Randomised aspirin assignment and risk of adult-onset asthma in the Women's Health Study

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ABSTRACT

Background: Randomised data in men show a small but significant reduction in the risk of adult-onset asthma among those given aspirin. The results from an observational study in women suggest that frequent use of aspirin decreases the risk of adult-onset asthma, but randomised data in women are lacking. A study was undertaken to test the effect of 100 mg aspirin or placebo on alternate days on the risk of adult-onset asthma in the Women's Health Study.

Methods: A randomised, double-blind, placebo-controlled clinical trial of aspirin and vitamin E was performed in apparently healthy women with no indication or contra-indication to aspirin therapy and no history of asthma at study entry. Female health professionals self-reported an asthma diagnosis on yearly questionnaires.

Results: Among 37 270 women with no reported history of asthma prior to randomisation and during 10 years of follow-up, there were 872 new cases diagnosed with asthma in the aspirin group and 963 in the placebo group (hazard ratio 0.90; 95% CI 0.82 to 0.99; p = 0.027). This apparent 10% lower relative risk of incident adult-onset asthma among those assigned to aspirin was significantly modified by body mass index, with no effect in women with a body mass index of $\geq 30 \text{ kg/m}^2$. The effect of aspirin on adult-onset asthma was not significantly modified by age, smoking status, exercise levels, postmenopausal hormone use or randomised vitamin E assignment.

Conclusions: In this large randomised clinical trial of apparently healthy adult women, administration of 100 mg aspirin on alternate days reduced the relative risk of a newly reported diagnosis of asthma.

Trial registration number: NCT00000479

Over the last few decades asthma has become a growing public health problem and the prevalence and incidence of this respiratory disease continues to increase.[1] Among the explanations for this increase are obesity and weight change,[2 3] dietary factors,[4] changes in early life antigen exposure,[5] environmental or geographical factors,[6] but it may also be the result of increased diagnostic testing. The increase in the incidence of asthma in children was accompanied by a change in the pattern of use of antipyretic and analgesic drugs. The use of aspirin decreased because of its linkage with Reye's syndrome and, as a result, the use of other analgesic drugs increased,[7] leading to the suggestion that aspirin may be associated with a decreased risk of asthma.[8]

In adult men, the results from a post hoc analysis of the Physicians' Health Study showed that apparently healthy men who were randomised to receive 325 mg aspirin every other day had a statistically significant relative risk reduction of 22% in newly reported diagnoses of adult-onset asthma during a mean follow-up period of 5 years.[9] In a large observational prospective cohort study, adult women who self-selected for frequent use of aspirin had a decreased risk of incident asthma compared with never users after adjusting for a variety of potential confounding factors.[10]

The Women's Health Study (WHS), a large-scale, randomised, double-blind, placebo-controlled trial of aspirin and vitamin E, provided a unique opportunity to test whether low-dose aspirin decreases the risk of a newly reported diagnosis of adult-onset asthma in apparently healthy women with no indication or contraindication for aspirin therapy.

METHODS

The WHS was a randomised, double-blind, placebo-controlled, two-by-two factorial trial testing the risks and benefits of low-dose aspirin (100 mg every other day; Bayer AG, Leverkusen, Germany) and vitamin E (600 IU every other day; National Source Vitamin E Association, Washington, DC, USA) in the primary prevention of cardiovascular disease and cancer.

