




# The Role of Heat Shock Protein 70 kDa in Asthma

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**Abstract:** Asthma is a complex chronic disorder of the airways, affecting immune and structural cells and inducing both protein and tissue remodeling. Heat shock proteins 70 kDa (HSP70s) are highly conserved members of the stress-induced family, possessing precisely described chaperone activity. There is growing evidence of a tight relationship between inflammatory diseases of different origins and the elevation of local HSP70 expression and secretion. Although extracellular HSP70 does not serve as a common marker of asthma, elevated HSP70 levels have been detected in the peripheral blood serum and sputum of patients with asthma, as well as in the bronchoalveolar lavage fluid of mice with induced allergic airway inflammation. Possessing diverse immunomodulating properties, extracellular HSP70 can manifest different activities in airway inflammatory processes and asthma, acting either as a pro-inflammatory trigger, or an anti-inflammatory agent. This review will discuss the effects and possible mechanisms concerning HSP70 implication in airway inflammation regulation in asthma. We examine ATPase and chaperone activities of HSP70 as potential modulators of immune responses in asthma. Given the crucial role of a chronic inflammatory response in asthma, understanding the effects of HSP70 on immune and structural cells may reveal new perspectives for the therapeutic control of inflammation.

**Keywords:** heat shock protein 70 kDa, asthma, allergic airway inflammation, immune cells, ATP

## Introduction

Asthma is a chronic inflammatory disease that includes several diverse phenotypes traditionally divided into “allergic” and “nonallergic” forms.<sup>1</sup> Recently, based on molecular mechanisms, asthma phenotypes were combined into two basic endotypes: “T2-high” and “Non-T2”.<sup>2</sup> The first is associated with the T helper 2 (Th2)-mediated response, and the other with a non-Th2-mediated response.<sup>2</sup> T2-high and allergic asthma are also associated with eosinophilia and can be referred to as “eosinophilic,” while non-allergic asthma is mostly associated with neutrophil-mediated inflammation. However, cases of non-allergic eosinophilic asthma have also been reported.<sup>1–3</sup>

Each endotype is characterized by different cytokine sets. Thus, allergic asthma is mediated by IL-4, IL-5, IL-9, and IL-13, while the dominant cytokines in non-allergic asthma are IL-1b, IL-8, IL-17, and IL-23.<sup>3,4</sup> However, some features are common to both endotypes. The initiation of inflammation is driven by tissue cytokines (TSLP, IL-25, and IL-33), and both allergic and non-allergic asthma switch to chronic inflammatory processes, which manifest with structural cell damage, followed by airway remodeling.<sup>1–4</sup> Besides the typical cytokines associated with asthma, some damage-associated molecular patterns (DAMP), including the 70-kDa heat shock protein (HSP70), reportedly play an essential role in asthma.<sup>5</sup>

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The HSP70 is a family of chaperone proteins, that consists of 13 members in humans: HspA1A, HspA1B, HspA1L, HspA2, HspA5, HspA6, HspA7, HspA8, HspA9, HspA12A, HspA12B, HspA13, HspA14.<sup>6</sup> Eight of them are expressed constitutively, but HspA1A, HspA1B, HspA5, HspA6, and HspA14 are stress-inducible.<sup>6</sup> Each HSP70 member has a definite intracellular location, and a number of these proteins were also found in the extracellular compartment, mostly in exosomes.<sup>6</sup> All HSP70 members had some alternative names that were used before the unique nomenclature adoption.<sup>7</sup> Hereafter the mentioned above HSP70 family members of humans and other species are referred to HSP70 if otherwise not purposely indicated by the authors.

Even though circulating HSP70 does not serve as a specific marker of asthma, it has been observed at high levels in the sputum and serum of patients with asthma, as well as in the bronchoalveolar lavage (BAL) fluid of mice in a model of induced allergic airway inflammation.<sup>8,9</sup> Several reports demonstrate both the pro-inflammatory and anti-inflammatory activity of HSP70 during the course of inflammation.<sup>10–14</sup> However, a lot is yet to be understood. This review will discuss the effects of HSP70 in the context of asthma and airway inflammation.

## HSP70 as a Potential Trigger of Asthma

Several members of the HSP70 family detected in other species are reported as allergens (Table 1). This relation to allergens was determined, based on the presence of antibodies to these proteins in the peripheral blood sera of patients with allergies. Most HSP70 allergens are listed as inhaled allergens, except Mala s 10 from *Malassezia sympodialis* yeast, which colonizes the skin, and Aed a 8 from

the *Aedes aegypti* mosquito. Antibodies to Aed a 8 have been identified in allergic individuals with asthma and/or rhinitis, and those sensitized to mosquitoes.<sup>15</sup> Despite the high homology among proteins of the HSP70 family, none is reported as an auto-allergen.<sup>16</sup> Recently, HSP70 from Chinese birch pollen extract was found to be the dominant protein that induced a strong specific IgE-response in a mouse model of birch pollen airway inflammation, suggesting it may play a role in Chinese birch pollen allergy.<sup>17</sup>

The antibodies that recognize recombinant human HSP70 have been found in the plasma of patients with asthma. Moreover, correlations were observed among asthma severity, increased IgE and IL-4 levels, and IgG to self (human) HSP70.<sup>18</sup> Based on these findings, Hosseini and coauthors<sup>19</sup> performed a clinical trial, in which they claimed that anti-HSP70 is a novel risk factor for asthma. They showed that anti-Hsp70 antibodies (IgG, IgA, and IgM) were elevated in the peripheral blood of placebo-treated patients with asthma. Signs of HSP70 involvement in allergic diseases have also been identified in patients with allergic dermatitis. In particular, serum Mala s 10-specific IgE, derived from patients with allergic eczema or allergic dermatitis, showed high levels of cross-reactivity to self HSP70.<sup>20</sup>

Thus, in accordance with the information above, upon inhalation, some exogenous proteins from the HSP70 family can promote allergic airway inflammation in sensitive people. Moreover, the data about the cross-reactivity of anti-HSP70 antibodies to self HSP70 in atopic patients<sup>20</sup> give evidence that under certain conditions, exogenous HSP70 can trigger reactivity to self HSP70. One possible reason for this is the highly conserved structural sequences of exogenous allergic proteins of the HSP70 family and self HSP70 proteins.

