Resource Summary Report

Generated by NIF on Aug 30, 2026

Parabase Genomics

RRID:SCR_012336
Type: Tool

Proper Citation

Parabase Genomics (RRID:SCR_012336)

Resource Information

URL: http://www.scienceexchange.com/facilities/parabase-genomics

Proper Citation: Parabase Genomics (RRID:SCR_012336)

Description: THIS RESOURCE IS NO LONGER IN SERVICE. Documented on May 2, 2024. Parabase Genomics is a molecular diagnostic company offering whole exome sequencing services to physicians and researchers with a specialty in the identification of rare variants associated with Mendelian Disorders. Our sample processing and variant analysis pipelines are highly optimized for calling rare variants that are normally missed by less optimized workflows. At Parabase Genomics, we have combined a team of innovators and experts in genomics, pediatrics and targeted sequencing that are not available at other service organizations. Our team is also unique in that we can deliver both research and well as CLIA/CAP certified exomes with insurance reimbursement.

Abbreviations: Parabase Genomics

Resource Type: commercial organization, service resource

Availability: THIS RESOURCE IS NO LONGER IN SERVICE

Resource Name: Parabase Genomics

Resource ID: SCR_012336

Alternate IDs: SciEx_11838

Record Creation Time: 20220129T080309+0000

Record Last Update: 20260829T182417+0000

Ratings and Alerts

No rating or validation information has been found for Parabase Genomics.

No alerts have been found for Parabase Genomics.

Data and Source Information

Source: SciCrunch Registry

Usage and Citation Metrics

We have not found any literature mentions for this resource.