#### **ASTHMA**

# Written action plans for asthma: an evidence-based review of the key components

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Received 22 June 2003 Accepted 8 October 2003 **Background:** Written action plans for asthma facilitate the early detection and treatment of an asthma exacerbation. Several versions of action plans have been published but the key components have not been determined. A study was undertaken to determine the impact of individual components of written action plans on asthma health outcomes.

**Methods:** Randomised controlled trials (n = 26) that evaluated asthma action plans as part of asthma self-management education were identified. Action plans were classified as being individualised and complete if they specified when and how to increase treatment (n = 17), and as incomplete (n = 4) or non-specific (n = 5) if they did not include these instructions.

**Results:** For individualised complete written action plans the use of 2–4 action points and the use of both inhaled (ICS) and oral (OCS) corticosteroid consistently improved asthma outcomes. Action points based on personal best peak expiratory flow (PEF) consistently improved health outcomes while those based on percentage predicted PEF did not. The efficacy of incomplete action plans was inconclusive because of insufficient data. Non-specific action plans led to improvements in knowledge and symptoms.

**Conclusion:** Individualised written action plans based on personal best PEF, using 2–4 action points, and recommending both ICS and OCS for treatment of exacerbations consistently improve asthma health outcomes. Other variations appear less beneficial or require further study. These observations provide a guide to the types of variations possible with written action plans, and strongly support the use of individualised complete written action plans.

Exacerbations of asthma usually occur gradually over several days to weeks or on a background of chronic poor asthma control.  $^{1/2}$  This provides an opportunity for early intervention with corticosteroids and  $\beta$  agonists which act to reverse airflow obstruction and reduce the severity of the exacerbation. A written action plan facilitates the early detection and treatment of an exacerbation and is therefore an essential part of the self-management of exacerbations.  $^3$ 

Asthma is characterised by episodes of expiratory airflow obstruction which occur in response to multiple stimuli. The frequency and severity of these episodes varies greatly, both between and within individuals. Since all individuals with asthma are susceptible to exacerbations of asthma, it follows that all those with diagnosed asthma need to know how to manage these episodes. This instruction in self-management can be formalised as a written action plan, and all asthmatics are candidates for such a plan. This contrasts with the need for inhaled anti-inflammatory therapy which only becomes necessary when the frequency and/or severity of exacerbations are sufficiently great. At present, however, there is a paradoxical situation where most patients are prescribed regular inhaled corticosteroids (ICS) yet only a few either have<sup>4 5</sup> or use<sup>6</sup> a written action plan.

A recent systematic review of asthma self-management education conducted in adults<sup>3</sup> identified 17 randomised controlled trials (RCTs) which evaluated written action plans compared with usual care. When a patient was provided with a written action plan and instructed in its use in the context of self-monitoring and a review of asthma medications and severity, there were highly significant improvements in asthma health outcomes. The risk of being admitted to hospital for asthma fell by over 40% and presentations to the emergency department with asthma fell by over 20% (relative risk (RR) 0.78, 95% CI 0.67 to 0.91). Since action plans are

highly effective when part of a planned self-management programme and many versions are available, it is reasonable to try to determine the key components of asthma action plans that make a difference to subjects with asthma. This study addresses this question.

#### **METHODS**

#### **Definitions**

#### Written action plan

An action plan is a set of instructions prescribed to a patient with asthma for use in the management of deteriorating asthma. An individualised written action plan is tailored to the patient's underlying asthma severity and treatment. It is further characterised by being a written plan which informs the subject about when and how to modify medications and how to access the medical system in response to worsening asthma. For the purpose of this review, written action plans were further classified as follows:

- Individualised complete written action plans contained each of the following four components of an action plan<sup>7</sup>:
  - when to increase treatment (action point);
  - how to increase treatment;
  - for how long;
  - when to seek medical help.
- Incomplete individualised action plans comprised a second form of asthma self-management. These plans were individualised and specified when to increase treatment, but the use of early intervention with ICS was not specified.
- Non-specific action plans provided general information about the management of deteriorating asthma.

