Research Article

Interferon characterization associates with asthma and is a potential biomarker of predictive diagnosis

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Interferon (IFN) plays a role in immune and inflammation responses. However, the effect of IFN in asthma is still not fully clear. The present study was conducted to better understand the role of IFN signatures in asthma. Blood samples from case–control studies (study 1: 348 asthmas and 39 normal controls and validation study 2: 411 asthmas and 87 normal controls) were enrolled. The single-sample gene set enrichment analysis (ssGSEA) method was used to quantify the levels of 74 IFN signatures. Gene Ontology analysis and pathway function analysis were performed for functional analysis and a protein–protein interaction (PPI) network was constructed. The area under the curve (AUC) value was used to evaluate the diagnostic ability. In our work, IFN-γ response-DN, negative regulation of IFN-γ secretion, IFNG pathway, negative regulation of response to IFN-γ, and type 1 IFN biosynthetic process showed higher levels in asthma. Functional analysis demonstrated that pathway and biological process involved in IFN signaling pathway, regulation of type 1 IFN production and response to IFN-γ. Hub IFN-related genes were identified, and their combination as biomarker exhibited a good diagnostic capacity for asthma (AUC = 0.832). These findings offered more insight into the underlying mechanism of how IFN signatures affected asthma. The use of the easy-to-apply IFN-related genes might serve as a promising blood-based biomarker for early diagnosis of asthma.

Introduction

Asthma, a major global public health issue, is a frequent chronic respiratory disease. This disease is characterized by variable and recurring airflow obstruction, airway inflammation, and bronchial hyperresponsiveness [1,2]. Asthma affects approximately 4.3% of the global population resulting in an increased burden on families and healthcare systems. The prevalence of asthma is increasing [3,4]. Asthma is considered as a multifactorial and heterogeneous disorder that is mainly influenced by genetic and environmental risk factors [5,6]. However, the exact and underlying mechanisms of asthma are still largely uncertain. In addition, it is not always easily diagnosed [7]. Therefore, it is necessary to further explore the potential mechanisms and find useful biomarkers of diagnosis for asthma.

The recognition that asthma is primarily associated with IgE production-related T helper type 2 (Th2) and recruitment of mast cells and eosinophils [8]. In recent years, studies suggest that T helper type 1 (Th1) cytokines are responsible for the pathogenesis of asthma. Th1 responses result in the production of interferon (IFN)-γ [9,10]. Toll-like receptor 9 (TLR9) activation is involved in AHR and airway inflammation. IFN-γ is essential for TLR9-driven suppression in asthma [11]. IFN signaling has a role in the regulation of peripheral T-cell functions [12]. IFN stimulates the expression of the endoplasmic reticulum stress-related genes and inflammation and may serve a new target for the treatment of asthma [13]. However, the role and underlying molecular regulatory of IFN signature remain largely unclear in asthma.
This comprehensive study was first conducted to determine the role of 74 IFN signatures in blood specimens of patients with asthma. In addition, we also explored IFN-related hub genes and their biological mechanisms, and we explored whether IFN could be a novel and useful biomarker in diagnosing asthma. This work could provide a more effective treatment approach and diagnostic test for asthma.

**Materials and methods**

**Study 1**

Expression profiling data from the microarray datasets for asthma and full-disease annotation were performed from Gene Expression Omnibus. The corresponding expression datasets with >50 participants in blood samples were included in the present work. In addition, obese asthma subjects were removed from further evaluation. Non-obese healthy subjects were used as a control group. Whole blood samples for microarray data from Affymetrix, we obtained GSE110551 and GSE137394 datasets. They were performed from the same platform GPL570 [HG-U133_Plus_2] Affymetrix Human Genome U133 Plus 2.0 Array. A robust multiarray averaging (RMA) method was adopted to normalize and process data. Moreover, the expression levels of all available genes were carried out by the log2 transformation for the quantification scale. When a gene symbol was mapped to multiple probes, the expression values of those probe sets were averaged. We integrated these two datasets to improve credibility and increase sample sizes because they were from the same platform and normalized method. Batch effects from these two datasets were corrected using the 'ComBat' algorithm of the 'sva' package [1,14]. Finally, we identified 348 cases with asthma and 39 healthy control subjects, which was considered as study 1 (discovery cohort).

