



Studies search

User guide for OmnibusX web tools

Introduction

Recent advances in multi-omics technologies have significantly expanded the scope of biological research by enabling detailed investigation of cellular states, molecular mechanisms, and tissue-level organization. Techniques such as bulk RNA sequencing, single-cell RNA sequencing (scRNA-seq), single-cell ATAC sequencing (scATAC-seq), and spatial transcriptomics each provide complementary views of gene expression, chromatin accessibility, and spatial context across biological systems.

However, the exponential growth in data generation introduces new challenges in accessing needed datasets and extracting human-understandable information. Traditional databases focus on indexing limited metadata like study titles, abstracts, and categorical tags. Crucially, they lack the detailed annotation efforts from authors and the comprehensive expression profiles of cells, which are key to revealing biological mechanisms. Consequently, researchers often rely on keywords to navigate to studies relevant to their research, then manually download each study for further integration to discern common patterns. This process not only demands substantial computational resources but also significant coding effort. Addressing this need, our database offers an advanced search functionality that encompasses not just traditional study information but also extends to detailed author annotations and comprehensive expression profiles. This capability sets our platform apart from conventional databases, which typically limit searches to basic metadata like titles and abstracts. With our enhanced search function, researchers can efficiently sift through extensive datasets to find studies that are directly relevant to their specific scientific queries.

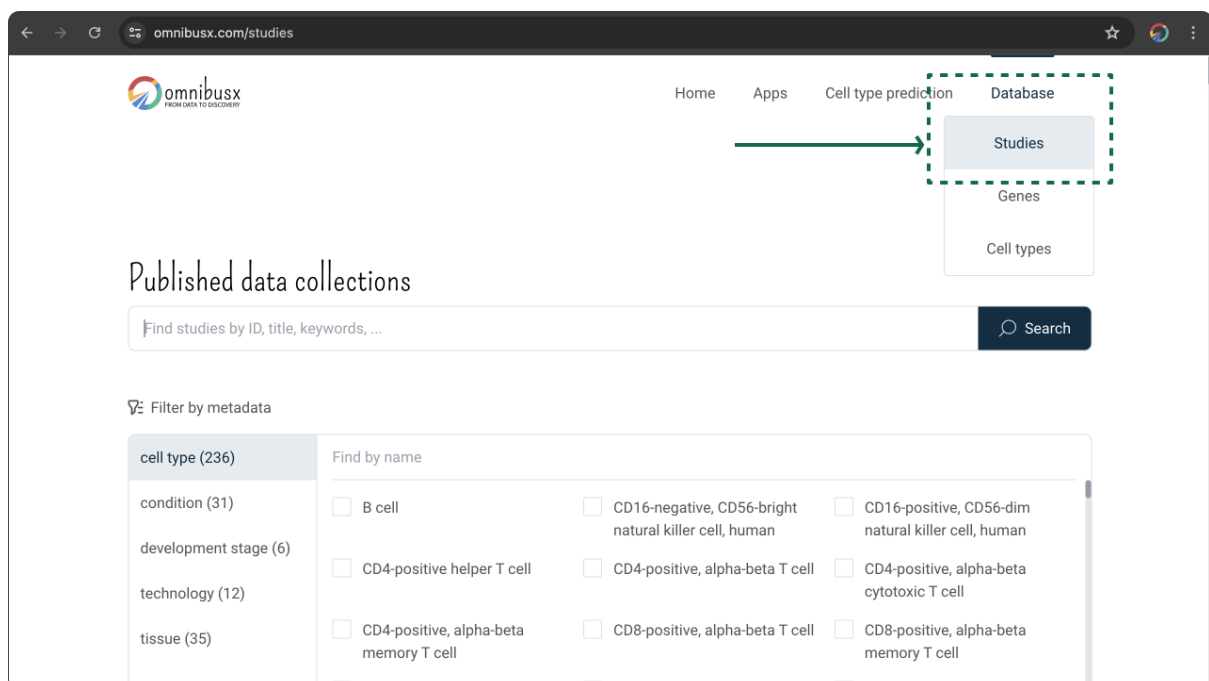
Beyond dataset discovery, analyzing these data types remains computationally challenging. Researchers often rely on disparate software tools, command-line

