GENETICS

CytoTalk: De novo construction of signal transduction networks using single-cell transcriptomic data

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Single-cell technology enables study of signal transduction in a complex tissue at unprecedented resolution. We describe CytoTalk for de novo construction of cell type–specific signaling networks using single-cell transcriptomic data. Using an integrated intracellular and intercellular gene network as the input, CytoTalk identifies candidate pathways using the prize-collecting Steiner forest algorithm. Using high-throughput spatial transcriptomic data and single-cell RNA sequencing data with receptor gene perturbation, we demonstrate that CytoTalk has substantial improvement over existing algorithms. To better understand plasticity of signaling networks across tissues and developmental stages, we perform a comparative analysis of signaling networks between macrophages and endothelial cells across human adult and fetal tissues. Our analysis reveals an overall increased plasticity of signaling networks across adult tissues and specific network nodes that contribute to increased plasticity. CytoTalk enables de novo construction of signal transduction pathways and facilitates comparative analysis of these pathways across tissues and conditions.

INTRODUCTION

Single-cell RNA sequencing (scRNA-seq) technologies are increasingly being used to characterize the heterogeneity of a complex tissue. Beyond cataloging cell types and transcript abundance, it is critical to understand how different cell types interact with one another to give rise to the emergent tissue complexity. Signal transduction is the primary mechanism for cell-cell communication. scRNA-seq technology holds great promise for studying cell-cell communication at much higher resolution. Using scRNA-seq data, several methods have been developed to infer ligand-receptor pairs that are active between two cell types. Skelly et al. (1) and Kumar et al. (2) predict ligand-receptor pairs if the two genes are expressed in the two respective cell types. Zhou et al. (3) and Vento-Tormo et al. (4) identify ligand-receptor pairs whose expression is specific to the cell types considered. Signaling pathways are highly dynamic, and cross-talk among them is prevalent. Because of these two features, simply examining expression levels of ligand and receptor genes cannot reliably capture the overall activities of signaling pathways and interactions among them (5, 6). As a step forward, Wang et al. (7) developed SoftSC and Browaeys et al. (8) developed NicheNet to identify both ligand-receptor pairs and genes downstream of them. However, these methods are based on known annotations of signaling pathways. To our knowledge, no method currently exists to perform de novo prediction of full signal transduction pathways emanating from the ligand-receptor pairs.

Here, we describe the CytoTalk algorithm for de novo construction of a signaling network (union of multiple signaling pathways) between two cell types using single-cell transcriptomic data. The algorithm first constructs an integrated network consisting of intracellular and intercellular functional gene interactions. It then identifies the signaling network by solving a prize-collecting Steiner forest (PCSF) problem based on appropriately defined node prize (i.e., cell-specific gene activity) and edge cost (i.e., probability of functional interaction between two genes). The objective of the PCSF problem is to find an optimal subnetwork in the integrated network that includes genes with high levels of cell type–specific expression and close connection to highly active ligand-receptor pairs. We benchmark the performance of CytoTalk using high-throughput spatial transcriptomic data and scRNA-seq data with perturbation to the receptor genes in a signaling pathway. To gain new insight into the heterogeneity of signaling pathways across tissues and developmental stages, we apply CytoTalk to perform a comparative analysis of signaling networks between macrophages and endothelial cells across human adult and fetal tissues. A software package implementing the CytoTalk algorithm has been deposited at GitHub (https://github.com/tanlabcode/CytoTalk).

RESULTS

Wiring of signaling pathways is highly cell type dependent

A hallmark of signal transduction pathways is their high level of cell type–specific wiring pattern. Single-cell transcriptome data allow us to examine the cell type–specific activity of individual signaling pathways beyond just ligand and receptor genes. To this end, we examined the canonical fibroblast growth factor receptor 2 (FGFR2) signaling pathway in two tissues, mammary gland and skin. Four canonical downstream pathways are known to signal from FGFR2 (9), including Janus kinase and signal transducer and activator transcription proteins (JAK-STAT), protein kinase C (PKC), mitogen-activated protein kinase (MAPK), and phosphoinositide 3-kinase and protein kinase B (PI3K/AKT) pathways. For mammary gland, we studied FGFR2 signaling between fibroblasts and luminal epithelial cells or basal cells (10, 11). For skin, we studied FGFR2 signaling between keratinocyte stem cells and basal cells (12). Using published scRNA-seq data (13) for each tissue type, we computed an expression specificity score, preferential expression measure (PEM) (14, 15) (see Materials and Methods), for each pathway gene in each cell type (Fig. 1, A to C). We found that the four canonical subpathways downstream of the same receptor (FGFR2) exhibit notable difference in cell type–specific

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activities not only within the same tissue but also across different tissues. In the mammary gland, the PI3K/AKT and PKC pathways are the most active pathways for signaling between fibroblasts and luminal epithelial cells and basal cells, respectively. In contrast, the JAK-STAT pathway is the most active pathway for signaling between keratinocyte stem cells and basal cells in the skin. To systematically evaluate the extent of cell type–specific wiring of signaling pathways, we examined all manually annotated signaling pathways in the Reactome database (16). For each pathway, we computed its cell type–specific activity score using the same mammary gland and skin scRNA-seq datasets (13). We found that most of the pathways exhibit high degree of cell type–specific activities (Fig. 1, D and E). This is true even for the same cell type across different tissues, such as basal cells from mammary gland versus basal cells from skin (Fig. 1F).