The WHS randomised 39 876 female health professionals between April 1993 and January 1996; of these, 19 934 were assigned to active aspirin and 19 942 to placebo. The results of the trial[11–13] and methods used to identify and recruit the study population have been described in detail previously.[14 15] In brief, invitational letters were mailed to 1 757 247 female health professionals throughout the United States and Puerto Rico, of whom 453 787 returned the initial study questionnaire. Women were considered eligible if they indicated willingness to participate, were aged ≥ 45 years with no previous history of coronary heart disease, cerebrovascular disease, cancer (except non-melanoma skin cancer) or other major chronic illness. Eligible women also had no history of side effects to any study medications, were not taking aspirin or non-steroidal anti-inflammatory drug medications more than once a week (or were willing to forego use of these medications during the trial), were not taking anticoagulants or corticosteroids, and were not taking individual supplements of vitamin A, E, or β-carotene more than once a week. Of the women who returned the initial study questionnaire, 65 169 were willing and eligible to participate and enrolled in a 3-month run-in period employing all placebo medications, designed to identify a group of participants with a higher likelihood of good long-term compliance. Of these, 39 876 were compliant
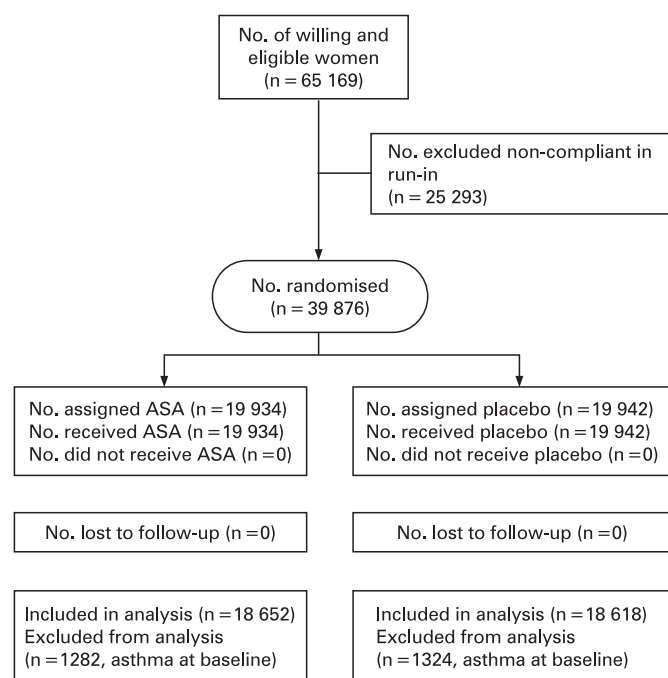
Figure 1 Flow diagram of the progress through the phases of the Women's Health Study.

during the run-in phase and were randomised into the trial (fig 1).

Baseline information was collected by mailed questionnaires that asked about many anthropometric, demographic, clinical and lifestyle variables. Twice in the first year and yearly thereafter, follow-up questionnaires were sent out asking about compliance with study medication, development of side effects, newly developed diseases that required medical attention including asthma, dates of diagnoses, and personal characteristics and habits.

Ascertainment of asthma

On each of the follow-up questionnaires, the participating female health professionals were specifically asked about newly diagnosed diseases including asthma and the date of diagnosis of these diseases. Asthma was not a prespecified end point of the WHS.

Statistical analysis

All reports of asthma from enrolment until the end of the randomised trial on 31 March 2004 were included. A total of 2606 participants (1282 in the aspirin group and 1324 in the placebo group) who indicated an asthma diagnosis prior to randomisation were excluded, leaving 37 270 participants without a history of asthma at baseline. Continuous baseline measurements were compared using the Student t test and categorical variables were compared using the χ^2 test according to aspirin or aspirin-placebo assignment. Following the intention-to-treat principle, we compared the cumulative incidence of adult-onset asthma over the trial follow-up period. A Cox proportional hazards model was used to analyse the association between randomised aspirin assignment and time from randomisation to date of the reported asthma diagnosis. Furthermore, we assessed whether the effect of aspirin on adult-onset asthma was modified by age (<55, $55–64$, $\geqslant65$ years), smoking status (never, past, current), body mass index (<25, $25–29.9$, $\geqslant30$ kg/m^2), exercise (rarely, <1/week, $1–3$ times/week, $\geqslant4$ times/week),

Table 1 Baseline characteristics according to randomisation to aspirin treatment in the Women's Health Study (N = 37 270)

Characteristics	Aspirin (N = 18 652)	Placebo (N = 18 618)	p Value
Mean (SD) age (years)	54.7 (7.1)	54.7 (7.1)	0.83
Mean (SD) BMI (kg/m^2)	26.0 (5.0)	25.9 (5.0)	0.39
BMI $\geqslant30$ kg/m^2 (%)	17.7	17.6	0.71
History of hypertension (%)	25.7	25.4	0.51
History of diabetes (%)	2.7	2.4	0.15
Smoking status (%)			
Never	51.2	51.2	0.61
Past	35.7	35.5	
Current (<15 cigs/day)	4.8	5.0	
Current (15+ cigs/day)	8.2	8.4	
Exercise (%)			
Rarely	38.3	38.1	0.51
<1/week	19.5	20.3	
1–3/week	31.4	31.1	
$\geqslant4$/week	10.9	10.5	
Alcohol consumption (%)			
Rarely	44.8	45.1	0.79
1–3/month	13.4	12.8	
1–6/week	31.6	31.8	
$\geqslant1$/day	10.2	10.3	
Postmenopausal	55.0	54.4	0.23
Postmenopausal hormone use			
Never	50.0	50.4	0.27
Past	9.0	9.2	
Current	41.0	40.4	

BMI, body mass index.

postmenopausal hormone use (never, past, current) and randomised vitamin E assignment. Statistical significant effect modification was tested by contrasting main effect models with models that also included interaction term indicators using the likelihood ratio test.

