ABSTRACT An integrative analysis focused on multi-tissue transcriptomics has not been done for asthma. Tissue-specific DEGs remain undetected in many multi-tissue analyses, which influences identification of disease-relevant pathways and potential drug candidates. Transcriptome data from 609 cases and 196 controls, generated using airway epithelium, bronchial, nasal, airway macrophages, distal lung fibroblasts, proximal lung fibroblasts, CD4+ lymphocytes, CD8+ lymphocytes from whole blood and induced sputum samples, were retrieved from Gene Expression Omnibus (GEO). Differentially regulated asthma-relevant genes identified from each sample type were used to identify (a) tissue-specific and tissue-shared asthma pathways, (b) their connection to GWAS-identified disease genes to identify candidate tissue for functional studies, (c) to select surrogate sample for invasive tissues, and finally (d) to identify potential drug candidates via connectivity map analysis. We found that inter-tissue similarity in gene expression was more pronounced at pathway-functional level than at gene level with highest similarity between bronchial epithelial cells and lung fibroblasts, and lowest between airway epithelium and whole blood samples. Although public-domain gene expression data are limited by inadequately annotated per-sample demographic and clinical information which limited the analysis, our tissue-resolved analysis clearly demonstrated relative importance of unique and shared asthma pathways. At the pathway level, IL-1b signaling and ERK signaling were significant in many tissue types, while insulin-like growth factor and TGF-beta signaling were relevant in only airway epithelial tissue. IL-12 (in macrophages) and Immunoglobulin signaling (in lymphocytes) and chemokines (in nasal epithelium) were the highest expressed pathways. Overall, the IL-1 signaling genes (inflammatory) were relevant in the airway compartment, while pro-Th2 genes including IL-13 and STAT6 were more relevant in fibroblasts, lymphocytes, macrophages and bronchial biopsies. These genes were also associated with asthma in the GWAS catalog. Support Vector Machine showed that DEGs based on macrophages and epithelial cells have the highest and lowest discriminatory accuracy, respectively. Drug (entinostat, BMS-345541) and genetic perturbagens (KLF6, BCL10, INFB1 and BAMBI) negatively connected to disease at multi-tissue level could potentially repurposed for treating asthma. Collectively, our study indicates that the DEGs, perturbagens and disease are connected differentially depending on tissue/cell types. While most of the existing literature describes asthma transcriptome data from individual sample types, the present work demonstrates the utility of multi-tissue transcriptome data. Future studies should focus on collecting transcriptomic data from multiple tissues, age and race groups, genetic background, disease subtypes and on the availability of better-annotated data in the public domain.
in disease vs. control tissues can be used to explore the connectivity between disease, gene expression and perturbagens (candidate therapeutic agents) (Lamb et al. 2006). Previous gene expression studies have utilized biological samples from bronchial and alveolar macrophages, upper and lower airway epithelial cells and peripheral whole blood or other circulating blood cells to identify genes differentially expressed in asthma compared to healthy controls (Tsitisou et al. 2012; Kicic et al. 2010; Choy et al. 2011). Although these studies produced lists of differentially expressed genes (DEGs), there tends to be little or no overlap among the lists of genes obtained using different tissues/cells suggesting disease-associated genes are likely to show tissue-specific expression patterns (Lage et al. 2008; Reverte et al. 2008). Most gene expression studies so far reported tissue combined approaches. Such an approach largely ignored the inherent gene expression heterogeneity of tissues/cells, and highly expressed genes rather than biologically relevant genes (but are too small to be detected in tissue-combined analysis) may be lost and potentially remain under-defined. Therefore, a tissue/cell-based gene expression analysis is necessary to track real driver genes and reduce potential confounders produced due to highly expressed genes from tissue/cell types. Thus, the analyses of genome-wide gene expression data originating from multiple tissue types would be helpful to understand tissue-specific or tissue-shared differentially regulated asthma genes. As the availability of genome-wide gene expression datasets has grown across major publicly data repositories, there is an increased interest to systematically mine these resources to identify novel patterns (Ramasamy et al. 2008).

In this study, we analyzed gene expression datasets available from GEO that were generated using multiple tissue/sample types (e.g., blood, lymphocytes, upper and lower airway epithelial cells, lung biopsies, fibroblasts, macrophages and induced sputum) obtained from asthma patients and matched controls in order to obtain a tissue-resolved perspective of differentially regulated asthma-relevant genes and biologic pathways. This study has potential to (i) identify biomarkers from easily accessible non-invasive samples (e.g., induced sputum) as surrogates for invasive, difficult to collect samples, (ii) connect GWAS-identified candidate genes to genome-wide gene expression results in asthma and (iii) identify potential therapeutic compounds for asthma based on tissue-specific and tissue-shared gene expression. In the current study we have used DEGs identified from different tissue/ sample types to select candidate perturbagens via Connectivity Map (CMAap), which currently covers > 1300 compounds connected with 7000 expression profiles (Wang et al. 2015). This approach can identify drugs that affect the expression of common genes and identify candidate that could be potentially repurposed for systemic management of asthma (Ravindranath et al. 2015). However, how DEGs identified from different tissue/ sample types can potentially modify the list of perturbagens identified from CMAP analysis remains to be elucidated. Since DEGs vary across asthma-relevant tissue types, a better understanding of tissue-specific/-shared DEGs underlying asthma may lead to the development novel drug candidates for to treat asthma.

