Research Article

Differences in the alveolar macrophage proteome in transgenic mice expressing human SP-A1 and SP-A2

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ABSTRACT

Surfactant protein A (SP-A) plays a number of roles in lung host defense and innate immunity. There are two human genes, SFTP A1 and SFTP A2, and evidence indicates that the function of SP-A1 and SP-A2 proteins differ in several respects. To investigate the impact of SP-A1 and SP-A2 on the alveolar macrophage (AM) phenotype, we generated humanized transgenic (hTG) mice on the SP-A knockout (KO) background, each expressing human SP-A1 or SP-A2. Using two-dimensional difference gel electrophoresis (2D-DIGE) we studied the AM cellular proteome. We compared mouse lines expressing high levels of SP-A1, high levels of SP-A2, low levels of SP-A1, and low levels of SP-A2, with wild type (WT) and SP-A KO mice. AM from mice expressing high levels of SP-A2 were the most similar to WT mice, particularly for proteins related to actin and the cytoskeleton, as well as proteins regulated by Nrf2. The expression patterns from mouse lines expressing higher levels of the transgenes were almost the inverse of one another – the most highly expressed proteins in SP-A2 exhibited the lowest levels in the SP-A1 mice and vice versa. The mouse lines where each expressed low levels of SP-A1 or SP-A2 transgene had very similar protein expression patterns suggesting that responses to low levels of SP-A are independent of SP-A genotype, whereas the responses to higher amounts of SP-A are genotype-dependent. Together these observations indicate that in vivo exposure to SP-A1 or SP-A2 differentially affects the proteomic expression of AMs, with SP-A2 being more similar to WT.

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Received Jan 25, 2013; Accepted Apr 28, 2013; Published Jul 19, 2013;
INTRODUCTION

Surfactant protein-A (SP-A) is a member of the collectin family with an N-terminal collagen-like domain and a C-terminal carbohydrate recognition domain (CRD) that binds to many macromolecules, pathogens, and allergens [1,2]. Most mammals possess a single SP-A gene, but humans and primates have two functional SP-A genes (SFTPA1 and SFTPA2). The human SP-A monomer is a 35kDa protein of 248 amino acids. In the coding region, the gene-specific differences of SP-A1 and SP-A2 consist of four amino acids [3,4].

SP-A influences host defense function in a variety of ways. It recognizes and binds to pathogen-associated molecular patterns. These interactions are complex and may involve binding sites in addition to the CRD. Direct interaction with pathogens constitutes one aspect of its host defense function, but SP-A also aids in the clearance of particulates, allergens, and debris [5-9]. SP-A also influences the alveolar macrophage (AM) by regulating many cytokines, including TNF-α, IL-1β, and others [10-20], and cell surface molecules, such as CD11b (CR3), TLR2 and TLR4, the mannose receptor, scavenger receptor A, and CD14 [21-25]. Also, SP-A helps maintain redox balance [26-30], enhances phagocytosis by AM [31-34], contributes to bacterial killing [35-37], affects the development of dendritic cells [38], and links innate and adaptive immunity [39]. Despite these diverse functions, gaps remain in our knowledge of how SP-A influences host defense and the cell types it affects, especially under basal or unstimulated conditions.

Several studies have identified functional differences between SP-A1 and SP-A2 in innate immune functions (including many of those mentioned above), and in several surfactant-related functions. These included cytokine production [12,18,20], modulation of surfactant secretion [40], phagocytosis by AM [33,34,41], and other surfactant structural characteristics [42-45]. Both SP-A1 and SP-A2 are required to form tubular myelin, an extracellular form of surfactant [46]. Differences in the structure and posttranslational modification of SP-A1 and SP-A2 have been observed [47], and it is likely that some of these structural differences are responsible for functional differences [44]. It is of interest that differences in the SFTPA1 and SFTPA2 expression, as assessed by the ratio of SP-A1 to total SP-A, have been reported in human bronchoalveolar lavage (BAL) based on age and lung health [48,49].

The AM, the primary effector cell for lung innate immunity, exhibits a unique phenotype [50] that is influenced by SP-A [21-25,31,32], although the extent of this effect is not fully understood. Previously, by administering exogenous SP-A to SP-A KO mice, we demonstrated that SP-A significantly alters the AM proteome, making it more like that of wild type mice [51]. In this study we hypothesized that the AM proteome is differentially affected by the in vivo presence of SP-A1 or SP-A2 in the alveoli of the lungs of SP-A humanized transgenic (hTG) mice. This hypothesis was investigated by generating various SP-A hTG mouse lines to use in a proteomics study with the two-dimensional difference gel electrophoresis (2D-DIGE) experimental design in order to study the AM proteome among the various mouse lines.

Running Title: The macrophage proteome in humanized transgenic SP-A mice

Key Words: surfactant, lung, 2D-DIGE, collectin, host defense
MATERIALS AND METHODS

Animals and Cells

For this study, we used wild-type C57BL/6 mice obtained from the Jackson Laboratory (Bar Harbor, ME). SP-A KO mice were propagated in the animal core facility of the Pennsylvania State University College of Medicine. Humanized TG SP-A1 and SP-A2 mice that each carried an SP-A1 or SP-A2 variant were generated on the SP-A KO C57BL/6 background as previously described [46]. In the normal human lung SP-A1 and SP-A2 are expressed in type 2 alveolar epithelial cells. The transgenes contain the promoter of SP-C, which is only expressed in type 2 cells. The inclusion of this promoter in the transgenes assures that they too are expressed only in type 2 cells.

