# Fenics **BIO**

#### Product data sheet

Raji/GFP stable cell line Catalog Number: CL-1022 Storage: Liquid nitrogen Components: 1 vial contains ~2 x10<sup>6</sup> cells in Cell freezing medium

### **Product description**

Raji/GFP cells are derived from the human Raji B lymphoblastoid cell line by stably integration of a constitutive GFP expression construct. Raji cell line was generated from human Brukitt's lymphoma, has been widely used in cancer research and drug development. Raji/GFP cells stably express GFP, can be used for in vitro assays and in vivo imaging.

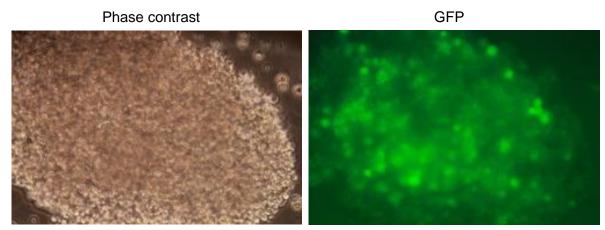


Figure 1. GFP expression in Raji/GFP stable cell line

### **Cell line description**

Organism: Homo sapiens (human) Tissue: Peripheral blood Cell Type: B lymphoblast Morphology: Lymphoblast Culture Properties: Suspension Disease: Burkitt's lymphoma Biosafety Level: 2

#### Medium

- Complete culture medium: RPMI-1640, 10% fetal bovine serum (FBS) 1 μg/mL of puromycin may be added to the culture medium. Puromycin should not be added until a culture has been well established from the thawed cells.
- 2. Freeze medium: Fetal bovine serum (FBS), 6% DMSO

## **Culture procedure**

#### Thawing of frozen cells

- 1. Thaw the frozen cryovial by gentle agitation in a 37 °C water bath in 1-2 minutes.
- 2. Remove the cryovial from the water bath as soon as the contents are thawed, and decontaminate by wiping with 70% ethanol.
- 3. Transfer the thawed cell suspension to a centrifuge tube containing 10 ml of Complete culture medium, centrifuge at 500 g for 5 minutes.
- 4. Remove the medium by aspiration, resuspend the cells with 2 ml of the Complete culture medium by gently pipetting up and down.
- 5. Transfer the cells to a T-25 suspension cell culture flask.
- 6. Place the cells in a 37°C incubator with 5% CO2.

#### Sub-culturing

Cultures can be maintained by the addition of fresh medium or replacement of medium. Alternatively, cultures can be established by centrifugation with subsequent resuspension at 3 to  $5 \times 10^5$  viable cells/ml. Maintain cell density between  $3 \times 10^5$  and 2 to  $3 \times 10^6$  viable cells/ml. Do not allow the cell density to exceed  $3 \times 10^6$  cells/ml.

Renew or add fresh medium every 2-3 days.

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