

Resource Summary Report

Generated by [SPARC SAWG](#) on Sep 17, 2026

[y\[1\] w\[*\]; Mi{Trojan-GAL4.2}Pxn\[MI01492-TG4.2\]/TM3, Sb\[1\] Ser\[1\]](#)

RRID:BDSC_66850

Type: Organism

Proper Citation

RRID:BDSC_66850

Organism Information

URL: <https://n2t.net/bdsc:66850>

Proper Citation: RRID:BDSC_66850

Description: Drosophila melanogaster with name y[1] w[*]; Mi{Trojan-GAL4.2}Pxn[MI01492-TG4.2]/TM3, Sb[1] Ser[1] from BDSC.

Species: Drosophila melanogaster

Notes: Donor: Gene Disruption Project; Donor's Source: Hugo J. Bellen, Baylor College of Medicine

Affected Gene: GAL4, Pxn, Sb, Ser, w, y

Genomic Alteration: Chromosome 1, Chromosome 3

Catalog Number: 66850

Database: Bloomington Drosophila Stock Center (BDSC)

Database Abbreviation: BDSC

Availability: available

Alternate IDs: BDSC_66850, BDSC:66850, BL66850

Organism Name: y[1] w[*]; Mi{Trojan-GAL4.2}Pxn[MI01492-TG4.2]/TM3, Sb[1] Ser[1]

Record Creation Time: 20240911T223023+0000

Record Last Update: 20260912T204236+0000

Ratings and Alerts

No rating or validation information has been found for y[1] w[*]; Mi{Trojan-GAL4.2}Pxn[MI01492-TG4.2]/TM3, Sb[1] Ser[1].

No alerts have been found for y[1] w[*]; Mi{Trojan-GAL4.2}Pxn[MI01492-TG4.2]/TM3, Sb[1] Ser[1].

Data and Source Information

Source: [Integrated Animals](#)

Source Database: Bloomington Drosophila Stock Center (BDSC)

Usage and Citation Metrics

We found 4 mentions in open access literature.

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

McLaughlin CN, et al. (2026) Endocytome profiling uncovers cell-surface protein dynamics underlying neuronal connectivity. *Neuron*, 114(11), 1935.

McLaughlin CN, et al. (2025) Endocytome profiling uncovers cell-surface protein dynamics underlying neuronal connectivity. *bioRxiv* : the preprint server for biology.

Gupta HP, et al. (2025) Decoding neuronal wiring by joint inference of cell identity and synaptic connectivity. *bioRxiv* : the preprint server for biology.

Marcogliese PC, et al. (2022) Drosophila functional screening of de novo variants in autism uncovers damaging variants and facilitates discovery of rare neurodevelopmental diseases. *Cell reports*, 38(11), 110517.