All procedures involving animals used protocols that were approved by the Institutional Animal Care and Use Committee at the Pennsylvania State University College of Medicine. All mice were maintained in facilities under pathogen-free conditions or in barrier containment facilities.

AM were obtained by BAL [51]. Cells were washed and counted. A cytospin was prepared, stained, and a differential cell count performed. In all cases AM constituted at least 95% of the cells obtained. The cell-free BAL fluid was frozen for later analysis. All groups consisted of at least 4 mice.

Characterization of humanized TG mice

Southern Blot and Hybridization

To study the copy number of the transgenes in the SP-A1 and SP-A2 hTG mice, total DNA was extracted and purified from the mouse tails and human lung tissue using the DNeasy Blood & Tissue Kit (Qiagen, Valencia, CA). Eight μg of pure DNA from each sample were digested overnight at 37°C with the restriction enzyme EcoRI and the digestions were subjected to 0.7% agarose gel electrophoresis and transferred to a Nylomembrane for Southern blot analysis. Two DNA probes (0.8 and 1.3 kb) amplified from human SP-A cDNA of SP-A2 (1A3) were labeled with digoxigenin (PCR DIG Probe Synthesis Kit, Roche, Indianapolis, IN) and used for hybridization to detect the SP-A transgenes by Southern blot. The 0.8kb probe was generated using primers 2058/2059 and contains the coding region of exons I-IV, and the 1.3kb probe was generated with primers 2058 and 2060 and contains the coding region (exons I-IV) plus a partial 3'UTR sequence which is part of exon IV. The sequences of the probes are listed in Supplementary Table 1. Hybridization was done overnight at 48°C in DIG Easy Hyb Granules solution (Roche, Indianapolis, IN). The membranes were then washed and detected with anti-DIG AP using chemiluminescence and exposed to X-ray film.

Quantitative Real Time PCR

Quantitative PCR (qPCR) was conducted in 10 ml reactions (384-well plates) consisting of 10 ng of mouse genomic DNA, TaqMan SP-A1, SP-A2, or mouse reference assays, and TaqMan master mix (Applied Biosystems, Foster City, CA, USA). Custom-designed TaqMan® Assays were purchased from Applied Biosystems (Foster City, CA, USA) for human SP-A1 (assay Hs01921510) and human SP-A2 (assay Hs00359837). The mouse Trfc TaqMan® Copy Number Reference Assay (Applied Biosystems, Foster City, CA, USA) was used as reference. Gene amplification was achieved using the following protocol: denaturing for 10 min at 94°C, and 40 cycles of 94°C for 15 sec and 60°C for 60 sec. Four replicates were run per reaction. These were monitored with the 7700 Real-Time PCR System (Applied Biosystems, Foster City, CA, USA). A standard curve was also amplified using serial dilutions of the vector used to generate the transgenic mice, ranging from 0.1 to 1000 copies of the transgene. The
mass of one copy of the plasmid was calculated by the formula described by Joshi et al [52].

Copy number determination

Transgene copy number was calculated by the standard curve method [52] as well as with the Copy Caller software (Applied Biosystems, Foster City, CA, USA), assuming the most frequent number of transgene was at least one, since no calibration sample was available [53].

Confirmation of orientation of integrated transgene

Endpoint PCR was performed in 400 ng of mouse genomic DNA with the Fast Start High Fidelity PCR system (Roche Diagnostics, Indianapolis, IN) and specific primers listed in Supplementary Table 1. Primer pairs 1636/1634 were used to detect head to tail concatemers, 2114/1636 to detect tail to tail, and primer 1634 was used to detect head to head concatemers. The following protocol was used: denaturing for 2 min at 94°C, and 40 cycles of denaturing at 94°C for 30 sec, annealing at 59°C for 30 sec, and extension at 72°C for 1 min. PCR products were resolved on 1% agarose and confirmed by sequencing.

SP-A assessment

Initial characterization of BAL fluid was done to confirm the presence of the expected SP-A protein. Blots were prepared as described subsequently and immunostained with either an antibody specific to SP-A1 [48] or an antibody specific for SP-A2 (Aviva Systems Biology, San Diego, CA). Because these two antibodies differed considerably in their relative affinities, these were used for positive identification of SP-A1 and SP-A2 transgenes and not for quantitation.

To determine the amount of secreted hSP-A in the lungs of hTG mice, lungs were lavaged and lavage samples (25 µl of BAL/sample) were subjected to gel electrophoresis (12.5% SDS-PAGE) under reducing conditions. Human SP-A in the BAL fluid was detected by Western blot analysis using a specific SP-A antibody that recognizes both SP-A1 and SP-A2, as described previously [40]. Briefly, proteins were then transferred to nitrocellulose and SP-A detected using a rabbit antibody (IgG) to human SP-A (1:20,000) and goat anti-

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**Supplementary Table 1. Primers used in this study**

| Primer | Sequence (5’- 3’) | Comments |
|--------|-------------------|----------|
| 1634   | CCTTGCCCCGTGCCTATGAG | Human SP-C 5’-flanking region - antisense |
| 1636   | TGAGCAATCTGGCAGCAAGCTA | Human SP-A 3’-UTR - sense |
| 2058   | CTGGAGGCTCTGTTGTTGGG | Human SP-A 5’-UTR – sense |
| 2059   | TCAGAACTCACAGATGGTCA | Human SP-A coding region - antisense |
| 2060   | CTGCCACAGACACCTCAGAGT | Human SP-A 3’-UTR - antisense |
| 2114   | TCTTACATGTCTGATCCCCGG | SV40 sequence – sense |

**Supplementary Table 1.** The sequences of the probes used for the Southern blot and hybridization (Figure 1) and their specificities are listed.
rabbit IgG (horseradish peroxidase-conjugated) secondary antibody. Following enhanced chemiluminescent detection, blots were exposed to X-ray film and densitometry performed on the films using a Bio-Rad GS-800 Densitometer and Quantity One Software.