**Table 1** HSP70 Allergens of Different Species

HSP70 Allergens	Species	References
Alt a 3	<i>Alternaria alternata</i> (mold)	Breitenbach & Simon-Nobbe 2002 <sup>113</sup>
Pen c 19	<i>Penicillium citrinum</i> (mold)	Shen et al 1997 <sup>114</sup>
Mala s 10	<i>Malassezia sympodialis</i> (skin colonizing yeast)	Andersson et al 2004 <sup>20</sup>
Der f 28	<i>Dermatophagoides farina</i> (American house dust mite)	Aki et al 1994, <sup>115</sup> An et al 2013 <sup>116</sup>
Der p 28	<i>Dermatophagoides pteronyssinus</i> (European house dust mite)	Liu et al 2018 <sup>117</sup>
Tyr p 28	<i>Tyrophagus putrescentiae</i> (mold mite)	Cui et al 2016 <sup>118</sup>
Cor a 10	<i>Corylus avellana</i> (European hazel)	Gruehn et al 2003 <sup>119</sup>
Aed a 8	<i>Aedes aegypti</i> (yellow fever mosquito)	Cantillo et al 2017 <sup>15</sup>
Cla h Hsp70	<i>Cladosporium herbarum</i> (mold)	Breitenbach & Simon-Nobbe 2002, <sup>113</sup> Zhang et al 1996 <sup>120</sup>
Blo t Mag29	<i>Blomia tropicalis</i> (dust mite)	Simon-Nobbe et al 2008 <sup>121</sup>

**Note:** In accordance with the database AllFam (<http://www.meduniwien.ac.at/allfam/>).<sup>16</sup>

## Elevation of HSP70 Expression and Secretion in Asthma

More than 20 years ago, the increased expression of HSP70 in airway epithelial cells and alveolar macrophages of people with asthma was demonstrated by immunostaining *in vitro*.<sup>21</sup> Further, the elevated expression of HSP70 mRNA was observed in peripheral blood mononuclear cells (PBMC) of patients with asthma during acute episodes.<sup>22</sup> Later, the local and systemic elevation of extracellular HSP70 levels were shown to be correlated with the severity of inflammation in asthma.<sup>8</sup> Furthermore, an association between increased plasma HSP70 levels and airway neutrophilia has been reported.<sup>8</sup> In addition, a correlation between sputum HSP70 level and lymphocyte number has been detected in patients with asthma.<sup>8</sup>

Recently, Huang and coauthors<sup>5</sup> compared the release of DAMP from structural airway cells and infiltrated immune cells in the sputum of patients with asthma, chronic obstructive pulmonary disease (COPD), and those with both asthma and COPD. Compared with the control group, the levels of HSP70 were elevated in the sputum of all three experimental groups (asthma group, COPD group, as well as the asthma-COPD overlap syndrome group). Interestingly, multiparametric analysis indicated a correlation between the percentage of neutrophils and the concentrations of HSP70.<sup>5</sup> Elevated levels of extracellular HSP70 in the airways have also been reported in other studies using animal models of allergic airway inflammation. Thus, HSP70-rich exosomes have been identified in the BAL fluid of mice that received the olive pollen allergen 24 h after intranasal allergen application.<sup>23</sup> Interestingly, in this study, to detect HSP70 in exosomes authors used antibodies to Hsc70 (HspA8 according to the unique nomenclature)<sup>7</sup> that is known to be constitutive, but not stress inducible.<sup>23</sup> Elevated BAL HSP70 production has also been observed in an ovalbumin-induced asthma model in mice. The kinetics of extracellular HSP70 levels indicated significant elevations at 24 and 48 h after the allergen challenge, compared to control mice.<sup>9</sup>

In summary, the elevated expression (of both the mRNA and protein) and secretion of HSP70 are associated with asthma, and are mostly detected in the acute phase of inflammation or exacerbation.

## Effects of HSP70 in Airway Inflammation

Both pro-inflammatory and anti-inflammatory effects have been reported for HSP70 in airway inflammation.<sup>10,14</sup> The effects of HSP70 can vary based on the origin (self or

exogenous) of the protein and the microenvironment, such as receptors that can bind HSP70, and cells that express those receptors. The study design, such as specific methods of HSP70 expression stimulation or the pathological conditions during HSP70 application, can also influence the effects of HSP70. Moreover, some conditions can affect both inflammation and HSP70 expression independently.<sup>9–14,24–29</sup>

Traditionally, it has been suggested that intracellular HSP70 are cytoprotective chaperones, which when secreted to the extracellular space, possess pro-inflammatory properties and serve as alarmins.<sup>24</sup> Thus, besides the potential immunogenicity and allergenicity of exogenous HSP70, extracellular self HSP70 can also exhibit pro-inflammatory effects during airway inflammation, which has been demonstrated *in vitro* using both primary cell cultures and various cell lines. The application of HSP70 to primary human lung fibroblasts can induce IL-8 production.<sup>25</sup> Moreover, using small interfering RNA, researchers have demonstrated that HSP70 are implicated in the induction of IL-8 production, in response to other stimuli, such as cigarette smoke extract.<sup>25</sup>

The treatment of human bronchial epithelial cells (16HBE14o-) with recombinant Hsp72 (HspA1A)<sup>7</sup> induces a dose-dependent increase in *IL-8* gene expression; and pretreatment of cells with polymyxin B to prevent the side effects of endotoxins that contaminate recombinant Hsp72 does not alter the effects of Hsp72.<sup>26</sup> Wheeler and coauthors<sup>27</sup> showed that recombinant Hsp72 treatment leads to increased IL-8 and TNF- $\alpha$  production by both primary human neutrophils and differentiated (by the addition of DMSO) HL-60 cells. The authors also pretreated cells with polymyxin B to prevent the effects of endotoxins. Similarly, elevated expression of TNF- $\alpha$  and KC – the murine functional homolog of human IL-8 – has been observed in murine bone marrow neutrophils treated with human recombinant Hsp72.<sup>27</sup> Using THP-1 cells differentiated by phorbol-myristate-acetate (PMA) activation into macrophage-like cells, Hulina and coauthors<sup>28</sup> showed dose-dependent and time-dependent production of IL-1 $\alpha$  and IL-8, in response to treatment with recombinant human HSP70 (both HspA1A and HspA1B)<sup>7</sup> containing low levels of endotoxin contamination. Later, the same group demonstrated the production of IL-6 and IL-8 by the human lung epithelial cell line NCI-H292, after incubation with extracellular recombinant HSP70.<sup>29</sup>

Some in vivo experiments have demonstrated the pro-inflammatory activity of HSP70. In one study, BALB/c mice were challenged with a single intratracheal inhalation of recombinant human HSP72, and euthanized 4 h later.<sup>27</sup> The Hsp72 inhalation significantly elevated the production of TNF- $\alpha$  and KC. This in turn, attracted neutrophils to the airways and subsequently, increased levels of myeloperoxidase were detected in the murine BAL fluid, 4 h after Hsp72 application.<sup>27</sup> However, these in vivo experiments did not permit any distinguishing between the effects of Hsp72 and those of endotoxin contaminants. As is well known from the standard ovalbumin-induced asthma mouse model, endotoxin contamination can affect the immune response to antigens. Thus, lipopolysaccharide (LPS)-free ovalbumin administration can induce IL-4, IL-5, and IL-13 secretion and eosinophil infiltration, while LPS-rich ovalbumin application can also increase IL-1b, TNF- $\alpha$ , and KC, as well as the infiltration of both eosinophils and neutrophils.<sup>30</sup>

