

Novel Blood-based Transcriptional Biomarker Panels Predict the Late-Phase Asthmatic Response

Amrit Singh^{1,2,3}, Casey P. Shannon², Young Woong Kim^{1,2}, Chen Xi Yang^{1,2}, Robert Balshaw⁴, Gabriela V. Cohen Freue⁵, Gail M. Gauvreau⁶, J. Mark FitzGerald^{7,8}, Louis-Philippe Boulet⁹, Paul M. O'Byrne⁶, and Scott J. Tebbutt^{1,2,8}

¹Centre for Heart Lung Innovation, St. Paul's Hospital, ³Department of Pathology and Laboratory Medicine, ⁵Department of Statistics, and ⁸Division of Respiratory Medicine, Department of Medicine, University of British Columbia, Vancouver, British Columbia, Canada; ²Prevention of Organ Failure Centre of Excellence, Vancouver, British Columbia, Canada; ⁴Centre for Healthcare Innovation, University of Manitoba, Winnipeg, Manitoba, Canada; ⁶Department of Medicine, McMaster University, Hamilton, Ontario, Canada; ⁷Vancouver Coastal Health Research Institute, Vancouver General Hospital, Vancouver, British Columbia, Canada; and ⁹Quebec Heart and Lung Institute, Laval University, Quebec City, Quebec, Canada

ORCID IDs: 0000-0002-7475-1646 (A.S.); 0000-0002-5687-3156 (C.P.S.); 0000-0002-3470-6728 (Y.W.K.); 0000-0002-8033-4769 (C.X.Y.); 0000-0002-2455-8792 (R.B.); 0000-0003-4526-0175 (G.V.C.F.); 0000-0002-6187-2385 (G.M.G.); 0000-0002-5367-5226 (J.M.F.); 0000-0003-3485-9393 (L.-P.B.); 0000-0003-0979-281X (P.M.O'B.); 0000-0002-7908-1581 (S.J.T.).

Abstract

Rationale: The allergen inhalation challenge is used in clinical trials to test the efficacy of new treatments in attenuating the late-phase asthmatic response (LAR) and associated airway inflammation in subjects with allergic asthma. However, not all subjects with allergic asthma develop the LAR after allergen inhalation. Blood-based transcriptional biomarkers that can identify such individuals may help in subject recruitment for clinical trials as well as provide novel molecular insights.

Objectives: To identify blood-based transcriptional biomarker panels that can predict an individual's response to allergen inhalation challenge.

Methods: We applied RNA sequencing to total RNA from whole blood ($n = 36$) collected before and after allergen challenge and generated both *genome-guided* and *de novo* datasets: genes, gene-isoforms (University of California, Santa Cruz, UCSC Genome Browser), Ensembl, and Trinity. Candidate biomarker panels were validated using the NanoString platform in an independent cohort of 33 subjects.

Measurements and Main Results: The Trinity biomarker panel consisting of known and novel biomarker transcripts had an area under the receiver operating characteristic curve of greater than 0.70 in both the discovery and validation cohorts. The Trinity biomarker panel was useful in predicting the response of subjects that elicited different responses (accuracy between 0.65 and 0.71) and subjects that elicit a dual response (accuracy between 0.70 and 0.75) upon repeated allergen inhalation challenges.

Conclusions: Interestingly, the biomarker panel containing novel transcripts successfully validated compared with panels with known, well-characterized genes. These biomarker–blood tests may be used to identify subjects with asthma who develop the LAR, and may also represent members of novel molecular mechanisms that can be targeted for therapy.

Keywords: gene expression profiling; biomarkers; asthma

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Correspondence and requests for reprints should be addressed to Scott J. Tebbutt, Ph.D., Department of Medicine, University of British Columbia Centre for Heart Lung Innovation, Room 166-1081 Burrard Street, Vancouver, BC, V6Z1Y6 Canada. E-mail: scott.tebbutt@hli.ubc.ca.

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At a Glance Commentary

Scientific Knowledge on the

Subject: Attenuation of the allergen-induced late-phase asthmatic response (LAR) is used to determine the efficacy of novel asthma therapies.

Recruitment, for clinical trials, of subjects who develop the late response is performed using an allergen inhalation challenge. However, only a subgroup of individuals with mild asthma develop the LAR.

What This Study Adds to the

Field: This study identified and validated novel RNA transcripts that are predictive of the LAR. These biomarkers may be used to enrich clinical trials with subjects who are susceptible to developing the LAR.

The allergen-induced late-phase asthmatic response (LAR), occurs reproducibly in a subgroup of individuals with mild allergic asthma after allergen inhalation challenge (1), and shares many characteristics with chronic asthma, such as prolonged airway contraction, persistent airway inflammation, mucus hypersecretion, and airway remodeling (2, 3). Although inhaled corticosteroids are effective in abolishing the LAR (4), they do not alter the natural course of the disease (5). Airway inflammation (e.g., presence of epithelial shedding, T cells, and eosinophils in mucosal biopsies) is already present in subjects with mild disease, and worsens in subjects with persistent asthma (6). Given the heterogeneity of the disease within an individual and over time, reproducible biomarkers that can help risk stratify individuals may help improve diagnosis, selection of therapy, and prevention of overtreatment (7–11).

The purpose of this study was to identify transcriptional biomarker panels (combinations of single molecules) using whole blood that could predict a subject's response to allergen inhalation challenge, before and 2 hours after challenge (Figure 1). Following the Institute of Medicine guidelines for the development of clinical biomarker tests (12), biomarker panels were identified in a discovery cohort with rigorous clinical phenotypic characterization. The biomarker panels

were developed using RNA-sequencing (RNA-Seq) data, and the performance of the computational model was assessed using cross-validation. The identified biomarker candidates were then transferred to the more clinically relevant NanoString platform and the biomarker panel formulas were locked down. Finally, the performance of these biomarker panels was assessed in an independent external validation cohort. Some of the results of these studies have been previously reported in the form of abstracts (13–15).

Methods

The institutional review boards of the participating institutions—University of British Columbia, McMaster University, and Université Laval—approved this study.

Subjects

Written informed consents were obtained from participants undergoing allergen inhalation challenges. Subjects were nonsmokers with mild atopic asthma, free of other lung diseases and any cardiovascular disease. All subjects had physician diagnosed, clinically stable asthma, with a baseline FEV₁ of 70% or greater of predicted value, and baseline provocative concentration of methacholine that caused a 20% drop in FEV₁ (PC₂₀) less than 16 mg/ml. All subjects developed the early asthmatic response (at least 20% fall in FEV₁ within 2 hours after allergen inhalation). Exclusion criteria included the use of inhaled corticosteroids, and use of other asthma medication, with the exception of infrequently inhaled β_2 -agonist, which was withheld for 8 hours before spirometry measurements (additional details can be found in References 16–22).

