

# Technical Data Sheet: NCI-H1650-GAS-Luc2

ATCC <sup>®</sup> Number	CRL-5883-GAS-LUC2™
Organism	Homo sapiens
Tissue/Disease Source	Lung/ Adenocarcinoma; Bronchoalveolar carcinoma; Stage 3B
Product Description	NCI-H1650 cell line (ATCC <sup>®</sup> CRL-5883 <sup>™</sup> ) is commonly used for immuno-oncology and lung cancer research. This luciferase reporter cell line was derived from parental line CRL-5883 by stably expressing firefly luciferase gene (luc2) under control of a gamma-activated site (GAS) promoter through lentiviral transduction and single cell cloning. The cells, upon stimulation with interferon gamma (IFN-γ), express high levels of enzymatically active luciferase protein, which can be detected via in vitro bioluminescence assays. This reporter cell line is useful for monitoring the activity of IFN-γ-induced GAS signal transduction pathway <del>s</del> .
Application	Enabling sensitive and quantitative assessment of signal transduction makes this reporter cell line ideal for in vitro bioluminescence assays to study immune response in cell lines overexpressing B7-H3 (CD276), development of new drugs, and safety evaluation of new chemicals and drugs.

#### In vitro activation of luciferase expression by IFN- $\gamma$ and T Cell-conditioned media



## Figure 1. In vitro activation of luciferase expression by IFN- $\gamma$ and T Cell-conditioned media. Luciferase expression from NCI-H1650-GAS-Luc2 cells upon signaling activation by (A) IFN- $\gamma$ stimulation (0.01 – 1,000 ng/ mL), (B) conditioned-media stimulation from checkpoint matched non-

activated and activated primary CD8+ T cells. N=3 in all experiments. \*, P < 0.05.

## NCI-H1650 GAS-Luc2 : NK = 2:1





Figure 2. In vitro activation of bioluminescence in co-culture assay. NCI-H1650-GAS-Luc2 cells were co-cultured with 2:1 (E:T) ratio of primary CD56+ NK cells for (A) 24 and (B) 48 hours in the presence of a B7-H3 antibody or an isotype control. Different concentrations of B7-H3 mAb were added to block the B7-H3 checkpoint ligand. N=3 in all experiments.

### Cell Morphology



Doubling time = 56 hours

NCI-H1650-GAS-Luc2 (CRL-5883-GAS-LUC2™)



Doubling time = 48 hours

Figure 3: Cell morphology of NCI-H1650 parental and NCI-H1650-GAS-Luc2. Cells were maintained in ATCC recommended culture conditions. Cell morphology was observed under microscopy and images were captured by digital camera.

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