#### Action point

Action plans prescribe a level of symptoms or lung function that determines when to activate the action plan. This is termed an action point. There are several variations in action points. Action points may be based on symptoms or peak expiratory flow (PEF). PEF based action points may be based on predicted or personal best PEF. The number of action points in any written action plan can also vary, generally between two and four different levels.

#### Literature search

To determine the impact of the individual components of written action plans on asthma health outcomes, RCTs that evaluated asthma action plans were identified. An initial broad search was conducted for RCTs on asthma education as part of a wider systematic review of asthma self-management education.<sup>3</sup> Studies were identified from the Cochrane Airways Group Clinical Trials Register which is derived from Medline, Embase, Cinahl, hand searched respiratory journals, and meeting abstracts. Bibliographies of included studies were also searched. These articles were examined to identify those that included asthma action plans as part of the intervention. The asthma action plans were classified using the criteria described above.

#### Analysis of data

Data were extracted independently by two reviewers. The relative risk (RR) with 95% confidence intervals (CI) was calculated for dichotomous outcomes. The RR is the probability of experiencing an outcome when treated compared with the probability of experiencing that outcome if untreated, with values of <1 indicating a favourable treatment effect. For continuous outcomes using different units of measure, a standardised mean difference (SMD) and 95% CI was calculated using a fixed effects model. Significance was accepted at p<0.05. The pooled results were tested for heterogeneity using a  $\chi^2$  test with appropriate degrees of freedom. All outcomes were analysed according to the variations of the action plan and compared with a usual care control group. The variations analysed were: number of action points, personal best or percentage predicted PEF, "traffic light" action plan, or use of ICS or oral corticosteroids (OCS) alone.

#### **RESULTS**

Twenty six RCTs comparing action plans with usual care were identified (table 1), 17 of which used individualised complete written action plans.  $^{8-28}$  In 15 trials education, self-monitoring and regular medical review as well as provision of written action plans were incorporated into the self-management programmes.  $^{8-23}$   $^{25-28}$  Four trials  $^{29-32}$  used incomplete individualised action plans and five used non-specific action plans.  $^{33-37}$ 

#### Individualised complete written action plans Action points

Full details of the individual action points for the written action plans are shown in table 2. The number and level of action points for when to increase treatment varied, but each trial gave some instruction on increasing treatment. Fifteen trials set their first action points at 70–85% of the personal best or predicted PEF value.<sup>8–15</sup> 17–27</sup> Action plans that used a personal best PEF were used in nine trials and six used percentage predicted PEF (table S1, figs S1–S4 available online only at www.thoraxjnl.com/supplemental). When compared with usual care in a meta-analysis (five and four trials respectively), both types of action plan reduced hospital admissions (RR 0.66 for personal best and 0.46 for percentage predicted; fig 1), whereas only the action plan based on

personal best PEF reduced emergency room visits (RR 0.78; fig S2). Similarly, only the personal best written action plan led to improvement in airway calibre (SMD: PEF 0.56; fig 2).

The number of action points provided in the individualised plans ranged from two to four. Eight studies used written action plans with four action points (six that could be used in a meta-analysis), whereas seven studies used written action plans with less than four action points (three that could be used in a meta-analysis). The use of two or three action points was consistently beneficial (figs 1 and 2; also table S2 and figs S5–S7 available online only at www.thoraxjnl.com/supplemental), as was the use of four action points.

Action points were presented as a "traffic light" system<sup>38</sup> in four trials.<sup>8</sup> <sup>10–14</sup> <sup>18</sup> <sup>22</sup> This was not consistently better than a conventional action plan presentation (fig 3; also table S3 and figs S8–S10 available online only at www.thoraxjnl.com/supplemental).