interfaces, and complex scripting workflows, which can pose significant barriers for researchers without programming expertise. These fragmented pipelines also raise challenges for reproducibility. To overcome this, OmnibusX provides a unified platform for real-time data analysis. OmnibusX integrates widely adopted open-source packages with proprietary analytical modules into a graphical interface that does not require programming. By supporting end-to-end workflows for multiple omics technologies within a single system, OmnibusX eliminates common barriers in multi-omics analysis, enhances reproducibility, and empowers a broader range of researchers to conduct robust, data-driven investigations.

This guide will provide you with detailed instructions on how to utilize the Studies Search to its full potential, facilitating a smoother and more effective research process.

Studies search

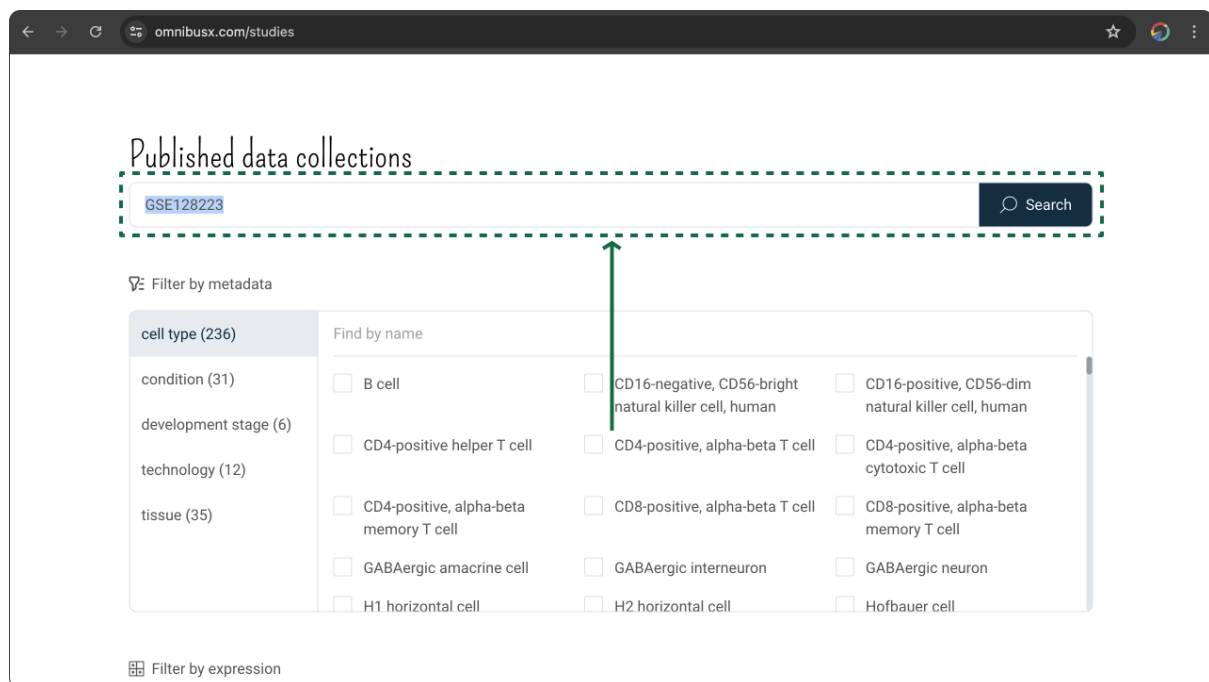
You can access the search tool directly at <https://omnibusx.com/studies>.



1. Queries

1.1. Free text search

To perform a free text search, enter your keyword in the text input and press **Enter** or click on the **Search** button. Your search will encompass all available text information related to a study, including titles, abstracts, author names, repository access IDs, and particularly, all annotations provided by the authors.