**MATERIALS AND METHODS**

**Gene expression datasets description**

Data used in this study were retrieved from the publicly accessible Gene Expression Omnibus (GEO) database at the NCBI (http://www.ncbi.nlm.nih.gov/geo/) (Barrett et al. 2013; Edgar et al. 2002). We used the query terms “human [organism] AND asthma AND 2000/01:2019/06[Publication Date]” to retrieve datasets from transcriptome studies comparing samples from multiple tissues (Figure 1) asthma patients with those of healthy individuals. Each individual dataset underwent consistent handling similar to when it was uploaded to GEO by the original study groups (preserving original normalization protocols, to maintain consistency with published reports). The following information was extracted from each study: (1) GEO accession, (2) sample type, (3) platform, (4) numbers of asthmatic and control individuals (Table 1). In total, 568,930 probes covering 25,000 genes were extracted for the analysis. The following asthma transcriptome data from the sample types were considered:

**Nasal epithelial cells (NECs):** Nasal epithelial cells were obtained by nasal brushing. Six asthmatics and 6 healthy controls were used (GSE44037). NECs are of particular interest for studies in children due to their easy accessibility during clinical visits (Poole et al. 2014).

**Airway epithelium:** Airway epithelial gene expression data were obtained from GSE4302 (asthmatics= 10, controls = 10) and GSE18965 (asthmatics= 9, controls = 7). Subjects using inhaled medication or tobacco smoke have been removed from our analysis.

**Bronchial biopsy:** Data were obtained from asthma (n = 4) and control (n = 4) subjects (GSE15823) and asthma (n = 4) and control (n = 4) subjects (GSE41649). Data related to treatment outcome was not included. The samples typically contained mucosal and epithelial/ sub-epithelial tissue (0.3 mm² to 0.5 mm²) (Laprise et al. 2004; Choy et al. 2011; Chamberland et al. 2009; Labonté et al. 2008). Bronchial epithelium is a source of cytokines and chemokines which plays a key role in recruitment of inflammatory cells into tissue from the circulation.

**Proximal airway fibroblasts:** We used asthmatic (n = 8) and control (n = 4) subjects (GSE27335) involving fibroblast samples obtained by bronchoscopy. Proximal airway fibroblasts are known to synthesize more collagen and eotaxin-1 than distal fibroblasts. Fibroblasts play key roles via an IL-13 induced mechanism involving TGF-β1 and MMPs leading to and lung remodeling in asthma (Ingram et al. 2011).

**Distal airway fibroblasts:** Matched case-control distal lung fibroblast pairs were isolated from above-mentioned subjects’ distal lung from GSE27335 (asthmatics= 8, controls = 4). Distal lung fibroblasts exhibit a distinct phenotype, proliferate faster and express higher levels of α-smooth muscle actin compared to proximal airway fibroblasts. Distal airways demonstrate distinctly different fibrotic property compared to the proximal segments. Excessive deposition of extracellular matrix is particularly prominent in the proximal parts compared to proximal airways.

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Alveolar macrophages (AMs): Two GEO datasets GSE22528 and GSE2125 have been used for the present study which allowed us to utilize transcriptome data containing 10 cases and 223 controls, and 30 cases and 547 controls respectively. To generate transcriptome data, AM cells from broncho-alveolar lavage fluid of asthmatics and control subjects were used to perform homogeneity checks by flow cytometry prior to RNA isolation and microarray experiments performed by the investigators. Alveolar macrophages constitute a unique subset of pulmonary macrophages, which serve as a first line of defense against foreign invaders to the lung tissue.

Blood and lymphocytes: Dataset GSE69683, which includes asthmatics (n = 399; severe and non-severe) and non-asthmatics (n = 101) excluding smokers, were used for this analysis. GSE 31773 was also used to identify asthma associated DEGs from CD4+ and CD8+ lymphocytes. This dataset was generated using CD4+ and CD8+ T cells isolated from peripheral blood of asthmatics (N = 18) and controls (N = 8) by negative selection and magnetic cell separation procedures followed by purity assessments using flow cytometry. Blood contains plasma and corpuscles, each paying multiple roles in inflammation. Peripheral blood mononuclear cells (PBMC) consist of lymphocytes (T cells, B cells, NK cells) and monocytes. TH2 cells represent a phenotype of T lymphocytes that upon activation generate IL-4, IL-5 and IL-13 cytokines in both peripheral blood and bronchial mucosa of asthma patients, leading to local tissue eosinophilia, generation of specific IgE antibody responses and airway hyperresponsiveness/ inflammation.

Induced sputum: We obtained transcriptomic data from GSE76262 which was generated using sputum cells collected from asthmatics (n = 118) and matched healthy controls (n = 21). Induced sputum is an easily accessible non-invasive approach for obtaining cellular (epithelial cells, eosinophils, lymphocytes, neutrophils and macrophages) and non-cellular (cytokines, chemokines and other secreted products/ inflammatory mediators) components from the lower respiratory tract.

Analysis workflow
The workflow for data analysis is illustrated in Figure 2. Genome-wide transcription data obtained from the seven different tissues/cell sample types include airway epithelial, bronchial biopsy, nasal epithelial, alveolar macrophages, distal lung fibroblasts, proximal lung fibroblasts, CD4+ lymphocytes, CD8+ lymphocytes, whole blood and induced sputum. Tissue-specific and tissue-combined analyses were performed to identify differentially expressed genes. The transcriptome datasets underwent careful quality control. If a single GEO dataset contained data from multiple tissues/ sample types (e.g., GSE31773 contains CD4+ and CD8+ cells and GSE27335 contains proximate and distal macrophages), such datasets were further divided based on tissue/sample types. This generated 13 datasets, each with cases and controls, for downstream analysis (Table1). Finally, data from subjects on asthma medication that could potentially change the gene expression were also removed.