Methacholine and Allergen Inhalation Challenge

Methacholine and allergen challenge were performed on triad visits. Methacholine inhalation tests were performed on Days 1 (pre-methacholine) and 3 (post-methacholine) in order to determine airway hyperresponsiveness, defined as the allergen-induced shift ([PC₂₀]_{pre}/[PC₂₀]_{post}). Allergen inhalation challenge was performed on Day 2 using allergen extracts in doubling doses until a

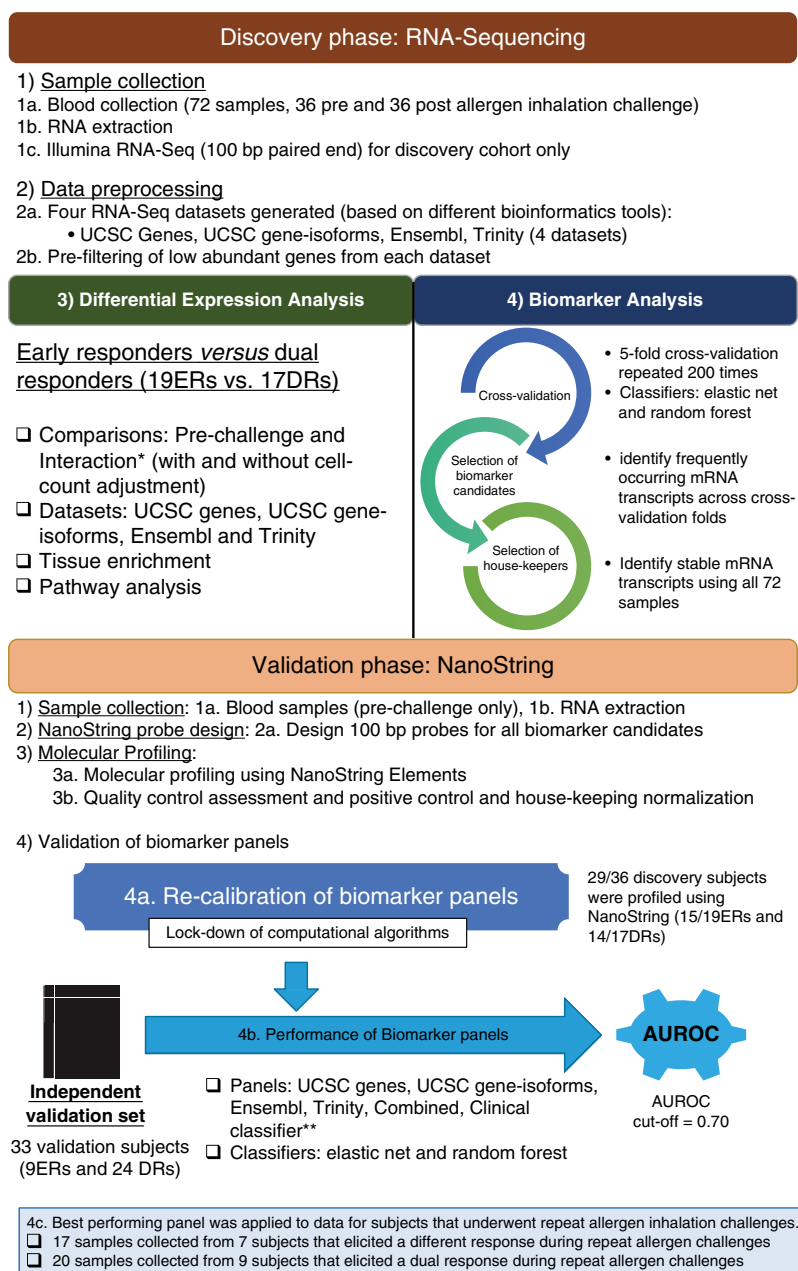
drop in FEV₁ of at least 20% was achieved; subsequently, FEV₁ was measured at regular intervals up to 7 hours after challenge. All subjects demonstrated an FEV₁ drop of 20% between 0 and 2 hours (early asthmatic response) after allergen inhalation challenge. Participants who demonstrated a maximum drop in FEV₁ greater than or equal to 15% between 3 and 7 hours after allergen challenge (LAR) were classified as dual responders (DRs). Subjects who did not meet the 15% drop in FEV₁ during the LAR were classified as early responders (ERs).

Cohorts

The discovery cohort consisted of 36 subjects. For all subjects of the discovery cohort, blood samples were collected for molecular analysis immediately before and 2 hours after allergen inhalation challenge. For the validation cohort, pre-challenge blood samples from 33 subjects were used for molecular analyses. Furthermore, 17 pre-challenge samples were collected from 7 subjects who elicited a different response during repeat allergen inhalation challenges, whereas 20 pre-challenge samples were collected from 9 subjects who elicited only the dual response during repeat allergen inhalation challenges. Standard operating protocols were used at each participating center for all blood collection procedures (*see* the online supplement for blood collection and processing details).

RNA-Seq

Total RNA from blood samples of the discovery cohort was purified and sent to Génome Québec (Centre d'Innovation Genome Quebec et Université McGill) for whole-transcriptome sequencing (eight samples per lane). Ribosomal RNA and globin-depleted stranded cDNA libraries were sequenced using an Illumina HiSeq 2000 as 100-bp paired end reads. All RNA-Seq files passed quality-control metrics based on FastQC standards. After trimming (Seqtk, version 1.0), the left and right concatenated files were used by the Trinity software (version r20131110) (23) to construct a *de novo* assembly of the blood transcriptome (reverse forward-stranded library type was specified). The abundance estimates of Trinity contigs were estimated using RSEM (RNA-Seq by Expectation Maximization, version 1.2.11) (24) using the Bowtie aligner. The abundance



estimates of University of California, Santa Cruz (UCSC) genes and gene-isoforms were estimated using RSEM (version 1.2.19) with the Bowtie2 (version 2.2.4) aligner. STAR (Spliced Transcripts Alignment to a Reference, version 2.5.0a), a fast RNA-Seq alignment software (25), was also used to align the paired end reads to the human genome using annotations from the GENCODE project (GENCODE release v21) (26). Feature counts (27) in the Subread/Rsubread package (version 1.5.0) was used to estimate the abundance of Ensembl transcripts. See the online supplement for complete details on RNA-Seq data preprocessing and normalization.

nCounter Elements TagSets

Oligonucleotide probes (100 bp) were designed for all biomarker candidates using the nDesign portal (<http://www.nanostring.com/>) and with the help of the bioinformatics team at NanoString Technologies. For uncharacterized candidates, the sequences were selected based on coverage plots of RNA-Seq data and provided to NanoString Technologies (see the online supplement for probe sequences). RNA samples from both the discovery and validation cohorts as well as repeated challenges were profiled for biomarker candidates using custom nCounter Elements TagSets (NanoString Technologies). All NanoString data were assessed for various quality-control metrics (28) and normalized according to their data analysis guide (see the online supplement for complete details).