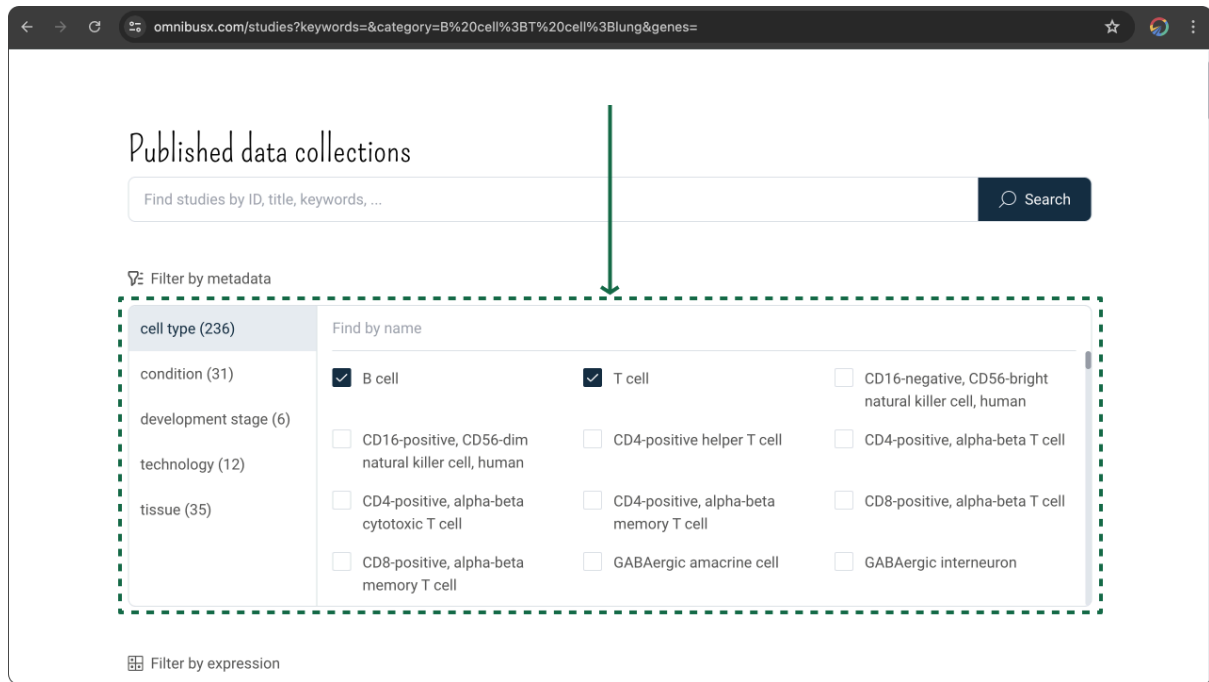


This function is especially useful for quickly locating a study by using well-known information such as the title or ID, or discovering novel annotations related to emerging cell populations that may not yet have official classifications.