Probe level analysis
Each dataset was downloaded from GEO for further tissue-specific analysis. Differentially expressed genes were identified for each individual dataset at the probe level using limma R packages from the Bioconductor project (https://www.bioconductor.org/). Genes in each dataset were arranged by statistical significance. Among the top DEGs based on p-value, those showing a fold-change of 1.5 (up- or down-regulated) were considered for further analysis.

Gene-level overlap between tissue/cell types
The probe IDs are mapped to NCBI generated annotations, which is available from the UCSC genome browser site. Annotations such as gene symbol and gene title are derived by extracting stable sequence identification information from the platform and periodically querying against the Entrez Gene and UniGene databases to generate consistent and up-to-date annotation. Then, the transcripts showing transcript IDs that did not have a corresponding gene symbol were exported to DAVID (http://david.abcc.ncifcrf.gov/home.jsp) for gene ID conversion (Huang et al. 2009a, 2009b).

Support vector machine (SVM) model
SVM is one of the most popular supervised learning methods for analyzing data and recognizing patterns. The objective of the support vector machine algorithm is to find a hyperplane that classifies data points (potentially case vs. control). Given labeled training data (supervised learning), the algorithm generates an optimal hyperplane that can be used to categorize new datapoints. SVM therefore represents a powerful technique...
Table 1 Summary of transcriptomics datasets used for the present study. NCBI Gene Expression Omnibus (GEO) accession number, tissue/cell sample type, platform, sample size (excluding smokers and subjects using inhaled corticosteroids) have been shown for each study.

| Sample type          | GEO ID     | Tissues/cell types             | Platform   | Case | Control |
|----------------------|------------|--------------------------------|------------|------|---------|
| I                    | GSE4302    | Airway epithelial cells        | Affy HG-U133 | 10   | 10      |
| II                   | GSE18965   | Airway epithelial cells        | Affy HG-U13A | 9    | 7       |
| III                  | GSE15823   | Endobronchial biopsies         | HG_U95     | 4    | 4       |
| IV                   | GSE41649   | Endobronchial biopses          | HG-U133    | 4    | 4       |
| V                    | GSE22528   | Alveolar macrophages           | Affy HG-U133 | 4   | 4       |
| VI                   | GSE2125    | Alveolar macrophages           | Affy HG-U133 | 15  | 15      |
| VII                  | GSE27335   | Proximal airway fibroblasts    | Affy HG-U133 | 8   | 4       |
| VIII                 | GSE27335   | Distal lung fibroblasts        | Affy HG-U133 | 8   | 4       |
| IX                   | GSE31773   | CD4+ T-cells                  | Affy HG-U133 | 12  | 8       |
| X                    | GSE31773   | CD8+ T-cells                  | Affy HG-U133 | 12  | 8       |
| XI                   | GSE44037   | Nasal epithelial cells         | Affy HG-U133 | 6   | 6       |
| XII                  | GSE69683   | Peripheral Blood               | Affy HG-U133 | 411 | 84      |
| XIII                 | GSE76262   | Induced sputum (whole)         | Affy HG-U133 | 118 | 21      |

Pathway and regulatory network analysis
Differentially expressed genes by tissue/cell types between asthma patients and normal controls were further investigated using Ingenuity Pathway Analysis (IPA) software (Qiagen, USA). The IPA methodology compares proportional representation of genes from a defined test set in a canonical pathway (a known, well-characterized pathway), compared to the proportional representation of the pathway genes in the entire set of known genes. The p-value is calculated using a right-tailed Fisher exact test and indicates the likelihood of the pathway association under the random model. Pathway/networks with a score >2 have >99% confidence that the genes included in the network are not generated by chance. The score represents a numerical value to rank networks according to their level of relevance to the 'Network Eligible' molecules in the dataset considering the number of 'Network-Eligible' molecules within the network and its size, as well as the total number of 'Network-Eligible' molecules analyzed and the total number of molecules in the Ingenuity Knowledge Base that could potentially be included in networks. Finally, the networks are ordered according to their score, with the highest scoring network displayed at the top. The network Score is based on the hypergeometric distribution and is calculated with the right-tailed Fisher’s Exact Test.

Linking DEGs with asthma genes identified from GWA Studies
There were 25 asthma studies resulting 38 genomic regions from GWAS Catalog (accessed July 2019). (Hindorff et al. 2009). Inclusion of asthma GWAS catalog-based associations was limited to those studies with P values of less than 5×10⁻⁸ (http://www.ebi.ac.uk/gwas/). Asthma-relevant genes were also identified from published literature using literature lab (Accumanta, USA) (Fibbo et al. 2007). Gene level expression of asthma-relevant genes was averaged over cases or controls in each dataset and correlation coefficients were calculated across datasets, which was used to run hierarchical clustering to identify groups of tissue/cell types that cluster more closely than others.

Connectivity Map (CMap) analysis using tissue-specific DEGS
Next, we used Connectivity Map (CMap) analysis approach to explore potential drugs targeting asthma by systematically mining functional connections between asthma disease, DEGs, and perturbagens (Lamb et al. 2006). Differentially expressed genes were divided into two parts, one for upregulation and the other for downregulation. The CMap analysis was performed through the web interface CLUE (https://clue.io/), a cloud-based platform was used to analyze perturbation-driven gene expression signatures, following a standard protocol described by Wang et al. (Wang et al. 2019; Subramanian et al. 2017). Briefly, the “Query” and “Touchstone” applications of CLUE were used to identify reciprocal connectivity (negative correlation) between asthma and perturbagens where up- and down-regulated genes were used inversely as inputs to identify negative correlation (Wang et al. 2019) (Lex et al. 2014). CMap instance was measured by an enrichment score, which ranged from -1 to 1, and a permutation P-value. For the current analysis, any connectivity score of below -0.85 or above 0.85 was considered for this analysis (Subramanian et al. 2017).