Using an alternative source of signaling pathway annotations that include gene signatures for Myc, Ras, E2F3, Src, and β-catenin pathways (17), we found that all pathways exhibit differential activities across cell types in mammary gland and skin (fig. S1). Together, these results highlight the need for analytical tools for de novo construction of complete signaling pathways (instead of ligand-receptor pairs) using single-cell transcriptomic data.

**Overview of the CytoTalk algorithm**

CytoTalk is designed for de novo construction of a signal transduction network between two cell types (Fig. 2; Materials and Methods), which is defined as the union of multiple signal transduction pathways. It first constructs a weighted integrated gene network composed of both intracellular and intercellular functional gene-gene interactions. Intracellular functional gene interactions are computed and weighted using mutual information between two genes. Two intracellular networks are connected via cross-talk edges (i.e., known ligand-receptor interactions). Ligand-receptor pairs with higher cell type–specific expression but lower correlated expression within the same cell type (thus more likely to be involved in cross-talk rather than self-talk) are assigned higher cross-talk weights. Nodes in the integrated network are weighted by a combination of their cell...
type–specific gene expression and closeness to the ligand/receptor genes in the network. We use a network propagation procedure to determine the closeness of a gene to the ligand/receptor genes. With the integrated network as the input, we formulate the identification of signaling network as a PCSF problem (18, 19). The rationale for using the PCSF algorithm is to find an optimal subnetwork that includes genes with high levels of cell type–specific expression and close connection to high-scoring ligand-receptor pairs. The objective function of the PCSF algorithm is designed to capture these two types of information/weights (see Materials and Methods for details). This optimal subnetwork is defined as the signaling network between the two cell types. The statistical significance of the identified signaling network is computed using a null score distribution of signaling networks generated using degree-preserving randomized networks.

Performance evaluation using spatial transcriptomic data
To evaluate the performance of CytoTalk, we applied it to scRNA-seq datasets on mouse visual cortex (20) and olfactory bulb (21), respectively. Both datasets cover the following cell types: neurons, astrocytes, microglia cells, endothelial cells, oligodendrocytes, and oligodendrocyte precursor cells. In the visual cortex dataset, neurons were further classified into two subtypes, glutamatergic (Glut) neurons and GABAergic (GABA) neurons. On average, 6358 and 836 genes were detected per cell in the visual cortex and olfactory bulb datasets, respectively. Among the covered cell types, neurons, endothelial cells, astrocytes, and microglia cells are known to signal to each other (22–29). We identified signaling networks between five pairs of these cell types, including neuron–endothelial cell (NeuronEndo for the olfactory bulb or GlutEndo and GABAEndo for the visual cortex), neuron-astrocyte (NeuronAstro or GlutAstro and GABAstro), neuron-microglia (NeuronMicro or GlutMicro and GABAMicro), astrocyte-endothelial cell (AstroEndo), and endothelial cell–microglia cell (EndoMicro), respectively. The eight predicted cell type pair–specific signaling networks in the visual cortex consist of 567, 445, 692, 332, 643, 628, 577, and 403 genes and involve 24, 22, 42, 29, 32, 28, 39, and 45 ligand-receptor interactions (cross-talk edges), respectively (table S1). For the olfactory bulb, the five predicted signaling networks consist of 627, 542, 604, 764, and 688 genes and involve 21, 19, 31, 42, and 39 ligand-receptor interactions, respectively (table S2).
To systematically evaluate the performance of CytoTalk, we used two matched sequential fluorescence in situ hybridization (SeqFISH+) datasets for the visual cortex and olfactory bulb (32), respectively, covering the same set of cell types as the scRNA-seq datasets (20, 21). On average, these two datasets provide a spatially resolved abundance of 5826 transcripts (3344 genes) per cell in the visual cortex and 1284 transcripts (844 genes) per cell in the olfactory bulb, respectively. For each pair of cell types under study, we divided cell pairs into close and distant groups based on their physical distance determined using the SeqFISH+ data (fig. S6 and Materials and Methods). We then calculated the mutual information of the SeqFISH+ expression among signaling pathway genes across the two groups of cell pairs (Fig. 3G). We reasoned that cells that are close are more likely to signal to each other. Therefore, bona fide signaling pathway genes are expected to have higher mutual information of spatial expression in these cells than cells that are distant from each other.

