#### **Feature Showcase:**

# High-viability, high-yield, and highly homogeneous single-cell suspensions were obtained, demon-strating excellent quality for downstream applications.

#### Cell viability analysis by flow cytometry

Single-cell suspension from adult mouse whole brain tissue was prepared, with cell viability >90% as determined by 7AAD staining assay.



### Preservation of surface antigens

#### Flow cytometry analysis of neural cell populations

Single-cell suspension from adult mouse whole brain was fractionated into distinct neural subpopulations using surface antigen markers (CD171/CD31/CD11b/O4/ACSA-2) via fluorescence-activated cell sorting (FACS).



### The prepared single-cell suspension meets all requirements for downstream primary cell culture applications.

#### Observation of cell morphology/status by optical microscopy

The single-cell suspension isolated from adult mouse whole brain exhibited excellent culture viability, with adherent cells still observable on day 7 of in vitro culture.





# **Gentle Tissue Enzymatic Digestion Kit (Series) Product Brochure**

www.rwdstco. com

## Introduction

Cell models can be divided into cell lines and primary cell models. For a long time, scientists have relied heavily on immortalized cell lines for various studies. This is because cell lines are generally easier to obtain, with many commercially available. Compared to cell lines, primary cells most closely resemble and reflect in vivo growth characteristics, making them suitable for experiments such as drug sensitivity tests and cell differentiation studies.

The critical first step in primary cell research is preparing a high-quality single-cell suspension. Tissue dissociation into single cells can be challenging due to factors such as tissue type, species, sample age, processing environment, and other variables, requiring tailored dissociation protocols to achieve optimal results.

#### Common methods of tissue dissociation:

Enzyme-free Mechanical cutting Enzyme digestion & Mechanical cutting

Combined with enzymatic digestion, the cell yield obtained is higher and more adaptable to different types of tissue dissociation compared to simple mechanical methods.





Tissue-specific optimization of composite enzymatic formulation, targeting extracellular matrix, aims to preserve cell structure integrity as much as possible, ensuring the acquisition of single-cell suspension with high activity, high yield, and surface antigen protection.

Note: \* It can be operated in conjunction with the RWD single-cell suspension dissociator, or manual dissociation.

# Order Information

www.rwdstco. com



- ity Tumor Tissue Enzymatic Digestion Kit (Mouse)
- ity Tumor Tissue Enzymatic Digestion Kit (Human)
- ity Brain Tumor Enzymatic Digestion Kit
- ity Neonatal Brain Enzymatic Digestion Kit (Mouse and Rat)
- ity Adult Brain Enzymatic Digestion Kit (Mouse and Rat)
- High Activity General Tissue Enzymatic Digestion Kit
- High Activity Intestine Tissue Enzymatic Digestion Kit (Mouse)
- $High \ Activity \ Whole \ Skin \ Enzymatic \ Digestion \ Kit \ (Mouse)$
- High Activity Adipose Tissue Enzymatic Digestion Kit (Mouse and Rat)
- High Activity Muscle Tissue Enzymatic Digestion Kit (Mouse and Rat)
  - vity Hepatocyte Extraction Kit (Mouse and Rat)
  - vity Umbilical Cord Enzymatic Digestion Kit (Human)
  - ity FFPE Tissue Enzymatic Digestion Kit

ency	Nuclei	Extraction	Kit
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sues	Type of cells	Specifications
	Tumor cells and immune cells	50T
s of	Neural stem cells (NSCs), astrocytes, oligodendrocytes, microglia, endothelial cells, and neural progenitor cells (NPCs)	50T
of mouse	Astrocytes, oligodendrocytes, microglia, endothelial cells, and neurons	50T
ı, lung,	Non-parenchymal cells, including immune cells, endothelial cells, macrophages, monocytes, epithelial cells, and fibroblasts	50T
	Tumor cells and immune cells	25T
etc.	Enables efficient debris removal across all routine tissue types.	50T
mina	Mouse lamina propria immune cells	50T
e brain	Tumor cells and immune cells	50T
	Immue cells, Macrophage, fibroblasts, Langerhans cells	25T
ipose	Stromal vascular fraction (SVF), adipose-derived mesenchymal stem cells (ADSCs), endothelial cells	25T
uscle	Myoblasts, muscle satellite cells	50T
2	High-quality mononuclear cell suspensions can be reliably obtained from routine tissue specimens through optimized dissociation protocols.	25T
er	Hepatocytes, hepatic stellate cells, etc.	15T
cord	Mesenchymal stem cells (MSCs), immune cells, and endothelial cells	25T