Data availability
The data supporting this work is publicly available from NCBI GEO (Gene Expression Omnibus): https://www.ncbi.nlm.nih.gov/gds?
Figure 2 Major steps to identify and analyze tissue specific asthma gene expression data. Individual datasets were obtained from GEO. Differentially expressed asthma-relevant genes identified from each tissue type were used to find tissue/ cell-specific genes and networks, and discriminatory gene-sets to classify samples into normal vs. asthma classes and to predict asthma disease state. In addition, differentially expressed genes identified from each sample/ tissue type were linked with GWAS catalog data and Connectivity Map resources to identify novel drug candidates.

term=asthma. Supplemental material available at figshare: https://doi.org/10.25387/g3.11935959.

RESULTS

Tissue-combined and tissue-resolved approaches
Tissue/ cellular samples used in this study included airway epithelial cells, bronchial biopsy, alveolar macrophages, peripheral blood, purified peripheral blood CD4+ lymphocytes, CD8+ lymphocytes, proximal lung fibroblasts, distal lung fibroblasts, induced sputum and nasal epithelium samples (Table 1). In an initial tissue-aggregated assessment (tissue-combined approach), top significant transcripts showed an inability to discriminate between normal and asthma subjects (Supplementary Figure S1A) by Principal Component Analysis (PCA) indicating the importance of tissue- specific analysis in asthma. Tissue-resolved analysis approach was therefore followed subsequently. Tissue-specific differentially expressed signature genes (number ranged from 126 to 1060) were then obtained for each of the seven tissues. Significant genes showing absolute fold change values of 1.5 or higher were considered for further analysis.

When the data were analyzed tissue by tissue (tissue-resolved approach), a clear distinction between normal and asthma subjects was observed. The lowest differentially expressed gene count was observed in bronchial biopsy tissues (173 DEGs) while the highest was observed in the nasal biopsy tissue (1320 DEGs). PCA showed clear separation of asthmatics from controls in tissue specific analysis (Supplementary Figure S1 B-D). Jaccard similarity index was used to determine similarity between various tissues, nasal and airway epithelial cells show the highest (0.61) while CD4+ lymphocytes and Bronchial biopsies (0.01) showed the lowest Jaccard similarity index, respectively. Differentially expressed genes were identified from datasets representing multiple sample types consisting of tissues (e.g., airway epithelial, bronchial, nasal) and isolated cells (e.g., CD4+ lymphocytes, CD8+ lymphocytes, macrophages, isolated fibroblasts). Figure 3A and 3B demonstrate the overlap of DEGs identified from tissue and cellular samples, respectively. Differentially expressed genes identified from different tissue types were compared to obtain tissue-specific and tissue-shared genes and pathways relevant for asthma as described below:

Airway epithelial samples: In airway epithelial tissue, increased expression of the chloride channel calcium-activated, family member 1 (CLCA1) (6.2 fold), peristin (4.4 fold), and serine peptidase inhibitor clade B member 2 (SerpinB2) (also known as plasminogen activator inhibitor-2) (3.5 fold) was observed. Originally described as a calcium-activated chloride channel, CLCA1 likely functions indirectly in chloride transport and has previously been reported to be up-regulated in asthma. Peristin is an integrin ligand and extracellular matrix protein with roles in cell adhesion, cell motility, and matrix remodeling. SerpinB2 is a member of the serpin class of proteases and functions to inhibit plasminogen activation and to promote fibrin formation and deposition. Among the other most differentially expressed genes in lung airway epithelial brushings were three mast cell markers: carboxypeptidase A3 (3.4 fold induced), tryptase /b (2.2 fold induced), and tryptase a/B (2.1 fold induced).

The transforming growth factor beta (TGF-β) signaling pathway seems to play a major role in the manifestation of asthma, most notably in the airway remodeling process. TGF-β stimulates fibroblasts to synthesize and secrete proteins of the extracellular matrix in patients with severe asthma may induce smooth-muscle hypertrophy (Ohno et al. 1996; Cohen et al. 2000). In addition to the TGF-β pathway, extracellular signal regulated kinase (ERK) plays a major role in airway smooth muscle activity by virtue of its upstream activation of cyclin D1 promoter activity which is required for DNA synthesis in airway smooth muscle cells (Ramakrishnan et al. 1998; Zhou and Hershenson 2003). Consistent with this observation that ERK is critical for signaling of airway smooth muscle cell activity in...
asthma, bronchoalveolar lavage fluid from asthma patients has been shown to augment ERK activation, increase cyclin D1 protein abundance, DNA synthesis, and proliferation of cultured human airway smooth muscle cells (Naureckas et al. 1999). One very significant finding is the upregulation of the estrogen pathway in the airway epithelium of asthma patients. Estrogen receptors are found on numerous immunoregulatory cells and estrogen’s actions skew immune responses toward allergy. The role of estrogen receptor in asthma is being increasingly recognized. For example, the G-protein-coupled estrogen receptor agonist has been found to suppress airway inflammation through IL-10 in a mouse model of asthma.

Lymphocytes: DEGs unique to CD4+ and CD8+ cells as well as common DEGs were analyzed. The top up-regulated genes were related to bacterial resistance, solute transport and calcium signaling. While asthma involved pathways like NFAT in regulation of the immune response and other lymphocyte activation pathways including signaling through IL-4, IL-8, IL-3 and IL-9 in CD8+ cells, the most prominent pathway in CD4+ cells was VDR/RXR activation signaling. Activation of ERK pathway was also noted to be the top common pathway in lymphocytes, followed by Immunoglobulin synthesis and inflammatory cytokine synthesis pathways.