Proteomics

For the proteomic study we used humanized transgenic mice expressing SP-A1 and SP-A2 at high levels and two mouse lines that expressed SP-A1 and SP-A2 at low levels, in addition to KO mice and WT mice. The SP-A1 mouse lines were 6A2 T1 (subsequently referred to as 6A2) that expressed high levels of SP-A1, and a mouse expressing low levels of SP-A1 that is designated 6A4-LE. The SP-A2 mice consisted of two high expressing mouse lines, 1A0-T10 and 1A0 T13, subsequently called 1A0 mice, and a low expressing mouse line designated 1A0-LE. The T stands for transgene and the number next to it indicates the particular mouse line expressing a given SP-A variant.

To prepare AM for 2D-DIGE frozen AM pellets were lyophilized until completely dry and resuspended in 25 µL of standard cell lysis buffer (30 mM Tris-HCl, 2 M thiourea, 7 M urea, 4% CHAPS, pH 8.5). Protein determinations were done using the Bio-Rad Protein Assay (Bio-Rad, Hercules, CA) and the concentration of protein was adjusted to 1mg/ml for labeling with either Cy3 or Cy5. Twenty micrograms of protein was used for each sample. These procedures and the subsequent 2D-DIGE were performed as we have described in detail [51]. Following electrophoresis gels were scanned using a Typhoon scanner and Gel images were imported into the Progenesis SameSpots v4.0 program (Nonlinear Dynamics) for analysis. For identified proteins having multiple isoforms, the normalized volumes of all isoforms of a given protein were added together and statistical analysis was performed on the totals using Microsoft Excel.

Normalization of data from combined 2D-DIGE experiments

Data were normalized between three independent 2D-DIGE experiments using the KO AM samples present in each experiment. The mean KO AM value was determined for each protein for each of the three experiments. These means were then used to calculate a correction factor for each protein and used to normalize the data for all experiments.

Generation of heat maps

Heat maps were generated in Excel (Microsoft). Means of the normalized values for each protein were subject to conditional formatting with the highest value assigned a red color, middle values a yellow color, lowest values a green color, and intermediate values colored with the appropriate shades depending upon where they fall within the range for that protein. All heat maps included values for lines containing the two highly expressed transgenes, the transgenes expressed at low levels, the WT mice, and the KO mice. For each analysis one group of AM was considered the index group and its proteins were arranged so that they went from high to low expression levels for that group.

RESULTS AND DISCUSSION

SP-A hTG mice and transgene copy number

First, we confirmed the presence of the transgene by Southern Blot (Figure 1) as described in Materials and Methods. Two specific probes were used for validation purposes. The absence of the transgene in both WT and SP-A KO mice is shown in lanes 1 and 2 (negative controls). Genomic DNA obtained from human lung tissue was used as a positive control (lane 11). All genomic DNA from humanized TG mice showed a
positive signal indicating that they contained human SP-A transgene sequences, with several bands and differences in the band intensities being observed among samples. Although these may indicate differences in the copy number of the transgenes, comparison of Southern blot band intensities does not always give accurate and quantitative results [53]. Therefore, we next estimated the copy number by Real
Time PCR, as described in Materials and Methods, and by absolute (standard curve) (not shown) and relative (Copy Caller) quantification (Table 1). Both approaches resulted in similar, if not equal, copy number.

Next we determined transgene orientation, and concatemer occurrence by end point PCR, with primers described in Supplementary Table 1. Supplementary Figure 2 shows specific amplification of head to tail concatemers, consistent with southern blot analysis and our previous work [46]. These were further purified, and confirmed by DNA sequencing. The reference numbers for the variants on NCBI are: 6A² (GenBank: HQ021434.1); 6A⁴ (GenBank: HQ021436.1); 1A⁰ (GenBank: HQ021421.1); 1A³ (GenBank: HQ021430.1).

No bands were detected in negative controls of one copy (human DNA, plasmid DNA of 1A³-T4 and 6A⁴-T3) or controls without the transgene (WT, KO). The presence of head to tail concatemers appeared to result in a band(s) of approximately 6.6kb size shown in Figure 1. The 1A⁰-T13, although it was shown to have multiple copies by PCR, these apparently are not concatemers (Fig. 1, lane 5) because it lacks the intense 6.6kb band. Also no bands were obtained with specific primers to detect head to head or tail to tail concatemers (data not shown). TG mouse lines with a high copy number of a given transgene tended to have a higher level of SP-A expression, although the level of expression and the copy number did not always have a linear relationship (data not shown). This may reflect different sites of integration.