The action plans were based on PEF in 10 trials<sup>10-17</sup> <sup>19-21</sup> <sup>25</sup> <sup>26</sup> <sup>28</sup> and on either PEF or symptoms in six trials.<sup>8</sup> <sup>10</sup> <sup>22-24</sup> <sup>27</sup> A previous systematic review found symptom based action plans to be equivalent to PEF based action plans.<sup>39</sup>

#### Treatment instruction

The instructions on how to increase corticosteroid treatment included the use of ICS and OCS in 13 trials and the use of OCS alone in four trials. ICS doses were increased by doubling the dose in 11 trials<sup>9-15</sup> 18-24 27 and by increasing or commencing ICS in two trials.<sup>8</sup> 25 The use of both ICS and OCS in the individualised plans was consistently beneficial (figs 1 and 2; also table S4 and figs S11–S14 available online only at www.thoraxjnl.com/supplemental). No comparison could be made with the use of OCS alone (four studies) because of insufficient data. No studies used action plans based on ICS alone.

#### Incomplete/non-specific plans

The efficacy of incomplete action plans was inconclusive as there were too few studies reporting data that could be used in a meta-analysis. Non-specific action plans led to an improvement in knowledge, 33 36 37 symptoms, 35 36 and reduced healthcare use in some 35 but not other studies. 29 Incomplete data reporting precluded meta-analysis.

#### **DISCUSSION**

This study has identified the key components of asthma action plans and examined the variations possible in preparing these plans. By using meta-analyses of data from RCTs, we have compared the different variations for their effect on key asthma outcomes such as hospital admissions, emergency room visits, and lung function. The results (table 3) give clear recommendations for preparing action plans and highlight areas needing further research.

#### Predicted or personal best PEF action points

Individualised action plans can be based upon the predicted PEF or the personal best PEF for that individual. Action points based on personal best PEF consistently improved health outcomes, whereas those based on percentage predicted PEF did not. With personal best PEF as the basis of the action plan, there were reductions in hospital admissions, emergency room visits, and improvement in PEF. This suggests that action points based on personal best PEF may perform better than those based on percentage predicted PEF. The likely reasons for this difference relate to individual variability in PEF results. However, significant differences between percentage predicted and personal best PEF could not be determined because of an overlap of the confidence intervals.

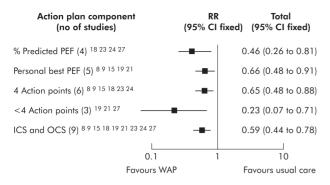
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		Treatment instructions					
Study	Individualised	When to increase treatment	How to increase treatment	How long to increase treatment	When to get help		
Individualised written o	action plans						
Cote et al <sup>®</sup>	Yes	Yes	Yes	Yes	Yes		
Cowie et al <sup>e</sup> *†	Yes	Yes	Yes	Yes	Yes		
Gallefoss et al <sup>10-14</sup>	Yes	Yes	Yes	Yes	Yes		
Ghosh <i>et al</i> 15*	Yes	Yes	Yes	Yes	Yes		
Grampian <sup>16</sup>	Yes	Yes	Yes	NS	Yes		
Hayward <i>et al</i> 17	Yes	Yes	NS	NS	Yes		
Heard <i>et al</i> 18†	Yes	Yes	Yes	Yes	Yes		
Ignacio-Garcia <sup>19</sup>	Yes	Yes	Yes	Yes	Yes		
Jones <i>et al</i> <sup>20</sup> *	Yes	Yes	Yes	Yes	Yes		
Lahdensuo <i>et al</i> 21	Yes	Yes	Yes	Yes	Yes		
Levy et al <sup>22</sup> *	Yes	Yes	Yes	Yes	Yes		
Moudgil <i>et al</i> <sup>23</sup> ‡	Yes	Yes	Yes	Yes	Yes		
Perneger <i>et al</i> ²⁴	Yes	Yes	Yes	NS	Yes		
Schermer <i>et al<sup>25</sup></i>	Yes	Yes	Yes	Yes	Yes		
Sommaragua <i>et al<sup>26</sup></i>	Yes	Yes	Yes	NS	Yes		
Yoon et al <sup>27</sup> §	Yes	Yes	Yes	Yes	Yes		
Zeiger <i>et al</i> <sup>28</sup>	Yes	Yes	Yes	NS	NS		
Incomplete action plan	S						
Bailey <i>et al</i> <sup>29</sup>	Yes	NS	NS	NS	NS		
George et al <sup>so</sup>	?	NS	NS	NS	NS		
Knoell <i>et al</i> <sup>31</sup>	Yes	NS	NS	NS	NS		
Kotses <i>et al</i> <sup>32</sup>	Yes	NS	NS	NS	NS		
Non-specific action pla	ans						
Abdulwadud <i>et al<sup>33</sup></i>	No	NS	NS	NS	NS		
Bailey <i>et al</i> ³⁴	No	NS	NS	NS	Yes		
Kotses et al <sup>35</sup>	No	NS	NS	NS	NS		
A II , B6	No	NS	NS	NS	NS		
Mulloy <i>et al<sup>s6</sup></i> Snyder <i>et al<sup>s7</sup></i>	No	NS	NS	NS	NS		