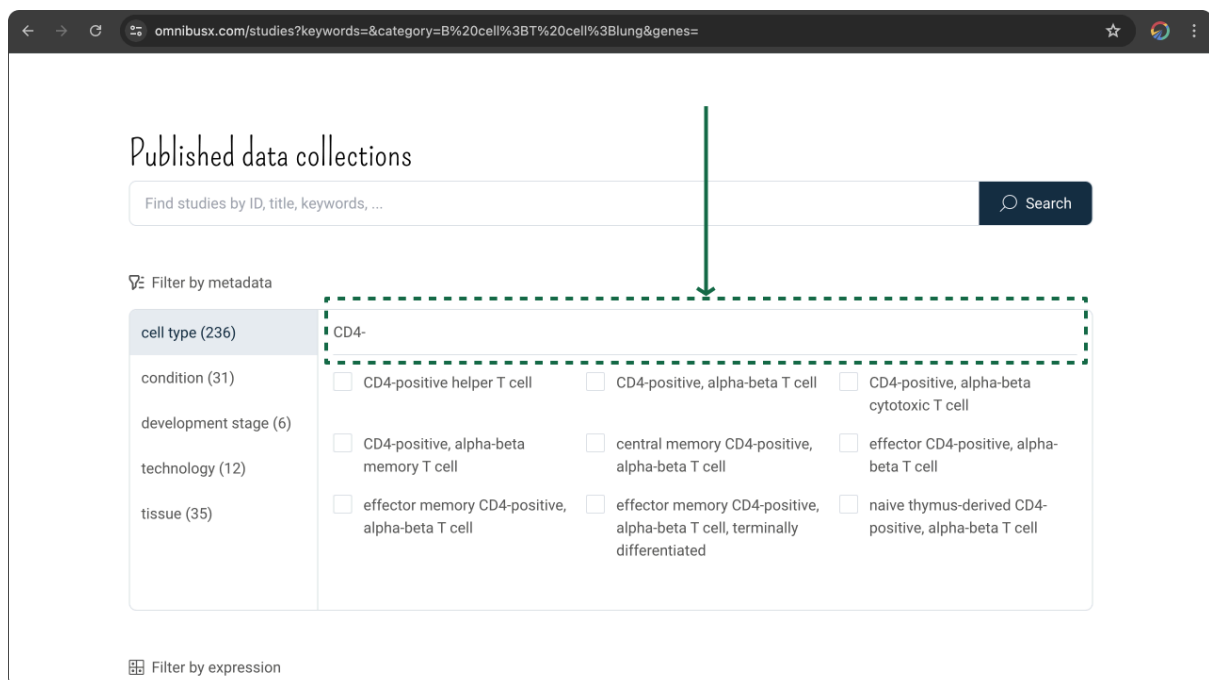
1.2. Controlled vocabularies search

Free text search might lead to missed results due to variations in naming conventions (e.g., CD4 T cell, CD4-positive T cell, CD4+). To accommodate naming variances, OmnibusX has extensively mapped different terminologies to standardized controlled vocabularies. You can utilize this feature by clicking on the needed categories from the **Filter by metadata** panel.

Selecting multiple categories from the same group (e.g., cell types like B cell or T cell) will apply an **OR** search filter, returning studies containing any of the selected categories. Selecting categories from different groups (e.g., tissue type and cell type) will apply an **AND** search filter, returning studies that meet all selected criteria (e.g., studies from lung tissues that contain B cells or T cells).