Statistical Analyses

Differential expression analysis was performed using the limma (linear models for microarray and RNA-Seq data) R-library (version 3.32.4; 29). Two comparisons were considered: ERs versus DRs before challenge and ERs versus DRs in the interaction (post-minus pre-challenge levels). A *t* test was used to compare ERs and DRs for all quantitative clinical and demographics variables (e.g., cell counts, age) when the data were deemed to be normally distributed (assessed using the Shapiro-Wilk normality test and normal Q-Q plots). The Wilcoxon rank-sum test was used when the normality assumption was not met. Tissue enrichment analysis was performed using the sear R-library (version 0.1), whereas gene set analysis was

Figure 1. Overview of analysis. The study was divided into a discovery and validation phase. Unbiased whole-genome expression profiling was performed using RNA sequencing, and predictive biomarker panels of the late asthmatic response were identified before challenge and at the interaction (after challenge normalized to before challenge). Selected transcripts were profiled using the NanoString platform on all samples. The biomarker panels were recalibrated using the discovery samples and applied to the additional samples to determine the classification performance. *Interaction: for each mRNA transcript, the expression of the pre-challenge samples was subtracted from the corresponding post-challenge sample before comparing ERs and DRs. **Clinical classifier consisted of baseline FEV₁, PC₂₀ before challenge, total leukocyte counts, and white blood cell-count frequencies (neutrophils, lymphocytes, monocytes, eosinophils, and basophils). AUROC = area under the receiver operating characteristic curve; DR = dual responder; ER = early responder; PC₂₀ = provocative concentration of methacholine that caused a 20% drop in FEV₁; UCSC = University of California, Santa Cruz.

performed using Enrichr (30). When applicable, the Benjamini-Hochberg correction procedure for multiple testing (31) was used, and a false discovery rate (FDR) of 0.15 was used to determine statistical significance.

Elastic net (32) and random forest (33) were used to identify biomarker panels that could distinguish ERs from DRs for each RNA-Seq dataset: UCSC genes, UCSC gene-isoforms, Ensembl, and Trinity. Classification performance of biomarker panels for each dataset was estimated using a 5-fold cross-validation repeated 200 times (200×5 -fold cross-validation), using an elastic net and random forest classifier. For each cross-validation run, the area under the receiver operating characteristic curve (AUROC) was computed and averaged over 200 repeats for each dataset. For both the discovery and validation phase an

AUROC cut-off of 0.70 was used as per recommended guidelines for clinical biomarker implementation (34). Biomarker selection was based on the frequently occurring (stable) transcripts in biomarker panels identified within cross-validation folds. To select stable biomarker candidates, the most frequently occurring transcripts among the 1,000 panels (5-fold \times 200 repeats) for both elastic net and random forest applied to each dataset were determined. The overlap between the top 100 ranked transcripts based on elastic net and random forest were identified for each dataset and annotated, from which a subset of candidates was selected to be transferred to the NanoString platform. Housekeeping transcripts per dataset were identified using the geNorm algorithm (35) in the NormqPCR R-library (version 1.16.0). Complete details on

the statistical methodology used in this study can be found in the online supplement.

Results

Discovery and Validation Cohorts

The discovery cohort consisted of 36 subjects (Table 1): 19 were classified as ERs and 17 were classified as DRs. The post-challenge PC_{20} was significantly lower in DRs compared with ERs, resulting in a significantly increased allergen-induced shift in DRs compared with ERs. No significant changes between ERs and DRs were identified for total leukocyte counts and specific white blood cell frequencies before challenge (Table 1). In the discovery cohort, white blood cell count frequencies were also not significantly different after

Table 1. Subject Demographics of the Discovery Cohort

Clinical Variable	Isolated ERs (n = 19)	DRs (n = 17)	P Value
Female, %	68	65	
Height, cm*	168.19 \pm 8.65	168.53 \pm 9.26	0.91
Weight, kg†	62.70 (58.55–78.25)	71.00 (63.35–87.15)	0.23
Age, yr†	28.00 (21.00–34.50)	23.00 (21.00–37.00)	0.65
Baseline FEV ₁ , L/s*	3.29 \pm 0.79	3.27 \pm 0.81	0.95
Predicted FEV ₁ , L/s†	3.43 (3.15–4.23)	3.53 (3.19–4.13)	0.91
% drop in FEV ₁ during the EAR†	30.40 (25.45–40.15)	34.80 (30.70–44.10)	0.12
% drop in FEV ₁ during LAR*	6.51 \pm 5.86	34.75 \pm 7.66	2.86×10^{-14}
Pre-methacholine PC ₂₀ , mg/ml†	5.25 (1.88–11.15)	2.24 (0.59–4.56)	0.11
Post-methacholine PC ₂₀ , mg/ml†	5.71 (1.00–9.50)	0.54 (0.23–2.26)	0.004
AIS‡	1.89 \pm 1.53	3.71 \pm 1.91	0.01
Allergen			
Cat	9	4	
Fungus	1	1	
Grass	3	2	
HDM	5	7	
Horse	0	1	
Ragweed	1	2	
Site			
Laval	11	7	
McMaster	8	9	
UBC	0	1	
Blood cell counts and frequencies before challenge			
Leukocytes, $\times 10^9$ cells/L*	6.38 \pm 1.44	5.87 \pm 1.35	0.31
Neutrophils, %*	0.57 \pm 0.08	0.53 \pm 0.09	0.19
Lymphocytes, %*	0.31 \pm 0.08	0.32 \pm 0.09	0.70
Monocytes, %*	0.08 \pm 0.01	0.09 \pm 0.02	0.23
Eosinophils, %†	0.03 (0.02–0.05)	0.04 (0.03–0.07)	0.27
Basophils, %†	0.01 (0.004–0.01)	0.0 (0.0–0.01)	0.36

Definition of abbreviations: AIS = allergen-induced shift; DRs = dual responders; EAR = early asthmatic response; ERs = early responders; HDM = house dust mite; LAR = late-phase asthmatic response; PC₂₀ = provocative concentration of methacholine that caused a 20% drop in FEV₁; UBC = University of British Columbia.

Grass refers to grass mix, orchard grass, or timothy grass. HDM refers to *Dermatophagoides farinae* or *D. pteronyssinus*.

*Variable is assumed to be normally distributed. Descriptive statistics are presented as mean \pm SD. A *t* test was used to compare ERs and DRs.

†Variable is assumed to not be normally distributed. Descriptive statistics are presented as median (25–75th percentiles). A Wilcoxon rank-sum test was used to compare ERs and DRs.

‡[PC₂₀]_{pre} divided by [PC₂₀]_{post}.

Table 2. Subject Demographics of the Validation Cohort

Clinical Variable	Isolated ERs (n = 9)	DRs (n = 24)	P Value
Female, %	56	46	
Height, cm*	168.11 ± 10.21	171.15 ± 8.91	0.43
Weight, kg*	69.00 ± 11.87	72.21 ± 11.55	0.51
Age, yr [†]	31.00 (27.00–36.00)	25.50 (21.75–41.00)	0.35
Baseline FEV ₁ , L/s*	3.05 ± 0.51	3.50 ± 0.81	0.14
Predicted FEV ₁ , L/s*	3.67 ± 0.84	3.68 ± 0.62	0.98
% drop in FEV ₁ during the EAR*	31.92 ± 9.32	36.39 ± 8.83	0.21
% drop in FEV ₁ during LAR [†]	7.20 (4.90–11.10)	23.75 (17.50–30.88)	1.39 × 10 ^{−5}
Pre-methacholine PC ₂₀ , mg/ml [‡]	5.13 (0.64–5.45)	1.58 (0.69–4.08)	0.49
Post-methacholine PC ₂₀ , mg/ml [‡]	5.11 (2.18–7.98)	0.76 (0.34–1.55)	0.04
AIS* [‡]	1.35 ± 1.13	2.92 ± 1.69	0.04
Allergen			
Cat	7	9	
Fungus	0	0	
Grass	2	4	
HDM	0	8	
Horse	0	1	
Ragweed	0	2	
Site			
Laval	6	9	
McMaster	1	12	
UBC	2	3	
Blood cell counts and frequencies before challenge			
Leukocytes, ×10 ⁹ cells/L*	5.13 ± 1.25	6.12 ± 1.52	0.10
Neutrophils, %*	0.58 ± 0.10	0.51 ± 0.08	0.04
Lymphocytes, %*	0.29 ± 0.08	0.37 ± 0.07	0.02
Monocytes, %*	0.07 ± 0.02	0.08 ± 0.01	0.62
Eosinophils, % [†]	0.03 (0.02–0.06)	0.04 (0.02–0.04)	0.70
Basophils, % [†]	0.00 (0.002–0.006)	0.01 (0.005–0.01)	0.17

For definition of abbreviations, see Table 1. Grass refers to grass mix, orchard grass, or timothy grass. HDM refers to *Dermatophagoides farinae* or *D. pteronyssinus*.