**Study 2**

Among a very large study population \( (n=498) \), we collected the expression data GSE69683 using blood samples. This dataset was from the GPL13158 platform [HT_HG-U133_Plus_PM] Affymetrix HT_HG-U133+ PM Array Plate. No obese participants were observed in this study cohort. This expression dataset from asthma patients and healthy control population was normalized and processed using the RMA method. Then, we underwent the log2 transformation for data. When multiple probes mapped to the same gene, we used the mean value of those probe sets for the corresponding gene. Eventually, a total of 411 cases with asthma and 87 healthy control subjects were identified in study 2 (validation cohort).

**The levels of IFN**

IFN-related signatures were collected from The Molecular Signatures Database (https://www.gsea-msigdb.org/gsea/msigdb/index.jsp). We obtained 74 IFN signatures such as IFN signaling, IFN-\( \gamma \) pathway, IFN-\( \alpha \) response, IFN-\( \gamma \) response, IFN-\( \beta \) response, production of type I IFN, IRF3-mediated activation of type I IFN, IFN-\( \nu \) and IFN-\( \beta \) secretion etc. (Supplementary Table S1). To quantify the levels of IFN signatures, we used the single-sample gene set enrichment analysis (ssGSEA) method [15,16]. The enrichment levels calculated by ssGSEA method were applied to represent the relative abundance of each IFN in each sample.

**Differentially expressed IFN genes and protein–protein interaction network**

IFN-related genes were summarized from IFN signatures (Supplementary Table S2). To identify differentially expressed IFN-related genes, we performed the Wilcoxon’s test between the asthma group and the normal control group. The criteria for determining significant genes were set as adjusted \( P \)-value <0.05. Metascape (http://metascape.org/) is an effective tool to provide a systematic analysis resource for functional enrichment, interactome analysis, gene annotation, and membership search [17]. We uploaded significant IFN-related genes as the enrichment background for pathway analysis and biological process enrichment analysis. The enrichment terms with a \( P \)-value <0.01 was used as statistical significance. In addition, the protein–protein interaction (PPI) analysis was also carried out from Metascape. The network consisted of the subset of proteins that formed the interactive correlation with at least one other member. If this network included between 3 and 500 proteins, we utilized the Molecular Complex Detection (MCODE) algorithm [17,18] to identify densely connected networks and obtain the best-scoring components. According to the degree of connectivity, the PPI network identified hub genes. The PPI network was also visualized and modified using the Cytoscape software.
Figure 1. IFN signatures in asthma and control groups in study 1

(A) IFNG pathway; (B) IFN-γ response-DN; (C) IFN-β biosynthetic process; (D) negative regulation of IFN-γ secretion; (E) negative regulation of response to IFN-γ; (F) positive regulation of IFNα production; (G) type 1 IFN biosynthetic process; (H) LRR FLII interacting protein-1 LRRFIP1 activates type I IFN production; (I) regulation of IFNA signaling; (J) response to IFNG-DN.
Statistical analysis
To conduct difference comparisons of IFN signatures and IFN-related genes between asthma and normal control groups, Wilcoxon’s signed-rank test was applied. Receiver operating characteristic (ROC) curve is a common tool used to evaluate the discrimination accuracy of a diagnostic test [19], ROC analyses of hub genes were plotted in the current work. The area under the curve (AUC) values were quantified to determine the diagnostic accuracy to distinguish asthma and healthy control subjects using the pROC R package. All statistical P-values were two-sided, \( P < 0.05 \) was considered statistically significant. All data processing was performed using R software (version: 3.5.3; R Foundation for Statistical Computing, Vienna, Austria).

