactamase activity are described below and are available from Chroma Technologies (800-824-7662, [www.chroma.com](http://www.chroma.com)).

**Chroma Set # 41031**

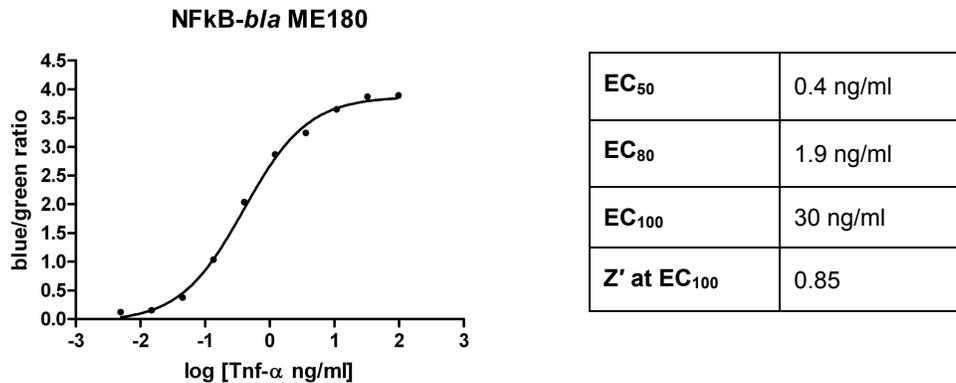
Excitation filter: HQ405/20x (405 ± 10 nm)

Dichroic mirror: 425 DCXR

Emission filter: HQ435LP (435 long-pass)

Filter sizes vary for specific microscopes and need to be specified when the filters are ordered. For epifluorescence microscopes, a long-pass dichroic mirror is needed to separate excitation and emission light and should be matched to the excitation filter (to maximally block the excitation light around 405 nm, yet allow good transmission of the emitted light).

### 6.3 Representative Data



**Figure 1. Dose response of NFκB-*bla* ME-180 cells to TNF-α.** NFκB-*bla* ME-180 cells were treated with agonist TNF-α over the indicated concentration range in a 384-well format. Cells were incubated for 5 hours with agonist and 0.5% DMSO and then loaded with LiveBLAzer™-FRET B/G Substrate (CCF4-AM) for 2.5 hours. Fluorescence emission values at 460 nm and 530 nm were obtained using a standard fluorescence plate reader and the 460/530 ratios were plotted against the concentration of the agonist.

## 7.0 DETAILED CELL HANDLING PROCEDURES

### 7.1 Thawing Method

1. Place 14 ml of Growth Medium without Blasticidin into a T75 flask.
2. Place the flask in a humidified 37°C/5% CO<sub>2</sub> incubator for 15 minutes to allow medium to equilibrate to the proper pH and temperature.
3. Remove the vial of cells to be thawed from liquid nitrogen and thaw rapidly by placing at 37°C in a water bath with gentle agitation for 1–2 minutes. Do not submerge vial in water.
4. Decontaminate the vial by wiping with 70% ethanol before opening in a Class II biological safety cabinet.
5. Transfer the vial contents drop-wise into 10 ml of Growth Medium without Blasticidin in a sterile 15-ml conical tube.
6. Centrifuge cells at 200 × g for 5 minutes.
7. Aspirate supernatant and resuspend the cell pellet in 1 ml of fresh Growth Medium without Blasticidin.
8. Transfer contents to the T75 tissue culture flask containing pre-equilibrated Growth Medium without Blasticidin and place flask in a humidified 37°C/5% CO<sub>2</sub> incubator.
9. At first passage, switch to Growth Medium with Blasticidin.

### 7.2 Propagation Method

1. Pass or feed cells at least twice a week. Maintain cells between 10% and 70% confluence. Do not allow cells to reach confluence.
2. To passage cells, aspirate medium, rinse once in PBS, add Trypsin/EDTA (3 ml for a T75 flask, 5 ml for a T175 flask, and 8 ml for T225 flask) and swirl to coat the cells evenly. Cells usually detach after ~2-5 minutes exposure to Trypsin/EDTA. Add an equal volume of Growth Medium to inactivate Trypsin.
3. Verify under a microscope that cells have detached and clumps have completely dispersed.
4. Centrifuge cells at 200 × g for 5 minutes and resuspend in Growth Medium in a new flask.

### 7.3 Freezing Method

1. Harvest the cells as described in Section 7.2. Count the cells, then spin cells down and resuspend in 4°C Freezing Medium at 2 × 10<sup>6</sup> cells/ml.
2. Dispense 1.0 ml aliquots into cryogenic vials.
3. Place in an insulated container for slow cooling and store overnight at –80°C.
4. Transfer to liquid nitrogen the next day for storage.

## 8.0 REFERENCES

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## 9.0 PURCHASER NOTIFICATION

### Limited Use Label License No. 28: CMV Promoter

The use of the CMV promoter is covered under U.S. Patent Nos. 5,168,062 and 5,385,839 owned and licensed by the University of Iowa Research Foundation and is sold for research use only. Commercial users must obtain a license to these patents directly from the University of Iowa Research Foundation (UIRF), 214 Technology Innovation Center, Iowa City, Iowa 52242. For further information, please contact the Associate Director of UIRF, at 319-335-4546.

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### Use of Genetically Modified Organisms (GMO)

**Information for European Customers** The NFκB-*bla* ME-180 cell line is genetically modified with the plasmid pLenti-*bsd*/NFκB Vector. As a condition of sale, use of this product must be in accordance with all applicable local legislation and guidelines including EC Directive 90/219/EEC on the contained use of genetically modified organisms.

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