RESULTS

Table 1 summarises the baseline characteristics of the 37 270 participants without a reported history of asthma at study entry according to randomisation to aspirin treatment. As expected in this large sample, the baseline characteristics were well balanced between the aspirin and placebo groups.

Over a mean follow-up period of 9.7 years (362 120 person-years), a total of 1835 participants reported new diagnoses of asthma which corresponded to a cumulative incidence of adult-onset asthma of 4.9%; 872 new diagnoses of asthma were reported in the aspirin group and 963 in the placebo group (table 2). This corresponded to a hazard ratio of 0.90 (95% confidence interval (CI) 0.82 to 0.99; p = 0.027) for newly reported adult-onset asthma, representing a 10% reduction in relative risk in favour of aspirin. The cumulative incidence of asthma in the aspirin and placebo groups over the follow-up period is shown in fig 2, indicating a constantly decreasing risk of adult-onset asthma in the aspirin group after approximately 6 years of follow-up.

There were no statistically significant differences in the observed effect of aspirin on adult-onset asthma by age, smoking status, levels of exercise, postmenopausal hormone intake or randomised vitamin E assignment (table 2). In contrast, body mass index significantly modified the effect of low-dose aspirin on reported asthma diagnosis (p = 0.002), indicating no apparent effect of aspirin on the incidence of asthma in women with a body mass index of $\geqslant 30$ kg/m² (hazard ratio 1.19, 95% CI 0.99 to 1.42). A similar lack of association was seen when we further categorised high body mass index values (data not shown). The effect of aspirin on the occurrence of asthma was also significantly modified by linear body mass index (p = 0.008).

DISCUSSION

In this large, randomised, double-blind, placebo-controlled trial in apparently healthy women, 100 mg aspirin every other day significantly reduced the relative risk of newly reported adult-onset asthma diagnosis by 10%. The magnitude of the reduction was similar across subgroups of age, smoking status, exercise, postmenopausal hormone intake and randomised vitamin E intake but was absent in those who were obese.

Our data are consistent with findings in men from a post hoc analysis of the randomised aspirin component of the Physicians' Health Study[9] which showed a statistically significant reduction of 20% in the relative risk of newly reported diagnoses of adult-onset asthma (hazard ratio 0.78, 95% CI 0.61 to 1.00, p = 0.045) in men randomly assigned to 325 mg aspirin every other day compared with men assigned to placebo. The smaller effect estimate in the current study may have been due to the lower aspirin dose in the WHS (100 mg every other day) compared with that in the Physicians' Health Study (325 mg every other day). However, gender differences in the association between

Table 2 Randomised aspirin assignment and report of newly diagnosed asthma in the Women's Health Study (N = 37 270)

	Aspirin group (N = 18 652) (181 467 person years)	Placebo group (N = 18 618) (180 652 person years)	Hazard ratio* (95% CI)	p Value
Newly diagnosed asthma (n)	872	963	0.90 (0.82 to 0.99)	0.027
Stratified analyses				
Age (years)				0.37†
<55	550	595	0.92 (0.82 to 1.03)	0.16
55–64.9	249	268	0.92 (0.78 to 1.10)	0.37
65+	73	100	0.73 (0.54 to 0.99)	0.04
Smoking status				0.97†
Never	419	459	0.91 (0.80 to 1.04)	0.16
Past	322	357	0.89 (0.77 to 1.04)	0.13
Current	131	146	0.91 (0.72 to 1.15)	0.44
BMI (kg/m²)				0.002†
<25	336	406	0.83 (0.71 to 0.95)	0.01
25–29.9	258	319	0.80 (0.68 to 0.95)	0.01
30+	265	224	1.19 (0.99 to 1.42)	0.06
Exercise				0.32†
Rarely	347	364	0.95 (0.82 to 1.10)	0.45
<1/week	198	205	1.00 (0.83 to 1.22)	0.97
1–3/week	240	294	0.81 (0.68 to 0.95)	0.01
$\geqslant$4/week	87	100	0.84 (0.63 to 1.12)	0.24
Postmenopausal hormone use				0.89†
Never	387	429	0.91 (0.79 to 1.04)	0.16
Past	86	92	0.96 (0.71 to 1.29)	0.77
Current	396	438	0.89 (0.77 to 1.01)	0.08
Vitamin E				0.57†
Active	428	485	0.88 (0.77 to 1.00)	0.05
Placebo	444	478	0.93 (0.81 to 1.05)	0.25

CI, confidence interval; BMI, body mass index

*Calculated from Cox proportional hazard models using the placebo group as reference.

