## Case study 5: analysis of an unknown sample

The analysis is based on an unknown sample obtained from either mice or human tissues. After tissue dissociation, cells were sorted by FACS and scRNA-seq libraries were prepared with the Smart-Seq2 protocol/platform.

The starting point is a digital count matrix with mouse or human genes as features.

Analyze the dataset with the methods that you prefer and try to cover the following points.

- 1) Quality control and filtering. Quality control and filtering (mitochondrial genes might be missing from the count matrix, but you can use spike-in RNAs, all starting with "ERCC", as an alternative quality control). ERCC: External RNA Controls Consortium.
- 2) Normalization, identification of variable features, scaling (normal procedure or sctransform)
- 3) Dimensionality reduction
- 4) Clustering
- 5) Identification of marker genes
- 6) Cell cycle analysis
- 7) Annotation

Based on your analysis, on the clusters of cells and marker genes you identified, what is your hypothesis on the sample you are analyzing?

8) Extra: try to integrate and analyze 2 different samples.