*Variable is assumed to be normally distributed. Descriptive statistics are presented as mean ± SD. A *t* test was used to compare ERs and DRs.

[†]Variable is assumed to not be normally distributed. Descriptive statistics are presented as median (25–75th percentiles). A Wilcoxon rank-sum test was used to compare ERs and DRs.

[‡][PC₂₀]_{pre} divided by [PC₂₀]_{post}.

challenge (normalized to before challenge, also called the interaction) between ERs and DRs. The validation cohort consisted of 33 subjects (Table 2): 9 were classified as ERs and 24 were classified as DRs. Significant changes between ERs and DRs with respect to pre-challenge neutrophil and lymphocyte frequencies were also identified (Table 2).

Differential Expression Analysis Reveals a Strong Discriminatory Signal between ERs and DRs before Challenge

At an FDR cut-off of 0.15, thousands of transcripts were identified to be differentially expressed between ERs and DRs before challenge with or without adjusting for total leukocyte counts (Figure 2A). Specifically, 1,177 UCSC genes (125 up in DRs, and 1,052 down in DRs), 683 UCSC gene-isoforms (69 up and 614 down), 1,032 Ensembl (305 up and 727

down), and 866 Trinity contigs (30 up and 836 down) were differentially expressed before challenge with or without adjustment for cell counts (Figure 2B). However, at the same FDR cut-off, no significant transcripts between ERs and DRs were identified in the interaction comparison (Figure 2A). Annotation of all differentially expressed transcripts resulted in 2,051 unique gene symbols, which were used for tissue and pathway enrichment analysis. Tissue enrichment analysis attributed the differentially expressed genes before challenge to T cell–specific genes, namely CD4 and CD8 T cells (Figure 2C). Enrichr identified 85 significant BioCarta pathways, 24 KEGG pathways, and 62 WikiPathways. Figure 2D depicts the overlap between the top 10 significant pathways with a pathway membership similarity of greater than 20%. Significantly enriched pathways included the T-cell receptor signaling

pathway, IL-2 receptor β chain in T-cell activation, and IL-1, TNF, TGF-β, and B-cell receptor signaling pathways (Figure 2D). Additional details regarding dataset preprocessing can be found in the online supplement.

Biomarker Analysis of RNA-Seq Data Identifies RNA Transcripts That Are Predictive of the LAR

Given that the discriminatory signal between ERs and DRs was stronger before challenge, we focused our efforts on identifying risk biomarker panels that could discriminate ERs from DRs before allergen inhalation challenge. Using a 200 × 5-fold cross-validation, estimates of the test performance of each of the RNA-Seq dataset classifiers as well as the clinical classifier (based on baseline FEV₁, before methacholine PC₂₀, total leukocyte counts, and white blood cell count frequencies [neutrophils, lymphocytes, monocytes,

