doi: 10.1111/j.1365-2222.2008.02982.x

Clinical and Experimental Allergy, **38**, 957–967

ORIGINAL PAPER **Basic Mechanisms in Allergic Disease**

Human innate immune responses to hexamethylene diisocyanate (HDI) and HDI–albumin conjugates

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Clinical and Experimental Allergy

Summary

Background Isocyanates, a leading cause of occupational asthma, are known to induce adaptive immune responses; however, innate immune responses, which generally precede and regulate adaptive immunity, remain largely uncharacterized.

Objective The aim of the study was to identify and characterize the cellular, molecular and systemic innate immune responses induced by hexamethylene diisocyanate (HDI).

Methods Human peripheral blood mononuclear cells (PBMCs) were stimulated *in vitro* with HDI–albumin conjugates or control antigen, and changes in phenotype, gene and protein expression were characterized by flow cytometry, microarray, Western blot and ELISA. Cell uptake of isocyanate was visualized microscopically using HDI–albumin conjugates prepared with fluorescently labelled albumin. *In vivo*, human HDI exposure was performed via a specific inhalation challenge, and subsequent changes in PBMCs and serum proteins were measured by flow cytometry and ELISA. Genotypes were determined by PCR.

Results Human monocytes take up HDI–albumin conjugates and undergo marked changes in morphology and gene/protein expression *in vitro*. The most significant (P-values 0.007–0.05) changes in mircoarray gene expression were noted in lysosomal genes, especially peptidases and proton pumps involved in antigen processing. Chemokines that regulate monocyte/macrophage trafficking (MIF, MCP-1) and pattern-recognition receptors that bind chitin (chitinases) and oxidized low-density lipoprotein (CD68) were also increased following isocyanate–albumin exposure. *In vivo*, HDI-exposed subjects exhibited a drastic increase in the percentage of PBMCs with the same HDI–albumin responsive phenotype characterized *in vitro* (HLA-DR$^+$/CD11c$^+$ with altered light scatter properties). An exposure-dependent decrease ($46 \pm 11\%$; $P<0.015$) in serum concentrations of chitinase 3-like-1 was also observed in individuals who lack the major (type 1) human chitinase (due to genetic polymorphism), but not in individuals possessing at least one functional chitinase-1 allele.

Conclusions Previously unrecognized innate immune responses to HDI and HDI–albumin conjugates could influence the clinical spectrum of exposure reactions.

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Keywords albumin, asthma, cathepsin, CD68, chitinase, exposure, innate, isocyanate, macrophage, MIF, monocyte

Submitted 5 April 2007; revised 28 February 2008; accepted 29 February 2008

Introduction

Isocyanates, highly reactive chemicals used to make polyurethane, are a leading cause of occupational asthma world-wide [1]. The natural history, clinical symptoms and pathology of isocyanate asthma are similar to common environmental asthma, suggesting that the disease is caused by a similar immune-mediated process [2, 3]. However, prototypic pathologic indices of common environmental asthma differ from isocyanate asthma. T-helper type 2 (Th2) cells, which are crucial to common environmental asthma and proliferate vigorously in response to causative allergens *in vitro*, exhibit comparatively minimal responses to isocyanate [3–6]. Allergen-specific IgE, atopy and elevated total serum IgE, which are strongly associated with common environmental asthma, are not generally associated with isocyanate asthma [7, 8]. Thus, in contrast to common allergens, the cell types and

effector molecules stimulated by isocyanate exposure remain unclear, including those that promote the development of pathogenic hypersensitivity responses vs. apparent 'tolerance'. The characteristic differences between isocyanate and common environmental asthma have led to the hypothesis that (low molecular weight) isocyanate chemicals elicit immune responses fundamentally distinct from those induced by common environmental allergens, which are generally (high molecular weight) proteins [2, 8, 9].

Uncertainty regarding the immunologic basis of isocyanate asthma is due in part to isocyanate's unique chemical reactivity, which presents a theoretical and technical obstacle to mechanistic studies [10]. Isocyanates react rapidly with proteins *in vitro* and *in vivo*, and isocyanate–protein conjugates are believed to represent the major immunogenic form of the chemical [6, 9, 11, 12]. In exposed humans, albumin in the airway fluid is a major target for isocyanate, and isocyanate–albumin conjugates are the only known 'isocyanate antigen' specifically recognized by rearranged antigen receptors of the adaptive immune system (i.e. IgG) [9, 13, 14]. However, as mentioned above, antigenic isocyanate–albumin conjugates do not stimulate vigorous *in vitro* Th2 cellular responses [4, 7]. Instead, limited recent studies suggest that human monocyte responses may play an important role in clinical isocyanate exposure outcome [15, 16].

Monocytes are crucial cells of the innate arm of the immune system that circulate in an immature form, and differentiate into 'tissue macrophages', dendritic cells and alveolar macrophages [17–20]. Monocytes possess the capacity to modulate immune responses by secreting chemokines and processing/presenting exogenous and 'self'-antigens [21–23]. The functional responses of monocytes (and other cells of the innate immune system) are also governed, in part, by germline-encoded pattern-recognition receptors, evolved to detect specific molecular motifs [24].

The present study investigates human monocyte responses to a major commercial isocyanate, hexamethylene diisocyanate (HDI), primarily used in protective spray coatings (e.g. automobiles, aircraft) [10]. Data were derived from *in vitro* experiments on human peripheral blood mononuclear cells (PBMCs), and clinical studies of human subjects before and after a specific inhalation challenge. The findings are discussed in the context of immune mechanisms that may modulate pathologic responses to exposure.

Materials and methods

Human subjects and samples

The study was approved by the Human Investigations Committee of Yale University, and written informed consent was obtained from all the participants. Thirty-one subjects were recruited prospectively from the SPRAY study and Yale's Occupational and Environmental Medicine Clinic in New Haven, CT, USA [25]. Peripheral blood was obtained by venipuncture and mononuclear cells were purified by density gradient centrifugation.