To clarify the role of Hsp70 in airway inflammation, Yombo and coauthors<sup>31</sup> recently demonstrated that HSP70 double knockout (Hsp70.1/3-/-) mice that lack inducible *HSP70* genes show significantly reduced Th2-mediated airway inflammation. Using the model of allergic airway inflammation induced by *Schistosoma mansoni* soluble egg antigen (SEA), those authors demonstrated that in the absence of inducible *HSP70* gene expression, the total BAL cell number and eosinophil level, as well as IL-4, IL-5, and IL-13 cytokine levels, are significantly reduced. Simultaneously, the levels of neutrophils and lymphocytes in the BAL fluid of HSP70.1/3-/- mice are significantly elevated in response to SEA administration, compared to those in the BAL fluid of wild type mice that received SEA.<sup>31</sup> Notably, sensitization and challenge with SEA did not modify *HSP70* gene expression in wildtype mice. Using bone marrow chimera and in vitro experiments, those authors showed that the loss of HSP70 is critical for Th2 response, and the absence of T cell activation leads to the attenuated airway inflammation observed in HSP70.1/3-/- mice.<sup>31</sup>

Using the same mouse strain (HSP70.1/3-/-) and a model of induced respiratory distress syndrome, a significant increase in lung injury and elevated levels of IL-6 and TNF- $\alpha$  were observed in the absence of inducible HSP70.<sup>32</sup> Thus, besides its pro-inflammatory effects, HSP70 can also display anti-inflammatory activity in the respiratory tract during inflammatory processes. Such activity has also been demonstrated in the murine

model of bleomycin-induced lung fibrosis, where the induction of increased levels of HSP70 inhibits the progression of inflammation and fibrosis.<sup>33,34</sup>

Prado and coauthors<sup>23</sup> detected increased levels of the constitutive form of Hsc70 (HspA8)<sup>7</sup> in BALF-derived exosome that were yielded from mice with induced tolerance to the allergen. Moreover, preventive intranasal immunization of mice with HSP70-rich exosomes blocked the development of olive pollen allergen (Ole e 1)-induced allergic airway inflammation, and particularly, significantly reduced IL-5, eosinophilia, and allergen-specific IgE.<sup>23</sup> Using a mouse model of ovalbumin-induced allergic airway inflammation, the direct protective effect of oropharyngeal HSP70 application in the acute phase of inflammation has been demonstrated.<sup>9</sup> Particularly, the application of murine HSP70 prevents airway eosinophilia and suppresses the allergen-induced elevation of both the total BAL cell number and the levels of pro-allergic cytokines, such as IL-4, IL-5, and IL-13.<sup>9</sup> Notably, in this study, airway inflammation was induced with ovalbumin grade 5, which is contaminated with endotoxin, and therefore provoked mixed eosinophilic and neutrophilic inflammation. Interestingly, the administration of HSP70 during the acute phase of inflammation in mice induced the retention of neutrophils in the BAL fluid, compared with mice affected by induced airway inflammation that did not receive HSP70.<sup>9</sup> In vitro, HSP1A1 prevented neutrophil necrosis or NETosis and facilitated a proapoptotic neutrophil state.<sup>9</sup>

Thus, as it is shown above, several in vivo and in vitro models demonstrate both pro-inflammatory and anti-inflammatory effects of HSP70 in the airways. In the case of pro-inflammatory effects, the following events are suggested: HSP70 stimulates the production of IL-1 alpha, IL-6, and IL-8 by the epithelial cells and alveolar macrophages, which attract neutrophils. The neutrophils in turn produce TNF- $\alpha$  and IL-8 (or KC in mice), which act as autocrine stimulators of the neutrophil-mediated response. Interestingly, in the case of anti-inflammatory activity, HSP70 also affects neutrophils, by preventing their hyperactivation. Regarding allergic inflammation, inducible HSP70 (among which HspA1A, HspA1B, HspA6, HspA7 and HspA14)<sup>7</sup> contributes to direct Th2 activation and the maintenance of a pro-allergic response, including eosinophilia, and elevated levels of IL-4, IL-5, and IL-13. However, exogenously supplied HSP70 down-regulates eosinophilia, and levels of pro-allergic cytokines and allergen-specific IgE. To address the question about



the dualistic effects of HSP70 in airway inflammation, further, we will focus on the molecular and cellular mechanisms of HSP70 action.

## HSP70 Binding Receptors and Their Role in Asthma

One key point in establishing the concept of the immunomodulatory effects of HSP70 is defining the receptor responsible for the recognition of HSP70 by immune or structural cells during airway inflammation. As HSP70 can be involved in interactions with a receptor in a membrane-bound or an extracellular form, in complex with a peptide and in ATP- and ADP-binding states, definition of the unique receptor is quite problematic. Calderwood et al<sup>35</sup> proposed that HSP70 mediates its activity by interacting with several cell surface receptors that can bind HSP70 or the HSP70-peptide complex. To date, these HSP70 binding receptors are represented as CD91, several scavenger receptors that belong to the family of C-type lectin receptors (such as LOX-1 and SCARF1), C-type lectin-like receptor NKG2D, as well as the sialic acid-binding immunoglobulin-type lectins, Siglec-5 and Siglec-14.<sup>35–38</sup> Furthermore, a heterodimeric cluster of TLR2/TLR4 and CD14, with and without the contribution of SCARF1, is reportedly involved in the binding of HSP70.<sup>39,40</sup> A recent study also provided the basis for the interaction between HSP70 and the receptor of advanced glycation end products (RAGE).<sup>41</sup>

With the exception of RAGE, little is known about the role of the above-mentioned receptors in airway inflammation and asthma. At the same time, RAGE is reportedly the “major mediator of pulmonary inflammatory responses.”<sup>42</sup> It is constitutively highly expressed in the lungs of healthy adults, on the airway epithelial, endothelial, and smooth muscle cells, as well as on immune cells, including macrophages, dendritic cells (DCs), eosinophils, T cells, and B cells.<sup>42</sup> The levels of endogenous and soluble RAGE are increased in the sputum samples of patients with asthma.<sup>42</sup> Furthermore, RAGE KO mice produce normal amounts of IL-4, in response to the house dust mite (HDM) but show attenuated IL-5 and IL-13 responses. In addition, RAGE promotes the expression of IL-33 and subsequent accumulation of ILC2s in the lungs of mice at the onset of allergic inflammation.<sup>42</sup> Based on the facts above, we can conclude that although RAGE can mediate the pulmonary allergic inflammatory response, it is not known, whether HSP70-RAGE interaction triggers this

response. The pro-inflammatory activity of RAGE is evidently driven by binding with the RAGE ligand, HMGB1, the expression of which is also elevated in asthma.<sup>42,43</sup> Interestingly, the expression of HMGB1 in asthma is negatively regulated by HSF1, the transcriptional factor responsible for the expression of HSP70.<sup>43</sup> Thus, although RAGE expression is associated with asthma, there is no direct proof that binding of HSP70 to RAGE can exacerbate asthmatic inflammation. Conversely, there is evidence that HSP70 expression can restrict the pro-inflammatory activity of RAGE by downregulating of HMGB1.<sup>42,43</sup>

Expression of CD91 has been observed on some structural cells, such as endothelial and smooth muscle cells and a wide range of immune cells, including DCs, monocytes, macrophages, and B and T lymphocytes.<sup>44</sup> Interestingly, CD91 has been identified as a negative regulator of DC-mediated Th2 immune responses during HDM-induced eosinophilic airway inflammation.<sup>45</sup> Whether HSP70-CD91 interaction involves in such suppression of the allergic immune response is currently unknown, however, it can be a hypothesis for the further research of the mechanisms of immunomodulatory activity of HSP70.

