Resource Summary Report

Generated by FDI Lab - SciCrunch.org on Apr 28, 2025

Anti-digoxigenin

RRID:AB_514496 Type: Antibody

Proper Citation

(Sigma-Aldrich Cat# 11333089001, RRID:AB_514496)

Antibody Information

URL: http://antibodyregistry.org/AB_514496

Proper Citation: (Sigma-Aldrich Cat# 11333089001, RRID:AB_514496)

Target Antigen: digoxinenin

Host Organism: sheep

Clonality: polyclonal

Comments: This is a Roche product, offered by Sigma.

Antibody Name: Anti-digoxigenin

Description: This polyclonal targets digoxinenin

Antibody ID: AB_514496

Vendor: Sigma-Aldrich

Catalog Number: 11333089001

Record Creation Time: 20231110T034746+0000

Record Last Update: 20240725T011342+0000

Ratings and Alerts

No rating or validation information has been found for Anti-digoxigenin.

No alerts have been found for Anti-digoxigenin.

Data and Source Information

Source: Antibody Registry

Usage and Citation Metrics

We found 29 mentions in open access literature.

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

Ku J, et al. (2024) Alternative polyadenylation determines the functional landscape of inverted Alu repeats. Molecular cell.

Kasal MR, et al. (2023) Lon degrades stable substrates slowly but with enhanced processivity, redefining the attributes of a successful AAA+ protease. Cell reports, 42(9), 113061.

Horio T, et al. (2023) Regulation of RNG105/caprin1 dynamics by pathogenic cytoplasmic FUS and TDP-43 in neuronal RNA granules modulates synaptic loss. Heliyon, 9(6), e17065.

Galupa R, et al. (2023) Enhancer architecture and chromatin accessibility constrain phenotypic space during Drosophila development. Developmental cell, 58(1), 51.

McKie SJ, et al. (2022) Topoisomerase VI is a chirally-selective, preferential DNA decatenase. eLife, 11.

Cui H, et al. (2022) Protocol to image and quantify nucleocytoplasmic transport in cultured cells using fluorescent in situ hybridization and a dual reporter system. STAR protocols, 3(4), 101813.

Gandara L, et al. (2022) Developmental phenomics suggests that H3K4 monomethylation confers multi-level phenotypic robustness. Cell reports, 41(11), 111832.

Janissen R, et al. (2022) High-throughput single-molecule experiments reveal heterogeneity, state switching, and three interconnected pause states in transcription. Cell reports, 39(4), 110749.

Kuijpers L, et al. (2022) Characterizing single-molecule dynamics of viral RNA-dependent RNA polymerases with multiplexed magnetic tweezers. STAR protocols, 3(3), 101606.

Janissen R, et al. (2021) Induced intra- and intermolecular template switching as a therapeutic mechanism against RNA viruses. Molecular cell, 81(21), 4467.

Domsch K, et al. (2021) The Hox Transcription Factor Ubx Ensures Somatic Myogenesis by Suppressing the Mesodermal Master Regulator Twist. Cell reports, 34(1), 108577.

Bera SC, et al. (2021) The nucleotide addition cycle of the SARS-CoV-2 polymerase. Cell reports, 36(9), 109650.

Kurihara M, et al. (2020) Genomic Profiling by ALaP-Seq Reveals Transcriptional Regulation by PML Bodies through DNMT3A Exclusion. Molecular cell, 78(3), 493.

Fukuda S, et al. (2020) The Biogenesis of SRP RNA Is Modulated by an RNA Folding Intermediate Attained during Transcription. Molecular cell, 77(2), 241.

Djabrayan NJ, et al. (2019) Metabolic Regulation of Developmental Cell Cycles and Zygotic Transcription. Current biology: CB, 29(7), 1193.

Johnson HE, et al. (2019) Signaling Dynamics Control Cell Fate in the Early Drosophila Embryo. Developmental cell, 48(3), 361.

Beier KT, et al. (2019) Topological Organization of Ventral Tegmental Area Connectivity Revealed by Viral-Genetic Dissection of Input-Output Relations. Cell reports, 26(1), 159.

Seol Y, et al. (2019) Homology sensing via non-linear amplification of sequence-dependent pausing by RecQ helicase. eLife, 8.

Ohno M, et al. (2019) Sub-nucleosomal Genome Structure Reveals Distinct Nucleosome Folding Motifs. Cell, 176(3), 520.

Le TT, et al. (2019) Synergistic Coordination of Chromatin Torsional Mechanics and Topoisomerase Activity. Cell, 179(3), 619.