For Glut neurons and endothelial cells in the visual cortex, there are 2812 and 916 cell pairs in the close and distant groups, respectively (Materials and Methods). We found that genes in signaling pathways predicted by CytoTalk have significantly larger mutual information of spatial expression in close GlutEndo cell pairs than distant cell pairs (one-sided Kolmogorov-Smirnov test P = 2.0 × 10−100), thus providing support to our predictions (Fig. 3, G and H, and fig. S7A). In comparison, predicted pathway genes by other methods show no or less significant difference in mutual information between close and distant cell pairs (Fig. 3H and fig. S7A), suggesting that those predictions have higher false-positive rates.

For the predicted signaling networks of the other 12 cell type pairs, we also found that CytoTalk predictions have consistently larger mutual information in close cell pairs than distant cell pairs, except for the NeuronMicro signaling network in the olfactory bulb. In comparison, predictions by other methods do not show significant separation in mutual information between close and distance cell pairs (Fig. 3, H and I, and fig. S7). Together, these results demonstrate that CytoTalk has substantial improvement over existing methods.

**Performance evaluation using scRNA-seq data with receptor gene perturbation**

To further evaluate the accuracy of CytoTalk, we applied it to an scRNA-seq dataset in which the transcriptomes of wild-type and receptor gene knockout cells were measured (34). The dataset covers 13 cell types in the mouse lung, including T and B cells, neutrophils, basophils, monocytes, macrophages, endothelial cells, alveolar type I (AT1) and type II (AT2) cells, club cells, smooth muscle cells, fibroblasts, and pericytes. On average, 2627 transcripts (1143 genes) were detected per cell. The authors found a novel signaling pathway involving interleukin 33 (IL33) secreted by AT2 cells and IL1 receptor–like 1 (IL1RL1) on basophils (34). Using this dataset, we first asked whether the IL33-IL1RL1 interaction between AT2 cells and basophils can be predicted. We found that the three methods (CytoTalk, NicheNet, and SoptSC) can identify the IL33-IL1RL1 interaction (table S1). We then evaluated the prediction accuracy using receiver operating characteristic (ROC) curve. To this end, we used the differentially expressed genes (DEGs; Materials and Methods) between IL1RL1-knockout and wild-type basophils as the ground truth. We found that predictions by CytoTalk have a higher area under the ROC curve (Fig. 4A). Furthermore, the downstream pathway genes predicted by CytoTalk tend to be more significantly

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Fig. 3. Performance evaluation of the CytoTalk algorithm using spatial transcriptomic data. (A and B) Venn diagrams for genes in the microglia-, astrocyte-, and endothelial cell–related signaling networks predicted by CytoTalk in the visual cortex (A) and olfactory bulb (B). (C to E) Representative signaling pathways uniquely predicted by CytoTalk for GlutMicro (C), EndoMicro (D), and AstroEndo (E) in the visual cortex. (F) Pathway genes in (C) to (E) are enriched for nervous system–related Gene Ontology (GO) biological processes and pathways in the Reactome (REAC) and KEGG databases. Shades of blue in the heatmap are inversely proportional to the enrichment P values (> 0.05) are indicated in yellow. (G) Schematic illustration of the procedure and rationale for using SeqFISH+ data to evaluate predicted signaling networks. (H and I) Performance evaluation using mutual information of spatial expression of predicted pathway genes across cell pairs in close and distant groups. Cell pairs were categorized into the two distance groups using SeqFISH+ data. Cells in the same FOV were considered as close, whereas cells from two most distant FOVs were considered as distant. Mutual information of predicted pathway genes was computed using the SeqFISH+ data and compared between the close and distant cell pair groups. Statistical significance between the two distributions of mutual information was computed using one-sided Kolmogorov-Smirnov test for predicted pathways in the visual cortex (H) and olfactory bulb (I). Each dot in the boxplot represents the performance on a given cell type pair. Detailed results are illustrated in fig. S7. Dashed line, significant P value of 0.05.
genes with significant differential entropies have larger tissue specificity associated with their diverse functions across tissues. We found several that transcriptomes differ more across adult organs than across fetal tissues (Wilcoxon rank sum test found that the significant genes have larger differential entropies among adult tissues than fetal tissues (Wilcoxon rank sum test). P = 0.03; Fig. 5C and table S2). Second, we found that significant genes are less evolutionary conserved than nonsignificant genes (Wilcoxon rank sum test P = 0.05; Fig. 5D and table S2). This anticorrelation between tissue specificity and sequence conservation was also observed in previous studies (40, 41). Third, we found that the ligand-receptor pairs that are most closely related to the significant genes have larger differential cross-talk scores (see Materials and Methods) across tissues (Fig. 5E) than ligand-receptor pairs related to nonsignificant genes (Wilcoxon rank sum test P = 0.03; Fig. 5F). Last, to further investigate the functions of significant genes that are shared by all three tissues, we extracted their associated pathways (Materials and Methods) from the predicted signaling networks. We found that they are enriched for tissue-specific functions and tissue-specific disease phenotypes (Fig. 5G), probably reflecting their functional adaptation across tissues (39). Together, these results highlight the heterogeneity of signaling pathways between the same cell type pair in different tissues, which may play an important role in tissue adaptation of these cell types.

