

Technical Data Sheet:

U-87 MG-Luc2

ATCC® Number	HTB-14-LUC2™
Organism	<i>Homo sapiens</i>
Tissue/Disease Source	Glioma
Product Description	U-87 MG-Luc2
Application	Excellent signal/background ratio and stable Luciferase expression make this cell line ideal for in vivo 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.

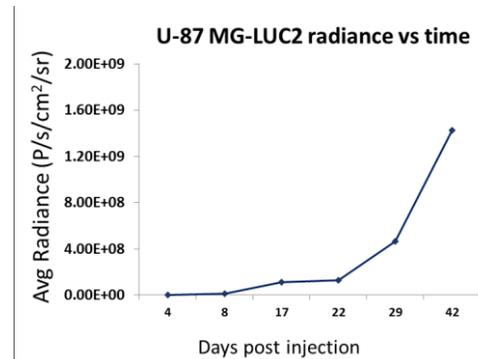
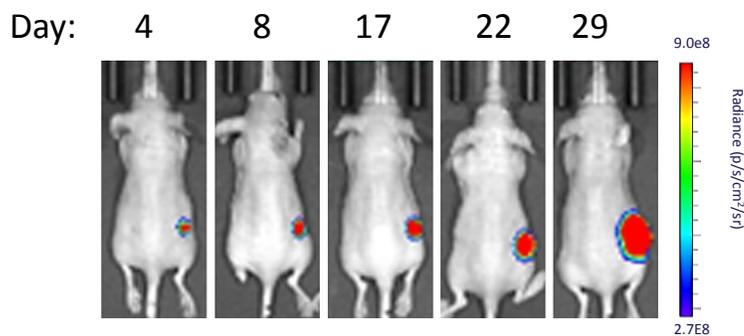
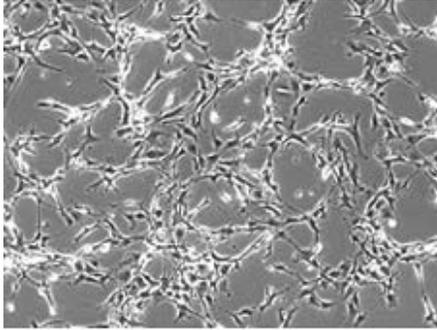


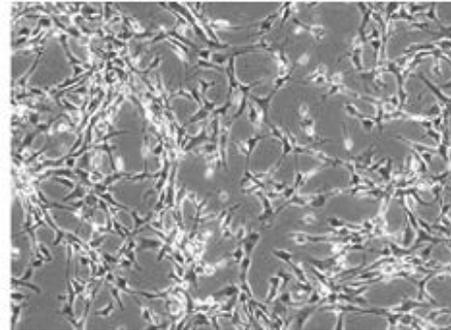
Figure 1: *In vivo* detection of luciferase activity of U-87 MG-Luc2. U-87 MG-Luc2 cells (1×10^6) were injected subcutaneously into the dorsal region near the thigh of female nude mice. Tumor growth was monitored weekly using a Xenogen IVIS Spectrum. *In vivo* bioluminescence imaging demonstrated the progression of tumors.

U-87 MG (HTB-14™)



Doubling Time = 29.0 hours

U-87 MG-Luc2 (HTB-14-LUC2™)



Doubling Time = 23.0 hours

Figure 2: Cell Morphology of U-87 MG parental and U-87 MG-Luc2. Cells were maintained in ATCC recommended culture conditions. Cell morphology was observed under Nikon™ microscopy and images were captured by Zeiss® digital camera.

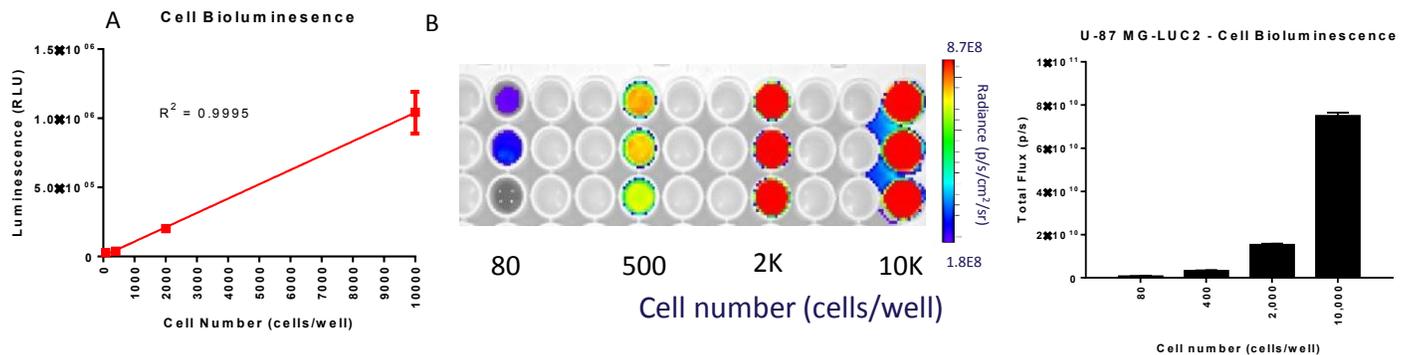


Figure 3: Linearity of luminescence and of *in vitro* quantification of luciferase activity of KRAS Mutant-A375 Isogenic-Luc2. Cells were seeded in a 96-well plate at indicated cell numbers per well, and Bright-Glo (Promega) was added to the indicated wells. The luminescence of the plate was read within 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.