



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Lieferung & Zahlungsart

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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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**Description**

Recombinant Raji cells constitutively expressing firefly (*Photinus pyralis*) luciferase under the control of a CMV promoter.

**Background**

The Raji line was established from a Burkitt's lymphoma patient. Raji cells constitutively express B cell antigens CD19, CD20, and CD22, and offer a physiologically relevant platform to evaluate cancer-directed immunotherapies such as Chimeric Antigen Receptor (CAR) T-cells. The Firefly Luciferase – Raji Recombinant Cell Line makes an excellent target for CAR-T or NK cells targeting CD19, CD20, or CD22. The signal generated by the firefly luciferase reporter is proportional to Raji cell numbers and facilitates the quantification of Raji cells killing upon co-culture with CAR-T or NK cells.

**Application**

1. Use as an internal control in CAR-T or NK co-culture killing assays
2. *In vitro* and *in vivo* Bioluminescence Imaging

**Materials Provided**

Components	Format
2 vials of frozen cells	Each vial contains $2 \times 10^6$ cells in 1 ml of 10% DMSO

**Parental Cell Line**

Raji, a human B lymphoblastoid cell line derived from a patient with Burkitt's lymphoma, suspension

**Mycoplasma Testing**

The cell line has been screened to confirm the absence of Mycoplasma species.

**Materials Required but Not Supplied**

These materials are not supplied with the cell line but are necessary for cell culture and cellular assays. BPS Bioscience's reagents are validated and optimized for use with this cell line and are highly recommended for best results. Media components are provided in the Media Formulations section below.

*Materials Required but Not Supplied*

Name	Ordering Information
Thaw Medium 2	<a href="#">BPS Bioscience #60184</a>
Growth Medium 2D	<a href="#">BPS Bioscience #79639</a>
96-well Tissue Culture-treated White Clear-bottom Assay Plate	Corning #3610
ONE-Step™ Luciferase Assay System	<a href="#">BPS Bioscience #60690</a>
Luminometer	

## Storage Conditions



Cells are shipped in dry ice and should immediately be thawed or stored in liquid nitrogen upon receipt. Do not use a  $-80^{\circ}\text{C}$  freezer for long term storage. Contact technical support at [support@bpsbioscience.com](mailto:support@bpsbioscience.com) if the cells are not frozen in dry ice upon arrival.

## Media Formulations

For best results, it is *highly recommended* to use these validated and optimized media from BPS Bioscience. Other preparations or formulations of media may result in suboptimal performance.



Note: Thaw Media do *not* contain selective antibiotics. However, Growth Media *do* contain selective antibiotics, which are used for maintaining the presence of the transfected gene(s) over passages. Cells should be grown at  $37^{\circ}\text{C}$  with 5%  $\text{CO}_2$ . BPS Bioscience's cell lines are stable for at least 15 passages when grown under proper conditions.

### Media Required for Cell Culture

*Thaw Medium 2 (BPS Bioscience #60184):*

RPMI1640 medium supplemented with 10% FBS, 1% Penicillin/Streptomycin.

*Growth Medium 2D (BPS Bioscience #79639):*

RPMI1640 medium supplemented with 10% FBS, 1% Penicillin/Streptomycin plus 200  $\mu\text{g/ml}$  of Hygromycin B.

## Cell Culture Protocol

### Cell Thawing

1. Swirl the vial of frozen cells for approximately 60 seconds in a  $37^{\circ}\text{C}$  water bath. As soon as the cells are thawed (it may be slightly faster or slower than 60 seconds), quickly transfer the entire contents of the vial to a tube containing 10 ml of pre-warmed Thaw Medium 2 (**no Hygromycin B**).  
**Leaving the cells in the water bath at  $37^{\circ}\text{C}$  for too long will result in rapid loss of viability.**
2. Immediately spin down the cells at  $300 \times g$  for 5 minutes, remove the medium and resuspend the cells in 5 ml of pre-warmed Thaw Medium 2 (**no Hygromycin B**).
3. Transfer the resuspended cells to a T25 flask and incubate at  $37^{\circ}\text{C}$  in a 5%  $\text{CO}_2$  incubator.
4. Cells should be passaged before they reach the density of  $2 \times 10^6$  cells/ml. At first passage and subsequent passages, use Growth Medium 2D (**contains Hygromycin B**).

### Cell Passage

Dilute the cell suspension into new culture vessels at no less than  $0.2 \times 10^6$  cells/ml of Growth Medium 2D (**contains Hygromycin B**). The sub-cultivation ratio is approximately 1:5 to 1:10 once or twice a week, so cells are maintained between  $0.2 \times 10^6$  cells/ml and  $2 \times 10^6$  cells/ml.

### Cell Freezing

1. Spin down the cells at  $300 \times g$  for 5 minutes, remove the medium and resuspend the cell pellet in  $4^{\circ}\text{C}$  Freezing Medium (BPS Bioscience #79796, or 10% DMSO + 90% FBS) at a density of  $\sim 2 \times 10^6$  cells/ml.
2. Dispense 1 ml of cell aliquots into cryogenic vials. Place the vials in an insulated container for slow cooling and store at  $-80^{\circ}\text{C}$  overnight.
3. Transfer the vials to liquid nitrogen the next day for storage.