We noticed that, among the significant genes shared by all three tissues, IFITM3 and CD74 are known to have important antiviral activities in endothelial cells (42) and macrophages (43), respectively. They are also the top genes with differential entropy in the endothelial (first ranked) and macrophage (second ranked) signaling networks, respectively. To investigate whether there exist functional cross-talk pathways between them, we extracted the shortest paths between IFITM3 and CD74 from the predicted signaling networks of the three tissues (Fig. 5H). We found that the majority of the genes (84%) on the shortest paths are different among the tissues. They are enriched for both common and tissue-specific virus-related biological processes and pathways, suggesting that different signaling pathways may be activated in response to a given virus in different tissues (Fig. 5I). Consistent with the fact that the lung is the main target organ of coronavirus infection, only the shortest-path genes in the lung are enriched for the term host-pathogen interaction of human corona viruses. This result implicates a possible role of the

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**Fig. 5. Performance evaluation of the CytoTalk algorithm using scRNA-seq data with receptor gene perturbation.** Pathways predicted by CytoTalk, NicheNet, and SopSC were evaluated by comparing to DEGs between receptor gene knockout and wild-type cells. DEGs were identified by DESeq2 using a Benjamini-Hochberg (BH)-adjusted P value cutoff of 0.05. (A) ROC curve. DEGs between IL1RL1 knockout and wild-type basophils are considered true positives. True-positive rate and false-positive rate were computed using the DEGs. Numbers at each point on the curve indicate the network distance of predicted genes to the receptor (IL1RL1). For instance, 1st means a predicted gene that is a first-order neighbor of the receptor. (B) Distribution of P values of differential expression for all genes in the predicted downstream pathways. Shown are violin plots of −log10(BH-adjusted P value). P values for comparing distributions were computed using one-sided Wilcoxon rank sum test.

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**Entropy of signaling pathways across human tissues**

Our analysis in Fig. 1 suggests that wiring of signaling pathway is highly cell type dependent. To better understand the heterogeneity of signaling pathways, we applied CytoTalk to perform a comparative analysis across tissues and developmental stages. Endothelial cells and macrophages are two common cell types in many mammalian tissues, and their interaction plays a critical role in the tissue adaptability of macrophages and tissue homeostasis (35, 36). We analyzed scRNA-seq datasets of these two cell types from human kidney, heart, and lung, at both fetal and adult stages (table S1) (37). On average, we found that 52 and 47% of the predicted pathway genes are shared among the three tissues at both fetal and adult stages, respectively (Fig. 5A). For shared pathway genes, we calculated their local Shannon entropy (38) that measures the uncertainty of signaling interaction patterns associated with a given pathway gene in a tissue. Genes with higher signaling entropies have more uniform signaling interaction probabilities with their interacting partners in the signaling network. We next identified genes that exhibit significantly large differential entropies across the three tissues (Materials and Methods).

These genes represent the most dynamic nodes of the signaling networks across tissues.

We identified 18 and 28 significant genes (empirical P < 0.1; table S2) in fetal and adult tissues, respectively. We found that the significant genes have larger differential entropies among adult tissues than fetal tissues (Wilcoxon rank sum test P = 6.8 × 10^{-5}; Fig. 5B). This is reminiscent of a previous observation that transcriptomes differ more across adult organs than across fetal organs (39).

We hypothesized that genes with differential entropies are associated with their diverse functions across tissues. We found several lines of evidence supporting this hypothesis. First, we observed that genes with significant differential entropies have larger tissue specificity scores (Materials and Methods) than nonsignificant genes (Wilcoxon rank sum test P = 0.03; Fig. 5C and table S2). Second, we found that significant genes are less evolutionary conserved than nonsignificant genes (Wilcoxon rank sum test P = 0.05; Fig. 5D and table S2). This anticorrelation between tissue specificity and sequence conservation was also observed in previous studies (40, 41). Third, we found that the ligand-receptor pairs that are most closely related to the significant genes have larger differential cross-talk scores (see Materials and Methods) across tissues (Fig. 5E) than ligand-receptor pairs related to nonsignificant genes (Wilcoxon rank sum test P = 0.03; Fig. 5F). Last, to further investigate the functions of significant genes that are shared by all three tissues, we extracted their associated pathways (Materials and Methods) from the predicted signaling networks. We found that they are enriched for tissue-specific functions and tissue-specific disease phenotypes (Fig. 5G), probably reflecting their functional adaptation across tissues (39). Together, these results highlight the heterogeneity of signaling pathways between the same cell type pair in different tissues, which may play an important role in tissue adaptation of these cell types.