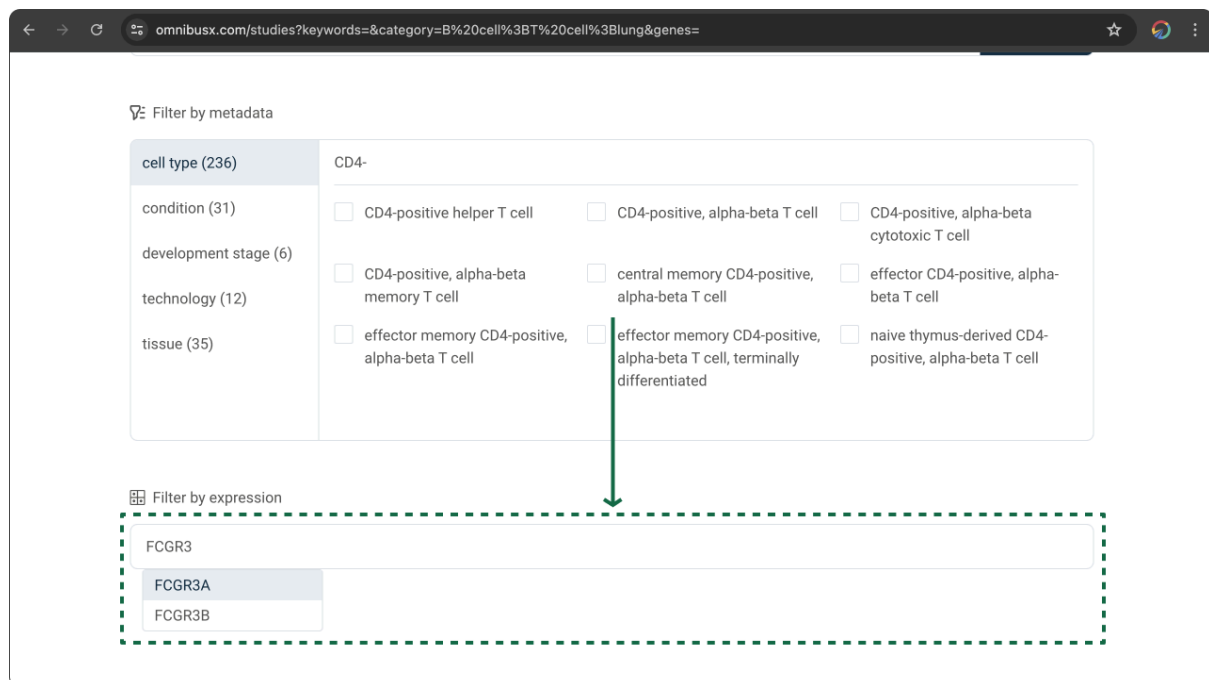


You can quickly find and select your target categories by typing the name into the filter input box above the metadata panel.



1.3. Expression search

This advanced feature of OmnibusX allows you to efficiently locate studies that report the expression of specific genes, saving significant time that would otherwise be spent downloading and processing datasets to verify gene expression. To use this feature, input the gene name into the search box in the **Filter by expression** section. As you type, OmnibusX will suggest related genes which you can select directly.



2. Results

Once your search is executed, all matching studies will be displayed in a tabular format. The number above the table provides a summary of the total search results. The table enables quick access to basic information for each study, including the title, reference URL, data deposited URL, and the number of cells. For each dataset associated with a study, all queries matching your search criteria will be listed for easy review. Original author annotations are displayed rather than controlled vocabularies to highlight potentially novel annotations that may be of

interest. You can click the **Explore** button associated with each study to delve deeper into detailed data and further investigate the study's specifics.

The screenshot shows the omnibusx.com website with search results for the query "peripheral%20blood%3BT%20cell%3BCD8-positive%2C%20alpha-beta%20T%20cell%3BCD8-positiv...". The interface includes a header with navigation links, a search bar, and a results table. Annotations with arrows point to various elements:

- Matched results**: Points to the top summary bar showing 14 studies, 21 datasets, and 3,013,122 cells.
- Sort results**: Points to the "Total cells" column header.
- Basic information**: Points to the first study entry, which includes the title, DOI, deposited data status, and cell count.
- Matched queries**: Points to the "Dataset 1" section of the first study, which lists tissue (blood), cell type (CD8+ Memory/Effector T), and expressed genes (LAG3).
- Explore**: Points to the "Explore" button next to the first study.
- View detail**: Points to the "Explore" button next to the second study, with an upward arrow indicating the flow from the study list to the detailed view.

3. Explore

On selecting the **Explore** option for a specific study, you will be presented with comprehensive information about that study. This includes:

- **Title, Authors, and Abstract**: Fundamental details providing a quick overview of the study's scope and purpose.

omnibusx.com/studies/348da6dc-5bf6-435d-adc5-37747b9ae38a

Immunophenotyping of COVID-19 and influenza highlights the role of type I interferons in development of severe COVID-19

Jeong Seok Lee, Seongwan Park, Hye Won Jeong, Jin Young Ahn, Seong Jin Choi, Hoyoung Lee, Baekgyu Choi, Su Kyung Nam, Moa Sa, Ji-Soo Kwon, Su Jin Jeong, Heung Kyu Lee, Sung Ho Park, Su-Hyung Park, Jun Yong Choi, Sung-Han Kim, Inkyung Jung, Eui-Cheol Shin