We used some of these mice to study the impact of chronic SP-A variant exposure on the AM phenotype using a proteomic approach. SP-A expression in the mouse lines used in the present study was first assessed by antibodies specific to SP-A1 and SP-A2 to confirm the presence and specificity of SP-A1 and SP-A2 expression in each transgenic line, as described in Methods and Materials. Because the affinities of these antibodies were different we could not use these for quantitative determinations. Thus, we measured total SP-A levels in the BAL of each mouse with an SP-A antibody recognizing both SP-A1 and SP-A2. Total SP-A levels were then measured on 25 μl of BAL/sample. Figure 2

### Table 1. Transgene copy number for SP-A1 and SP-A2 hTG mice

| SP-A1 mice | Copy # |
|------------|--------|
| *6A² T1    | 2      |
| 6A⁴ T1     | 3-4    |
| *KO        | 0      |
| *WT        | 0      |

| SP-A2 mice | Copy # |
|------------|--------|
| 1A⁰ T7     | >10    |
| *1A⁰ T10   | >10    |
| *1A⁰ T13   | >10    |
| 1A³ T2     | 1-2    |
| 1A³ T4     | 1-2    |
| *KO        | 0      |
| *WT        | 0      |

Table 1. T stands for transgene and the numbers next to the T indicate the particular mouse line that expressed a given SP-A variant. The numbers shown in the right hand column depict the copy caller values. The asterisks (*) indicate mouse lines used in the proteomic study. Note that the 1A⁰ mice used in the study were obtained from two different lines (1A⁰ T10 and 1A⁰ T13) that had high levels of expression. The table does not include the low expressors, 1A³-LE and 6A⁴-LE mouse lines used in the present study.
shows a box plot in which the mean values from the immunostained SP-A bands from the 6A² (SP-A1) and 1A⁰ (SP-A2) mice used in this study are plotted on a log scale. Under these conditions levels from the mice referred to as low expressors (6A⁴-LE and 1A⁰-LE) were barely detectable. The Western blot from which these data were obtained is depicted in Supplementary Figure 2. With this protein load the bands for the LE mice are not visible.

Supplemental Figure 2 Western blot of BAL fluid. The figure shows an immunoblot of a gel on which 25µl of BAL fluid from each mouse has been loaded. The gel was immunostained with an antibody that recognizes both SP-A1 and SP-A2. Note that in the 1A⁰ bands, dimeric and trimeric forms of SP-A are visible in addition to the monomer, which is the only band visible in the 6A² samples.

Supplemental Figure 3. Reference gel and protein list. This figure shows an image of the reference gel with identified proteins circled and numbered. The legend contains the names of each identified protein and the functional groups each has been assigned to. These include: actin-related/cytoskeletal proteins; Nrf2-regulated proteins; protease balance/chaperone function proteins; regulation of inflammation proteins; and regulatory/differentiative process proteins. Some proteins are

Proteomics

Overview: We have previously studied the expression of 76 proteins in AM from SP-A KO mice after administration of exogenous SP-A [51]. The reference gel and list of identified proteins are shown in Supplementary Figure 3. These proteins were identified by MALDI-ToF/ToF and in cases where multiple spots were determined to represent different isoforms of a given protein, the values were added together and statistical analysis was done on the sum of the spots. The identified proteins were grouped into five functional groups [51] based on a review of the literature. These groups included:
included in more than one group and some others are not in any group.

In this study we compared SP-A KO mice or WT mice with several mouse lines containing human SP-A transgenes to determine the effects of endogenous expression of the human SP-A transgenes on the AM proteome. The high-expressing SP-A1 (6A²) and SP-A2 (1A⁰) lines used in the proteomic study are marked with an asterisk in Table 1. The two low expressors (LE) for SP-A1 (6A⁴-LE) and SP-A2 (1A⁰-LE) are not shown in Table 1. Mouse lines included those that expressed the human SP-A1 gene, one expressing the 6A⁴ variant at low levels and another expressing the 6A² variant at higher levels, and those expressing the human SP-A2 gene (1A⁰ variant) at low levels and at high levels. In summary, from here on, the SP-A1 and SP-A2 variants used in the proteomics study are referred to as 6A², 6A⁴-LE, and 1A⁰, 1A⁰-LE, respectively.

**Significant changes from KO:** We first compared the transgenic mice to SP-A KO mice in order to determine how the presence of human SP-A in the transgenics altered the proteome of the alveolar macrophage (Table 2). All of the proteins with significantly changed levels are listed in Supplementary Table 2A. As one might anticipate, both of the low expressing (LE) mouse lines had many fewer proteins that changed significantly from KO levels than did lines expressing higher levels. There were 11 proteins that were significantly different in 6A⁴-LE mice than in KO and 8 proteins that were significantly different in the 1A⁰-LE mice. Interestingly, in both 1A⁰-LE and 6A⁴-LE there were four proteins that changed significantly and in similar directions (increased or decreased) including cathepsin D precursor, chitinase 3-like 3 precursor, nucleophosmin 1, and the 4-hydroxylase beta polypeptide precursor. The fact that these changes occurred in both hTG lines suggests that while these proteins may be human SP-A responsive, the responses are not genotype-specific.

AM from mice expressing higher levels of SP-A1 and SP-A2 had many more significant differences from KO, 27 in SP-A1 (6A²) and 19 in SP-A2 (1A⁰), suggesting some degree of dose dependency as compared to the “low expressing” lines. However, the greater number of significant changes vs KO in the 6A² mice as compared to the 1A⁰ mice suggests that SP-A dose dependency (based on the differences in SP-A levels between the 1A⁰ and 6A² lines (Figure 2)) did not have a major influence on the differences between these mouse lines and that these differences were largely variant-specific rather than dose-specific. As in the lower expressing mouse lines, there were some proteins (n=10) that were significantly different from KO in both lines. However, of these ten, while five were similarly regulated in both mouse lines (actin-related protein 2/3 complex, subunit 5; heat shock protein 5 precursor; keratin complex 2, basic, gene 8; prolyl 4-hydroxylase, beta polypeptide precursor; protein synthesis initiation factor 4A), there were five significantly changed proteins that were increased relative to KO mice in one mouse line and decreased in the other (ferritin heavy chain; gelsolin precursor; high mobility group 1 protein; protein disulfide isomerase associated 6; Rab GDP dissociation inhibitor beta). This implied that although the expression of the nine proteins was human SP-A-

| Transgenic line      | vs. KO | vs. WT |
|----------------------|--------|--------|
| 6A⁴-LE (SP-A1)       | 11     | 31     |
| 1A⁰-LE (SP-A2)       | 8      | 34     |
| 6A² (SP-A1)          | 26     | 47     |
| 1A⁰ (SP-A2)          | 19     | 17     |

Table 2. The table summarizes the specific differences seen in each comparison of transgenic mice to either KO or WT mice.
dependent, the directionality of the response (increase vs decrease) was genotype-dependent.