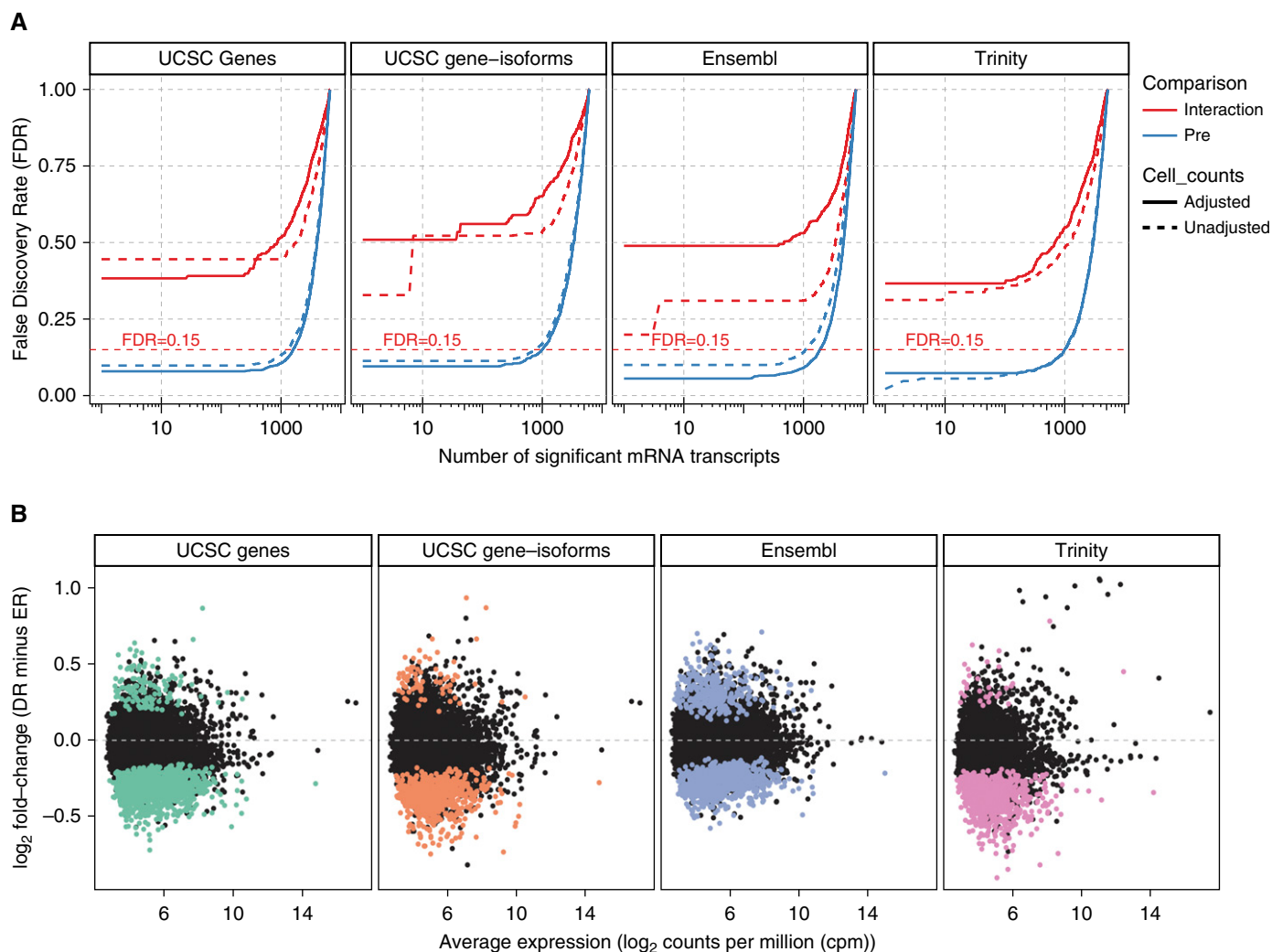


Figure 2. Differential expression analysis. (A) Benjamini-Hochberg false discovery rate (FDR) versus the number of significant mRNA transcripts, before challenge and at the interaction (before minus after), with and without adjusting for total leukocyte counts, for each dataset. (B) Minus-average plots depicting the fold change versus the average expression of mRNA transcripts across all pre-challenge samples, for each dataset. The colored points represent differentially expressed transcripts (comparing early responders [ERs] and dual responders [DRs] before challenge) common between the with- and without-adjustment for total leukocyte count comparisons, whereas the black points represent nondifferentially expressed transcripts. (C) Tissue enrichment of common (with and without adjusted for total leukocyte counts) differentially expressed transcripts between ERs and DRs before challenge from all datasets. (D) Top 10 significantly enriched pathways from the BioCarta, KEGG, and WikiPathways databases using the entire list of differentially expressed genes from all datasets. Connected pathways have a Jaccard similarity of greater than 20%. UCSC = University of California, Santa Cruz.

eosinophils, and basophils]) were determined (Figure 3A). The performance of the Trinity biomarker panel met an AUROC cut-off of 0.70 with average classification performances of AUROC of 0.71 (elastic net) and AUROC of 0.74 (random forest). The clinical panel had the lowest classification performance (AUROC of ~50%).

Setting the tuning parameter ($\alpha = 0.9$) resulted in approximately 13–17 transcripts per elastic net biomarker panel (an equivalent-sized random forest classifier was chosen based on the

importance score ranking) within the cross-validation folds. Biomarker panel transcripts across the 1,000 panels ($200 \times 5\text{-fold} = 1,000\text{-fold}$) were tallied for both the elastic net and random forest panels for each dataset (see METHODS for details). The overlaps between the top 100 ranked transcripts in the elastic net and random forest panels were determined for each dataset and annotated, from which a subset of candidates was selected to be transferred to the NanoString platform: 33 UCSC genes, 35 UCSC gene-isoforms, 35 Ensembl transcripts, and 14 Trinity contigs

(see Figure 3B for an overlap between transcripts selected from each dataset). The RNA biomarker transcripts retained their discriminative ability in separating ERs from DRs 2 hours after allergen inhalation challenge, based on principal component analysis (Figure 2C).

Nine housekeeping transcripts were identified across three datasets: UCSC genes, Ensembl, and Trinity (Figure 2D). However, because the Trinity housekeeping contigs were well above the expression of all other selected transcripts, they were removed from the final NanoString assay design,

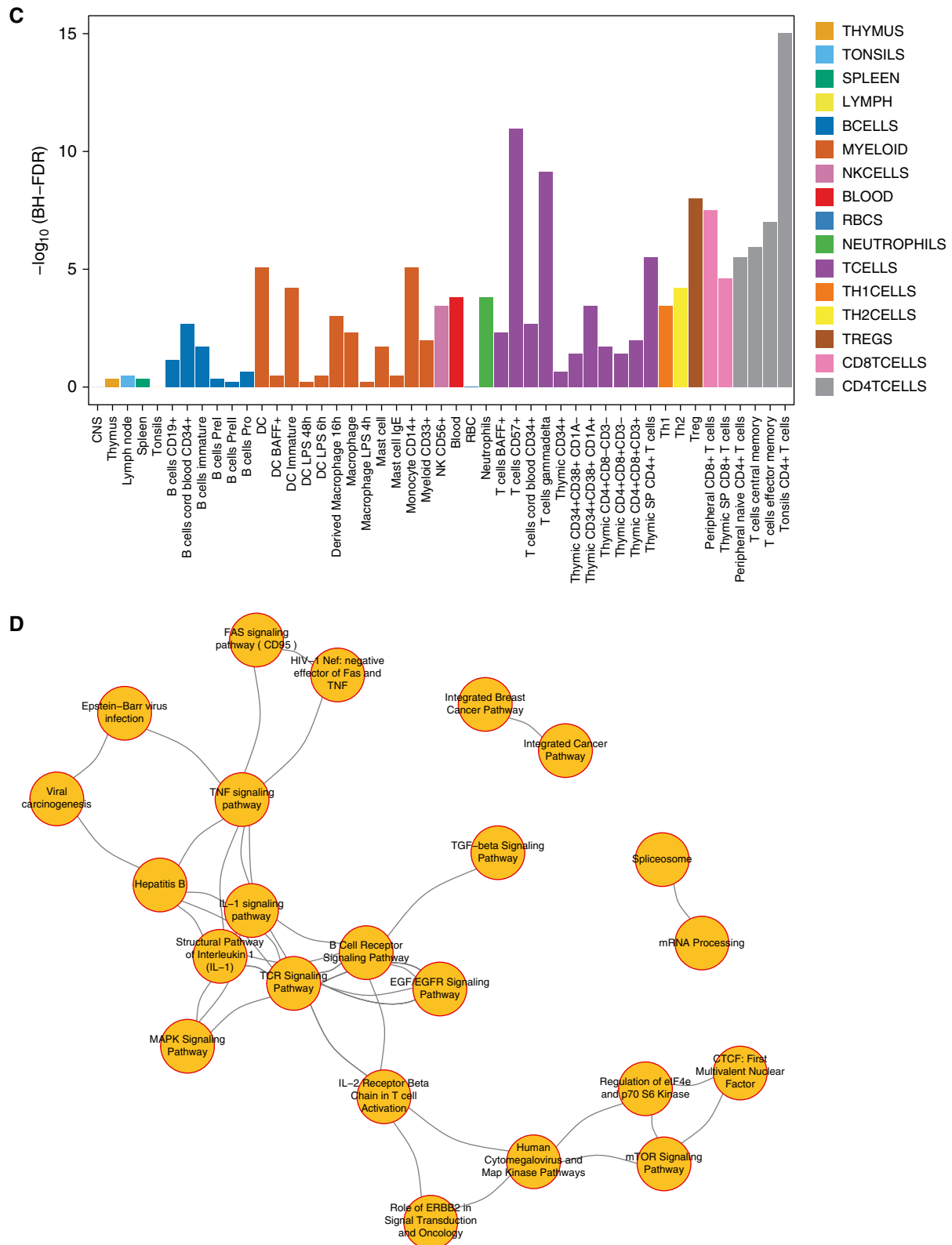


Figure 2. (Continued).