†p Value for modification from contrast of a model that included main effects only with a model that also included interaction term indicators using the likelihood ratio test.
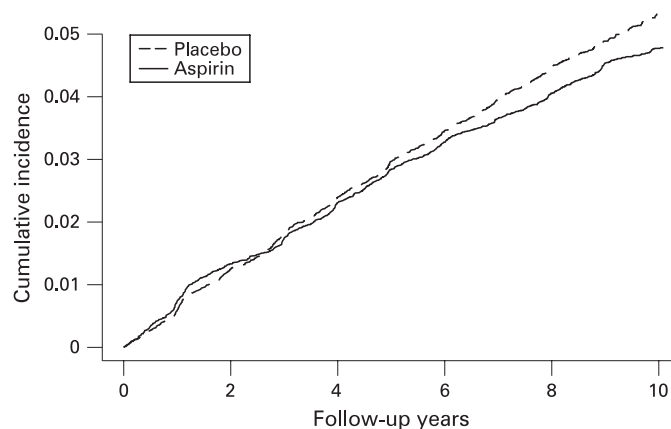
Figure 2 Cumulative incidence of adult-onset asthma during the randomised component of the Women's Health Study.

aspirin and adult-onset asthma may also account for the reduced effect in women.

Our results are also consistent with previous findings from a large prospective observational cohort study of female registered nurses in which women who self-selected for frequent aspirin use (15 days or more per month) had a 40% lower risk of newly reported physician diagnosis of adult-onset asthma than women who did not take aspirin.[10] The stronger effect size may be due to higher aspirin doses or to residual confounding. However, when the attributable incidence rate from the randomised data of the WHS (5.3/10 000 women per year), the randomised data of the Physicians' Health Study (5.9/10 000 men per year)[9] and the observational data from the Nurses' Health Study (4.4/10 000 women per year)[10] are compared, the results are very similar.

Among individuals with existing asthma, aspirin can acutely precipitate bronchospasm in the subset of patients with aspirin-intolerant asthma. This intolerance to aspirin affects 4–11% of patients with asthma.[16–18] The aetiology of aspirin-intolerant asthma is complex and not fully understood, but most evidence points towards an abnormality of arachidonic acid metabolism.[19] However, most patients with asthma tolerate aspirin well, and some patients with asthma improve when challenged with aspirin or other non-steroidal anti-inflammatory drugs.[20–22]

The precise biological mechanism by which long-term low-dose aspirin use may reduce the risk of asthma is currently unknown and should be the target of future studies. Several mechanisms that relate to immune-mediating effects of aspirin, which act via cyclooxygenase (COX)-dependent[23] and COX-independent pathways, can be envisaged. The effect of inhibiting COX with regard to asthma is probably dependent on the balance between proinflammatory prostaglandin D_2, prostaglandin $F_{2\alpha}$, thromboxane A_2 and anti-inflammatory prostaglandin E_2 production.[23] Aspirin irreversibly inhibits the isoenzyme COX-1 which can lead to a reduction in prostaglandin E_2 and subsequently to a reduction in the synthesis of interferon-γ of Th1 lymphocytes.[24 25] Consistent with this mechanism, aspirin inhibits the activation of NF-κB[26] and interleukin (IL)4 gene expression in T cells. Therapeutic doses of aspirin also inhibit IL4- and IL13-induced activation of STAT6 via non-COX dependent pathways.[27] Conversely, aspirin promotes STAT1 activation in interferon-γ signalling.[28] Aspirin therefore seems to inhibit Th2 immune responses while promoting Th1 responses. Since asthma has been associated with cytokine production of the Th2 phenotype,[29] a shift towards a non-asthmatic Th1 phenotype may lead to reduced susceptibility to asthma.

An additional non-COX activity of aspirin—but not of COX-2 inhibitors—is its potent induction of lipoxin A_4 and 15-epi-lipoxin A_4.[30] These products of arachadonic acid metabolism have profound anti-inflammatory properties that block airway hyperresponsiveness and pulmonary inflammation in murine models[31] and in vivo,[32] and are currently seen as novel targets for asthma therapies.