Note: It is recommended to expand the cells and freeze at least 10 vials at an early passage for future use.

**Validation Data**

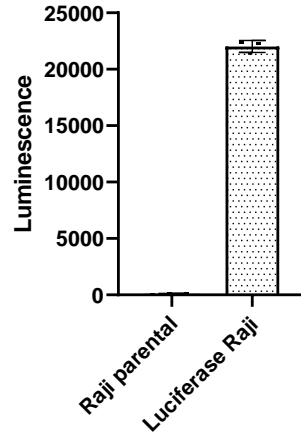


Figure 1. Luciferase activity in Firefly Luciferase Raji recombinant cells. Firefly Luciferase Raji recombinant cells were seeded into a 96-well plate at 5000 cells/well in 50 µl Thaw Medium 2, and the luciferase activity was measured using the ONE-Step luciferase assay system (BPS Bioscience, #60690).

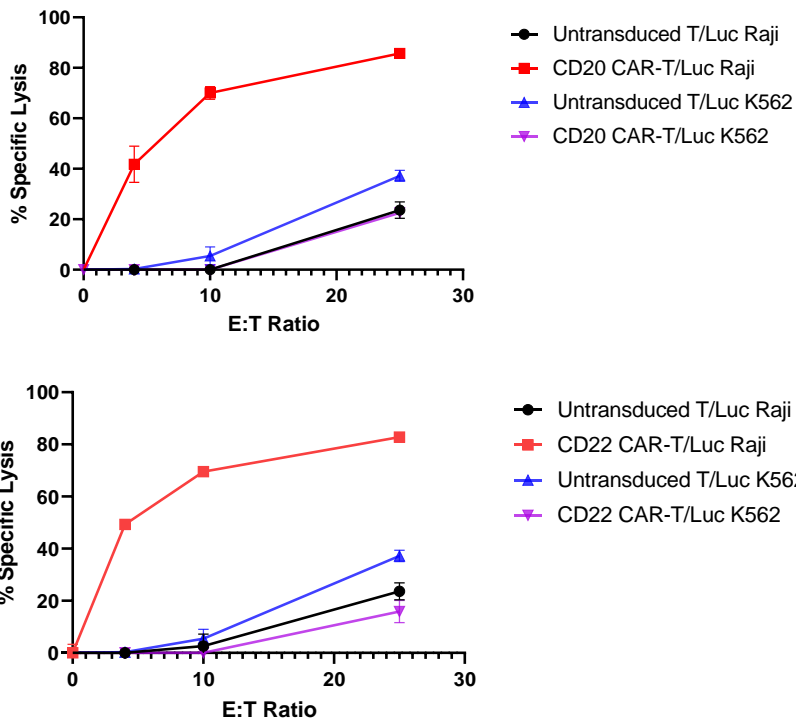


Figure 2. Anti-CD20 or anti-CD22 CAR-T cells cytotoxicity against Firefly Luciferase Raji recombinant cell line in co-culture assays. Firefly Luciferase Raji recombinant cells were seeded into a 96-well plate at 5000 cells/well in 50 µl of Thaw Medium 2, and were cocultured overnight with 50 µl of anti-CD20 CAR-T cells (prepared using anti-

CD20 CAR lentivirus, BPS Bioscience #78606) or anti-CD22 CAR-T cells (prepared using anti-CD22 CAR lentivirus, BPS Bioscience #78608) at various effector-to-target ratios. The luciferase activity was measured using the ONE-Step luciferase assay system (BPS Bioscience #60690). Target only wells determined the maximum signal (Lmax). The blank was determined from medium only wells and was subtracted from all values. Percent specific lysis was calculated as:  $1-(L)/(L_{max})$ . Firefly Luciferase K562 cells (BPS Bioscience #78621) which do not express endogenous CD20 or CD22, were used as a negative control.

**License Disclosure**

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**Troubleshooting Guide**

Visit [bpsbioscience.com/cell-line-faq](https://bpsbioscience.com/cell-line-faq) for detailed troubleshooting instructions. For all further questions, please email [support@bpsbioscience.com](mailto:support@bpsbioscience.com).

**Related Products**

<i>Products</i>	<i>Catalog #</i>	<i>Size</i>
Firefly Luciferase K562 Cell Line	78621	2 vials
Firefly Luciferase - CHO Recombinant Cell Line	79725	2 vials
RPMI 8226 Recombinant Cell Line	79834	2 vials
CD19 CAR Lentivirus	78600	50 µl
CD20 CAR Lentivirus	78606	50 µl
CD22 CAR Lentivirus	78608	50 µl
BCMA CAR Lentivirus	78603	50 µl
CD19/ Firefly Luciferase - CHO Recombinant Cell Line	79714	2 vials
CD22/ Firefly Luciferase - CHO Recombinant Cell Line	79715	2 vials
CD19/CD20/ Firefly Luciferase - CHO Recombinant Cell Line	78186	2 vials
BCMA/CD20/ Firefly Luciferase - CHO Recombinant Cell Line	78185	2 vials