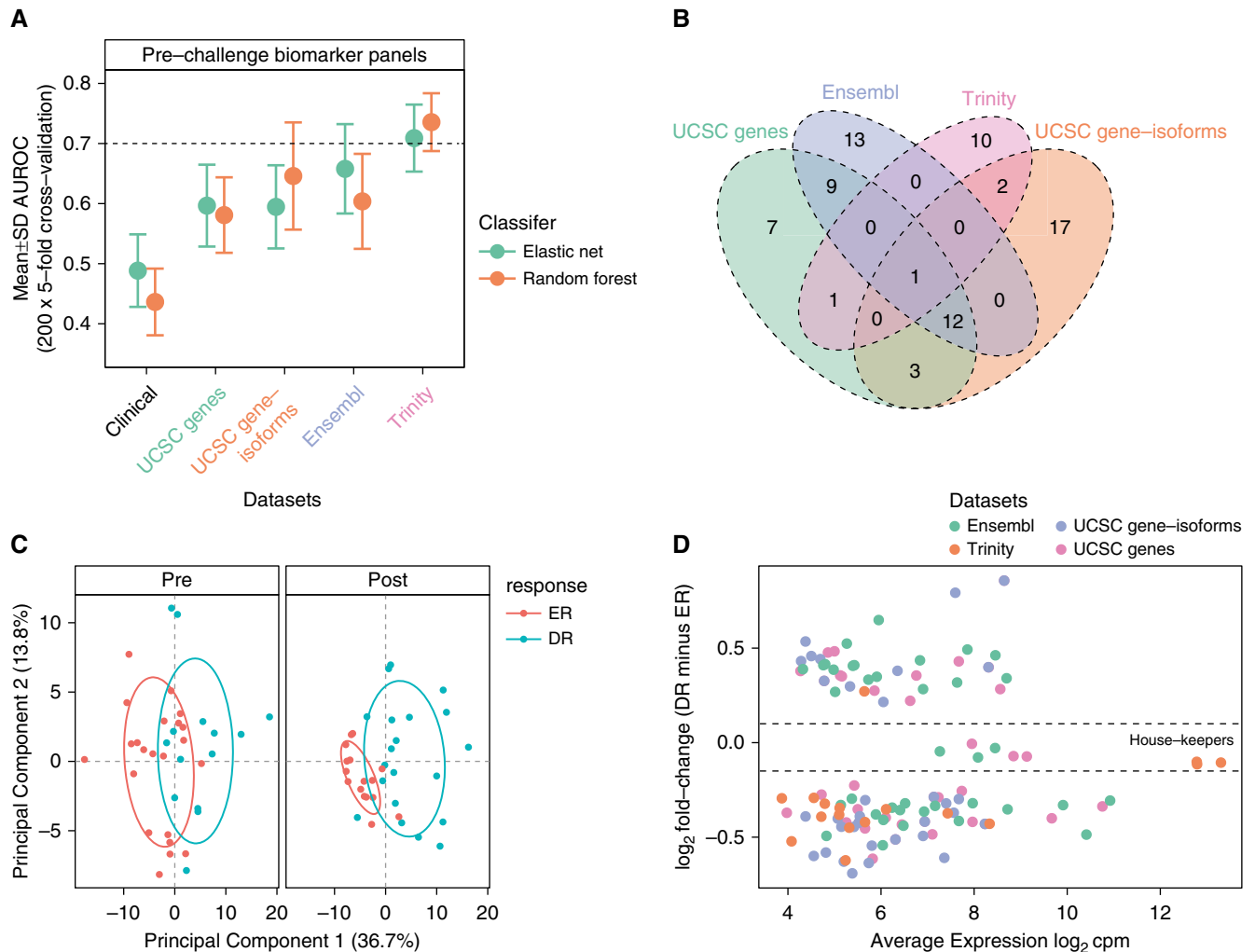


Figure 3. Discovery of pre-challenge biomarker panels. (A) Classification performance of pre-challenge biomarker panels based on a 5-fold cross-validation repeated 200 times (200 \times 5-fold cross-validation) using an elastic net and random forest classifier. (B) Overlap between the selected mRNA biomarker transcripts from each dataset; each set of transcripts serves as the resulting biomarker panel for each dataset. (C) Principal component analysis plot of all samples (before and after) using all biomarker transcripts in B. The ellipses (with a 68% confidence interval) depict a separation between the early responders (ERs) and dual responders (DRs) before challenge, which is sustained at 2 hours after challenge. (D) Minus-average plot of all biomarker transcripts as well as housekeeping transcripts (within dashed lines). Note: complete clinical data were present for 17/19 ERs and 15/17 DRs. AUROC = area under the receiver operating characteristic curve; UCSC = University of California, Santa Cruz.

resulting in six housekeeping gene candidates: *MED13*, *TOR1AIP2*, and *WAC* (Ensembl dataset) and *ARPC4*, *TMBIM6*, and *RHOA* (UCSC gene dataset), all of which were transferred to the NanoString platform. A NanoString assay can reliably measure up to two million counts per sample. However, if some transcripts are highly abundant (e.g., globin in a whole-blood sample), they may saturate the assay such that most counts would be attributed to the highly expressed targets. Therefore, the highly abundant Trinity housekeeping transcripts were not selected for downstream NanoString analyses.

Panel Recalibration and Validation Using the NanoString Platform

Six housekeeping probes as well as 75 probes corresponding to the 117 biomarker candidates (due to gene redundancy across datasets) were designed. Transcripts were annotated to their corresponding gene symbol, and a probe was selected from the nDesign portal (www.nanostring.com). Custom probes were designed for Trinity contigs that mapped either to intronic regions of known genes or to uncharacterized loci of the human genome, based on visualizing the nucleotide coverage of each transcript, and sent to NanoString technologies, which selected

100 bp from this sequence as the final probe sequence (see Table E1 in the online supplement).

All samples of the discovery and validation cohorts (Figure 4A) were profiled using the NanoString platform for all biomarker and housekeeping probes. Quality-control assessment and normalization based on positive spike-in controls and housekeeping probes were performed (see the online supplement). Five biomarker panels (UCSC genes, UCSC gene-isoforms, Ensembl, Trinity, and Combined [all transcripts]) and a clinical panel were locked down (fixed weights of transcripts in the biomarker

