# Resource Summary Report

Generated by FDI Lab - SciCrunch.org on Apr 2, 2025

## H1.1

RRID:CVCL_C813
Type: Cell Line

### Proper Citation

(RRID:CVCL_C813)

### Cell Line Information

**URL:** https://web.expasy.org/cellosaurus/CVCL_C813

**Proper Citation:** (RRID:CVCL_C813)

**Sex:** Male

**Comments:** From: University of Wisconsin; Madison; USA.

**Category:** Embryonic stem cell

**Name:** H1.1

**Synonyms:** WA01.1, WAe001-A-1, WICELLe001-A-1

**Cross References:** hPSCreg:WAe001-A-1, SKIP:SKIP001997, Wikidata:Q54871907

**ID:** CVCL_C813

**Record Creation Time:** 20250131T200124+0000

**Record Last Update:** 20250131T201236+0000

### Ratings and Alerts

No rating or validation information has been found for H1.1.

No alerts have been found for H1.1.

### Data and Source Information

**Source:** Cellosaurus

## Usage and Citation Metrics

We found 1 mentions in open access literature.

**Listed below are recent publications.** The full list is available at FDI Lab - SciCrunch.org.

Chia PH, et al. (2018) A homozygous loss-of-function CAMK2A mutation causes growth delay, frequent seizures and severe intellectual disability. eLife, 7.