**Significant changes from WT:** We also compared the transgenics to WT mice. In these comparisons, the greater the similarity between a transgenic mouse line and WT, the lower the number of significant differences. Changes may reflect differences in the effects of endogenous mouse SP-A (in WT mice) vs those produced by the expressed human SP-A transgene or differences in the levels of SP-A regardless of type. The benchmark for these analyses is the comparison between WT and SP-A KO mice where there were 21 significant differences out of the 76 identified proteins.

**SP-A2:** We first compared the mice expressing the 1A⁰ variant to WT. The AM of mice expressing higher amounts of SP-A expressed 17 proteins at levels significantly different from those in WT mice. The proteins undergoing significant changes are listed in **Supplementary Table 2B.** This is in sharp contrast to the mice expressing very low amounts of 1A⁰-LE. In the 1A⁰-LE mice there were 34 significantly different proteins as compared to WT. These results suggested that higher levels of expression of 1A⁰ resulted in an AM phenotype that was more similar to that of WT mice compared to that observed with low expressors. In fact, nine of the 17 significantly different proteins in the 1A⁰/

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**Figure 2.** *Values from SP-A immunostained Western blots from mouse BAL.* Identical aliquots from the BAL of mice used for the proteomic study were subject to electrophoresis, immunostained with an antibody to human SP-A (both SP-A1 and SP-A2), and scanned. Means values (dark lines) from the scans are graphed on a log scale with a box plot and individual data points are shown. The immunoblot from which these values were derived is shown in Supplementary Figure 2.
WT comparison were also significantly different in the 1A0-LE/WT comparison, suggesting that the threshold levels of SP-A2 responsible for the effects on AM protein expression were quite low.

**SP-A1:** We then compared each of the two mouse lines expressing SP-A1 (6A2-LE and 6A2) to WT. AM from the transgenic mice that had high amounts of SP-A1 (6A2) expressed 47 proteins at levels significantly different from WT (see Supplementary Table 2B). In the 6A2-LE mice there were 31 significant differences from the WT, indicating that the similarity to WT was greater with lower levels of SP-A1. Of these changes, 24 were present in both groups (high and low SP-A1), although 2 of the 24 were differentially expressed in the two lines. This is in contrast to the findings with 1A0-LE and 1A0 (see above) where the degree of similarity of 1A0 to WT was increased with higher 1A0 levels (i.e. fewer significant differences between 1A0 and WT). This much higher number of significant changes between SP-A1 and WT indicates the SP-A1 transgenes are far less effective, regardless of level, than SP-A2 transgenes in terms of restoring a WT phenotype. Alternatively, it may be that SP-A2 regulates many macrophage proteins in a manner similar to that of native mouse SP-A, whereas SP-A1 may be less effective or may alter the expression of proteins that are not affected by either native mouse SP-A or human SP-A2.

**Heat maps**

**Overview:** In an effort to gain some insight into the functional significance of the changes in protein expression we generated a series of heat maps comparing protein expression in each of the hTG mouse lines, SP-A KO mice, and WT mice. In these maps the highest level of expression among all mouse lines was encoded with a bright red color, the lowest level of expression with a bright green and intermediate levels with yellow. All values were ranked and assigned shades of red, yellow, or green depending on where they fell within the spectrum of expression for each protein.

We organized the map by arranging all 76 proteins in the WT mice on the basis of highest to lowest levels and then compared WT to the other groups (Figure 3). Although this map is very complex there are a couple of interesting features. The first of these is that the map for the 1A0 mice resembles, to some degree, that of the WT mice. This is apparent in the high abundance of high expressing (red) proteins in the top half of the figure and an abundance of low expressing (green) proteins in the bottom portion. The second prominent feature is that the 6A2 mice are probably the most different from WT in terms of protein expression. In the 6A2 mice in the bottom half of the heat map there is a large number of high expression levels (red) corresponding to the lowest levels (green) in the WT. A similar discordance between WT and 6A2 is seen in the top half where highly expressed proteins (red) in the WT correspond to the lowest expressed (green) in the 6A2. The differences between 1A0 and 6A2 mice are further accentuated when a heat map is generated in which expression levels are stratified according to values obtained in the 1A0 mice (Supplementary Figure 4). Thus the observation from this heat map is that the 1A0 is most similar to the WT and 6A2 is the most different.

