Resource Summary Report

Generated by RRID on Sep 4, 2026

NUP98 Polyclonal Antibody

RRID:AB_2757247
Type: Antibody

Proper Citation

ABclonal Cat# A0530, RRID:AB_2757247

Antibody Information

URL: http://antibodyregistry.org/AB_2757247

Proper Citation: ABclonal Cat# A0530, RRID:AB_2757247

Target Antigen: NUP98

Host Organism: rabbit

Clonality: polyclonal

Comments: Applications:WB

Antibody Name: NUP98 Polyclonal Antibody

Description: This polyclonal targets NUP98

Target Organism: mouse, human

Antibody ID: AB_2757247

Vendor: ABclonal

Catalog Number: A0530

Record Creation Time: 20231110T033310+0000

Record Last Update: 20260829T095434+0000

Ratings and Alerts

No rating or validation information has been found for NUP98 Polyclonal Antibody.

No alerts have been found for NUP98 Polyclonal Antibody.

Data and Source Information

Source: Antibody Registry

Usage and Citation Metrics

We found 12 mentions in open access literature.

Listed below are recent publications. The full list is available at RRID .

Shao J, et al. (2024) An achievement has been made in establishing an induced pluripotent stem cell line (SDPHi005-A) from a healthy Chinese male donor. Stem cell research, 77, 103393.

Guan J, et al. (2024) Establishment of iPS cell line (SDQLCHi061-A) from a patient with carbamoylphosphate synthetase I deficiency due to CPS1 mutation. Stem cell research, 76, 103353.

Guan J, et al. (2024) Establishment of iPS cell line (SDQLCHi080-A) from a patient with GM1 gangliosidosis due to GLB1 mutation. Stem cell research, 81, 103545.

Shao J, et al. (2021) Non-integrating episomal vectors induced pluripotent stem cell line (SDPHi001-A) from a healthy female individual. Stem cell research, 56, 102540.

Zhang H, et al. (2021) Generation and characterization of an induced pluripotent stem cell line SDQLCHi018-A from a congenital myasthenic syndrome patient carrying compound heterozygote mutations in RAPSN gene. Stem cell research, 51, 102160.

Ma Y, et al. (2020) An integration-free iPSC line (SDQLCHi017-A) derived from a patient with nemaline myopathy-2 disease carrying compound heterozygote mutations in NEB gene. Stem cell research, 43, 101729.

Zhang H, et al. (2019) Establishment of a human induced pluripotent stem cell line (SDQLCHi005-A) from a patient with mastocytosis carrying heterozygous mutation in KIT gene. Stem cell research, 40, 101565.

Ma Y, et al. (2019) Establishment of a human induced pluripotent stem cell line (SDQLCHi004-A) from a patient with nemaline myopathy-4 disease carrying heterozygous mutation in TPM2 gene. Stem cell research, 40, 101559.

Zhang H, et al. (2019) Generation of an induced pluripotent stem cell line (SDQLCHi003-a) from a patient with short rib-thoracic dysplasia syndrome type III carrying compound heterozygous mutations in DYNC2H1. Stem cell research, 39, 101505.

Zhang H, et al. (2019) An integration-free iPSC line (SDQLCHi013-A) derived from a patient with maple syrup urine disease carrying compound heterozygote mutations in BCKDHA gene. Stem cell research, 41, 101585.

Zhang H, et al. (2019) Establishment of a human iPSC line (SDQLCHi010-A) from a patient with optic nerve malformation carrying a heterozygous mutation in PAX6 gene. Stem cell research, 41, 101611.

Ma Y, et al. (2019) Generation of two induced pluripotent stem cell lines from patients with unbalanced translocation (3;22). Stem cell research, 40, 101545.