Macrophages: The two datasets generated using bronchial macrophage cells, showed that ERK was the main regulatory network. In an individual study, Madore et al. identified differentially expressed asthma genes related to stress and immune responses using airway macrophage cells obtained from asthma and healthy controls (Madore et al. 2010).

Fibroblasts: The fibroblast transcripts focused on the differences in gene expression between distal and proximal fibroblasts. Distal Lung Fibroblasts exhibited a higher basal activation of SMAD3 and MAPK8 compared to its proximal counterpart.

Nasal epithelial cells: Nasal epithelial cell samples revealed CCL chemokines as the most biologically relevant DEG candidates exhibiting higher expression in asthma patients and were associated with increased eosinophil and monocyte chemotaxis. In addition to prominent eosinophilic inflammation, increased IL1RL1 expression was also found.

Tissue-shared DEGs, the multi-tissue factors: The ATXN2 gene was significantly upregulated in airway epithelial brushing, bronchial biopsies and proximal as well as in distal lung fibroblasts. Interestingly, a variant of this gene was recently found to be associated with compromised lung function (Wain et al. 2017; Soler Artigas et al. 2015). In addition, upregulation of the integrin ITGB1 at the mRNA level has been demonstrated in airway smooth muscle cells in the asthma model group (Álvarez-Santos et al. 2016). The integrins are important cell surface adhesion receptors that recognize extracellular matrix components known to be altered in airway cells in asthma. Intrinsic differences specifically related to proliferation, differentiation, and migration contribute to the dysregulated bronchial epithelial cell response to injury.

Functional enrichment, pathway-level overlap and networks
In order to gain further insight into the functional significance of tissue-specific and/or shared DEGs, a gene set enrichment analysis based on gene ontology annotations was performed. Immune system process and cell adhesion were the GO terms that were overrepresented (8 genes each) within our list of DEGs. Interestingly, out of the 8 genes representing an immune system process, only 4 (ALOXA15, ITGB2, CD44 and EDNRA) had previously been associated with asthma. The differentially expressed genes between asthma and controls in each tissue type were used to identify gene networks by the IPA software application. Two gene networks were identified with a network score of >2. The highest scoring network
had a network score of 21 and was associated with the network functions of cellular movement, immune cell trafficking and inflammatory response. Figure 4A and 4B demonstrate the overlap of enriched pathways in tissue and isolated cellular samples in asthma.

Pathway analysis identified autophagy, VEGF (vascular endothelial growth factor) Signaling, Paxillin Signaling, Actin Cytoskeleton Signaling and ERK5 Signaling as the top canonical signaling pathways in lower airway epithelial samples. Interestingly, VEGF Signaling, has been designated as a very important pathway in asthma by inducing remodeling and enhancing TH2-mediated sensitization and inflammation in the lung (Lee et al. 2004). Paxillin and actin cytoskeletal signaling have been found to be important for bronchial tissue contraction and airway remodeling in asthma. However, thus far, the role of the Pax gene (encoding paxillin protein) in asthma has not been sufficiently explored. Analysis of the lymphocyte data using IPA pathway analysis tool indicated that activation of multiple pathways related to T-cell activation, inflammatory cytokine and immunoglobulin synthesis were important. In lymphocytes, IgE synthesis, cytokine signaling, and ERK-mediated cell proliferation have been found to be the major driving factors for asthma. For macrophages, pathway and network analysis identified IL-12 and ERK as the major hub genes or nodal genes indicating the role of sustained inflammation via the IL12 pathway and control of cell proliferation via the ERK pathway. For fibroblast samples, the up-regulation of extracellular matrix-associated molecules, actin binding and cytoskeletal protein molecules has been previously reported in proximal and distal lung fibroblasts respectively. Genome-wide gene expression differences between these two populations of regional lung fibroblasts might explain different responses to cell injury, cell regeneration, and subsequent airway remodeling in the lung. We noticed that top significant asthma-relevant networks are distinctly different between airway macrophages, proximal lung fibroblasts and distal lung fibroblasts of asthmatics indicating that although all of these samples were obtained from airway compartment they demonstrate different asthma-relevant genes and networks (Supplementary Figure S2).

Interestingly, although epithelial gene expression patterns differ between nasal and bronchial samples, they exhibit significant similarities at the functional level. For example, the genes CTSC, ELF1L2, IL13RA2, IL1R2, IRX4 had higher expression in nasal compared to bronchial epithelium, but both demonstrated activated functional clusters important for cell-mediated immune responses, hematological system development and function as well as immune cell trafficking.

We also compared the pathway-level overlap between samples to assess whether easily accessible sample types (such as nasal epithelium and induced sputum) could be used as surrogates for more invasive sample types (such as airway epithelial brushing and bronchial biopsy sample). Airway epithelium can be partially represented by the nasal samples, rather than from induced sputum which might partially represent bronchial samples at the pathway-level indicating the significance of easily accessible samples as surrogates for less accessible clinical samples (Supplementary Figure S3). Therefore, the nasal epithelial cells are easier to obtain and may act as a surrogate for airway epithelium (calculated Jaccard coefficient = 0.43) in asthma studies as previously indicated (McDougall et al. 2008).