Preparation of isocyanate antigens

HDI (Sigma; St Louis, MO, USA) was conjugated to endotoxin-free human albumin (alb^{Ø}) (Sigma) as described previously, with slight modifications to minimize processing steps that might introduce contaminants (i.e. endotoxin, trace metals, glycerol, etc.) [5]. Briefly, 100 μL of HDI was added to 25 mL of a 0.5% (w/v) solution of alb^{Ø} in tissue-culture grade phosphate-buffered saline (PBS), pH 7.2 (Gibco, Grand Island, NY, USA), and mixed end over end for 2 h. The isocyanate–albumin-conjugated reaction product (HDI–alb^{Ø}) was immediately sterile filtered (0.2 μm) for use in culture. FITC-labelled human albumin (alb^{FITC}) (Sigma) was used to generate fluorescently labelled isocyanate conjugates (HDI–alb^{FITC}), using an abbreviated reaction time (20 min) to prevent quenching of the fluorochrome signal. Adipoyl chloride (Sigma)-exposed, timellitic anhydride (TMA) (TCI America; Portland, OR, USA) and DNCB-conjugated albumin were prepared according to previously published methods, except using endotoxin-free albumin [26–28]. For HDI–alb^{Ø} and TMA–alb^{Ø} conjugates, the molar ratios of chemical : albumin were 36 : 1 and 38 : 1, respectively, based on chemical substitution analysis, performed as described previously [29]. Heat-denatured/aggregated albumin was prepared by placing samples of 0.5% alb^{Ø} in PBS, pH 7.2, in a rapidly boiling water bath for 10 min. Control antigen for all studies was 'mock-exposed' alb^{Ø}, which had been processed in the same manner as the HDI–alb^{Ø} (equal concentration, mixing, filtration, etc.), except omitting the HDI.

In vitro *culture and processing*

Human PBMCs were cultured at 2×10^6 cells/mL in RPMI 1640 media (Gibco) supplemented with 10% autologous serum and stimulated with 500 μg/mL of HDI–alb^{Ø} or control antigen or 5 μg/mL concanavalin A (ConA) (Sigma). An Olympus microscope (Nashua, NH, USA) was used to photograph cultures with a Pentax camera under automatic exposure settings. Cells were cytospun on slides with a Shandon Cytospin (Cheshire, UK) and stained with Diff-Quick (VWR; Bridgeport, NJ, USA), or RNA was isolated using RNAeasy columns from Qiagen (Valencia, CA, USA) and DNA was removed with a kit from Ambion (Austin, TX). In other experiments, cell proteins were solubilized with Cell Lytic-M^{TM} lysis buffer (Sigma) supplemented with a protease inhibitor cocktail (Sigma).

Flow cytometry

Cells were incubated with fluorochrome-conjugated monoclonal antibodies (mAbs) to HLA-DR, CD11c, CD3 and CD19, and data were acquired on a FACscan and analysed with Cell Quest software, from Becton-Dickinson (San Jose, CA, USA).

Immunofluorescence analysis

PBMCs were incubated with 50 μg/mL of alb^{FITC} or HDI–alb^{FITC} for varying amounts of time, washed, fixed with paraformaldehyde (Sigma) and settled onto poly-L-lysine (Sigma)-coated coverslips. In some experiments, cells were incubated with phycoerythrin-conjugated mAbs to CD3 or CD14 (Becton-Dickinson), or a lysosomal stain, Lysotracker-Red DND-99 (Molecular Probes, Eugene, OR, USA), before fixation. Coverslips were mounted with anti-fade mounting media and viewed under a Nikon Microphot-FXA microscope (Melville, NY, USA).

Microarray studies

DNA-free, total RNA, from 5-day PBMC cultures of individual subjects was transcribed to labelled cDNA with the MICROMAX™ TSA labelling and detection kit (Perkin-Elmer; Shelton, CT, USA), according to the manufacturer's instructions. Labelled cDNA was hybridized to a broad-spectrum 4.6 K cDNA microarray (Gene Expression Omnibus repository platform GPL993). Fluorescence hybridization data were acquired on an Axon GenePix 4000A scanner and processed using GenePix software (Molecular Devices Corp.; Sunnyvale, CA, USA). The fold increase in gene expression was calculated based on the ratio of the average median fluorescence intensity of duplicate spots after subtraction of the background signal. A twofold change in expression was statistically significant using Student's *t*-test. Hierarchical cluster analysis and gene ontology (GO) groupings were completed using GeneSpring software (Agilent Technologies, Palo Alto, CA, USA). Data were subjected to Lowess normalization before clustering. The significance of enrichment for genes in GO groups was determined using a Fischer exact test.

Protein studies

Cell proteins were Western blotted as described previously, using mAbs specific for CD68, cathepsin B and Annexin (Santa Cruz; Santa Cruz, CA, USA) [26]. Chitinase-1 enzyme activity levels in culture supernatants were determined as reported previously, based on spectrophotomtric measurements of cleavage of a fluorescent chitin analogue [30]. Chitinase 3-like-1, monocyte chemoattractant protein (MCP)-1 and macrophage migration inhibitory factor (MIF) concentrations, in serum or culture supernatants, were quantitated by ELISA using kits, respectively, from Quidel Corp. (San Diego, CA, USA), BD Biosciences-Pharmingen (San Diego, CA, USA) and R&D Systems (Minneapolis, MN, USA). The statistical significance of changes in chemokine concentrations or enzyme activity was determined using Student's *t*-test or ANOVA.

In vitro *respiratory tract isocyanate exposure*

Human subjects were exposed to 15–20 parts per billion (p.p.b.) of isocyanate for up to 2 h, in an isocyanate exposure chamber as described previously [31].

Genotyping

The Chit-1 genotype was determined by PCR using primers that span the polymorphic region, a 24 bp duplication in exon 10 [32]. PCR with the forward primer 5′-CTC CCT GCA CAG TCA GCT AT-3′ and the reverse primer 5′-TAG GAT GTT TGG CTC CTT GG-3′ amplify an 86 bp fragment corresponding to the normal allele and an 110 bp fragment corresponding to the polymorphic allele.