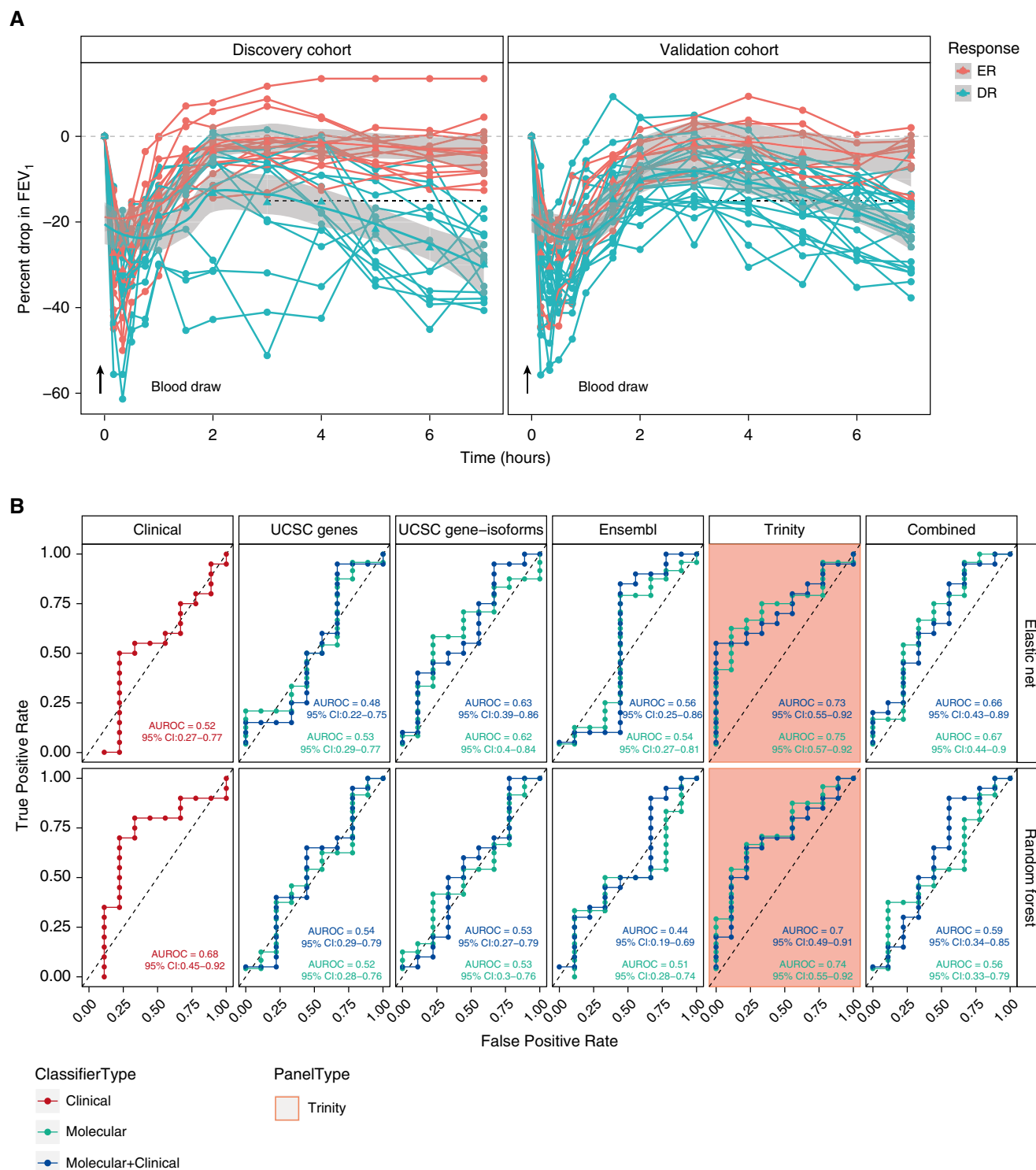


Figure 4. Validation of pre-challenge biomarker panels. (A) FEV₁ drop from baseline of subjects in the discovery and validation cohorts undergoing allergen inhalation. The LOESS (locally weighted scatterplot smoothing) curve was fitted to each group separately, with a 95% confidence interval. (B) Area under the receiver operating characteristic curve (AUROC) for all molecular panels (from each dataset) and in conjunction with clinical variables, such as baseline FEV₁, provocative concentration of methacholine that causes a 20% drop in FEV₁ before challenge, total leukocyte counts, and white blood cell count frequencies (neutrophils, lymphocytes, monocytes, eosinophils, and basophils). The Trinity panel (pink) achieved an AUROC of greater than or equal to 0.70 by itself and in conjunction with clinical variables. Notes: given the limited RNA quantity available for samples in the discovery cohort, molecular profiling was performed for 15/17 early responders (ERs) and 14/17 dual responders (DRs). Complete clinical data were present for 14/15 ERs and 12/14 DRs in the discovery cohort and 9/9 ERs and 20/24 DRs in the validation cohort. CI = confidence interval; UCSC = University of California, Santa Cruz.

panels) by fitting a model (elastic net and random forest) to NanoString data from discovery samples (15 ERs and 14 DRs; some samples were lost due to the lack of sufficient RNA material remaining for some samples). An “off the shelf” test was performed using an independent cohort of 9 ERs and 24 DRs for all biomarker panels, and the AUROC was determined (Figure 4B). The Trinity biomarker panels performed well using both elastic net (AUROC = 0.75; 95% confidence interval = 0.57–0.92) and random forest (AUROC = 0.74; 95% confidence interval = 0.55–0.92) biomarker panels (Figure 4B), with both panels having an accuracy of 0.72 (using a probability threshold of 0.5).

The Trinity Biomarker Panel and Its Application to Subjects Undergoing Repeated Allergen Inhalation Challenges

The Trinity biomarker panel consisted of 14 transcripts (both known and uncharacterized) that demonstrated the strongest discriminatory signal among all molecular panels, as well as the clinical panel. Figure 5A depicts the separation of the responder groups (using samples from both the discovery and validation cohort) along the first principal component. The first principal component is a linear combination of the 14 transcripts weighted by the contribution of each individual transcript, leading to responder group separation observed in Figure 5A. Figure 5B shows the contribution of each transcript that leads to the separation depicted in Figure 5A. Overall, the expression levels of the Trinity panel transcripts were higher in ERs compared with DRs (Figure 5C).

The Trinity panel-based elastic net and random forest classifiers were applied to 17 samples corresponding to 7 subjects who elicited a different response during repeated allergen inhalation challenges. The Trinity panel had an accuracy of 0.71 and 0.65 for the elastic net and random forest classifier, respectively (Figure 5D, upper panel). The Trinity panel-based elastic net and random forest classifiers were applied to subjects who elicited only the dual response upon repeated allergen challenges and achieved an accuracy of 0.70 and 0.75, respectively (Figure 5D, lower panel).

Discussion

We identified four predictive biomarker panels of the LAR, of which the Trinity biomarker panel, consisting of novel RNA transcripts, was validated using an independent cohort, achieving an AUROC of greater than 0.70. Interestingly, the Trinity panel demonstrated utility in predicting the response of subjects who underwent repeated allergen inhalation challenges. This may indicate that the Trinity biomarker panel was sensitive to molecular fluctuations within an individual that are predictive of the type of response a given subject will elicit on the day of the allergen inhalation challenge. These results may imply inherent differences in the underlying molecular mechanisms that can be detected in peripheral whole blood, which predispose individuals with asthma for the LAR.

All biomarker transcripts, with the exception of adenosine deaminase 2 (*CECR1*), were downregulated in DRs compared with ERs. *CECR1* is secreted by antigen-presenting cells, such as monocytes, and leads to the proliferation of T cells and macrophages (36). Other Trinity panel transcripts included those with multiple cellular functions, such as splicing factor 3b subunit 1 (*SF3B1*), which is involved in DNA damage response (37), and quaking homolog (*QKI*), which is involved in pre-mRNA splicing, protein translation, blood vessel development, and apoptosis (38). Downregulation of these transcripts, as well as caspase 8 (*CASP8*), another promoter of apoptosis, may suggest a reduced capacity of cellular maintenance in DRs compared with ERs. Reduced senataxin (*SETX*) levels have been shown to lead to viral resistance through inhibition of antiviral gene expression (39). Lysosomal trafficking regulator (*LYST*) has been shown to control Toll-like receptor-3- and -4-induced endosomal TRIF (TIR domain-containing adapter-inducing IFN- β) signaling pathways (40). Collectively, these results suggest a dysregulation of complex immune signaling networks that predispose an individual with asthma to the late-phase response upon allergen inhalation. A dampened antiinflammatory response was observed through reduced gene expression of the formyl peptide receptor (FPR) family, *FPR1_intron* and *FPR2*, in DRs compared with ERs. *FPR1* and *FPR2* are

G protein-coupled receptors expressed by phagocytic cells, such as monocytes, macrophages, and dendritic cells, which play roles in host defense against pathogens (41). Lower levels of receptor expression of *FPR1* and *FPR2* in DRs suggest a reduced capacity to bind antiinflammatory ligands, thus impairing the resolution of inflammation.