Initial versions of written action plans recommended that the same level of PEF should be applied to all patients in order to indicate when to increase treatment. This was

expressed as either percentage predicted or percentage personal best PEF. This commonly was set at 80% or 60% of the predicted peak flow value. Such a level would be

	When to inc	rease treatment	How to increase treatment			
Study	% Level	Traffic light	No of action point levels	PEF or symptom based	ICS	ocs
Cote et al <sup>8</sup>	85**	Yes	4	PEF/symptom	BDP 2000 μg	Yes
Cowie et al <sup>9</sup> *	70**	No	4	PEF/symptom	Double	Yes
Gallefoss et al <sup>10-14</sup>	80**	Yes	4	PEF	Double/triple	Yes
Ghosh et al <sup>15</sup> *	70**	No	4	PEF	Double .	Yes
Grampian <sup>16</sup>	NS	NS	NS	PEF	NS	Yes
Hayward <i>et al</i> <sup>17</sup>	70**	NS	2	PEF	NS	Yes
Heard et al <sup>18</sup> †	70††	Yes	4	NS†	Double	Yes
Ignacio-Garcia <sup>19</sup>	70**	No	3	PEF	Double	Yes
Jones <i>et al</i> <sup>20</sup> *	75**	No	4	PEF	Double	Yes
Lahdensuo <i>et al<sup>21</sup></i>	85**	No	2	PEF	Double	Yes
Levy et al <sup>22</sup> ¶	80††	Yes	3	PEF/symptom	Double	Yes
Moudgil et al <sup>23</sup> ‡	80††	No	4	PEF/symptom	Double	Yes
Perneger et al²⁴‡	80††	NS	4	PEF/symptom	Double	Yes
Schermer et al <sup>25</sup>	80**	NS	3	PEF	Double or commence	Yes
Sommaragua <i>et al</i> <sup>26</sup>	80††	No	3	PEF	No	Yes
Yoon <i>et al<sup>27</sup></i> §	80††	No	3	PEF/symptom	Double	Yes
Zeiger <i>et al</i> <sup>28</sup>	NS	NS	NS	PEF	NS	Yes



**Figure 1** Comparison of the effects of action plan components on hospital admissions for asthma. WAP=written action plan; PEF=peak expiratory flow; ICS=inhaled corticosteroids; OCS=oral corticosteroids.

inappropriate for someone with moderate asthma who could have daily variability of peak flow of up to 30%, and therefore their peak flow would regularly drop to 70% of the best during a stable period. Similarly, if there was any component of fixed airflow obstruction, the best peak flow may not reach 80% of the predicted value and the patient would be below their action point even when stable.

#### Number of action points

Action plans can use a variable number of action points. The studies in this review used two, three or four action points. Action plans with four action points were not consistently better than action plans with two or three action points (table S2 available online at www.thoraxjnl. com/supplemental). Using a greater number of action points leads to a narrower range of peak flow in each of the zones which may not be feasible to use in practice. These results argue in favour of a simpler approach using two or three action points. It may be that the improved precision obtained by using four action points is offset by the greater complexity of these plans which limits patient understanding and acceptance.