Both SCARF1 and LOX-1 are expressed on activated endothelium and antigen-presenting cells (APCs).<sup>38,40</sup> Recently, the role of C-type lectin receptors in asthma was highlighted.<sup>46</sup> Some HDM or fungal allergens are recognized by C-type lectin receptors, mostly dectin receptors and mannose receptors. However, no direct evidence exists of the implication of SCARF1 and LOX-1 in allergic airway inflammation and asthma. Nevertheless, the expression of LOX-1 on the endothelium affects (slow down) neutrophil turnover, which could be important in the inflammatory process in the airway.<sup>47</sup> These facts suggest the HSP70 interaction with SCARF1 or LOX-1 on neutrophils can be responsible for the neutrophil retention, which was associated with the inflammation resolution in experiments describing the protective effects of HSP70 in the asthma mouse model.<sup>9</sup> Moreover, both SCARF1- and LOX-1-mediated endocytosis of apoptotic cells is necessary step for the resolution of inflammation.<sup>38</sup>

Several Siglecs have been implicated in the pathogenesis of asthma, particularly Siglec-8 (Siglec-F in mice) in allergic asthma, and Siglec-9 in severe neutrophilic asthma.<sup>48–50</sup> Neutrophils also express Siglec-5 and Siglec-14, both of which reportedly bind HSP70. Siglec-5 and Siglec-9 are thought to be inhibitory, while Siglec-14 is an activator that induces the production of IL-8 and

TNF- $\alpha$  upon interaction with sialic acid.<sup>48–50</sup> In accordance with this concept, the regulation of airway inflammation through the interaction of HSP70 with Siglecs has been demonstrated. Particularly, HSP70 stimulation through Siglec-5 acts as an anti-inflammatory signal, while stimulation through Siglec-14 is pro-inflammatory.<sup>37</sup> The facts indicated above support the hypothesis that the effect of HSP70 in airway inflammation depends on both the presence of the certain cell type (e.g. neutrophils or eosinophils) and the current level of the certain Siglec expression on these cells. Such a hypothesis gives the mechanistic explanation of the multidirectional effects of HSP70 in inflammatory processes.

The effects of TLR activation in asthma, particularly TLR2 and TLR4, are dependent on many factors.<sup>51</sup> Thus, multiple studies of TLR4 activation in asthma mouse models have shown either protection against, or exacerbation of asthma after exposure to endotoxins (LPS).<sup>30,51</sup> The effect is dependent on LPS concentration and the route of exposure. In general, the administration of low doses of LPS together with the allergen induces a mixed Th2 and Th17 response, and the infiltration of both eosinophils and neutrophils to the airways; while the administration of high doses of LPS with the allergen induces a strong Th1 response.<sup>51</sup>

Simultaneously, endotoxin-free HSP70 – TLR4 interaction remains a matter of debate. Thus, Hulina et al<sup>28</sup> demonstrated that HSP70, contaminated with a very low dose of LPS, can modulate the effects of LPS and lipoteichoic acid (LTA) interactions with TLR4 and TLR2, respectively. However, other researchers have shown opposite responses to LPS-free HSP70. Thus, opposite to contaminated with LPS recombinant *Plasmodium falciparum* HSP70 obtained in *E. coli* XL1 Blue, LPS-free HSP70 obtained in *E. coli* ClearColi BL21 and in *Brevibacillus choshinensis*, did not stimulate production of IL-6 and IL-8 in murine bone marrow-derived DCs.<sup>52</sup> Furthermore, the addition of LPS-free HSP70 to LPS-activated human peripheral blood monocytes downregulates TNF- $\alpha$  and IL-6 secretion.<sup>53</sup> Notably, in the same study, the downregulation of TLR4 expression under LPS-free HSP70 exposure has not been demonstrated. Other researchers have identified HSP70 as an effector molecule downstream of TLR4 in the inflammatory signaling cascade, and have shown that KC production in response to TLR activation is significantly reduced in B6-*Hspa1a*/*Hspa1b*<sup>tm1Dix</sup>/NIEHS mice deficient in *HSpA1A* and

*HSPA1B* genes.<sup>54</sup> Thus, the controversial effects of TLR4 activation in asthma and controversial data about HSP70-TLR4 interactions require more detailed analysis. Additional studies are necessary to determine a single consistent hypothesis about the effects of HSP70-mediated TLR4 activation in asthma.

The natural killer (NK) cell receptor, NKG2D, and NK cells are activated in asthma.<sup>55</sup> NKG2D-deficient mice show no signs of allergic airway inflammation after exposure to the HDM allergen.<sup>56</sup> However, several other studies showed that blocking of NKG2D by antibodies or genetic depletion have not significant influence on the development of asthma in different mouse models.<sup>55</sup>

The data shown above suggest, that most of the HSP70-binding receptors have other ligands besides HSP70. Data showing that binding of HSP70 to its cognate or other immune receptors affects inflammation in asthma do not yet exist. The effects of activation of HSP70-binding receptors through interaction with HSP70 or other ligands in asthma depend on multiple factors such as the dose and nature of the ligand. The type of cell that expresses the HSP70-binding receptor is also very important in evaluating HSP70-mediated effects. Further studies are needed to explore whether HSP70 exerts its effects in asthma by binding to its cognate receptor or other immune receptors.

## Effects of HSP70 on Neutrophils

The neutrophil-mediated response is mostly associated with non-allergic asthma and correlates with asthma severity and resistance to corticosteroid therapy.<sup>57</sup> During the acute phase of allergic airway inflammation, as in any other inflammatory process, IL-8 secretion and subsequent neutrophil attraction is evident, as demonstrated by both clinical studies in human and experimental studies with mouse models of allergic airway inflammation.<sup>58,59</sup> In cases of allergic asthma, neutrophils can suppress the progression of asthma.<sup>60</sup>