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**Fig. 5. Plasticity of signaling networks across tissues and developmental stages.** Signaling networks between endothelial cells and macrophages in kidney, heart, and lung were predicted for both fetal and adult stages. (A) Venn diagrams for genes in the predicted signaling networks of the three tissues. (B to D) Distribution of differential entropies (B), tissue specificity scores (C), and sequence conservation scores (D) of the shared signaling network genes of the three tissues. The shared genes were stratified into significant and nonsignificant groups based on empirical P values. (E) Cross-talk scores of ligand-receptor pairs that are most closely related to the significant genes in the predicted networks. (F) Distributions of differential cross-talk scores between the three tissues for the ligand-receptor (LR) pairs that are most closely related to the significant and nonsignificant genes. P values for comparing distributions were computed using one-sided Wilcoxon rank sum test. (G) The neighborhood of significant genes shared by all three tissues in the predicted signaling networks is enriched for tissue-specific Gene Ontology biological processes and human disease phenotypes (HP). (H) The shortest paths between two important host antiviral genes, IFITM3 and CD74, in the predicted signaling networks of the three tissues. (I) Genes on the shortest paths in (H) are enriched for virus-related Gene Ontology biological processes and infection response pathways from Reactome, KEGG, and WikiPathways (WP) databases. Shades of blue in the heatmap are inversely proportional to the enrichment P values that are adjusted for multiple testing using the Benjamini-Hochberg method. Nonsignificant P values (> 0.05) are indicated in yellow.

**DISCUSSION**

We introduce a computational method, CytoTalk, for the construction of cell type–specific signaling pathways using single-cell transcriptomic data and known ligand–receptor interactions. Unlike previous methods using known pathway annotations (7, 8), CytoTalk constructs full signaling pathways de novo. Using CytoTalk, we performed a comparative analysis of signaling pathways across human tissues and developmental stages, revealing increased heterogeneity of signaling networks in adult tissues compared to fetal tissues and specific nodes in the network that exhibit significant changes in signaling entropy across tissues.

Systematic evaluation of predicted signaling pathways represents a major challenge due to the lack of gold-standard cell type–specific pathway annotations. Here, we propose two benchmarking strategies using single-cell spatial transcriptomic data and perturbation-based scRNA-seq data, respectively. Using spatial transcriptomic data, cells of two types can be stratified into close and distant cell pair groups based on their physical separation. The predicted signaling pathways can be validated by computing the mutual information of spatial expression for pathway genes across cell pairs. Using perturbation-based

**IFITM3 and CD74 signaling cross-talk in SARS-CoV-2 (COVID-19) infection,** which are also supported by several recent studies (43–45).
scRNA-seq data, especially data with ligand/receptor gene perturbation, DEGs between the perturbed and wild-type cells can be used to validate predicted pathways. Using these two benchmarking approaches, we demonstrated that CytoTalk substantially outperforms six existing methods that also use scRNA-seq data to characterize cell–cell communication.

In the current version of CytoTalk, the node prize is defined on the basis of cell type specificity of gene expression. Thus, CytoTalk may fail to identify signaling pathways whose genes have low expression specificity in the cell types under study. To address this issue, the node prize can be redefined by considering both absolute gene expression level and cell type specificity of gene expression. It is also well known that activity of a signaling pathway is regulated by post-translational modifications. With the rapid development of single-cell proteomic technologies (46), CytoTalk can be further improved by incorporating these data. The current version of CytoTalk uses intracellular networks constructed using scRNA-seq data. Other types of molecular interaction data such as protein–protein interactions and genetic interactions are becoming more abundant. The utility of these data types can be investigated in the future development of CytoTalk.

In summary, CytoTalk provides a much-needed means for de novo construction of full cell type–specific signaling pathways. Comparative analysis of signaling pathways will lead to a better understanding of cell–cell communication in healthy and diseased tissues.

MATERIALS AND METHODS
Construction of intracellular functional gene interaction network
We construct an intracellular gene coexpression network for each cell type by calculating the mutual information between all pairs of genes. Construction of cell-cell communication in healthy and diseased tissues.

The cross-talk score equals the product of the min-max–normalized expression score and non–self-talk score. If genes i and j are specifically expressed in cell types A and B, respectively, but not co-expressed in either cell type (likely involved in self-talk), then the Cross-talk score would be high, suggesting a high possibility of cross-talk between the two cell types.

Construction of an integrated network between two cell types
We construct an integrated network consisting of two intracellular networks connected by known ligand–receptor interactions. We collected 1942 manually annotated ligand–receptor interactions, including 1894 interactions from (31) and 48 interactions from (34, 52–56) (table S3). Note that both secreted and cell surface proteins could be ligands. For each ligand–receptor pair, if the ligand gene and the receptor gene are present in the two intracellular networks, then we connect them and denote the edge as a cross-talk edge. The cross-talk score is used as the edge weight as described above. Because of the difference in scale between mutual information value and cross-talk score, we separately normalize the edge weights of intracellular networks and cross-talk edges using z score transformation.