**Functional group heat maps:** In an attempt to try to simplify the complicated picture obtained following the comparison of all 76 proteins, we studied heat maps of proteins assigned to the 5 functional groups that we have defined in previous studies [51,54]. The protein names, accession numbers, and references supporting the assignment of each protein to a particular group have been previously published [51,54]. These groups
| Gel # | WT | 1A0 | 6A2 | 1A0-LE | 6A4-LE |
|-------|----|-----|-----|--------|--------|
| 1     | 3.474 | 3.470 | 1.982 | 2.841 | 2.648 |
| 2     | 1.855 | 1.853 | 1.383 | 1.814 | 1.678 |
| 3     | 2.119 | 1.781 | 0.849 | 1.862 | 1.130 |
| 4     | 2.129 | 1.721 | 1.281 | 1.439 | 1.161 |
| 5     | 1.428 | 0.920 | 0.792 | 0.778 | 0.771 |
| 6     | 0.707 | 0.516 | 0.380 | 0.390 | 0.308 |
| 7     | 0.087 | 0.058 | 0.058 | 0.020 | 0.048 |
| 8     | 0.028 | 0.026 | 0.026 | 0.026 | 0.026 |
| 9     | 0.022 | 0.022 | 0.022 | 0.022 | 0.022 |
| 10    | 0.020 | 0.020 | 0.020 | 0.020 | 0.020 |
| 11    | 0.018 | 0.018 | 0.018 | 0.018 | 0.018 |
| 12    | 0.016 | 0.016 | 0.016 | 0.016 | 0.016 |
| 13    | 0.014 | 0.014 | 0.014 | 0.014 | 0.014 |
| 14    | 0.012 | 0.012 | 0.012 | 0.012 | 0.012 |
| 15    | 0.010 | 0.010 | 0.010 | 0.010 | 0.010 |
| 16    | 0.008 | 0.008 | 0.008 | 0.008 | 0.008 |
| 17    | 0.006 | 0.006 | 0.006 | 0.006 | 0.006 |
| 18    | 0.005 | 0.005 | 0.005 | 0.005 | 0.005 |
| 19    | 0.004 | 0.004 | 0.004 | 0.004 | 0.004 |
| 20    | 0.003 | 0.003 | 0.003 | 0.003 | 0.003 |
| 21    | 0.002 | 0.002 | 0.002 | 0.002 | 0.002 |
| 22    | 0.001 | 0.001 | 0.001 | 0.001 | 0.001 |

**Figure 3**
represented important macrophage functions and included: actin-related/cytoskeletal proteins; Nrf2-regulated proteins; protease balance/chaperone function proteins; regulation of inflammation proteins; regulatory/differentiative proteins. In each of these heat maps (Figures 4-8), Panel A depicts levels of expression in 1A0 and 6A2 that are stratified by levels in WT mice, and Panel B depicts 1A0 and 6A2 levels arranged by levels of expression in the 1A0 mice, making the differences between these two hTG lines much more evident.

The impression obtained from examining the actin-related proteins (Figure 4A) was similar to that when we examined all proteins. Namely, the overall expression pattern was most similar to WT in the 1A0 mice and most different in the 6A2 mice. The difference between the 1A0 (SP-A2) and 6A2 (SP-A1) mice was further accentuated when we rearranged the heat map according to the levels of expression in the 1A0 mice (Figure 4B). The two protein expression patterns obtained in this case are nearly the inverse of one another. The changes in the actin-related/cytoskeletal proteins may indicate differences in motility and phagocytosis in the AM from the two hTG lines. Indeed, in a study from our laboratory studying the effects of SP-A1 and SP-A2 on phagocytosis in vitro we found that SP-A2 enhanced phagocytosis of Pseudomonas aeruginosa more than SP-A1 did [33,34].

Similar patterns, where the 1A0 resembled that of WT, were also seen among the Nrf-2 regulated proteins (Figure 5A) and the regulatory/differentiative proteins (Figure 6A). The 1A0 heat maps (Figures 5B and 6B) clearly demonstrated the contrast between 1A0 and 6A2 AM. The similarity between WT and 1A0 was less obvious in the protease balance/chaperone function (Figure 7A) and the regulation of inflammation (Figure 8A) groups, although in these two functional groups the markedly different patterns between 1A0 and 6A2 remained readily apparent (Figure 5B-8B). The marked differences in the proteomic profile of each of these functional groups between these two hTG lines lends further support to the idea that these two very similar protein variants have unique functional capabilities and that there are definite differences in phenotypes with respect to their associated functions. The variation in the pattern of proteins regulated by Nrf2 indicates that these two lines may have a very different response to oxidative stress. Similarly, the disparities in the two mouse lines in the protease balance/chaperone function protein group may result in distinctly different abilities to repair damage following injury. This is also likely to be the case with the differences in the proteins involved in the regulation of inflammation that predict altered responses to inflammatory stimuli. Indeed, when we tested the ability of SP-A1 and SP-A2 to induce proinflammatory cytokines in a macrophage cell line we found SP-A2 to be more effective than SP-A1 [18,20]. Confirming these

Figure 3. **Heat map of all proteins.** A heat map is shown for WT mice and all of the hTG lines used in this study. The protein order is arranged based on levels of expression in WT mice. Red bars have the highest levels, yellow are in the middle of the range, and green are lowest. Intermediate levels are colored with the appropriate shades. The column labeled “Gel#” indicates the number of a given protein in the reference gel (Supplementary Figure 3). The numbers in each bar indicate the average normalized volume for the spot/s representing each protein.
effects and the consequences of the above-described proteomic changes in vivo is an important undertaking for future studies. However, based on the distinct proteomic differences observed in the present study, we speculate that significant differences will be found in vivo between SP-A1 and SP-A2 in terms of their ability to regulate inflammatory processes, responses to oxidative stress, and other functions.