Results
IFN signatures in asthma
Blood samples were available for 348 asthma cases and 39 non-asthma healthy controls in study 1, which was used as the discovery set. Similarly, blood samples with larger study participants from 411 asthmatics and 87 healthy control subjects in study 2, which was applied for validation. We first summarized and evaluated 74 components of IFN signatures in asthma. Data from the discovery data demonstrated that a higher proportion of IFN-\( \gamma \) response-DN, IFN-\( \beta \) biosynthetic process, regulation of IFNA signaling, negative regulation of IFN-\( \gamma \) secretion, IFNG pathway, response to IFNG-DN, negative regulation of response to IFN-\( \gamma \), type 1 IFN biosynthetic process, and positive regulation of IFN\( \alpha \) production etc. was observed in asthma group than in healthy control group (all \( P \)-values <0.05) (Figure 1). We further validated these results in study 2 and found that five of these IFN signatures such as IFN-\( \gamma \) response-DN, negative regulation of IFN-\( \gamma \) secretion, IFNG pathway, negative regulation of response to IFN-\( \gamma \) and type 1 IFN biosynthetic process were consistent and significantly higher in asthma (all \( P \)-values <0.05) (Figure 2).
Differentially expressed IFN-related genes and function analysis

Using the Wilcoxon's test, IFN-related genes from IFN signatures were analyzed in asthma and normal control groups. Finally, 322 significant genes with an adjusted P-value < 0.05 were found in the study. We further performed a functional analysis for significant genes using Metascape. We confirmed that pathway and biological process analyses were mainly involved in regulation of innate immune response, IFN signaling pathway, regulation of type 1 IFN production and response to IFN-γ (Figure 3A).
Figure 4. Expression of 13 hub IFN-related genes showing lower in asthma than healthy control group in blood samples
Figure 5. Expression of eight hub IFN-related genes showing higher in asthma than healthy control group in blood samples

Hub IFN-related genes from PPI networks
The PPI network contained the interactions with at least one other member for genes. According to the MCODE algorithm, we selected the densest network and demonstrated that the hub genes (Figure 3B), such as CDK1, TUBB, RFC2, HSP90AB1, HSPA9, UBA52, GCH1, NUP85, PPP2CA, XRCC6, MAP3K1, EIF4E2, KPNA5, SEH1L, RANBP2, NUP98, NUP107, NUP214, GAPDH, NUP188, and NUP93.

Expression of hub IFN-related genes and diagnostic capacity
Figures 4 and 5 showed the expression level of the above 21 IFN-related genes. The results demonstrated that GCH1, HSP90AB1, HSPA9, KPNA5, NUP85, NUP93, NUP107, NUP188, RANBP2, SEH1L, TUBB, UBA52, and XRCC6 had lower expression levels in asthma, the remaining eight genes CDK1, EIF4E2, GAPDH, MAP3K1, NUP98, NUP214, PPP2CA, and RFC2 had higher expression levels in asthma. Additionally, we conducted ROC curve analyses to estimate the capability of these IFN-related genes for distinguishing between asthma and control blood samples. AUC values with >0.5 for these IFN-related genes were observed, and the combination of them showed the highest AUC value with 0.832 (Figure 6). Moreover, we used brochoalveolar lavage data (GSE67940: 73 asthmas and 31 normal
controls) and further observed that the combination of them also had the highest AUC value (AUC = 0.864) (Supplementary Figure S1A). These suggested that a pooled biomarker could improve the predictive accuracy for diagnosis in asthma.

**Discussion**

In the world, asthma affects a large population (>300 million). Although the treatment and management strategies have some advancements, this disease has no cure. The rate of death of this disease remains a crucial global challenge [7]. The precision medicine strategy for the diagnosis and management of asthma is still necessary [20]. Evidence suggests that Th2 cytokines are involved in the pathogenesis of asthma, currently, IL-5 and IL-4 antagonists also show promising efficacy. When considering TLR9 in asthmatic treatment, Th1 pathway activation indirectly/directly balances the overactivation of the Th2 pathway [21–23]. IFN therapy represents a new opportunity and an appropriate approach to treat asthma [13]. The role of IFN signaling is investigated in inflammation and asthma [12,13]. However,
the effect of IFN signaling in asthma is still not largely known. In the current work, we carried out a comprehensive analysis to explore the pattern of IFN signatures and their potential biological mechanisms in asthma.