Apart from known genes, uncharacterized transcripts that were constructed *de novo* were also present in the Trinity biomarker panel. These included intronic sequences in genes, such as *FPR1*, *IFRD1*, *TNFRSF10C*, and noncoding RNAs (LOC254896, LOC101927568, and LOC105378945). A previous study indicated that “dark matter” (unassigned functional role or unannotated) RNA represents a significant proportion of total RNA (42) of sequenced reads in normal and neoplastic tissues. The authors found that 50–65% of all nonribosomal, nonmitochondrial RNA was dark matter RNA, suggesting a greater discovery space for identifying novel transcripts associated with disease. Noncoding RNAs are known to regulate gene expression through mechanisms, such as chromatin remodeling, post-transcriptional regulation, and mRNA degradation (43). Given the unexplored territory of noncoding RNAs in the context of asthma pathogenesis, the transcripts identified in the present study may just be the low-hanging fruit, given the depth of sequencing. The functional role of these transcripts at present remains unknown. Given their reproducibility across platforms and cohorts, these transcripts may not be technical artifacts, but rather surrogates of the underlying biology. However, additional work will be required to delineate their biological function.

A prospective study will be required to demonstrate the utility of the Trinity biomarker panel. Briefly, the same inclusion/exclusion clinical criteria will be retained for future subjects. A blood sample will be collected from all subjects and the Trinity biomarker panel will be applied, such that each subject is assigned a probability score that describes the likelihood that the subject will elicit a dual response upon allergen inhalation challenge. Subjects with a score greater than or equal to 0.5 will be retained as potential candidates for entry into clinical trials for new asthma drugs. These subjects will then undergo

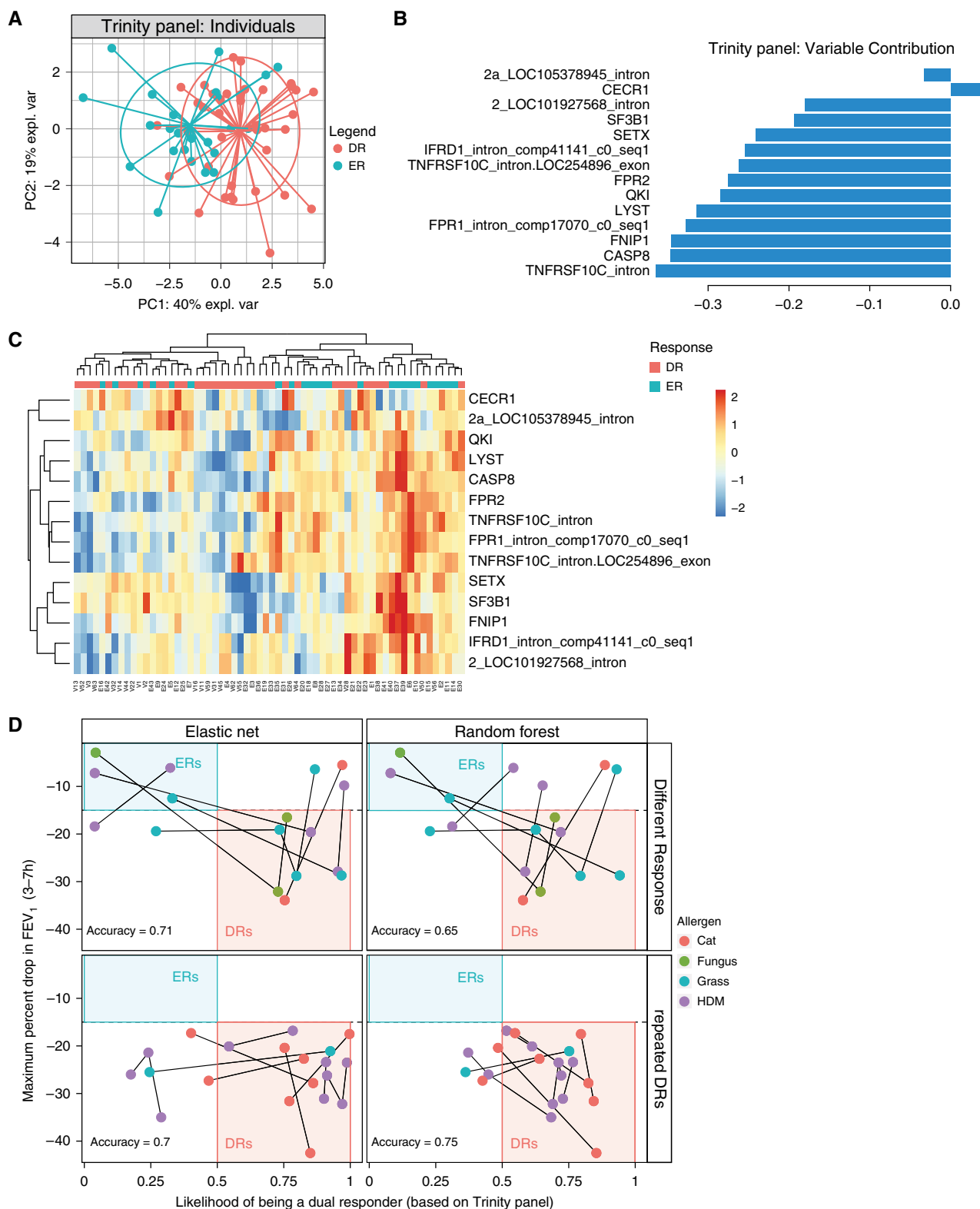


Figure 5. The Trinity biomarker panel. (A) Principal component analysis (PCA) plot of all samples (discovery + validation) using the transcripts of the Trinity biomarker panel. (B) Variable contributions that led to the separation between early responders (ERs) and dual responders (DRs) in the PCA. (C) Heatmap of the Trinity biomarker panel transcripts. All transcripts with the exception of CECR1 are upregulated in ERs compared with DRs. (D) Maximum drop in FEV_1 during the late asthmatic response and the Trinity biomarker panel–based predicted probabilities of repeatedly challenged subjects, using their baseline expression data before allergen inhalation challenge. HDM = house dust mite.

methacholine challenge tests as well as allergen inhalation challenges to determine their response to allergen challenge. This prospective study may demonstrate that fewer subjects are required for the screening process (allergen inhalation challenge) to obtain sufficient numbers of subjects for clinical trials. This study is not only a standard biomarker study, but also depicts the novelty of tapping into the world of noncoding RNA through the use of total RNA and *de novo* transcriptome assembly. Noncoding RNA may serve as novel predictive biomarkers, as well as provide additional insights into the biological mechanisms of asthmatic responses.

The present study uses a rigorous statistical approach in the identification and validation of whole-blood RNA biomarker panels that are predictive of the LAR before allergen inhalation challenge. It may be possible that the “state” of circulating immune cells may predispose individuals with asthma to the LAR. Additional studies that apply these panels to larger cohorts may provide appropriate estimates of positive and negative predictive values. These biomarker transcripts will be further investigated to elucidate the underlying molecular mechanisms of asthmatic responses, and may be used to develop novel asthma therapies. ■

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