Results

Characterization of isocyanate–albumin conjugates

The antigenicity of HDI–albumin conjugates used in the present study (HDI–$alb^{Ø}$) was confirmed by Western blot with serum from exposed workers (Fig. 1a). Serum IgG binding to HDI–$alb^{Ø}$ is associated with chemical conjugation, cross-linking and conformational changes in albumin, highlighted by reducing SDS-PAGE, native gel electrophoresis (Figs 1b and c) and previously published studies [33]. [Fig. 1 also shows other 'modified' albumins used in later experiments].

Isocyanate–albumin-induced changes in cell morphology

HDI–$alb^{Ø}$-induced cellular immune responses were studied *in vitro* using PBMCs from individuals participating in a study on HDI exposure health effects. Microscopic examination of cultures (Figs 2a and b) revealed that HDI–$alb^{Ø}$, but not control 'mock-exposed' $alb^{Ø}$, induced striking morphologic changes in PBMCs from all study subjects ($N = 31$). The HDI–$alb^{Ø}$-responsive cells could be purified by plastic adherence (Fig. 2c), and co-expressed high levels of monocyte/macrophage-associated cell membrane proteins (HLA-DR and CD11c) (Fig. 2d), but not T or B cell-specific markers CD3 (Fig. 3c) or CD19 (not shown). The morphologic changes observed under the light microscope were accompanied by a marked shift in side and forward light scatter in flow cytometry studies, consistent with changes in cell shape and complexity (Fig. 2d).

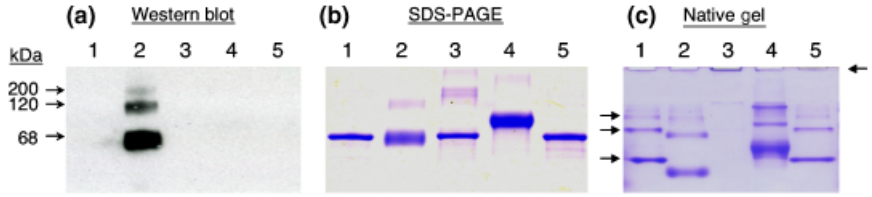

Fig. 1. Characterization of isocyanate–albumin conjugates. Pooled serum IgG from autobody shop workers was blotted (a) against control 'mock-exposed' endotoxin-free albumin (alb^{Ø}) (lane 1), HDI–alb^{Ø} (lane 2), heated/aggregated alb^{Ø} (lane 3), TMA–alb^{Ø} (lane 4) and adipoyl chloride–alb^{Ø} (lane 5). Parallel protein stains of SDS-PAGE (b) and native gels (c) highlight changes in albumin's conformation, as reflected by gel migration. The major band observed under SDS-PAGE is monomeric albumin, while fainter upper bands are likely 'multimers'. Albumin multimers that occur spontaneously (see arrows in lane 1, c), but not those chemically/physically induced (lanes 2–4, b and lane 2, a), are dissociated under disulphide-reducing conditions (lanes 2–4, b).
Note: Western blots with serum from unexposed human subjects were uniformly negative against the conjugates shown above. TMA, timellitic anhydride.