In our study, we first explored the role of 74 IFN signatures in blood specimens of patients with asthma and observed that five IFN signatures (IFN-γ response-DN, negative regulation of IFN-γ secretion, IFNG pathway, negative regulation of response to IFN-γ, and type 1 IFN biosynthetic process) exhibited a higher level in asthma group than in healthy control group. In addition, the analysis of biological processes and pathways was still involved in regulation of type 1 IFN production, response to IFN-γ, and IFN signaling pathway. Based on the MCODE algorithm, 21 hub IFN-related genes were observed in this work. In a large population-based case–control study (411 patients with asthma and 87 healthy control subjects), these 21 IFN-related genes were performed to evaluate whether their combination can improve diagnostic ability for asthma, which suggested a more effective blood-based biomarker for diagnosing asthma in clinic.

Numerous studies have suggested that the ssGSEA method can represent the relative abundance of gene set signature and explore the underlying molecular mechanism in tumor [24–26]. In the airway epithelium, IFN-λ plays key roles in the mucosal immune response to viral infection for asthma [27]. The expression of IFN-β and IFN-λ1/IL-29 was higher in asthma compared with healthy subjects [28]. In the current work, we first determined 74 IFN signatures in asthma. IFN-γ response-DN, negative regulation of IFN-γ secretion, IFNG pathway, negative regulation of response to IFN-γ, and type 1 IFN biosynthetic process displayed higher proportion in blood samples of asthma. Using brochoalveolar lavage data (GSE67940), we further found that some IFN signatures (i.e., IFN-γ response-DN) were also higher in asthma (Supplementary Figure S1). These findings suggested that some IFN signatures had pivotal roles in the pathogenesis of asthma. Asthma is an uncontrolled disease, with a substantial mortality and morbidity burden. Because under- and overdiagnosis and misdiagnosis are common in clinical practice for asthma, thus, asthma shows a large challenge for the diagnosis and management [7]. Here, we identified 21 hub IFN-related genes and pooled these genes in blood samples of asthma. We found that asthmatic subjects exhibited a superior AUC value of the combination of biomarkers compared with control subjects (AUC = 0.832). Because of the easy-to-use whole blood samples, our work revealed the use of IFN-related gene biomarkers can be regarded as a useful tool for early detection and intervention of asthma.

The present study had several limitations. First, our work was a retrospective design. Second, because of insufficient data on different groups (such as allergic asthma and non-allergic asthma and exacerbated asthma condition), we did not perform the relevant analysis for different types of asthmatic groups. Finally, the survival data were not available for asthma, additional studies are necessary to validate the correlation between the survival data and asthma for the selected IFN genes in the future.

In conclusion, we identified that some IFN signatures were associated with asthma. Based on the use of blood samples, the combination of IFN-related genes demonstrated a robust and effective diagnostic ability for early detection of asthma. These findings offered the first evidence to better understand the underlying mechanisms of how IFN pathways signatures affected asthma. However, further research is still necessary to elucidate the role of IFN signatures in asthma in the future.

Data Availability

All data were available in this manuscript.

Competing Interests

The authors declare that there are no competing interests associated with the manuscript.

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Author Contribution

Hairong Shu, Minqiao Zheng, and He Li contributed to the conception and design of this research. Hairong Shu, Yong Li, Hangyu Xu, Qing Yin, Jianxin Song, Minqiao Zheng, and He Li contributed to the drafting of the article and final approval of the submitted version. Hairong Shu, Yong Li, Hangyu Xu, Qing Yin, Jianxin Song, Minqiao Zheng, and He Li contributed to data analyses and the interpretation and completion of the figures and tables. All authors read and approved the final manuscript.
Ethics Approval
All procedures performed in studies involving human participants were in accordance with the ethical standards of the Gene Expression Omnibus Human Subjects Protection and Data Access Policies.

Informed Consent
Informed consent was obtained from all individual participants included in the study.

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Abbreviations
AHR, airway hyperreactivity; AUC, area under the curve; DN, down; IFN, interferon; IFNG, interferon gamma; IgE, immunoglobulin E; IL-4/5, interleukin-4/5; IL-29, interleukin-29; IRF3, interferon regulatory factor 3; MCODE, Molecular Complex Detection; PPI, protein-protein interaction; RMA, robust multiarray averaging; ROC, receiver operating characteristic; ssGSEA, single-sample gene set enrichment analysis; Th1/2, T helper type 1/2; TLR9, Toll-like receptor 9.

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