We found a statistically significant modification of the effect of aspirin on asthma by body mass index, indicating no effect of low-dose aspirin on adult-onset asthma among obese subjects (body mass index $\geqslant$30 kg/m²). Although subgroup analyses should be interpreted with caution, it is possible that there was less effect among obese women since respiratory symptoms in obese individuals may represent a different phenotype from asthma in non-obese individuals.[33] Furthermore, it can be envisaged that the effects of aspirin in obese women are reduced, potentially due to a larger volume distribution. A similar lack of effect in obese was also observed for the beneficial effect of aspirin on the risk of ischaemic stroke in the WHS which was not apparent in women who were obese.[11]

The strengths of this trial include its randomised, double-blind, placebo-controlled design, large number of participants and outcome events, as well as the high rates of follow-up and compliance. Several limitations should be considered when interpreting our results. Although this large trial had adequate size to detect a small to moderate benefit of low-dose aspirin on incident adult-onset asthma, the trial was not designed to test this specific hypothesis. The occurrence of incident adult-onset asthma was based on self-reports by the participating health care professionals and, thus, random misclassification is possible. We did not collect information on aspirin-intolerant asthma in the WHS. However, since the WHS was a trial of aspirin, participants are expected to tolerate aspirin well. Chronic obstructive pulmonary disease (COPD) can be misdiagnosed as adult-onset asthma, even in health professionals.[34] Owing to the randomised prospective design, however, such misclassification would be expected to result in an underestimation of effect. Supporting this line of reasoning, the effect of aspirin was somewhat stronger among never smokers and weaker among current smokers, who are more likely to develop COPD. In addition, when we censored participants who reported a diagnosis of COPD during follow-up, the results were similar (data not shown). Finally, the WHS was composed of apparently healthy female health professionals aged $\geqslant$45 years of whom the majority were white. Thus, generalisability to other populations may be limited.

In summary, 100 mg of aspirin every other day reduced the relative risk of newly reported diagnoses of adult-onset asthma in this large randomised trial of apparently healthy women. Although aspirin can worsen symptoms in some patients with asthma, our biologically plausible finding—along with similar results from a large randomised trial in men and observational cohort studies in women—suggests a small benefit of aspirin for the prevention of the development of asthma in adults. However, before public recommendations are provided, results from randomised trials are needed that are specifically designed to test whether low-dose aspirin reduces the risk of asthma.