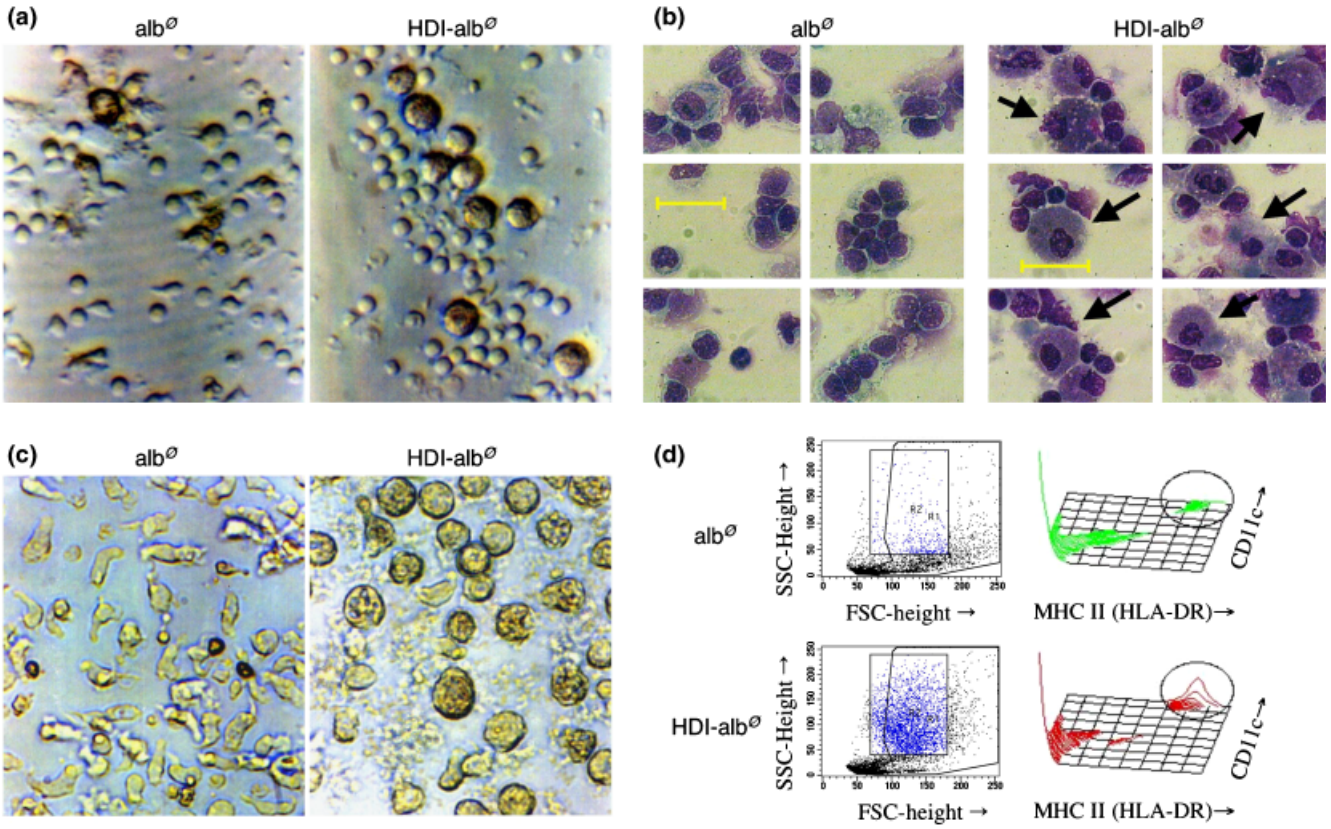

Fig. 2. Isocyanate–albumin-induced phenotypic changes in PBMCs. Cells were cultured with 500 μg/mL HDI–alb^{Ø} or control antigen (alb^{Ø}) for 3 days and photographed under the light microscope at × 400. (a): live cultures, (b): cytospins, (c): live cultures initiated with plastic adherent PBMCs, (d): flow cytometry light scatter and two-dimensional histogram staining for HLA-DR and CD11c of PBMCs. Cells with increased light side scatter are highlighted by the boxed region and correspond to the circled area of the histogram plot. SSC, side scatter; FSC, forward scatter, PBMCs, peripheral blood mononuclear cells; HDI, hexamethylene diisocyanate.

Isocyanate–albumin uptake

To better understand the marked cell morphology changes induced by isocyanate–albumin, we performed additional studies that traced the chemical–protein conjugate's cell uptake and subcellular distribution. HDI conjugates were prepared with fluorescently labelled albumin (alb^{FITC}), which could be visualized directly through fluorescence microscopy. As shown in Fig. 3a, HDI–alb^{FITC} uptake occurred within 30 min of culture, the time at which it was localized around the cell membrane. Over time (72 h), HDI–alb^{FITC} accumulated deeper inside the cytoplasm, with a staining pattern suggestive of localized sequestration. In contrast, little or no staining was observed in cultures incubated with control antigen, 'mock-exposed' alb^{FITC}. The cells that take up high levels of HDI–alb^{FITC}

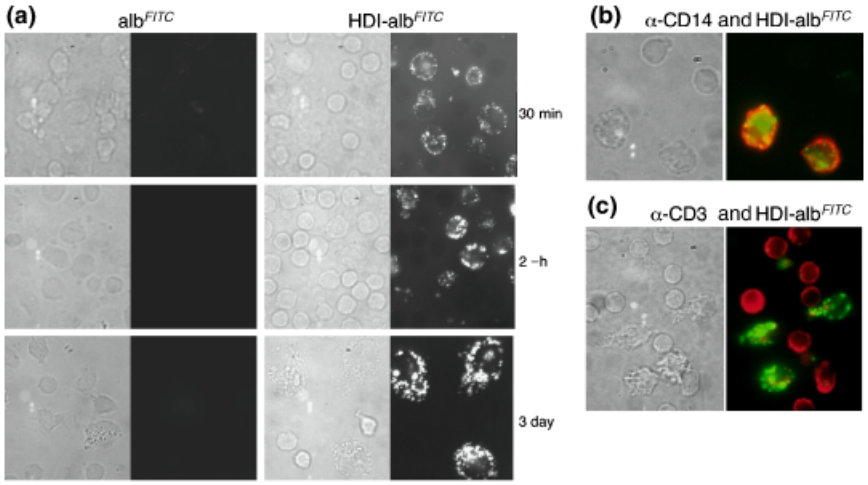

Fig. 3. Isocyanate–albumin is taken up and accumulates in cells of monocyte lineage. PBMCs were cultured with fluorescently labelled control 'mock-exposed' albumin (alb^{FITC}) or HDI–alb^{FITC}, and analysed under phase contrast (left frame) and fluorescence (right frame) microscopy at different time points shown (A). HDI–alb^{FITC}, but not alb^{FITC}, is taken up along the outer cell membrane in a subset of cells, and accumulates over time. Monocyte-derived cells (B), but not T cells (C), are the major cell types that take up HDI–alb^{FITC} (green), as shown by dual staining with PE (red)-labelled α-CD14 or α-CD3, at 2-h and 3-day time points, respectively. PBMCs, peripheral blood mononuclear cells; HDI, hexamethylene diisocyanate.

were further phenotyped as CD14$^+$/CD3$^-$, in double-stain experiments (Figs 3b and c).

Isocyanate–albumin– induced changes in gene expression

To help understand the molecular changes underlying the human PBMC response to HDI–alb^{Ø}, we profiled gene expression changes using a broad-spectrum, 4.6 k cDNA microarray. HDI–alb^{Ø}-induced genes were identified relative to control cultures stimulated with 'mock-exposed' alb^{Ø}, and further stratified through hierarchical clustering and GO. As shown (Fig. 4), HDI–alb^{Ø} induced substantial changes in gene expression of PBMCs from each of the four different subjects tested. The most significant changes (*P*-values 0.007–0.05) were consistently observed in lysosomal genes, especially proteases and proton pump subunits associated with antigen processing (Fig. 4b). Several of the HDI–alb^{Ø}-induced genes are also known to be highly dysregulated in genetic lysosomal storage diseases, including chitinase-1 (T94272), tartrate-resistant acid phosphatase (ACP5), and GM2 ganglioside-activating protein (GM2A) [34–36]. Consistent with the microarray data, the cellular lysosomal content was also elevated in cultures of PBCMs stimulated with HDI–alb^{Ø} (Fig. 5).

Isocyanate–albumin-induced changes in cellular proteins

To determine whether HDI–alb^{Ø}-induced gene expression profiles extend to the protein level, to validate specific microarray findings and to begin evaluating the specificity of the response for isocyanate, we performed additional Western blot experiments. Protein levels for two of the genes whose expression was most consistently elevated in microarray experiments, CD68 and cathepsin B (CTSB), were substantially increased by HDI–alb^{Ø}. CD68 was not induced by other stimuli tested, including mitogen or albumin exposed to adipoyl chloride, another reactive chemical of similar size and bi-functionality (Fig. 6a). The HDI–alb^{Ø}-induced cathepsin B protein differed in size from the lysosomally processed fully mature form [37] found in control cultures (Fig. 6a), supporting the gene expression and cytology data (above) suggestive of altered lysosomal activity.