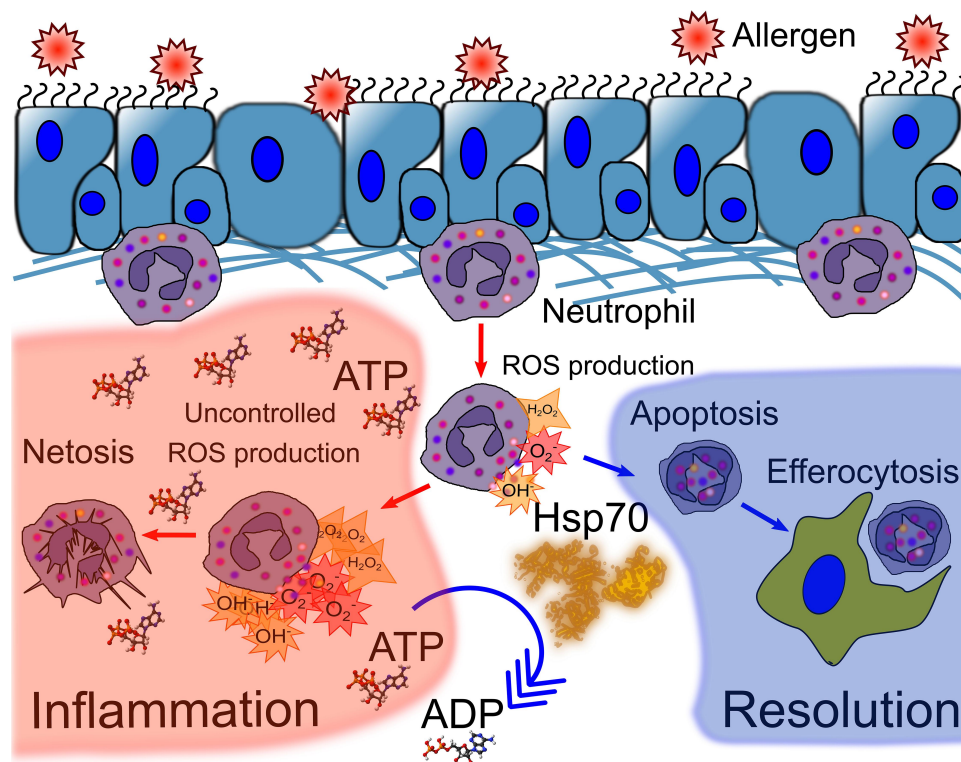
Besides the indirect neutrophil stimulation through alterations in the expression of neutrophil-attracting cytokines that were reported above,<sup>27</sup> some direct effects of HSP70 on neutrophils have been demonstrated in experiments that were unrelated to airway inflammation. Thus, the dose-dependent potential of stress-inducible Hsp72 (HspA1A)<sup>7</sup> to stimulate chemotaxis of human neutrophils has been previously reported in in vitro models.<sup>61</sup> Particularly, the participation of CD14 and TLR2 is reported to be associated with the mechanism of Hsp72-induced neutrophil chemotaxis.

Another mechanism that can be implicated in HSP70-mediated neutrophil recruitment from the bone marrow to the site of inflammation might be caused by the ability of HSP70 to interact with tetratricopeptide repeat (TPR) domain-containing protein co-chaperones.<sup>62</sup> One such protein, HSP interacting protein (Hip), can also interact with CXCR2, which is expressed on mature neutrophils and is responsible for neutrophil recruitment from the bone marrow to peripheral blood.<sup>63</sup> Moreover, the mutant Hip, which is unable to bind HSP70 (particularly, heat shock cognate protein 71 kDa (Hsc70) or HspA8),<sup>7</sup> induces attenuated chemotaxis of neutrophils.<sup>63</sup> Thus, the elevation of extracellular HSP70 concentration at the site of inflammation may affect the chemotactic activity of neutrophils.

Neutrophils recruited to the site of inflammation can neutralize the pathogen through one of the following strategies: phagocytosis, reactive oxygen species (ROS) production, or neutrophil extracellular trap (NET) formation. Furthermore, ROS hyperproduction, as well as uncontrolled NETosis, are known to be important factors of asthma complications, in response to allergen provocation; while neutrophil apoptosis is advantageous for the resolution of inflammation.<sup>64-66</sup> In vitro

studies have demonstrated that HSP70 prevented bacterial antigen-induced ROS production and pro-inflammatory cytokine secretion by both human peripheral blood and mouse bone marrow neutrophils.<sup>67,68</sup> Furthermore, HspA1A suppresses PMA-activated and fMLP-activated bone marrow neutrophil hyperactivation and prevents NETosis and necrosis.<sup>9</sup> Simultaneously, HspA1A facilitates spontaneous apoptosis in the primary culture of bone marrow neutrophils.<sup>9</sup> Recently, the contribution of ATP to the NETosis amplification was demonstrated.<sup>69</sup> Thus, the inherent ATPase activity of HSP70 can be considered as one of the potential mechanisms of NETosis prevention (Figure 1).

In addition, the indirect effects of HSP70 regarding the neutrophil-mediated immune response are noteworthy. Some of these effects have been described above and are related to the production of neutrophil-attractant cytokines, such as IL-8 (or KC) and IL-1 $\beta$  by epithelial cells or macrophages.<sup>27,28</sup> Furthermore, it was recently shown, that the Th2-associated cytokine, IL-4, interacts directly with IL-4R $\alpha$  on bone marrow neutrophils and blocks GM-CSF-induced neutrophil recruitment to the site of inflammation.<sup>70</sup> Moreover, neutrophil depletion can lead to a significant increase in Th2 cytokine



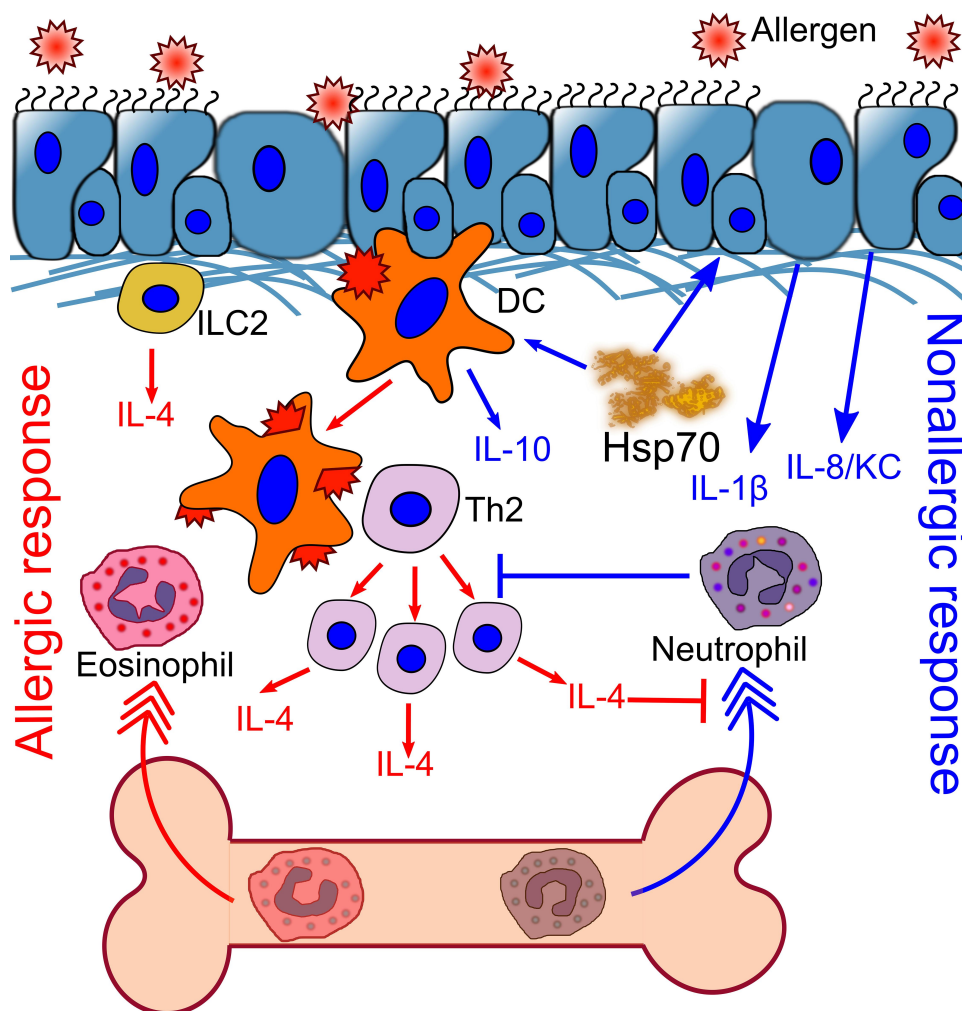
**Figure 1** HSP70 prevents neutrophil hyperactivation at the site of inflammation. Production of reactive oxygen species (ROS) is one mechanism of neutrophil defense against pathogens; however, the trigger of ROS production can also cause damage to host tissues or damage-associated molecular patterns (DAMPs). ATP is known to stimulate ROS production. Uncontrolled ROS production leads to NETosis and inflammation progression. HSP70, as an ATPase, can dephosphorylate ATP to ADP, which is less pyrogenic than ATP, and thus prevent uncontrolled ROS production. Moreover, HSP70 was involved in switching from necrosis and NETosis of neutrophils to apoptosis. Apoptosis and efferocytosis of apoptotic neutrophils are associated with inflammation resolution.