**Linking tissue-specific DEGs with asthma GWA studies**

To further assess the inter-tissue expression differences of asthma-relevant genes we analyzed the expression of asthma-associated genes identified from GWAS and from published asthma literature. GWA studies have identified many genetic variants associated with complex diseases like asthma. However, gene regulatory architectures differ between tissues, and the enrichments for regulatory variants among GWAS SNPs are more pronounced in disease-relevant tissues (Roadmap Epigenomics Consortium et al. 2015). Thus the relevance of significant GWAS loci in altering biological processes (such as up/down regulated genes, biological process, functions and networks) that result in risk for or protection from the disease remains to be a key question, which requires access to the disease tissues of interest (Koester and Insel 2016). Immunoglobulin genes and IL-1 signaling pathways, which are key regulatory pathways in asthma,
Predicting asthma risk using easily accessible surrogate clinical samples

Next, we attempted to ascertain whether more accessible clinical sample types (such as peripheral blood, airway macrophage) could be used to discriminate asthma from controls thereby potentially useful for predicting asthma. Prediction analysis using SVM was used to determine a minimal set of genes that could best discriminate asthmatics from controls. Representative asthma signature for peripheral blood, Airway Macrophage and Induced sputum that could discriminate asthma from control with > 80% predictive accuracy have been shown in Table 2. Interestingly, in our SVM model, the expression of only a set of 10 genes could discriminate cases from controls with 96% predictive accuracy.

Utilizing tissue-specific DEGs to identify candidate ASTHMA therapeutics

Asthma has been recognized as a systemic disease consisting of networks between various tissues showing inflammatory changes involving a broad spectrum of structural cells, adaptive and innate immune cells. Therefore, the DEGs identified from each tissue were used to “connect” asthma with candidate therapeutics using computational drug repurposing approach (Subramanian et al. 2017). For this, tissue-specific asthma-associated gene expression signatures were matched to previously characterized perturbagens that may potentially reverse the signature in multiple cell/tissue sample types. Table 3 shows perturbagens associated with three or more asthma tissue/sample types, occurrence (number of connected asthma tissues), and their association with allergy/asthma/lung diseases (if known). Perturbagens can be “genetic” (gene knock-down and over-expression; 20 candidates) or “pharmacologic compound” (e.g., entinostat, BMS-345541, calyculin, importazole and topotecan; 5 candidates), while pharmacologic compound could be sub-categorized into biologic or chemical drugs. The connectivity scores of perturbagens are presented in supplement S1.

We found that no L1000-characterized perturbagen is shared by all asthma tissue sample types. The candidate perturbagen shared by most sample types is KLF6 (5 sample types), followed by BCL10, HOXB13, IFNB1 and Entinostat (4 sample types each). While 20 other perturbagens (ATOX1, BAMBI, BMS-345541, CCNL1, CDCA8, DHX8, DTX2, KLF3, LASP1, LOXL1, PPP2R3C, PREB, PUF60, SORBS3, TRIP10, XPO7, YWHAZ, calyculin, importazole, topotecan) are associated with three different sample types, large numbers of perturbagens are connected to either two (171 perturbagens) or only one (1126 perturbagens) sample type. Several top-ranking perturbagens have been indicated by previous studies to be involved in allergy/asthma or other lung diseases (i.e., Chronic Obstructive Pulmonary Disease; COPD). Figure 6 depicts unique and shared perturbagens connected to asthma status in different asthma sample types. The blue connected dots (6A) indicate shared candidates between sample types (gray tracks) with the asterisk showing perturbagen connecting maximum samples, while the bar chart (6B) indicates the number of unique perturbagens for each sample type. Finally, figure 6C represents a clustered heatmap generated using the connectivity scores (shown in supplementary table s1) of perturbagens (25 significant perturbagens listed in table-3) depicting the connections between disease, candidate perturbagens and tissue-based gene signatures of asthma. Connectivity score values of perturbagens were generated by touchstone application of CLUE (www.clue.io). Further details about their characteristics have been listed in Supplementary S2. Analysis of functional significance of the perturbagens showed their roles in immune function, cellular transport function, apoptosis and inflammation, with several agents showing one or more overlapping functions. The results clearly demonstrated that the connectivity scores of perturbagens can vary depending on input source tissue used to identify asthma-relevant DEGs. For example, the genetic perturbagen can be relevant for distal fibroblasts, CD+ lymphocyte and nasal epithelium, but may not be relevant for proximal fibroblasts and CD4+ lymphocytes. Similarly, Macrophages, the chemical perturbagen entinostat may be relevant.
Table 2  SVM-generated classifiers (Asthma Gene Expression signatures) for sample types (blood, Macrophage, induced sputum) that are widely used to study the pathology of asthma. Percent predictive accuracy for each classifier has also been indicated

| Tissue Type                  | Asthma Gene Signature (for >80% predictive accuracy)                                                                 | % correct prediction |
|------------------------------|-------------------------------------------------------------------------------------------------------------------------|----------------------|
| Blood (GSE69683)             | MYD88, MYL12A, BOLA2, HBXIP, ACSL1, VPS24, GPM6B, F5, PDE7A, AB11FIP3, ZNF785, TCF4, RAB11A, GPR109B, GNAQ, DIS3L2, SLCT7ASP1/P2, TSR2, AMICA1, CR1, QRIC1, SERBP1, ZNF37A, DERL2, ZBTB20, CDC42SE1, NSMAF, KIAA0562, NT5C2, DAPPP1, ATP6V0E1, LOC203274, TMEM59, NAPEPLD, ANXA3, WIP2, ATAD5, LAR5, PTBP1, SPG7, DNAJC16, IL6ST, CLN8, SAT1, EVI2B, MRPS5, AQP9, RPP14, ZNF747, GAPT | 83                   |
| Macrophage Cells (GSE2125)  | KBTBD2, UPF3A, MLLT10, P5PC1, ST3GAL1, ARFRP1, TRIM14, EIF4G1, BM25, CDK11A/B, IL18R1, TLR7, GPR85, LIMA1, SPINT1, EIF1AY, PDCD6, MGAT4A, CSTA, LOC100129845 | 83                   |
| Induced Sputum (GSE76262)   | IL18R1, TLR7, GPR85, LIMA1, SPINT1, EIF1AY, PDCD6, MGAT4A, CSTA, LOC100129845 | 96                   |

for distal fibroblasts, CD8+ lymphocytes and nasal epithelium, but may actually have an opposing effect based on sputum and whole blood asthma signatures. Collectively our results indicate a critical role of tissue sample (used in generating gene expression data) in identifying connected drug candidates.