Isocyanate–albumin-induced changes in secreted proteins

We also evaluated isocyanate–albumin-induced changes in monocyte-secreted proteins of particular interest to asthma. The chemokine MIF, an important modulator of asthma severity [38], rapidly accumulated in the culture media of PBMCs stimulated with isocyanate–albumin. Compared with control cultures, MIF levels were increased >2-fold by 6 h (not shown) and continued to increase for at least 72 h (Fig. 6b). Another chemokine, MCP-1, which acts as an autocrine factor for monocytes (and induces histamine release from mast cells) [39] was variably increased at the protein level in culture supernatants; but, was consistently up-regulated at the message level based on RT-PCR (not shown). Chitinase-1, a soluble secreted pattern-recognition receptor, was significantly increased by isocyanate–albumin, based on enzyme activity in PBMC culture supernatants (Fig. 6c), consistent with the microarray data, and morphologic changes (Figs 1 and 2), reminiscent of Gauchers' cells, which are known to be a major source of chitinase-1 *in vivo* [34, 40].

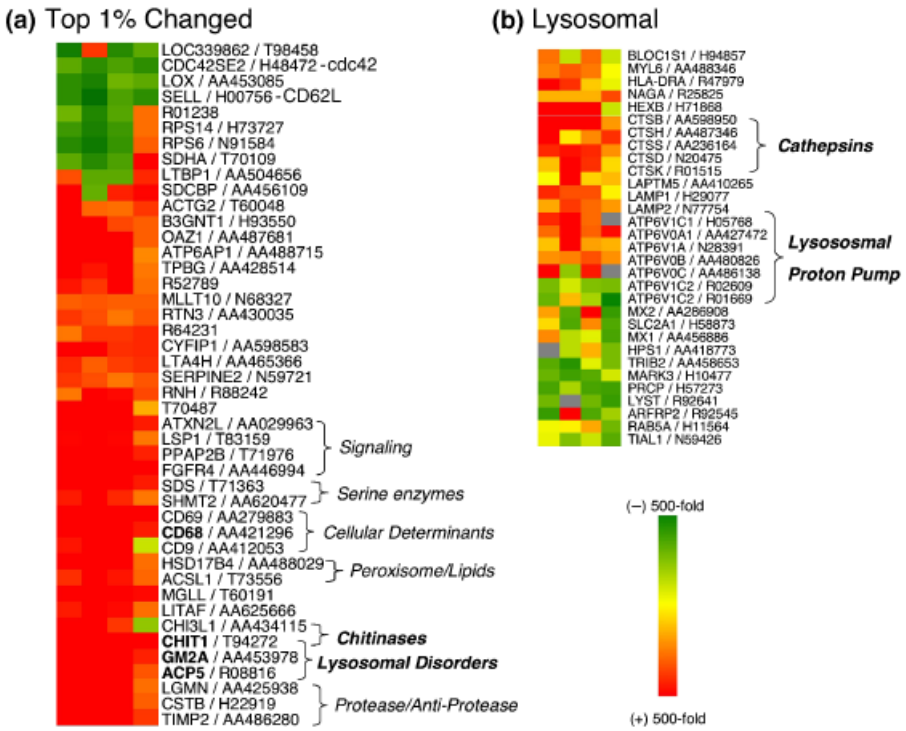

Fig. 4. Isocyanate–albumin-induced changes in gene expression. Cluster analysis of genes most up-/down-regulated in PBMCs cultured with HDI–albumin for 5 days. Colour and intensity represents the fold change in cDNA microarrays (4.6 k) comparing cultures stimulated with HDI–alb^{Ø} vs. control 'mock-exposed' alb^{Ø}. (a): top 1% of affected genes based on fold change in expression. (b): lysosome gene ontology subset, GO:0005764 – Functional Cluster-Cellular Component: cell: intracellular: cytoplasm: vacuole: lysosome; one-third (29/95) of genes (those > 2-fold change in expression) are shown. PBMCs, peripheral blood mononuclear cells; HDI, hexamethylene diisocyanate.

Specificity of the In vitro *human response to isocyanate–albumin conjugates*

Additional experiments were performed to determine whether *in vitro* responses to HDI–alb^{Ø} were HDI-dependent, or merely reflected biophysical/biochemical changes in albumin (solubility, conformation, aggregation), known to influence any proteins' immunogenicity [41]. Heat-denatured/aggregated albumin, as well as albumin conjugated with other reactive, asthma-causing chemicals (see Fig. 1), were compared with HDI–alb^{Ø} for their ability to induce chemokine and chitinase-1 secretion. As shown in Fig. 7, HDI–alb^{Ø}, but not heat-denatured/aggregated alb^{Ø} or TMA–alb^{Ø}, induced marked MIF, MCP-1 and chitinase-1 production. HDI–alb^{Ø} was also distinct from DNCB–alb^{Ø}, which induced MIF (consistent with previous reports) [28] but not chitinase-1 or MCP-1. Thus, the human cellular response to HDI–albumin exhibits HDI specificity and does not reflect non-specific consequences of biochemical/biophysical changes, or chemical conjugation.

Monocyte activation In vitro *in exposed subjects*

To begin to determine whether isocyanate exposure, *in vivo*, has effects on human monocytes, similar to those observed *in vitro*, we studied human subjects exposed to HDI under controlled laboratory settings. Human volunteers breathed for 2 h from a chamber, conditioned with 15–20 p.p.b. HDI, the short-term exposure limit recommended by the NIOSH. Following the exposure period, cells with the HDI responsive phenotype defined *in vitro* (Fig. 1d) were quantitated in the peripheral blood by flow cytometry. As shown (Figs 8a–c), HDI exposure induced a drastic, 3–5-fold increase in the percentage of HLA-DR^{+}/CD11c^{+} cells with altered light scatter in the peripheral blood of exposed individuals ($N = 5$), which returned to near-baseline levels by 24 h post-exposure.