production by ILC2 during the early stages of allergic airway inflammation and amplifies the progression of inflammation.<sup>60</sup> By reducing IL-4 levels in BAL fluid (as reported for the anti-inflammatory effects of HSP70),<sup>9,71</sup> HSP70 potentially prevents neutrophil retention in the bone marrow and facilitates neutrophil recruitment (Figure 2).

Thus, the indirect effects of HSP70 on the neutrophil-mediated immune response can be the link between the pro-inflammatory and anti-inflammatory activity of HSP70 in allergic airway inflammation. By recruiting neutrophils, exogenous HSP70 suppresses the Th2-associated immune response in inflammation (Figure 2). At the same time, although elevated levels of extracellular HSP70 induce the recruitment of neutrophils, HSP70 prevents both neutrophil hyperactivation and the further progression of inflammation (Figure 1).

## HSP70-Mediated Effects on APCs, Antigen Presentation, and T Cell Activation: Implications in Allergic Asthma

APCs play a pivotal role in asthma, particularly in allergic airway inflammation. Professional APCs include DCs, macrophages, and B cells. APCs recognize, internalize, and present peptides derived from allergenic proteins in complex with major histocompatibility complex (MHC) II, and in this way, activate adaptive immunity by inducing clonal allergen-specific Th2 expansion.<sup>72</sup> In the atopic phenotype, Th2 cells are key cells of the adaptive immune system responsible for the hyperimmune reaction.<sup>73,74</sup> Thus, the impact of HSP70 on APCs in allergic sensitization and allergic airway inflammation is of great interest.



**Figure 2** HSP70 induces neutrophil recruitment to the site of inflammation. Exogenous substance challenge induces neutrophil recruitment to the airways. However, inhaled allergen promotes IL-4 production by ILC2, and further by DC-activated Th2 cells. IL-4 concentration is increased locally and systemically. IL-4 blocks neutrophil recruitment, which promotes the recruitment of eosinophils and stimulates allergic airway inflammation. HSP70 activates the production of IL-1 $\beta$  and IL-8 by epithelial cells, which attract neutrophils to the periphery. Neutrophils block Th2 proliferation and allergic airway inflammation. HSP70 also prevents DC maturation and induces IL-10 production.



In the early studies that examined the potential of HSP70 to stimulate antitumor immunity, it has been shown that extracellular HSP70 has significant stimulatory effects on APCs and subsequently facilitates the adaptive immune response. Particularly, HSP70 induces APC maturation and the secretion of pro-inflammatory cytokines such as TNF- $\alpha$ , IL-1b, IL-12, IL-6, and GM-CSF.<sup>14,75</sup> These effects were served as a mechanistic explanation of antitumor effect of exogenous HSP70.<sup>76,77</sup> However, later, the results were not confirmed; opposite, the suppressive effect of HSP70 and the induction of the tolerogenic APC phenotype were demonstrated.<sup>13,78</sup>

The HSP70-mediated suppressive effects have been demonstrated by findings that demonstrate endotoxin-free recombinant mycobacterial, and human HSP70 prevents bone marrow-derived DC maturation and induces the tolerogenic phenotype in mature DCs.<sup>13,78</sup> In the presence of endotoxin-free HSP70, immature monocyte-derived DCs show reduced capacity to prime T cells, and activate T cell proliferation. The proliferation rates of both total pools of activated CD3<sup>+</sup>CD25<sup>+</sup> T cells and non-activated CD3<sup>+</sup>CD25<sup>-</sup> cells, including T helper CD4<sup>+</sup>CD25<sup>-</sup> and T cytotoxic CD8<sup>+</sup>CD25<sup>-</sup> cells, are significantly reduced in the presence of recombinant human endotoxin-free HSP70. Moreover, HSP70 inhibits the proliferation of T cells that have been pre-activated with IL-12 or phytohemagglutinin.<sup>13</sup>

Molecular mechanisms underlying the suppressive effects of HSP70 on the antigen-presenting activity of innate immune cells are still not completely understood. On one hand, HSP70–MHCII interaction does not affect the primary structure of the antigenic peptide, and therefore, cannot modulate specific T cell responses even in conditions of enhanced or suppressed antigen presentation.<sup>79</sup> On the other hand, the observed effects can be caused by the antigenic properties of HSP70, as exogenous or endogenous HSP70 protein can undergo internalization, following the processing and presentation of different HSP70-derived peptides in the context of MHCII. Peptide fragments of HSP70 have been shown to be expressed in complexes with both MHCII and MHCI.<sup>80</sup>

Notably, peptides of both extracellular and intracellular HSP70 pools can accumulate on MHCII, either as a result of cross-presentation or autophagy.<sup>10,80</sup> Thus, even when HSP70 enhances antigen presentation, the presentation of peptides from the extracellular HSP70 pool is simultaneously increased. As a result, the specificity of Th2

cells shifts. Alteration of Th2 response specificity is associated with the suppression of allergic inflammation during allergen-specific immune therapy.<sup>81,82</sup> Altogether, the data above suggest the following: antigen-presentation of HSP70 peptides might modulate allergic airway inflammation by preventing a part of the intracellular pool of MHCII from participation in allergen processing and presentation by occupying the MHCII complex. In this way, HSP70 reduces the number of allergen-specific Th2 clones, which, in turn, results in a reduced level of allergen-specific IgE. Thus, the immune-modulatory activity of HSP70 might be exhibited by switching the allergic response to tolerance.