Low expressors: It was interesting to note that although there were many differences in protein expression between AM from mice with higher levels of SP-A1 and SP-A2, a high degree of similarity was observed between mice expressing low levels of SP-A2.
Figure 5. **Heat map of Nrf2-regulated proteins.** Panel A depicts a heat map of the Nrf2-regulated proteins expressed in WT, 1A0, and 6A2 mice stratified according to WT levels. Color scheme is as described in Figure 3. The column labeled “Gel #” refers to the numbers in Supplementary Figure 3. Panel B depicts the same group of proteins arranged according to levels of expression in 1A0 mice.

**Figure 6.** **Heat map of regulatory/differentiative process proteins.** Panel A depicts a heat map of the regulatory/differentiative proteins expressed in WT, 1A0, and 6A2 mice stratified according to WT levels. Color scheme is as described in Figure 3. The column labeled “Gel #” refers to the numbers in Supplementary Figure 3. Panel B depicts the same group of proteins arranged according to levels of expression in 1A0 mice.
Figure 7. **Heat map of protease balance/chaperone function proteins.** Panel A depicts a heat map of the protease balance/chaperone function proteins expressed in WT, 1A<sup>0</sup>, and 6A<sup>2</sup> mice stratified according to WT levels. Color scheme is as described in Figure 3. The column labeled “Gel #” refers to the numbers in Supplementary Figure 3. Panel B depicts the same group of proteins arranged according to levels of expression in 1A<sup>0</sup> mice.

Figure 8. **Heat map of regulation of inflammation proteins.** Panel A depicts a heat map of the regulation of inflammation proteins expressed in WT, 1A<sup>0</sup>, and 6A<sup>2</sup> mice stratified according to WT levels. Color scheme is as described in Figure 3. The column labeled “Gel #” refers to the numbers in Supplementary Figure 3. Panel B depicts the same group of proteins arranged according to levels of expression in 1A<sup>0</sup> mice.
Figure 9. **Heat maps of functional groups.** Heat maps are shown for each functional group of proteins depicting the expression in KO, 1A₀-LE and 6A⁴-LE mice stratified according to KO levels. Color scheme is as described in Figure 3. The column labeled “Gel #” refers to the numbers in Supplementary Figure 3. Panel A shows the heat map for actin-related/cytoskeletal proteins. Panel B shows the heat map for Nrf2-regulated proteins. Panel C shows the heat map for protease balance/chaperone function proteins. Panel D shows the heat map for regulation of inflammation proteins. Panel E shows the heat map for regulatory/differentiative proteins.
found a range of 0.003 to 0.999 (mean 0.21; SD 0.241) [48].

In the present study we compared KO mice to 1A\(^0\) high expressing mice that had been stratified further within this high-expressing group into a higher subgroup and a lower subgroup, as well as the 1A\(^0\) low expressors. When this was done, although in some instances protein expression levels varied proportionally with SP-A levels, in most cases it did not, and only two of these cases involved significant changes from the KO values. It was interesting to note that in several of the examples where protein expression varied with 1A\(^0\) levels, the protein changes with 6A\(^2\) were in the other direction (decrease vs. increase or vice versa). In the only case where protein expression differences from KO in 1A\(^0\) and 6A\(^2\) mice appeared proportional to SP-A levels, the difference vs KO was significant in both 1A\(^0\) and 6A\(^2\) mice. This analysis led us to draw the conclusion that under these basal conditions most, if not all, of the effects observed occur after reaching a given threshold level rather than varying in proportion to SP-A levels. However, the possibility remains that in the presence of an insult (i.e oxidative stress) the much higher levels may better maintain the alveolar macrophage phenotype, since some of the SP-A could be oxidized, and this can reduce its functional activity [33,34].

It should be noted that despite the fact that the 6A\(^2\) mice had lower levels of SP-A than 1A\(^0\) mice did, they exhibited almost 50% more significant changes vs KO, further reinforcing that the different response patterns reflect primarily functional differences between 1A\(^0\) and 6A\(^2\) rather than differences between absolute amounts of SP-A. Further evaluation of the expression data for individual proteins in this study did not show that differences in specific protein levels between the 1A\(^0\) and 6A\(^2\) mice were due to levels of SP-A, but rather due to the specific variant present. This was also supported by the very different response patterns elicited by 1A\(^0\) and

(1A\(^2\)-LE) and SP-A1 (6A\(^4\)-LE) in the heat map stratified by expression patterns in the WT (Figure 3). As pointed out above, there were relatively few significant differences between KO (Supplementary Table 2A) and each of these two low expressing hTG mouse lines.

Both low expressing hTG lines were very different from KO, but very similar to one another (Supplementary Figure 5).

The similarity between these two lines is perhaps best appreciated in heat maps of the different functional groups after stratification of the heat maps based on expression levels in AM from SP-A KO mice (Figure 9A-E). As noted earlier, there were relatively few significant differences from the KO in these low expressing lines (8 in 1A\(^0\)-LE; 11 in 6A\(^4\)-LE). The similarities in the heat maps of the two low expressing hTG lines may indicate that a relatively low level of human SP-A expression, regardless of variant type, is adequate to produce some changes that seem to be independent of genotype, but that when higher levels of SP-A are expressed each variant produces distinct phenotypes.

