

# Resource Summary Report

Generated by [RRID](#) on Sep 9, 2026

## PC12

RRID:CVCL\_0481

Type: Cell Line

### Proper Citation

RRID:CVCL\_0481

### Cell Line Information

**URL:** [https://web.expasy.org/cellosaurus/CVCL\\_0481](https://web.expasy.org/cellosaurus/CVCL_0481)

**Proper Citation:** RRID:CVCL\_0481

**Description:** Cell line PC12 is a Cancer cell line with a species of origin Rattus norvegicus (Rat)

**Sex:** Male

**Disease:** Rat adrenal gland pheochromocytoma

**Defining Citation:** [PMID:682602](#), [PMID:1065897](#), [PMID:2187480](#), [PMID:2493070](#), [PMID:2759161](#), [PMID:6500171](#), [PMID:8282765](#), [PMID:9886495](#), [PMID:15541571](#), [PMID:16223334](#), [PMID:18005394](#), [PMID:22538498](#), [PMID:33171774](#), [PMID:34348336](#)

**Comments:** Characteristics: Responds reversibly to NGF by induction of a neuronal phenotype., Group: Space-flown cell line (cellonaut).

**Category:** Cancer cell line

**Organism:** Rattus norvegicus (Rat)

**Name:** PC12

**Synonyms:** PC-12, PC 12, PC12.1

**Cross References:** BTO:BTO\_0001009, CLO:CLO\_0008392, CLO:CLO\_0050821, EFO:EFO\_0001225, MCCL:MCC:0000379, CLDB:cl3876, CLDB:cl3877, CLDB:cl3878, CLDB:cl4922, CLDB:cl5245, Abcam:ab279978, AddexBio:C0032001/4927, ATCC:CRL-1721, BCRC:60048, BCRJ:0203, CCRID:1101RAT-PUMC000024, CCRID:3101RATSCSP517, CCRID:3101RATTCTCR3, CCRID:4201RAT-CCTCC000006, CCRID:5301RAT-KCB07036YJ, CCTCC:GDC0006, ChEMBL-Cells:ChEMBL3307976, ChEMBL-Targets:ChEMBL612556, CLS:500311, DSMZ:ACC-159, DSMZCellDive:ACC-159, ECACC:88022401, ICLC:ATL98004, IZSLER:BS TCL 91, JCRB:IFO50278,

JCRB:JCRB0733, KCB:KCB 200735YJ, KCB:KCB 200736YJ, KCB:KCB 93033YJ, KCLB:21721, Lonza:124, MeSH:D016716, NCBI\_Iran:C153, PubChem\_Cell\_line:CVCL\_0481, RCB:RCB0009, TOKU-E:2867, TOKU-E:3554, Ubigen:YC-C047, Wikidata:Q7118481

**ID:** CVCL\_0481

**Record Creation Time:** 20220427T221414+0000

**Record Last Update:** 20260905T182030+0000

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## Ratings and Alerts

No rating or validation information has been found for PC12.

No alerts have been found for PC12.

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## Data and Source Information

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

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## Usage and Citation Metrics

We found 3886 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [RRID](#).

Lee S, et al. (2026) Synthetic serum markers enable noninvasive monitoring of gene expression in primate brains. *Neuron*, 114(10), 1751.

Mohr J, et al. (2026) Multimodal Imaging Reveals Rapid Catecholamine Uptake and Release by Neutrophils. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), e24193.

Yang T, et al. (2026) "Membrane-Guided" Repair Strategy: Precision Delivery of GGT1 Degradar for Targeted Repair and Regeneration of Spinal Cord Neurons. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 13(42), e75554.

Zamora F, et al. (2026) Docosahexaenoic Acid Modulates Autophagy and Confers Neuronal Resilience under Hypoxia-Reoxygenation Stress. *Journal of molecular neuroscience* : MN, 76(2).

Shin N, et al. (2026) Neutrophil-Like Mitochondria Enable Blood-Brain Barrier Transmigration and Neuroprotection. *ACS nano*, 20(21), 15116.

Cui C, et al. (2026) Aggregation-Enhanced Piezoelectric Nanotransducers Facilitate Transgene-Free Wireless Neurostimulation under Low-Intensity Focused Ultrasound. *ACS nano*, 20(1), 835.

Zhang C, et al. (2026) Quantification of Single-Cell Cysteine Using an Electrochemical Nanosensor for Predicting Tumor Disulfidptosis Susceptibility. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 13(13), e23478.

Li L, et al. (2026) Targeting G3BP1 Condensate Topology Promotes Stress Granule Assembly via m6A-IGF2BP1 for Ischemic Stroke Rescue. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 13(6), e14703.

Wang W, et al. (2026) Nanotherapeutic Macrophage-Neuro Reprogramming Through Immunometabolic Crosstalk Mitigates Sepsis-Induced Lung Injury and Neurologic Damage. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 13(24), e20665.

Shi X, et al. (2026) 3D-Printed, Cellulose-Derived Scaffold Promotes Neuroregeneration and Functional Recovery after Spinal Cord Injury. *ACS biomaterials science & engineering*, 12(3), 1814.

Li ZT, et al. (2026) Natural Product Rengyolone Attenuates LPS-Induced Microglia Inflammation via Suppression of the TLR4/NF- $\kappa$ B Pathway. *Chemistry & biodiversity*, 23(5), e71314.

He ZNT, et al. (2026) H2S induces apoptosis of tumors with high IDO1 expression via NR4A1-BCL-2 and SOCS3 pathways. *Cellular & molecular biology letters*, 31(1).

Ying X, et al. (2026) Exercise-derived exosomal miR-151-3p: An innovative anti-inflammatory and antioxidant therapeutic for spinal cord injury. *Bioactive materials*, 65, 535.

Cidem A, et al. (2026) BLF stimulates neuronal differentiation via activation of p35/CDK5 signaling and AMPK-mediated mitochondrial regulation. *Neurochemistry international*, 192, 106106.

Zhou D, et al. (2026) Stepwise Regulation of Cellular Oxidative Stress via Conductive-Piezoelectric Integrated Microstructured Conduits for Enhanced Nerve Regeneration. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), 13(13), e16124.

Zhai Z, et al. (2026) Rationalizing nose-to-brain drug delivery: Machine learning-guided optimization and mechanistic elucidation of olfactory deposition for nasal sprays. *Journal of controlled release : official journal of the Controlled Release Society*, 392, 114667.

Nigri J, et al. (2026) Myofibroblasts Induce Neuroplasticity to Promote Pancreatic Inflammation and Cancer Progression. *Cancer discovery*, 16(5), 1014.

Pavan B, et al. (2026) A new microphysiological platform to study the permeation of neuroactive agents across intestinal and brain barriers. *Biofabrication*, 18(2).

Wu C, et al. (2026) Engineered exosomes mediated siNox4 delivery loaded polyzwitterionic hydrogel with electrophysiologic conductivity for synergistic treatment of spinal cord injury via suppressing neuronal ferroptosis. *Biomaterials*, 331, 124096.

Han KJ, et al. (2026) Molecular Mechanistic Studies on Caffeoylquinic Acid Derivatives From *Vaccinium dunalianum* Wight as Dual-Target Inhibitors of AChE and BChE With In Vitro Inhibitory Evaluation and Molecular Docking. *Chemistry & biodiversity*, 23(5), e03013.