In vitro *chitinase response to isocyanate exposure*

Based on our microarray findings, we further explored the potential *in vivo* effect of isocyanate exposure on chitinases, a family of innate immune proteins that bind chitin, a polysaccharide found in fungi, parasites and nematodes, but not in mammals [40, 42]. We focused on two different chitinases: chitinase-1, the major human chitinase, which possesses true chitin-degrading capabilities [40, 43], and chitinase 3-like-1 (HC-gp39, YKL-40), which binds, but cannot cleave chitin [44, 45]. An 'inactivating' polymorphism exists in chitinase-1 and individuals with two polymorphic alleles (~6% of the

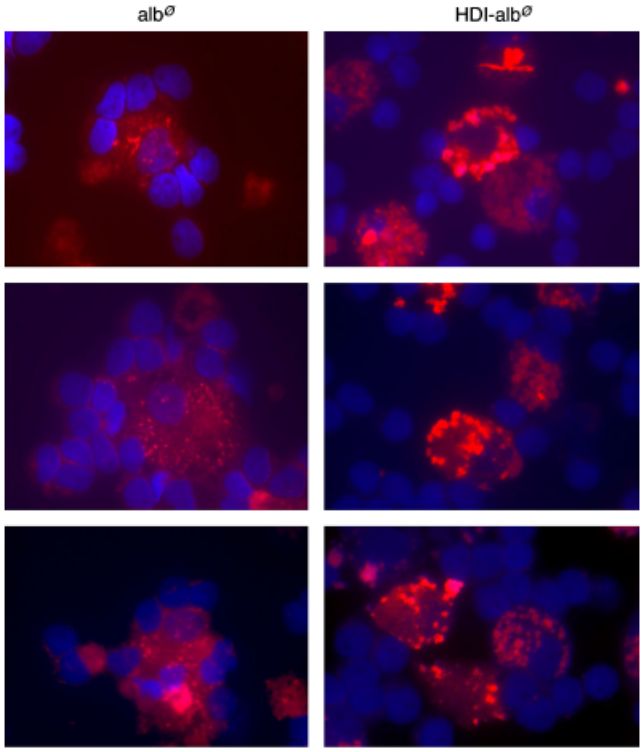

Fig. 5. Isocyanate–albumin-induced increases in lysosomal content. PBMCs cultured with 500 μg/mL HDI–alb$^{\emptyset}$ or control antigen (alb$^{\emptyset}$) for 5 days were incubated with a lysosomal membrane stain (red/pink) and counterstained with DAPI (blue) before settling onto poly-lysine-coated coverslips. Representative fields were photomicrographed at × 200. PBMCs, hexamethylene diisocyanate; HDI, hexamethylene diisocyanate.

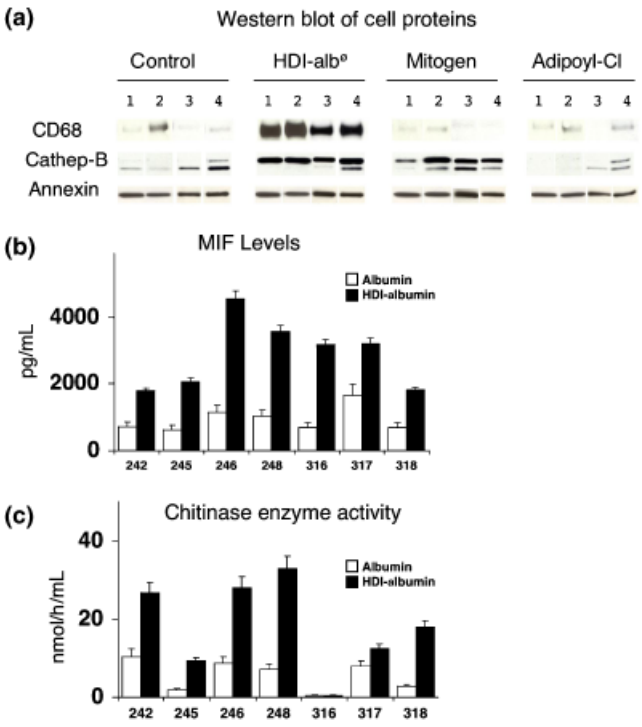

Fig. 6. Isocyanate–albumin-induced changes in intracellular and secreted proteins. (a) Presents a composite of representative Western blots of cultures of PBMCs from four different subjects (lanes 1–4). Equivalent amounts of cell protein from PBMCs cultured with 500 μg/mL of 'mock-exposed' alb$^{\emptyset}$ (control), HDI–alb$^{\emptyset}$, mitogen or adipoyl-chloride for 5 days were blotted with mAb against CD68, cathepsin B and annexin 1. (b): MIF levels (pg/mL) in the culture media of PBMCs from seven additional human subjects (identification numbers 242–318 shown on *X*-axis) similarly stimulated with 500 μg/mL of 'mock-exposed' alb$^{\emptyset}$ (open bars) or HDI–alb$^{\emptyset}$ (filled bars) for 72 h. (c): Chitotriosidase enzyme activity (nmol/h/mL) on day 5 of the same cultures. PBMCs, hexamethylene diisocyanate; HDI, hexamethylene diisocyanate; mAb, monoclonal antibody.

population) are deficient in the enzyme's activity throughout their body [32]. Chitinase 3-like-1 is thought to play a role in tissue remodelling, and serum levels have recently been associated with human asthma severity [46].

In the present study, chitinase-1 enzyme activity and chitinase 3-like-1 protein concentrations were measured in serum before and 24 h after a specific inhalation challenge with HDI. In the majority of subjects ($N = 6$), there was no significant change in chitinase-1 or chitinase 3-like-1 following exposure (Figs 9a and b). However, in a subset of subjects who were chitinase-1 deficient due to genetic polymorphism ($N = 4$), isocyanate exposure caused a $46 \pm 11\%$ ($P < 0.015$) decrease in serum concentrations of chitinase 3-like-1 (Fig. 9b).