Moreover, HSP70 can inhibit the proliferation of T lymphocytes both indirectly, by downregulating the stimulatory activity of APCs, and by direct influence on previously activated T cells.<sup>10,83</sup> In addition to the suppressive effects on APCs and T cell proliferation, recent studies on the implications of HSP70 in the development of autoimmune disorders have demonstrated the potential of HSP70 to elevate the number of regulatory T cells (T regs). Injections of HSP70 upregulate total Foxp3 expression and increase the percentage of CD4<sup>+</sup>CD25<sup>+</sup>Foxp3<sup>+</sup> cells in lymph nodes.<sup>83,84</sup>

Taken together, we can assume that under specific conditions, HSP70 promotes the differentiation of naïve T cells to CD4<sup>+</sup>CD25<sup>+</sup>Foxp3<sup>+</sup> inducible regulatory T cells and prevents the differentiation of naïve T cells to the Th2 phenotype.<sup>10,85</sup> Thus, HSP70 can suppress the activation and proliferation of T cells and inhibit the differentiation of naïve T cells to the Th2 phenotype, which makes it one of the key features of allergic airway inflammation.

## Chaperone Activity of HSP70 in Asthma

While extracellular HSP70 reveal both pro-inflammatory and anti-inflammatory activity, intracellular HSP70 are traditionally considered to be cytoprotective chaperones.<sup>24</sup> Functional chaperone activity of HSP70 is due to their ability to interact with the hydrophobic regions of denaturing (unfolded or misfolded) proteins.<sup>86</sup> The substrate-binding domain of HSP70 that can recognize denaturing proteins is located at the C-terminus of the molecule. This domain is represented by the 15 kDa  $\beta$ -sheet region, that contains the hydrophobic groove.<sup>62,87</sup> Interacting with hydrophobic regions of newly synthesized or damaged denatured proteins, HSP70s participate in the

stress-induced cellular responses, that can lead to the reparation of cellular homeostasis or cell death.<sup>88</sup>

As HSP70 are localized in the different cell compartments including the nucleus, mitochondria, cytosol,<sup>6</sup> they also administrate the translocation of the polypeptide chains into different cell organelles, including the endoplasmic reticulum (ER).<sup>88</sup> In the ER, HSP70 (particularly, HspA5)<sup>7</sup> are involved in the mechanisms of the ER stress and the unfolded protein response (UPR) that reportedly play a sufficient role in asthma and airway inflammation.<sup>89,90</sup> The ER stress was shown to contribute to such asthma features as airway remodeling and mucus hyperproduction.<sup>91</sup> Moreover, the overexpression of UPR sensors (that are localized in the ER membrane – inositol requiring enzyme (IRE) 1, activating transcription factor (ATF) 6, and PKR-like ER kinase (PERK)) was demonstrated in both ovalbumin-induced and HDM-induced animal asthma models as well as in the tissue samples of asthma patients.<sup>91,92</sup> In the absence of the ER stress, IRE1 $\alpha$ , ATF6, and PERK bind HspA5. During the unfolded protein accumulation in ER, HspA5 binds the unfolded proteins and dissociates from the interaction with the UPR sensors; the loss of the interaction triggers the UPR activation.<sup>90–92</sup>

In the case when the UPR is not sufficient to repair the damage, the fragment of ER can be degraded via macroautophagy.<sup>93</sup> HSP70 (particularly, Hsc70 or HspA8) plays an essential role in protein degradation through the ubiquitin–proteasome system (UPS) and in all three types of autophagy: chaperon-mediated, microautophagy and macroautophagy.<sup>88,94</sup> Co-expression of HSP70 and the autophagy marker LC3 was detected in the airway cell culture A549 under the exposure of carbon nanoparticles and reported as an indicator of chaperon-mediated autophagy.<sup>95</sup> In certain conditions, carbon nanoparticles can trigger or exacerbate airway inflammation in asthma, and chaperon-mediated autophagy can be potentially involved in inflammation regulation.<sup>95,96</sup>

The role of autophagy in asthma is widely discussed and thought to be associated with airway remodeling and severe asthma.<sup>97,98</sup> Single nucleotide polymorphisms in autophagy-related gene *Atg5* are associated with reduced lung function and airway remodeling.<sup>93,96</sup> Also, the expression of *Atg5* product was higher in lung tissue samples of asthma patients compared with nonasthmatic samples.<sup>97</sup> Furthermore, in the ovalbumin-induced mouse model of asthma, the treatment with *Atg5* short hairpin RNA inhibited autophagy together with asthma features

such as airway hyperreactivity and eosinophilia.<sup>98</sup> Activation of autophagy is also involved in the NET formation (that implication in asthma was described above) and in neutrophilic airway inflammation.<sup>65,96</sup>

Thus, the facts above indicate that intracellular HSP70 contribute to ER stress-induced UPR, UPS, and autophagy. All these processes are observed in asthma, particularly severe asthma. Taking into account elevated expression of HSP70 in airway epithelial and immune cells of people with asthma,<sup>21,22</sup> we can assume that the protective chaperone activity of intracellular HSP70 during the severe inflammation in asthma plays an essential role in the degradation of the damaged proteins and the removal of destroyed organelles from the cells.

## Potential Suppressive Effects of HSP70 ATPase Activity in Systemic and Local Allergic Airway Inflammation

One of the main functional characteristics of the HSP70 family is ATPase activity.<sup>86,99</sup> The nucleotide-binding domain (45 kDa) belongs to the highly conserved region of the primary amino acid sequence of HSP70 proteins. It is in close proximity to the N-terminus of the molecule and typically contains the actin-like ATPase protein superfamily (sugar kinase/heat shock protein 70/actin superfamily) ATP-binding center.<sup>100</sup> The ATP-binding center of HSP70 is mainly represented by proline, tyrosine, phenylalanine, arginine, and glutamine, which are positioned at the specific places relative to each other.<sup>101</sup> For different members of the HSP70 family, the positions can be shifted by one or two amino acid residues; however, the shift does not affect domain conformation.<sup>101</sup>

Moreover, HSP70 can bind to ATP with nanomolar affinity.<sup>99,102</sup> Consecutive interaction of HSP70 with ATP, followed by interaction with a peptide or denatured protein in the presence of co-chaperone and adapter proteins, induces the ATPase activity of HSP70, and hydrolysis of ATP to ADP.<sup>103</sup> The difference in peptide binding of ATP-bound and ADP-bound HSP70 isoforms is the basis of chaperone activity.<sup>88</sup> However, chaperone activity can also be manifested by HSP70 alone in the presence of co-chaperones. The most well-described co-chaperones are members of the J-protein family, which facilitate ATP dephosphorylation, and the nucleotide exchange factors (NEFs), which are responsible for the removal of ADP from the HSP70-nucleotide complex as well as the TPR

domain-containing proteins.<sup>62</sup> Both J-proteins and NEFs have binding sites in the region of the nucleotide-binding domain.<sup>62,87</sup>