De novo identification of signaling network between two cell types
We formulate the identification of a signaling network between two cell types as a PCSF problem (18, 19). Because the forest is a disjoint set of trees, the PCSF problem is a generalization of the classical prize-collecting Steiner tree (PCST) problem (57, 58). The individual signaling pathways are represented as trees, and the collection of (forest) represents the entire signaling network between two cell types.

We define edge costs and node prizes in the integrated network as follows. The z score–normalized edge weights of the integrated network are first scaled to the range of [0, 1]. The cost of edge e in
where \(\text{Relevance}^i\) is the relevance coefficient vector for all genes in the intracellular network at iteration \(i\). \(\text{Relevance}^0\) is the initial value of the relevance coefficient vector such that \(\text{Relevance}^0(i) = 1\) if gene \(i\) is a ligand or receptor. Otherwise, \(\text{Relevance}^0(i) = 0\). \(W\) is a normalized edge weight matrix for an intracellular network, which is defined as \(W = D^{-1/2}W^{1/2}\). Here, \(W\) is set to the original mutual information matrix, and \(D\) is defined as a diagonal matrix such that \(D(i, i)\) is the sum of row \(i\) of the matrix \(W\). This network propagation procedure is equivalent to a random walk with restart on the network. \(\alpha\) is a tuning parameter that controls the balance between prior information (known ligands or receptors) and network smoothing. Thus, node prize of gene \(i\) in the integrated network is defined as \(\rho(i) = \text{PEM}_i \times \text{Relevance}^i(i)\). This definition can capture both the cell type specificity and the closeness of this gene to the ligand or receptor gene in the network. To avoid extremely large node prizes for ligand or receptor genes, we used \(\alpha = 0.9\) in this study.

The PCSF algorithm identifies an optimal forest in a network that maximizes the total amount of node prizes and minimizes the total amount of edge costs in the forest. Although the PCSF problem is NP (nondeterministic polynomial time)-hard and often needs a high computational cost (18), we use a PCSF formulation established in (18, 59) and use a highly efficient PCST algorithm (60) to identify the PCSF. The objective function of the PCSF problem is defined as below

\[
\min c(F) + \beta \times \rho(F) + \omega \times k
\]

where \(F\) represents a forest (i.e., multiple disconnected trees) in the integrated network. \(c(F)\) denotes the sum of edge costs in the forest \(F\), and \(\rho(F)\) denotes the sum of node prizes of the remaining subnetwork excluding the forest \(F\) from the network. We modify the integrated network by introducing an artificial node and a number of artificial edges to the original network. The artificial edges connect the artificial node to all genes in the original network. The costs of all artificial edges are the same and are defined as \(\omega\), which influences the number of trees, \(k\), in the resulting PCSF. \(\beta\) is a parameter for balancing the edge costs and node prizes, which influences the size of the resulting PCSF. By tuning parameters \(\beta\) and \(\omega\), multiple PCSTs can be identified with the artificial node as the root node. For each identified PCST, a PCSF can be obtained by removing the artificial node and artificial edges from the PCST.

We identify the signaling network between two cell types by searching for the most robust PCSF across the full parameter space (see below; fig. S8). To further evaluate the statistical significance of the identified PCSF, we construct null distributions for the objective function and for the fraction of cross-talk edges in a PCSF using 1000 null PCSFs identified from randomized integrated networks (fig. S2). To generate the randomized networks, we separately shuffle the edges of the two intracellular networks while preserving the node degree distribution, node prizes, and cross-talk edges as the original integrated network.

**Parameter selection**

The main parameters of CytoTalk are \(\beta\) and \(\omega\) in the objective function of the PCSF problem. We first determined the optimal ranges of the two parameters based on the total number of genes in the input integrated network and the size and overlap of the resulting PCSFs. For the mouse visual cortex dataset, \(\beta\) values were tested from 1 to 60 with a step size of 1. For the mouse olfactory bulb and lung tissue datasets, \(\beta\) values were tested from 5 to 500 with a step size of 5. For the Human Cell Atlas dataset, \(\beta\) values were tested from 1 to 80 with a step size of 1. We found that the upper limit of the test values of \(\beta\) is inversely proportional to the total number of genes in the integrated network of a given cell type pair (fig. S8A). Using \(\beta\) values above this upper limit results in very large PCSFs (>2000 edges) (fig. S8B). On the basis of this observation, we suggest that \(\beta\) values can be tested from 1 to 100 if the total number of genes in a given cell type pair is above 10,000. However, if the total number of genes is below 5000, then it is necessary to increase the upper limit of test \(\beta\) value to 500 to ensure that the largest PCSF includes more than 2000 edges.