Discussion

The present studies demonstrate that human monocytes, important cells of the innate arm of the immune system, respond to hexamethylene diisocyanate in a manner that could modulate exposure outcome. *In vitro*, HDI–albumin conjugates were shown to be taken up by monocytes, and stimulate marked changes in morphology, phenotype and gene/protein expression. In clinical studies, respiratory

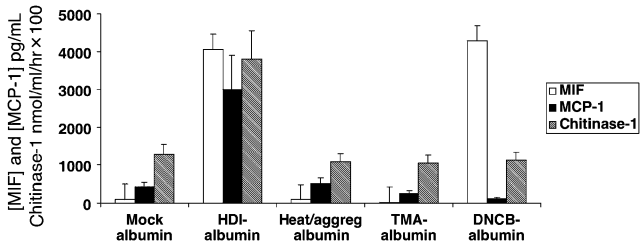

Fig. 7. Specificity of the human innate immune response to isocyanate–albumin. PBMCs from non-isocyanate-exposed individuals ($N = 4$) were cultured with 500 μg/mL of control 'mock-exposed' alb$^{\emptyset}$, HDI–alb$^{\emptyset}$, heated/aggregated alb$^{\emptyset}$, TMA–alb$^{\emptyset}$ and DNCB–alb$^{\emptyset}$. The concentrations of MIF (open bars) and MCP-1 (filled bars) in the culture supernatants at 48 h were measured by ELISA, and chitinase-1 enzyme activity (stripped bars) on day 5 was measured fluorometrically as described in the methods. PBMCs, hexamethylene diisocyanate; HDI, hexamethylene diisocyanate; TMA, timellitic anhydride; MCP, monocyte chemoattractant protein.

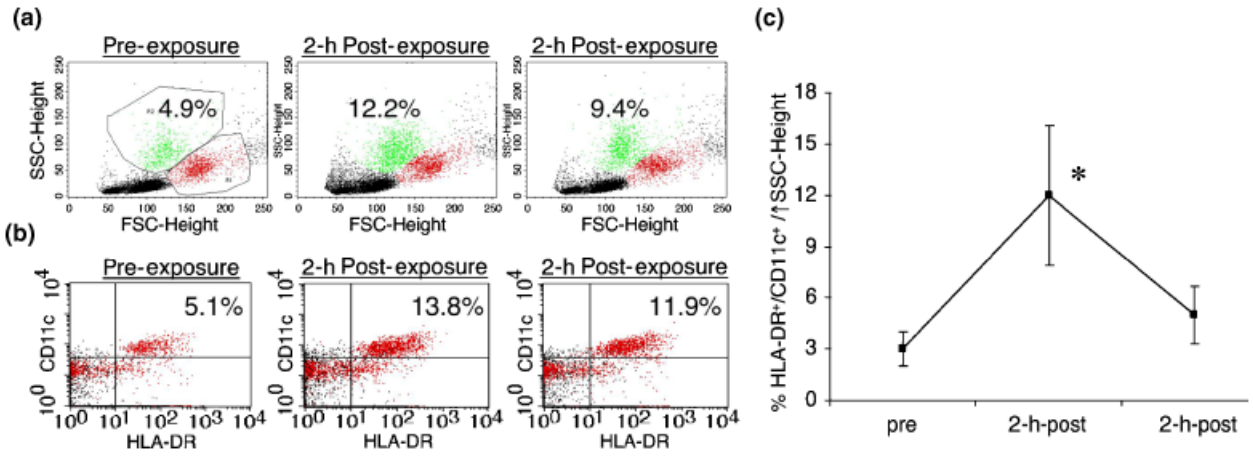

Fig. 8. *In vivo* monocyte/macrophage response to isocyanate exposure. The peripheral blood cells of human subjects exposed to 20 p.p.b. isocyanate (HDI) for 2 h were analysed by flow cytometry. (a) shows light scatter and (b) shows HLA-DR/CD11c dot plots from a representative subject pre-exposure, 2 and 24 h post-exposure. The percentages of cells in the gated region or quadrant are shown. (c) summarizes the percentage of HLA-DR$^+$/CD11c$^+$ cells with shifted light scatter (as gated) in the peripheral blood of five different subjects at each time point (mean ± SE). Pre-exposure and 2 h post-exposure values are significantly different ($^*P < 0.01$). HDI, hexamethylene diisocyanate.

tract exposure to HDI vapours increased the percentage of circulating HLA-DR$^+$/CD11c$^+$ blood cells with the HDI–albumin-responsive phenotype observed *in vitro*. Furthermore, a multi-gene (chitinase)/environment (isocyanate exposure) interaction was observed *in vivo*, which affects the serum concentrations of chitinase 3-like-1, a soluble pattern-recognition receptor recently associated with human asthma [46]. Together, the data describe previously unrecognized effects of HDI on the innate arm of the human immune system, which may provide novel insights into HDI's immunogenicity and may help explain unusual features of exposure-induced asthma.

The present findings are consistent with limited reports on human cellular responses to isocyanate published to date, including pronounced effects of cells and cytokines/chemokines other than the prototypic, adaptive, Th2 type driven by common environmental allergens. Lummus et al.[16] have shown that isocyanate–albumin conjugates stimulate PBMCs to produce chemokines *in vitro*, with profiles typical of monocyte-dominated responses (MCP-1, IL-8, TNF-α), while Bernstein et al.[15] further demonstrated that isocyanate–albumin-induced *in vitro* MCP-1 production by PBMCs was associated with clinical responses to inhalation challenge tests. The present data suggest that chemokine responses represent part of a much broader cellular reaction triggered by isocyanate, which may contribute to the chemical's potent (hyper)-sensitizing capability.

Innate immune responses to isocyanate, as described in this study, could influence exposure outcomes through several different, although not necessarily independent, pathways. Isocyanate-induced chemokine production may create a microenvironment that favours further development of pathologic immune responses. Two