REFERENCES

1. **Maziak W,** Behrens T, Brasky TM, et al. Are asthma and allergies in children and adolescents increasing? Results from ISAAC phase I and phase III surveys in Munster, Germany. Allergy 2003;**58**:572–9.
2. **Shore SA,** Fredberg JJ. Obesity, smooth muscle, and airway hyperresponsiveness. J Allergy Clin Immunol 2005;**115**:925–7.
3. **Camargo CA Jr,** Weiss ST, Zhang S, et al. Prospective study of body mass index, weight change, and risk of adult-onset asthma in women. Arch Intern Med 1999;**159**:2582–8.
4. **Shaheen SO,** Sterne JA, Thompson RL, et al. Dietary antioxidants and asthma in adults: population-based case-control study. Am J Respir Crit Care Med 2001;**164**:1823–8.
5. **Leynaert B,** Neukirch C, Jarvis D, et al. Does living on a farm during childhood protect against asthma, allergic rhinitis, and atopy in adulthood? Am J Respir Crit Care Med 2001;**164**:1829–34.
6. **Janson C,** Anto J, Burney P, et al. The European Community Respiratory Health Survey: what are the main results so far? European Community Respiratory Health Survey II. Eur Respir J 2001;**18**:598–611.
7. **Kogan MD,** Pappas G, Yu SM, et al. Over-the-counter medication use among US preschool-age children. JAMA 1994;**272**:1025–30.
8. **Varner AE,** Busse WW, Lemanske RF Jr. Hypothesis: decreased use of pediatric aspirin has contributed to the increasing prevalence of childhood asthma. Ann Allergy Asthma Immunol 1998;**81**:347–51.
9. **Barr RG,** Kurth T, Stampfer MJ, et al. Aspirin and decreased adult-onset asthma: randomized comparisons from the physicians' health study. Am J Respir Crit Care Med 2007;**175**:120–5.
10. **Barr RG,** Wentowski CC, Curhan GC, et al. Prospective study of acetaminophen use and newly diagnosed asthma among women. Am J Respir Crit Care Med 2004;**169**:836–41.
11. **Ridker PM,** Cook NR, Lee IM, et al. A randomized trial of low-dose aspirin in the primary prevention of cardiovascular disease in women. N Engl J Med 2005;**352**:1293–304.
12. **Cook NR,** Lee IM, Gaziano JM, et al. Low-dose aspirin in the primary prevention of cancer: the Women's Health Study: a randomized controlled trial. JAMA 2005;**294**:47–55.
13. **Lee IM,** Cook NR, Gaziano JM, et al. Vitamin E in the primary prevention of cardiovascular disease and cancer: the Women's Health Study: a randomized controlled trial. JAMA 2005;**294**:56–65.
14. **Buring JE,** Hennekens CH, for the Women's Health Study Research Group. The Women's Health Study: summary of the study design. J Myocardial Ischemia 1992;**4**:27–9.
15. **Rexrode KM,** Lee IM, Cook NR, et al. Baseline characteristics of participants in the Women's Health Study. J Womens Health Gend Based Med 2000;**9**:19–27.
16. **Kasper L,** Sladek K, Duplaga M, et al. Prevalence of asthma with aspirin hypersensitivity in the adult population of Poland. Allergy 2003;**58**:1064–6.
17. **Hedman J,** Kaprio J, Poussa T, et al. Prevalence of asthma, aspirin intolerance, nasal polyposis and chronic obstructive pulmonary disease in a population-based study. Int J Epidemiol 1999;**28**:717–22.
18. **Vally H,** Taylor ML, Thompson PJ. The prevalence of aspirin intolerant asthma (AIA) in Australian asthmatic patients. Thorax 2002;**57**:569–74.
19. **Hamad AM,** Sutcliffe AM, Knox AJ. Aspirin-induced asthma: clinical aspects, pathogenesis and management. Drugs 2004;**64**:2417–32.
20. **Szczeklik A,** Gryglewski RJ, Nizankowska E. Asthma relieved by aspirin and by other cyclo-oxygenase inhibitors. Thorax 1978;**33**:664–5.
21. **Kordansky D,** Adkinson NF Jr, Norman PS, et al. Asthma improved by nonsteroidal anti-inflammatory drugs. Ann Intern Med 1978;**88**:508–11.
22. **Resta O,** Foschino-Barbaro MP, Carnimeo N, et al. Asthma relieved by acetylsalicylic acid and nonsteroid anti-inflammatory drugs. Respiration 1984;**46**:121–7.
23. **Pang L,** Pitt A, Petkova D, et al. The COX-1/COX-2 balance in asthma. Clin Exp Allergy 1998;**28**:1050–8.
24. **Betz M,** Fox BS. Prostaglandin E2 inhibits production of Th1 lymphokines but not of Th2 lymphokines. J Immunol 1991;**146**:108–13.
25. **Snijdewint FG,** Kalinski P, Wierenga EA, et al. Prostaglandin E2 differentially modulates cytokine secretion profiles of human T helper lymphocytes. J Immunol 1993;**150**:5321–9.
26. **Kopp E,** Ghosh S. Inhibition of NF-kappa B by sodium salicylate and aspirin. Science 1994;**265**:956–9.
27. **Perez GM,** Melo M, Keegan AD, et al. Aspirin and salicylates inhibit the IL-4- and IL-13-induced activation of STAT6. J Immunol 2002;**168**:1428–34.
28. **Chen LC,** Kepka-Lenhart D, Wright TM, et al. Salicylate-enhanced activation of transcription factors induced by interferon-gamma. Biochem J 1999;**342**:503–7.
29. **Busse WW,** Lemanske RF Jr. Asthma. N Engl J Med 2001;**344**:350–62.
30. **Fiorucci S,** Santucci L, Wallace JL, et al. Interaction of a selective cyclooxygenase-2 inhibitor with aspirin and NO-releasing aspirin in the human gastric mucosa. Proc Natl Acad Sci USA 2003;**100**:10937–41.
31. **Levy S,** Volans G. The use of analgesics in patients with asthma. Drug Saf 2001;**24**:829–41.
32. **Bonnans C,** Vachier I, Chavis C, et al. Lipoxins are potential endogenous antiinflammatory mediators in asthma. Am J Respir Crit Care Med 2002;**165**:1531–5.
33. **Beuther DA,** Weiss ST, Sutherland ER. Obesity and asthma. Am J Respir Crit Care Med 2006;**174**:112–9.
34. **Barr RG,** Herbstman J, Speizer FE, et al. Validation of self-reported chronic obstructive pulmonary disease in a cohort study of nurses. Am J Epidemiol 2002;**155**:965–71.