Recent studies demonstrate that ATP plays an important role in the activation of APCs.<sup>104</sup> Release of ATP to the extracellular space, as a result of damage, stress, or cell death, has been implicated in the development of allergic airway inflammation.<sup>105</sup> In particular, a sharp increase in ATP levels in the BAL fluid of persons with asthma and in mouse models has been detected shortly after allergen provocation.<sup>106</sup> Extracellular ATP binding to the P2X7 receptor can activate different types of cells and increase the production of pro-inflammatory cytokines such as: IL-1, IL-6, IL-18, and TNF- $\alpha$  by macrophages; IL-1 $\beta$ , IL-18, TNF- $\alpha$ , and IL-23 by DCs; IL-8 and eosinophil cationic protein (ECP) by eosinophils; as well as histamine secretion by mast cells. Taken together, extracellular ATP upregulation can facilitate Th2-activation and hyperimmune reactions.<sup>107</sup> Direct neutralization of ATP through the administration of the ATP-hydrolyzing enzyme apyrase can significantly reduce inflammatory cell infiltration and conducting airway obstruction in induced allergic airway inflammation.<sup>106</sup>

Under non-inflamed steady-state physiological conditions, exoferments ectonucleoside triphosphate diphosphohydrolases (E-NTPD)s regulate extracellular ATP levels.<sup>108</sup> Both E-NTPD and HSP70 belong to the sugar kinase/heat shock protein 70/actin superfamily and possess similar conformations of the ATP-binding domain, which is sensitive to Ca<sup>2+</sup> or Mg<sup>2+</sup>.<sup>109</sup> The E-NTPDs manifest ATPase activity; however, owing to their membrane-bound location, they can only act on the cell surface, but not within the extracellular matrix, unlike HSP70.<sup>108,110</sup> The role of HSP70 in the regulation of extracellular ATP concentrations has not yet been determined in vivo. However, in vitro studies have shown a reduction in extracellular ATP levels in the presence of HSP70 in culture filtrates of bone marrow-derived DCs.<sup>111</sup> Taken together, increased extracellular HSP70 concentration at the site of allergic airway inflammation and ATPase activity of the protein support the hypothesis that HSP70 can downregulate extracellular ATP levels and suppress P2X7 activation by reducing the activation stimulus.

Thus, the neutralization of extracellular ATP (the pro-inflammatory stimulus activating APCs to promote allergic airway inflammation) by HSP70 from the extracellular

pool can be considered one of the molecular mechanisms associated with the anti-inflammatory potential of HSP70.

## Discussion

In the presented review, we surveyed the published results related to effects and potential regulatory mechanisms of HSP70 in asthma.

The literature data demonstrated that under certain conditions, some exogenous HSP70 can trigger specific IgE production and even autologous allergic-like response to self HSP70. Regarding these findings, it would be of great interest to investigate the potential of some exogenous HSP70, for example, Pen c 19, to induce allergic airway inflammation in the murine model. As it was shown, murine HSP70 (yielded from the homogenate of mouse liver and kidney) was not potent to induce eosinophilia and pro-allergic cytokine production when was injected and inhaled to mice in accordance with the scheme similar to ovalbumin-induced allergic airway inflammation.<sup>9</sup> Mouse model of Pen c 19-induced airway inflammation would help to investigate the mechanisms of cross-reactivity and self-protein tolerance. In fact, tolerance is of great importance as the elevated *HSP70* mRNA expression and HSP70 protein secretion were found in asthma patient cell and tissue samples.

Despite the clear implication of HSP70 in asthma, there is no a unique concept about the role of HSP70 in asthma. Both pro-inflammatory and anti-inflammatory activities of HSP70 are reported. The ambiguous effects are detected in experiments with animal models of allergic airway inflammation and even with in vitro cell cultures. One explanation of such effects comes from the findings of Fong et al<sup>37</sup> who demonstrated that HSP70 stimulation through Siglec-5 acts as an anti-inflammatory signal, while stimulation through Siglec-14 is pro-inflammatory. Thus, the current cell state, particularly the level of expression of certain receptor interacting with HSP70, can be a determinant in triggering the pro-inflammatory or anti-inflammatory effects.

The other explanation of the dualistic effects can be related to the HSP70 protein characteristics. While the traditional concept suggests intracellular HSP70 to be cytoprotective chaperones, and extracellular HSP70 act as alarmins, both of these forms are elevated in asthma, and some studies reported the suppressive effects of extracellular HSP70. Due to cell and tissue damage during the inflammatory processes, it is quite difficult to discriminate

the effects of physiologically released HSP70 (secreted as a component of exosomes) and effects of HSP70 that appeared in the extracellular space due to cell necrosis. Moreover, exosomal HSP70 can be represented with the different HSP70 family members at the different phases of allergic airway inflammation and in nonallergic response. Some experiments directed to precise characterization of the certain HSP70 function in the inflammatory processes in asthma could be of great importance. Such experiments should also take into account the state of the protein (ATP-bound or ADP-bound), and therefore, not only LPS contamination, but also the concentration of ATP, and  $\text{Ca}^{2+}$ , and  $\text{Mg}^{2+}$  in HSP70 preparations should be verified.

The estimation of ATP contamination in HSP70 preparations is of great importance in the studies of the ATPase activity of HSP70. Due to inherent ATPase activity, HSP70 can be involved in the neutralization of extracellular ATP, which is known to trigger the inflammatory response in both Th2- and T17-mediated asthma inflammation. Thus, the potential of the extracellular HSP70 to neutralize ATP and subsequently suppress the inflammation in asthma must be further in detail investigated. Additionally, the application of HSP70 inhibitors, particularly, ATP analogs and ATPase activity modulators, is very valuable in the investigation of the suppressive potential of ATPase activity in the inflammation in asthma. Such inhibitors can be applied in both animal models and in vitro experiments with asthma patient primary cell cultures and tissue samples.<sup>112</sup>

It can also be suggested, that some stimulatory HSP70 effects that are referred to as pro-inflammatory, in some situations, can prevent the inflammation. In particular, the observed in vitro pro-inflammatory HSP70 effects that are directed at structural (primarily epithelial) and innate immune cells, act through the stimulation of IL-1 alpha, IL-6, TNF- $\alpha$ , and IL-8 production, in vivo can trigger the neutrophil infiltration. In the case of allergic airway inflammation, stimulation of neutrophil response can suppress eosinophilia and Th2 proliferation. Thus, the neutrophil-mediated response and activation of HSP70 can downregulate allergic airway inflammation (Figure 2). Regarding this suggestion, the estimation of a shift in cytokine production under the HSP70 exposure in the mouse models of allergic airway inflammation should not be restricted to common pro-allergic cytokines (IL-4, IL-5, IL-13), but neutrophil-attracting cytokine (IL-1b, IL8 or KC, as well as GM-CSF) production should also be estimated.

## Conclusion

Many facts are confirming the involvement of HSP70 in the inflammatory processes in asthma, however, the mechanisms of this involvement require further study. In mouse models of airway inflammation, both intracellular and extracellular HSP70 exhibit the anti-inflammatory activity that characterizes the protein as a promising molecule for the development of new therapeutic approaches to managing asthma. However, some facts demonstrate a pro-inflammatory activity that HSP70 can display in different experimental models. In this review, we provided evidence that both pro-inflammatory and anti-inflammatory effects resulted from the unique mechanisms of HSP70 action. Further understanding of the molecular mechanisms of the HSP70-mediated regulatory functions in airway inflammation may reveal new therapeutic possibilities for the treatment of patients with asthma.

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## Disclosure

The authors report no conflicts of interest in this work.

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