

Technical Data Sheet: EL4-Luc2

ATCC [®] Number	TIB-39-LUC2™
Organism	Mus musculus
Tissue/Disease Source	Lymphoma
Product Description	 This luciferase expressing cell line was derived from EL4 cell line by transduction with a lentiviral vector encoding firefly luciferase gene (luc2) and subsequently through single cell cloning. Signal noise ratio: ≥ 1,000 Bioluminescence: ≥ 20,000 photons/cell/sec (subject to imaging and culture conditions) Confirmed to be murine pathogen-free
Application	Excellent signal/background ratio and stable luciferase expression make this cell line ideal for <i>in vivo</i> bioluminescence imaging of xenograft animal model to study human cancer and monitor activity of anti-cancer drug. It also can be used in cell-based assays for cancer research.

In vivo Bioluminescent Imaging

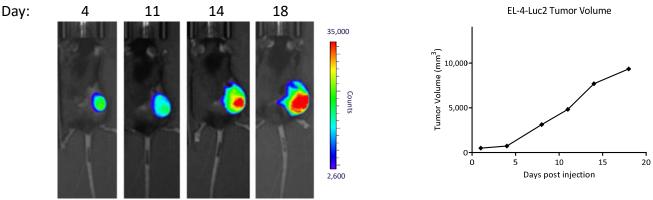
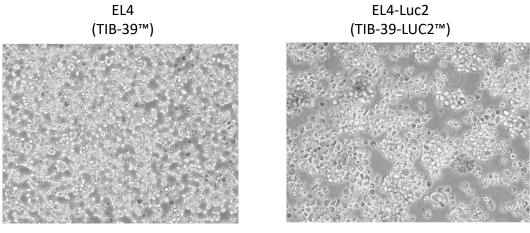


Figure 1. *In vivo* detection of luciferase activity of EL4-Luc2. EL4-Luc2 cells (3 x 10⁶) were injected subcutaneously into the dorsal region near the thigh of female C57BL/6N mice. Tumor growth was monitored weekly using a Xenogen IVIS Spectrum. *In vivo* bioluminescence imaging demonstrated the progression of tumors.

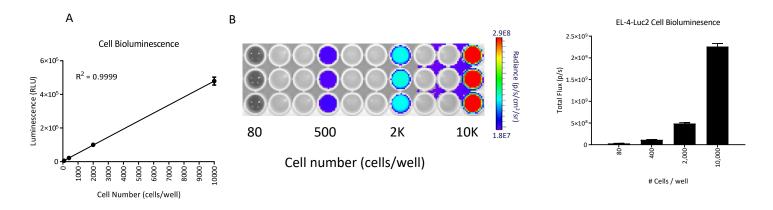
Cell Morphology



Doubling time = 12.6 hours

Doubling time = 12.3 hours

Figure 2. Cell morphology of EL4 parental and EL4-Luc2. Cells were maintained in ATCC recommended culture conditions. Cell morphology was observed under microscopy and images were captured by digital camera.



Luciferase Expression

Figure 3. Linearity of luminescence and of *in vitro* quantification of luciferase activity of EL4-Luc2. Cells were seeded in a 96-well plate at indicated cell numbers per well, and Bright-Glo (Promega) was added to the indicated wells. The luminescence of the plate was read within 10 minutes using a luminescence plate reader (A) and determined to have a linear correlation of bioluminescence intensity with cell numbers. (B) The plate was imaged using a Xenogen IVIS Spectrum to quantify that photons emitted per cell.

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