For all datasets, \(\omega\) values were tested from 0.1 to 1.5 with a step size of 0.1. We found using \(\omega\) values above this range resulting in PCSFs with little difference compared to existing PCSFs (fig. S8C). Subsequent optimal parameter selection was conducted using the above parameter ranges. For all PCSTs identified using the \(\beta\) and \(\omega\) ranges determined above, the occurrence frequency of each edge in a PCSF was computed to construct a background distribution of edge occurrence frequency. A \(P\) value for each PCSF was computed by comparing the edge occurrence frequency distribution of this PCSF to the distribution of all other PCSFs using one-sided Kolmogorov-Smirnov test. The PCSF with the minimum \(P\) value (red dot) was considered as the most robust signaling network predicted by CytoTalk (fig. S8D).

**Processing of scRNA-seq data**

For all scRNA-seq datasets used in this study (table S4), we only retained protein-coding genes based on annotations from the GENCODE database (61). For data from the Human Cell Atlas (37), following the authors’ data processing procedure, we removed genes expressed in fewer than three cells of a given type. For all other datasets, we removed genes expressed in less than 10% of all cells of a given type.

For identifying DEGs between \(IL1RL1\)-knockout and wild-type basophils, we first filtered out genes that have fewer than five sequencing reads in at least five cells. Then, we used the zinwave function in the zinwave R package (62) to model the zero inflation of the counts. The DESeq2 R package (63) was used to perform differential expression analysis. \(P\) values were adjusted for multiple testing using the method of Benjamini-Hochberg (64). A total of 619 DEGs with adjusted \(P < 0.05\) were identified (table S5).

**Processing of SeqFISH+ data**

Two SeqFISH+ datasets were downloaded from a published study (32), including cell type annotation, cell spatial location [i.e., field of view (FOV) index] data, and two cell-by-gene count matrices for 523 visual cortex cells and 2050 olfactory bulb cells, respectively (table S6). On the basis of the authors’ preprocessing procedure.


Mutual information of spatial expression of signaling pathway genes using SeqFISH+ data

Cell pairs consisting of two types were categorized into two groups based on their physical locations imaged by SeqFISH+, namely close and distant groups. SeqFISH+ can image mRNAs for thousands of genes in all single cells in each FOV. Five and seven FOVs were imaged in the visual cortex and olfactory bulb datasets, respectively (32). We considered two cells in the same FOV as a close cell pair, whereas two cells from two most distant FOVs were considered as a distant cell pair (fig. S6). Genes in predicted signaling pathways were intersected with the genes detected by SeqFISH+. Among these genes, we computed the mutual information of SeqFISH+ expression values between any gene pair (one from each cell type and are connected in the predicted signaling pathways) across individual cells of the two types.

Running of published methods

NicheNet (8) uses a prior ligand-target regulation model by integrating known signaling transduction and transcriptional regulatory interactions to predict a signaling network between two cell types. Given the scRNA-seq data of cell type A and B, we first defined two gene sets that are expressed in these two cell types, respectively. We used the same expression cutoff and only retained protein-coding genes for analysis as described above. NicheNet requires a pre-defined gene set of interest in cell type B as candidate target genes regulated by ligands in cell type A. This gene set was defined as the genes that are specifically expressed (i.e., PEM score of >0) in cell type B. Using these gene sets as the input, we predicted ligands, their signaling pathways, and target genes using predict_ligand_activities and get_ligand_signaling_path functions in the NicheNet R package. These identified signaling pathways were considered as a “cell type A to B” signaling subnetwork. Using another set of genes that are specifically expressed in cell type A as input, we also predicted a “cell type B to A” signaling subnetwork. By combining the two subnetworks, we obtained a final predicted signaling network between the two cell types.

SoptSC (7) also uses known pathway annotations to predict signaling pathways between two cell types. For comparison, we used mouse Reactome pathways and the same ligand-receptor pairs as used by CytoTalk. For each ligand-receptor pair, we computed a matrix of signaling probabilities between any two cells from cell types A and B using the LR_Interaction function in the SoptSC MATLAB package. Using this matrix, we computed two probabilities of signaling via the given ligand-receptor pair from each direction (A → B and B → A), respectively. On the basis of these signaling probabilities, we selected the same number of ligand-receptor pairs as predicted by CytoTalk and their known downstream pathways for performance comparison.

Different from NicheNet and SoptSC, the other four methods only predict active ligand-receptor pairs between two cell types. Among these four methods, Zhou’s and Skelly’s methods are similar, which consider the gene expression levels of a ligand and its receptor separately. On the basis of Zhou’s method (3), for each gene i, we calculated the mean $\mu_i$ and SD $\sigma_i$ of the gene expression values across all cell types. If the average expression values of the ligand gene in cell type $A$ and the receptor gene in cell type $B$ are both larger than $\mu_i + 2 \sigma_i$, then this ligand-receptor pair is predicted to be active by Zhou’s method. On the basis of Skelly’s method (1), for each ligand-receptor pair, if the ligand and the receptor genes are expressed in more than 20% of the cells of cell types $A$ and $B$, respectively, then this ligand-receptor pair is retained and considered to transmit a signal from cell type $A$ to $B$. We considered all retained ligand-receptor pairs from both directions ($A \rightarrow B$ and $B \rightarrow A$) as the final predictions by Skelly’s method.