Abstract

Although most severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)-infected individuals experience mild coronavirus disease 2019 (COVID-19), some patients suffer from severe COVID-19, which is accompanied by acute respiratory distress syndrome and systemic inflammation. To identify factors driving severe progression of COVID-19, we performed single-cell RNA sequencing using peripheral blood mononuclear cells (PBMCs) obtained from healthy donors, patients with mild or severe COVID-19, and patients with severe influenza. Patients with COVID-19 exhibited hyperinflammatory signatures across all types of cells among PBMCs, particularly up-regulation of the tumor necrosis factor/interleukin-1 β (TNF/IL-1 β)-driven inflammatory response as compared with severe influenza. In classical monocytes from patients with severe COVID-19, type I interferon (IFN) response coexisted with the TNF/IL-1 β -driven inflammation, and this was not seen in patients with milder COVID-19. We documented type I IFN-driven inflammatory features in patients with severe influenza as well. On the basis of this, we propose that the type I IFN response plays a pivotal role in exacerbating inflammation in severe COVID-19.

Datasets

1. Immunophenotyping of COVID-19 and influenza highlights the role of type I interferons in development of severe COVID-19

- **Annotations:** All available annotations from the authors are accessible. This section allows you to toggle between different fields from the summary table to explore various aspects of the study in greater depth.

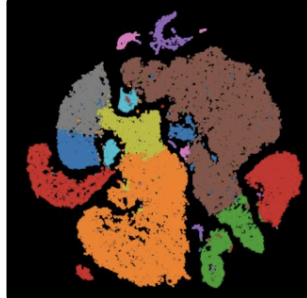
omnibusx.com/studies/348da6dc-5bf6-435d-adc5-37747b9ae38a

Datasets

1. Immunophenotyping of COVID-19 and influenza highlights the role of type I interferons in development of severe COVID-19

Metadata		
Sample ID	nCoV 2	4999 cells
Disease group	Flu 1	4895 cells
Comorbidity	Normal 2	4646 cells
Hospital day	nCoV 6	4526 cells
WBC per microL	Normal 3	4490 cells
Neutrophil per microL (%)	nCoV 1	4464 cells
Lymphocyte per microL	nCoV 11	4425 cells
	Normal 1	4331 cells

Preview



Analyze this study

Dataset 1

- **Real-time analysis:** Click the **Explore** button corresponding to any dataset to begin analyzing it immediately.

2. Coronal (WB_imputation_animal2) Explore

Subject ID	C57BL6J-2	1,900,313 cells
Cell type (subclass transfer)		
High quality transfer		
Tissue annotation (major brain region)		
Tissue annotation (CCF)		
Sample ID		

3. Sagittal (WB_imputation_animal3) Explore

Subject ID	C57BL6J-3	2,081,186 cells
Cell type (subclass transfer)		
High quality transfer		

For a detailed walkthrough of the analysis, please refer to our user guides:

- [Single-cell RNA-seq, scATAC-seq, and bulk RNA-seq analysis](#)
- [Spatial analysis](#)

Sagittal (WB_imputation_animal4) -- Molecularly defined and spatially resolved cell atlas of the whole mouse brain
Number of wells: 214,699

spatial RNA-seq

Import Export Dimensional reduction Quality control Information

Composition Basic charts DEG Enrichment Heatmap Editplot

Subcluster

QUERY GENES FIND MARKERS

rna

Input genes

Unit Color mode

Run AUCell

Cluster Predict cell type Annotate

Switch to scatter

NAME SIZE

brain	18,370
cerebellum	51,392
cortical subplate	7,953
hindbrain	1,188
hippocampal formation	33,238
neocortex	66,020
pallidum	904
striatum	14,056
thalamic complex	115
ventricular system of brain	2,403
white matter	19,060

Thank you!

We extend our heartfelt gratitude to all users of the OmnibusX Multi-Omics Database. Your engagement and feedback are invaluable to us and are what drive continuous improvement and innovation within our database. We are committed to supporting the scientific community by providing robust tools that facilitate groundbreaking research and discovery.

If you have suggestions, feedback, or would like to share how OmnibusX has assisted in your research endeavors, please do not hesitate to reach out to us at support@omnibux.com. Your stories inspire us, and your feedback helps us refine our tools to better serve your needs.