**DISCUSSION**

Tissue-specific gene expression plays a fundamental role in multi-cellular pathobiology (Song et al. 2013). However, until recently direct comparative analysis across multiple tissue types was not possible due to the limited number of gene expression data available from public domain. Herein, we studied tissue-specific DEGs in multiple datasets originating from independent studies using multiple asthma tissue/ cell sample types (bronchial epithelium, peripheral blood, CD4+ and CD8+ lymphocytes, nasal epithelial, induced sputum, airway fibroblast and airway macrophages). To our knowledge this is the first study to identify and compare tissue-specific asthma-relevant genes and pathways from publicly available gene expression data and to utilize them for better asthma management.

**Multi-tissue transcriptomic analysis can identify appropriate surrogate samples for invasive/ difficult to collect samples**

Although it is ideal to determine gene expression from target tissues (i.e., cells from lung tissue in asthma), this is challenging when considering the large number of samples required given the demands of statistical power. However, recently studies have demonstrated the value of focusing on surrogate target tissues/cells in predicting gene expression in tissues/cells that are challenging to access in large numbers (i.e., lung tissue), which have the potential to significantly move the field forward. For example, Poole and colleagues used whole-transcriptome sequencing (RNA-Seq) to demonstrate that the nasal airway epithelium mirrors the bronchial airway (Marenholz et al. 2006). Their study confirmed that children with asthma have an altered nasal airway transcriptome compared with healthy controls, and these changes are reflected by differential expression in the bronchial airway. Noninvasively obtainable biological samples showing considerable pathway-level overlap with difficult to obtain disease-relevant tissues could be used as surrogates for assessing disease risk and progression. For example, in experimental data indicated that nasal epithelial cells are easier to obtain and may act as a surrogate for bronchial epithelium in asthma studies (Thavagnetnam et al. 2014; McDougall et al. 2008). The accurate prediction of tissue-specific gene expression could thus provide useful information for biomarker development and drug targeting. Since nasal epithelial tissue and induced sputum are readily accessible from patients and controls, this study does suggest using these samples as a model system for asthma, as there is more pathway-level similarity to the upper airway and bronchial samples, respectively.

**Genes and networks interact across tissues and show connection to Canonical Pathways showing the hallmarks of asthma**

A striking finding of our study is that the top significant DEGs do not significantly overlap between tissues, rather participate in complex pathways and networks crucial for asthma manifestation. Several of these genes are involved in cellular processes such as cell migration, airway remodeling and mucus production that are crucial for asthma manifestation. For examples, significant DEGs in our list are involved biomarkers of Th2 response (POSTN), inflammation (NOS2, ALOX15) and mucus production (MUC5AC) corroborated with a previous metanalysis of airway gene expression in asthma.(Liu et al. 2018)

Pathway-level overlap between multi-tissue gene transcription data shows that ‘leukocyte adhesion and diapedesis’ and ‘Th1/Th2 activation’ are top significantly enriched pathways in all samples – both from the blood and the airway compartments (Figure 4a and 4b). The roles of these pathways in the installation (recruitment of inflammatory cells from blood to airways) and persistence of inflammation is well known. Activation of IL-1beta, IL-7 and ERK signaling pathways were common across a wide range of tissue types maintaining inflammation. Signaling via Calcium-sensitive chloride channels (CLCA1) plays critical part in mucus production and hyperresponsiveness in the airways. Our study also identified Acute Phase Response Signaling pathway and Nitric Oxide and Reactive Oxygen Species (ROS) pathway activation in nasal, airway epithelial, bronchial biopsy samples that might contribute to hyperresponsiveness, while chronic inflammation as well as pro-fibrotic factors (TGF-beta signaling pathway and factors release by infiltrating eosinophils) lead to sub-epithelial fibrosis with changes of the extracellular matrix composition, smooth muscle growth, Goblet cell hyperplasia and mucus secretion, collectively known as airway remodeling. Taken together, pathway-level overlap distinctly reflects molecular features of Hyperresponsiveness, inflammation and remodeling – three hallmarks of asthma.

**Multi-tissue transcriptomic analysis can be useful for identifying candidate tissues for functional validation of GWAS-identified genes**

Since GWAS identify loci rather than functional variants, most GWAS have provided limited insights into underlying mechanisms. (Gallagher and Chen-Plotkin 2018) Genome-wide association studies have nominated many genetic variants for common human traits, including diseases, but in many cases the underlying biological reason for a trait association is unknown.(Gallagher and Chen-Plotkin 2018;
The current study utilized connectivity map approach to identify the expression of asthma-relevant genes at the multi-tissue level. The overlap between GWAS-identified genes and tissue-resident memory lymphocytes in asthma (Vieira Braga et al. 2019). Therefore, their functional modifiers could modify/reverse asthma signature at the multi-tissue level.