A potential limitation of this study is the fact that the different SP-A variants were expressed at different levels. However, in normal and diseased humans SP-A levels in the BAL are quite variable as well [48,55,56]. The SP-A levels in BAL from normal healthy humans varied over an order of magnitude, and when samples from diseased humans were included the variation could exceed two orders of magnitude. In addition, the use of different antibodies and standards has resulted in variability in reported BAL SP-A levels among different studies [57]. Moreover, in a study involving 49 normal human lungs we measured SP-A1/total SP-A ratios and

found a range of 0.003 to 0.999 (mean 0.21; SD 0.241) [48].
At the other end of the spectrum, with the low expressing mouse lines, while we were unable to measure SP-A using the same gel parameters as with the high expressing group, the protein expression patterns in these mice was very different from that present in the SP-A KO group. This indicated that even very low levels of SP-A have a profound effect on protein expression, and this expression profile is very different from that seen when the proteins are expressed at higher levels.

Another potential limitation is whether the effects of human SP-A on mouse alveolar macrophages provides a good surrogate for the human lung. While many gaps remain in our knowledge of the mechanism of action of SP-A and the specific receptors involved, similarities in the effects of human SP-A on phagocytosis in human and rat alveolar macrophages [34,41] and mouse alveolar macrophages (our unpublished data) confirm that at least some of the SP-A effects are SP-A-specific, rather than species-specific. Specifically, in terms of phagocytosis, when human, rat, and mouse alveolar macrophages were incubated with bacteria in the presence of SP-A1 or SP-A2, although the macrophages responded to both SP-A1 and SP-A2, in all cases SP-A2-treated macrophages exhibited a higher phagocytic index than those treated with SP-A1. Further evidence is provided by the many studies we have published examining the effects of SP-A and the functional differences between the human SP-A variants [12,18,20,33,34,40,44] in a number of in vitro systems.

**Summary**

In the present study we compared the AM proteome of hTG mice to KO mice and were able to gain insight into changes in protein expression resulting from the presence of the SP-A transgenes. We studied the expression of 76 AM proteins using a 2D-DIGE experimental design. We found that even when SP-A was expressed at low levels the AM proteome differed markedly from that of SP-A KO mice. While there were relatively few significant differences from KO, the similarity in these responses led us to speculate that these low amounts of SP-A might be affecting the AM by acting through a high affinity receptor that was not variant-specific.

On the other hand, when we studied the mice expressing higher levels of SP-A there were more than twice as many significant changes, with greater numbers of changes occurring in the SP-A1 (6A^2) mice. The other difference between these mouse lines that became apparent with the study of heat maps was that these two mouse lines produced AM with dramatically different phenotypes. Examining both the overall protein expression pattern as well as the individual functional groups, clearly showed that the patterns of expression in the 1A^0 and 6A^2 mice were almost the inverse of one another (proteins at their highest levels in 1A^0 were at their lowest levels in 6A^2, and vice versa). Another very interesting feature of the 1A^0 expression pattern was its similarity to the WT pattern. These observations led us to speculate that when high levels of SP-A1 and SP-A2 are present they acted in a variant-specific manner, either through different receptors or by eliciting distinctly different responses through a single receptor.

Further studies are required to determine how these changes in the AM proteome in various humanized SP-A hTG mice affect AM function. On the basis of the similarities between WT and the SP-A2 hTG mice we would predict that SP-A2 transgenics would clear bacteria more effectively from their lungs than SP-A1 mice and that their AM would exhibit increased phagocytic function, as we have previously demonstrated in vitro with AM treated with exogenous SP-A2 [34,41]. However, both SP-A variants must be present in order for tubular myelin to be present in the alveolar space [46] indicating a divergence of function. Of relevance is the fact that the ratio of SP-A1 to total SP-A in humans varies...
as a function of lung disease and age [48,49]. We speculate that many aspects of lung function, particularly with respect to the AM, depend on the relative amounts of SP-A1 and SP-A2 in the alveolus.

CONCLUSIONS

1. AM protein expression patterns differed markedly from SP-A KO in hTG mouse lines generated on the SP-A KO background.
2. The AM protein expression patterns from mouse lines expressing low levels of SP-A1 and SP-A2 were similar to each other but different from KO indicating a lack of variant-specific effects at very low levels of SP-A.
3. SP-A1 and SP-A2 at higher levels exhibited very different AM protein expression patterns indicating variant-specific patterns at higher levels of expression of hSP-A.
4. AM protein expression in SP-A2 hTG mice was similar to WT and the pattern from SP-A1 mice was very different from WT.

ACKNOWLEDGEMENTS

The authors thank Susan DiAngelo for genotyping. Current address for Guirong Wang: Department of Surgery, SUNY Upstate Medical University, Syracuse, NY 13210.

This study was supported in part by R01 ES009882 from the National Institute of Environmental Health Sciences.

Supplementary Table 2A: Summary of all significant changes versus SP-A KO AM.

Supplementary Table 2B: Summary of all significant changes versus WT AM.

Supplementary Figure 4. Heat map of all proteins. A heat map is shown for WT mice, KO mice, and all of the transgenic lines used in this study. The protein order is arranged based on levels of expression in 1A0 mice. Red bars have the highest levels, yellow are in the middle of the range, and green are lowest. Intermediate levels are colored with the appropriate shades. The column labeled “gel#” indicates the number of a given protein in the reference gel (Supplementary Figure 3). The numbers in each bar indicate the average normalized volume for the spot/s representing each protein.

Supplementary Figure 5. Heat map of all proteins. A heat map is shown for WT mice, KO mice, and all of the transgenic lines used in this study. The protein order is arranged based on levels of expression in KO mice. Red bars have the highest levels, yellow are in the middle of the range, and green are lowest. Intermediate levels are colored with the appropriate shades. The column labeled “gel#” indicates the number of a given protein in the reference gel (Supplementary Figure 3). The numbers in each bar indicate the average normalized volume for the spot/s representing each protein.

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