

# Resource Summary Report

Generated by [FDI Lab - SciCrunch.org](#) on May 8, 2024

## GL261

RRID:CVCL\_Y003

Type: Cell Line

### Proper Citation

(DSMZ Cat# ACC-802, RRID:CVCL\_Y003)

### Cell Line Information

**URL:** [https://web.expasy.org/cellosaurus/CVCL\\_Y003](https://web.expasy.org/cellosaurus/CVCL_Y003)

**Proper Citation:** (DSMZ Cat# ACC-802, RRID:CVCL\_Y003)

**Description:** Cell line GL261 is a Cancer cell line with a species of origin *Mus musculus* (Mouse)

**Disease:** Mouse glioblastoma

**Defining Citation:** [PMID:16734735](#), [PMID:23248259](#), [PMID:23686484](#), [PMID:25894527](#), [PMID:30084848](#), [PMID:30524896](#)

**Comments:** Breed/subspecies: C57BL/6., Derived from sampling site: Brain., Omics: HLA class I peptidome analysis by proteomics., Omics: Deep exome analysis., Omics: Cell surface proteome., Transformant: ChEBI; CHEBI:34342; 3-methylcholanthrene (3-MC., Doubling time: ~100-120 hours (DSMZ=ACC-802).

**Category:** Cancer cell line

**Organism:** *Mus musculus* (Mouse)

**Name:** GL261

**Synonyms:** Glioma 261, GLIOMA 261, Glioma-261, GL-261

**Cross References:** BCRJ:0299, ChEMBL-Cells:CHEMBL4483160, ChEMBL-Targets:CHEMBL4483239, DSMZ:ACC-802, DSMZCellDive:ACC-802, NCI-DTP:Glioma 261, PRIDE:PXD000589, PRIDE:PXD008733, PubChem\_Cell\_line:CVCL\_Y003, Wikidata:Q27547979

**ID:** CVCL\_Y003

**Vendor:** DSMZ

**Catalog Number:** ACC-802

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## Ratings and Alerts

No rating or validation information has been found for GL261.

No alerts have been found for GL261.

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## Data and Source Information

**Source:** [Cellosaurus](#)

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## Usage and Citation Metrics

We found 454 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [FDI Lab - SciCrunch.org](#).

Kirschenbaum D, et al. (2024) Time-resolved single-cell transcriptomics defines immune trajectories in glioblastoma. *Cell*, 187(1), 149.

Schweiger MW, et al. (2024) Glioblastoma extracellular vesicles modulate immune PD-L1 expression in accessory macrophages upon radiotherapy. *iScience*, 27(2), 108807.

Rashidi A, et al. (2024) Myeloid cell-derived creatine in the hypoxic niche promotes glioblastoma growth. *Cell metabolism*, 36(1), 62.

Bai L, et al. (2023) Biomimetic three-dimensional glioma model printed in vitro for the studies of glioma cells and neurons interactions. *International journal of bioprinting*, 9(4), 715.

Tu L, et al. (2023) Incorporation of a TGF-?2-inhibiting oligodeoxynucleotide molecular adjuvant into a tumor cell lysate vaccine to enhance antglioma immunity in mice. *Frontiers in immunology*, 14, 1013342.

van Hooren L, et al. (2023) CD103+ regulatory T cells underlie resistance to radioimmunotherapy and impair CD8+ T cell activation in glioblastoma. *Nature cancer*, 4(5), 665.

Hou X, et al. (2023) Gut microbiota mediated the individualized efficacy of Temozolomide via immunomodulation in glioma. *Journal of translational medicine*, 21(1), 198.

Miao Y, et al. (2023) Anti-cancer effect of targeting fibroblast activation protein alpha in glioblastoma through remodeling macrophage phenotype and suppressing tumor

progression. *CNS neuroscience & therapeutics*, 29(3), 878.

Wang T, et al. (2023) Reversing T Cell Dysfunction to Boost Glioblastoma Immunotherapy by Paroxetine-Mediated GRK2 Inhibition and Blockade of Multiple Checkpoints through Biomimetic Nanoparticles. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 10(9), e2204961.

Turco V, et al. (2023) T cell-independent eradication of experimental glioma by intravenous TLR7/8-agonist-loaded nanoparticles. *Nature communications*, 14(1), 771.

Ye S, et al. (2023) Rapid and label-free histological imaging of unprocessed surgical tissues via dark-field reflectance ultraviolet microscopy. *iScience*, 26(1), 105849.

Li Y, et al. (2023) Glioma-derived LRIG3 interacts with NETO2 in tumor-associated macrophages to modulate microenvironment and suppress tumor growth. *Cell death & disease*, 14(1), 28.

Chen H, et al. (2023) A nitric-oxide driven chemotactic nanomotor for enhanced immunotherapy of glioblastoma. *Nature communications*, 14(1), 941.

Zhao Q, et al. (2023) FGL2-targeting T cells exhibit antitumor effects on glioblastoma and recruit tumor-specific brain-resident memory T cells. *Nature communications*, 14(1), 735.

Zhou S, et al. (2023) Reprogramming systemic and local immune function to empower immunotherapy against glioblastoma. *Nature communications*, 14(1), 435.

Datta M, et al. (2023) Losartan controls immune checkpoint blocker-induced edema and improves survival in glioblastoma mouse models. *Proceedings of the National Academy of Sciences of the United States of America*, 120(6), e2219199120.

Liu Y, et al. (2023) In Situ Nitric Oxide Gas Nanogenerator Reprograms Glioma Immunosuppressive Microenvironment. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 10(18), e2300679.

Chen Q, et al. (2023) Dynamic change in Siglec-15 expression in peritumoral macrophages confers an immunosuppressive microenvironment and poor outcome in glioma. *Frontiers in immunology*, 14, 1159085.

Qi Z, et al. (2023) Live-attenuated Japanese encephalitis virus inhibits glioblastoma growth and elicits potent antitumor immunity. *Frontiers in immunology*, 14, 982180.

Liu X, et al. (2023) Interaction between PD-L1 and soluble VEGFR1 in glioblastoma-educated macrophages. *BMC cancer*, 23(1), 259.