Perturbagens associated with up to 4 tissue types: Three genetic perturbagens (HOXB13, BCL10 and IFNB1) and one drug compound perturbagen (entinostat; associated with Macrophages, FibroDistal, CD8+, Nasal) are associated with four tissue types (Table 3). While BCL-10 can modulate T cell proliferation, exogenous IFN-β might confer resistance to rhinoviral infection in human mast cell rhinoviral infection model (Akoto et al. 2017). Both BCL-10 and IFNB1 are known to be associated with asthma status (Ramakrishnan et al. 2019; Klemm et al. 2006; Bhakta et al. 2018). Among the perturbagens associated with three different tissue types, we found four chemical drugs and 16 genetic perturbagen candidates that negatively correlated with asthma status, several of these are previously known for their involvement in asthma. Major functional groups include: the agents with anti-inflammatory function such as BMS-345541 (marketed by Merck KGaA), a cell-permeable quinoxaline which primarily binds IKK-2 and blocks NF-κB function, and dampsens airway inflammation.
inflammation and remodeling in mice model (Zhu et al. 2018). BAMBI (Bone Morphogenetic Protein and Activin Membrane-bound Inhibitor) has been described as a TGF-β type I pseudoreceptor, which regulates Treg/TH 17 differentiation (Onichtchouk et al. 1999). BAMBI knock out mice model of asthma demonstrated significantly reduced airway hyperresponsiveness, pulmonary inflammation, broncho-alveolar lavage fluid (BALF) eosinophil count, as well as diminished IL-4, IL-5, IL-13 and IL-6. In vitro data suggested that this effect is mediated by a potential regulatory role for BAMBI in TGF-β driven Treg differentiation (Vock et al. 2017). BAMBI also regulates macrophages inducing the differentiation of Treg through the TGF-β1 pathway (Vock et al. 2015). In addition to asthma, the enhanced plasma BAMBI level in COPD positively correlated with the increased plasma TGF-β1 levels and peripheral Th17/Treg ratio indicating the significance of TGF-beta/BAMBI pathway in pulmonary disease (Zhang et al. 2016). LASP1, a protein-coding gene that regulates cytoskeletal activity, is downregulated in sputum proteomics of asthmatics (Takahashi et al. 2018). The Role of lysyl oxidases in pathogenesis of pulmonary emphysema Lysyl oxidases regulate fibrillar collagen remodeling in idiopathic pulmonary fibrosis(Tjin et al. 2017) The copper-dependent lysyl oxidases play a role in the formation and accumulation of elastic fibers in the extracellular matrix and augments Pulmonary emphysema in COPD (Besiktepe et al. 2017). Increased fibrosis and remodeling potentially via peristin-mediated pathway. Most of the perturbagens identified dsRNA-induced changes in gene expression profiles of primary nasal and bronchial epithelial cells from patients with asthma, rhinitis and controls (Wagener et al. 2014). These genetic and chemical perturbagens could be regarded as novel drug candidates.

Finally, it is important to note that our analyses, and hence interpretations, are subject to limitations associated with the data available in the public domain. First, multi-origin transcriptome data from independent investigators that could not be adjusted for age, race, sex and asthma severity were used. We, therefore, could not adjust all confounders. However, to minimize heterogeneity, the datasets were analyzed individually to obtain up- and down-regulated genes. Furthermore, the following transcriptome data were removed: identifiable samples related to inhaled medication and tobacco use, nasal polyps (in case of nasal dataset), animal cells, cultured cell lines. Second, although considered as a state-of-the-art drug repurposing tool, the drug perturbation gene expression profiles in connectivity maps are derived from cancer cell lines, while the asthma transcriptomic data are in vivo data from human patients. This could contribute to noise in the connectivity results which should be subjected to further validation in addition to acquiring multi-tissue gene expression data from systemic asthma drugs including biologics (e.g., Omalizumab). In spite of these limitations, the availability of multiple data sets from public resources allowed us to explore DEGs from various tissues to identify asthma-relevant genes specific and shared tissue/cell types. Our analysis showed that tissue-specific, but not tissue-combined analysis can resolve DEGs in principal component analysis. Therefore, our transcriptome analysis was performed in a tissue-specific manner without combining gene expression data from multiple tissues. This approach was helpful to identify asthma signatures from each tissue/ sample type and to use for potential drug repurposing. The results from this study highlight the importance of tissue specific analysis of gene expression data and that this tissue/ cell-type based approach could be used to treat the disease at tissue specific or at systemic/multi-tissue level.

CONCLUSIONS

Our study utilized public domain gene expression data to capture an integrated multi-tissue snapshot of asthma-relevant genes and pathways. This is, to our knowledge, the first report utilizing multi-tissue transcriptomics for drug repurposing to manage asthma as depicted in Figure 7. The results indicated that (1) Asthma, although primarily affects the lungs, can be associated with tissue-specific DEGs and signatures as seen in other systemic diseases; (2) Major significant signature genes identified from each tissue/ sample type have previously-documented connections with asthma; (3) DEGs rarely overlap between tissues, but interact at the pathway level to manifest asthma. Top significant networks show distinct functional roles including bronchial hyperresponsiveness, inflammation and remodeling – three major hallmarks of asthma. Since asthma transcriptome signature depends on tissue (source of RNA), it is important to mention source tissue while presenting a gene signature; (4) Our study represents a model of tissue-resolved asthma transcriptome which can be used for (i) mapping asthma-relevant genes/ pathways in tissues and their functional interactions (ii) to find appropriate surrogate samples for difficult to obtain tissues (iii) to select appropriate
tissue for functional studies of GWAS-identified genes (iv) to use in rational drug development such as connectivity map analysis; (5) Although major networks show their functional relevance to asthma, limited per-sample demographic/clinical data are currently available in the public domain. Therefore, future studies should focus on collecting transcriptomic data from multiple sample types, age and race groups, genetic background, disease subtypes and better annotated data should be made available in the public domain. Taken together, while previous studies have used different sample types to identify asthma-relevant genes and pathways, the present work has demonstrated the utility of multi-tissue gene expression data as parts of a puzzle to obtain an integrated overview of asthma.

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