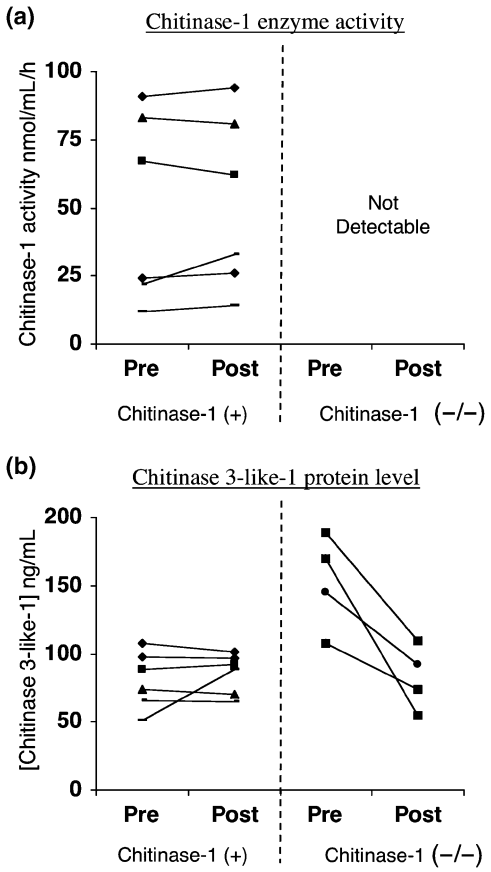

Fig. 9. *In vivo* effect of isocyanate exposure on chitinases. Chitinase-1 enzyme activity levels (a) and chitinase 3-like-1 protein levels (b) in study subjects' serum were measured pre- and (24 h) post-respiratory tract exposure. Each line represents measurements from a single individual, with at least one active ($^+$) chitinase allele ($N = 6$) or two copies ($^{-/-}$) of the non-active (polymorphic) allele ($N = 4$, the only four subjects in our study population with this genotype)

isocyanate–antigen induced chemokines, MIF and MCP-1, have well-characterized pro-inflammatory properties, and contribute to the airway pathology observed in common environmental asthma [38, 39]. Isocyanate-induced up-regulation of germline-encoded (innate immune) pattern-recognition receptors may also alter secondary 'signals' that influence antigenic responsiveness [24]. Two isocyanate–antigen-induced pattern-recognition receptors (CD68 and chitinases) bind ligands (oxidized LDL and chitin, respectively) that evoke potent immunostimlatory responses [40, 47, 48]. Isocyanate's effects on lysosomes, the cellular compartment where exogenous antigens are processed, might further modulate adaptive responses [23]. General disruption of lysososmal equilibrium underlies and/or contributes to the activity of some well-known immunomodulatory drugs/agents (e.g. lysosomotropic amines), as well as the classical concept of lysosomal stabilization/labilization as a mechanism of immune regulation [49, 50]. Recent studies have identified the specific lysosomal proteases and proton pumps induced by isocyanate antigen (e.g. cathepsin B, legumaine a.k.a. AEP) as important determinants of 'antigen fate', differentially expressed in functionally distinct antigen-presenting cells (macrophage vs. dendritic cells) [51, 52]. The present data suggest that lysosomal effects may represent a previously unappreciated mechanism by which isocyanates might stimulate the human immune system. Thus, innate immune responses to isocyanate could influence isocyanate's antigenicity in multiple ways that differ from other common allergens.

Isocyanate's effects on the innate arm of the human immune system may at first seem surprising, given the modern history (1930s) of isocyanate usage, and the prevailing theory that the innate immune system evolved to specifically detect pathogen-associated molecular patterns (PAMPs) [10, 53]. However, the data fit well with recent hypotheses suggesting an overlap between PAMPs and damage-associated molecular patterns (DAMPs), based on shared hydrophobic moitifs, as well as what is known about isocyanate's chemistry [53, 54]. Commercially used isocyanates are highly hydrophobic, comprised almost entirely of carbon and hydrogen [10]. Their bivalent nature and intra-molecular cross-linking ability are also known to alter the 'normal' structure of albumin, perhaps exposing stimulatory hydrophobic regions ('hypos') or creating other 'damage' moieties that might act as innate immune signals [33, 54]. Enzymatic or hydrolytic metabolism of isocyanate might yield derivatives similar to other immunomodulatory agents described above (lysosomotropic amines) [10, 50]. Additionally, isocyanate conjugation raises the isoelectric point of albumin, and increases its sensitivity to (acid) precipitation at lysosomal pH (unpublished observations). Thus, multiple characteristics of isocyanates could contribute individually or coordinately towards their ability to stimulate innate immune responses.

A major strength of the present study is the use of human subjects (PBMCs, serum, albumin), rather than laboratory animals, given well-recognized species-specific differences in gene polymorphisms/mutations and expression, as well as isocyanate responsiveness. For example, in contrast to humans, murine macrophages do not express chitinase-1 and lack the inactivating polymorphism of the gene. Furthermore, chitinase-1 over-producing cells, observed in Gauchers disease, a genetic human lysosomal storage disorder, exhibit striking morphologic similarities to HDI-responsive cells, but are not known to occur in other species. Chitinase 3-like-1, which has recently been linked to human asthma, possess several polymorphisms in humans; however, its expression, activity, disease association and genetics in other species are less understood [46]. Another strength of the present study was the use of a specific inhalation challenge for *in vivo* exposures, which allowed precise dosing and timing of sample acquisition in relation to exposure. The use of human subjects for the present study did, however, impose certain limitations, particularly for *in vivo* studies, which required intensive safety precautions, specialized well-calibrated equipment, physician and industrial hygienist oversight [55, 56]. Safety and ethical concerns precluded human subjects from repeated exposures, and limited the dosage, timing and available tissue samples/subjects. Thus, potential differences in human monocyte responses between subjects with different levels of exposure and/or isocyanate sensitivity could not be deduced from the present data. Future studies will address this provocative possibility, but will require substantial subject characterization and selective recruitment, beyond the scope of the present study.

In summary, we have identified and characterized the effects of HDI and HDI–albumin conjugates on the innate arm of the human immune system. The data highlight lysosomal responses, known to affect antigen processing, as well as a family of pattern-recognition receptors (chitinases), recently linked to asthma. These newly defined HDI-induced responses provide unique perspectives on the chemical's immunogenicity and suggest several mechanisms by which innate immune responses to exposure could modulate the development of allergy/asthma. Future studies will address whether qualitative individual variation in the innate immune response to HDI influences the outcome of occupational exposure and/or helps explain potential differences between isocyanate asthma and prototypical 'atopic'/IgE-dependent asthma.

Acknowledgements

This work was supported by grants from the National Institute of Health (HL62622, M01RR00125), National Institute of Occupational Safety and Health (OH3457) and National Institute of Environmental Health Sciences

(ES00355). The authors acknowledge Carole Holm, RN, Frank Walsh and Dr You-cheng Liu for help with patient recruitment and isocyanate chamber operation.

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