Kumar’s method (2) is different from the two methods above, which defines an interaction score for a given ligand-receptor pair as the product of the average expression of the ligand gene in cell type $A$ and the average expression of the receptor gene in cell type $B$. We selected the same number of ligand-receptor pairs as predicted by CytoTalk based on interaction scores as the final predictions by Kumar’s method.

CellPhoneDB (4) is a repository of curated ligand-receptor pairs and can be used for predicting ligand-receptor interactions based on their cell type specificity. Given an scRNA-seq gene expression matrix and cell type annotation data as the input, we used the cellphonedb function in the CellPhoneDB Python package to compute a P value for the likelihood of cell type specificity of each ligand-receptor pair. For two given cell types $A$ and $B$, we selected the ligand-receptor pairs with $P < 0.05$ from each direction ($A \rightarrow B$ and $B \rightarrow A$) as the final predictions by CellPhoneDB.

Differential signaling entropy

Given a signaling network and gene expression data, a local Shannon entropy for each gene in the network can be computed to measure the uncertainty of signaling interaction incident on a given gene. Following the definition in (38), signaling entropy of gene $i$ is defined as

$$S_i = -\frac{1}{\log k_i \prod_{j \in N_i} p_{ij}} \log p_{ij}$$

where $N_i$ represents the first-order neighbors of gene $i$ in the signaling network. $w_{ij}$ denotes the edge weight between genes $i$ and $j$ and is defined as $w_{ij} = \text{Expr}_i \cdot \text{Expr}_j$, where $\text{Expr}_i$ and $\text{Expr}_j$ are the normalized expression levels of gene $i$ and $j$, respectively. $p_{ij}$ is the probability of interaction between genes $i$ and $j$. Here, $k_i$ is the degree of gene $i$ in the signaling network, which serves as a normalization factor to scale the signaling entropy to the range of $[0, 1]$.

We computed signaling entropies for all genes in the predicted networks in kidney, heart, and lung at fetal and adult stages using the R package provided in (38). We then defined differential entropy for a given gene as the absolute value of the difference in signaling entropies between a pair of tissues. Note that a gene shared by all three tissues has three differential entropy values. We selected the maximum differential entropy value of that gene for determining the significance of differential entropy.

To identify genes with significantly large differential entropies, we constructed a null distribution of differential entropies based on randomized networks. To generate the randomized network for each of the three tissues at a developmental stage, we separately shuffled the edges of the two intracellular networks while preserving the node degree distribution, node prizes, and cross-talk edges as the original integrated network. We then identified 100 PCSFs from the randomized integrated networks for each tissue and computed signaling entropies of genes in these PCSFs. To generate the

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null distribution of differential entropy, we performed a total of 30,000 pair-wise comparisons of signaling entropies between all PCSFs from a pair of tissues. An empirical $P$ value of 0.1 was used as the cutoff to identify genes with significant differential entropies.

**Tissue specificity and conservation scores of genes**

Tissue specificity scores of genes are based on the Tau metric (65), which is defined as $\tau = \sum_{i=1}^{N} (1 - x_i)/(N - 1)$, where $N$ is the number of tissues and $x_i$ is the expression of gene $i$ in tissue $i$ normalized by its maximal expression across all tissues. Tau values for all genes were downloaded from (39). Conservation scores of genes were calculated on the basis of the phastCons score, which is the probability that a nucleotide belongs to a conserved element based on the multiple sequence alignment (66). We downloaded phastCons scores based on multiple alignments of 30 mammalian genomes from the University of California, Santa Cruz Genome Browser. Conservation score for a gene was computed as the average phastCons score of all exonic sequences of the gene. Both tissue specificity and conservation scores of the shared pathway genes across kidney, heart, and lung tissues are listed in table S2.

**Differential cross-talk score across tissues**

For each ligand-receptor pair, we computed three absolute differential values of cross-talk scores between the three human tissues (kidney, heart, and lung) and used the maximum of the three values as the differential cross-talk score for the ligand-receptor pair.

**Functional enrichment analysis of signaling pathways**

To investigate the functions of the signaling pathway associated with a given gene, we first extracted the 10th-order neighborhood of that gene from the predicted signaling network. Then, we performed enrichment analysis of the neighborhood using g:Profiler (67) and removed redundant Gene Ontology terms using REVIGO (68). Enrichment $P$ values were adjusted for multiple testing using the method of Benjamini-Hochberg (64).

**SUPPLEMENTARY MATERIALS**

Supplementary material for this article is available at https://advances.sciencemag.org/cgi/content/full/7/16/eabf1356/DC1

View/request a protocol for this paper from Bio-protocol.

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CytoTalk: De novo construction of signal transduction networks using single-cell transcriptomic data
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