#### Symptoms versus peak flow based action plan

Written self-management plans can use action points that are based on either symptoms, peak flow, or both. Several RCTs have compared these approaches and found them to be equivalent for both the number of people admitted to hospital for asthma (RR 1.17, 95% CI 0.44 to 3.12) and those attending the emergency department (RR 0.91, 95% CI 0.61 to 1.35).<sup>39</sup>

#### Presentation of written action plan

A number of different ways of presenting action plans have been developed to facilitate their use. These include placing written self-management guidelines on a plastic card the size of a credit card,40 using electronic diaries or computerised "expert" systems, or the "traffic light" system where the colours green, yellow and red are used to signal continue usual treatment (green), increase treatment (yellow), or to seek help urgently (red).38 RCTs were identified that examined the "traffic light" approach. Individual written action plans without a "traffic light" action plan showed consistent benefits. There were fewer studies using a "traffic light" action plan configuration. Some outcomes were beneficial whereas others were not. This probably represents a lack of power rather than a lack of efficacy (table S3 available online at www.thoraxjnl.com/ supplemental).

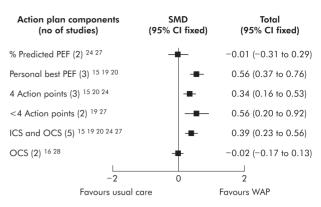


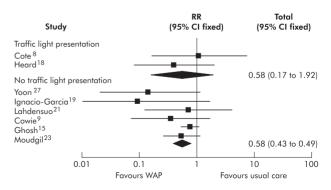
Figure 2 Comparison of the effects of action plan components on mean peak expiratory flow (PEF) in asthma. WAP=written action plan; ICS=inhaled corticosteroids; OCS=oral corticosteroids; SMD=standardised mean difference.

#### Treatment instruction: how to increase treatment

An exacerbation of asthma consists of a deterioration in both airflow obstruction and airway inflammation.41 42 It is therefore appropriate to recommend treatment with both bronchodilators and corticosteroids. A meta-analysis has confirmed the strong beneficial effect of OCS in severe exacerbations of asthma.43 The optimal treatment of mild exacerbations of asthma (forced expiratory volume in 1 second >60% predicted) is less clear. Current management practices include increased β<sub>2</sub> agonists, ICS, and OCS. A recent randomised trial conducted in a primary care setting compared fluticasone 1 mg twice daily with prednisone (40 mg daily and reducing). Treatment of a mild exacerbation with ICS had a comparable success rate to the use of ingested prednisone (48% v 48%). There was a surprisingly high rate of treatment failure in both groups (27% and 23%, respectively).44 This failure rate may reflect inadequate doses, inadequate duration of treatment, or non-eosinophilic exacerbations.45 Doubling ICS was found to be less effective than OCS in another trial. 46 The studies in this review support a combined approach where both ICS and OCS are used in the action plan.

#### Limitations

This review has used data derived from RCTs of asthma self-management programmes where written action plans were a key component. There are a number of limitations that need to be considered when reviewing these data. In some cases there were insufficient studies to allow a comparison and hence a type II error is possible. Where this is the case—for example, in the comparison of OCS with ICS and OCS—we



**Figure 3** Comparison of the effects of different action plan presentations (traffic light versus other) on hospital admissions for asthma. WAP=written action plan.

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Action plan variation	Result			
Action points				
Symptoms v PEF triggered	Equivalent			
Standard written instruction	Consistently beneficial			
Traffic light configuration	Not clearly better than			
2 2	standard instruction			
2–3 action points 4 action points	Consistently beneficial Not clearly better than			
4 delion points	Not clearly better than 4 points			
PEF based on	Consistently beneficial			
personal best PEF	consistent, consistent			
PEF based on %	Not consistently better			
predicted PEF	than usual care			
Treatment instruction				
Individualised WAP	Consistently beneficial			
using ICS and OCS	Consistently beneficial			
Individualised WAP	Insufficient data to evaluate			
using OCS only				
Individualised WAP	Insufficient data to evaluate			
using ICS only				

have been cautious in our interpretation of the data, reported the number of studies contributing to the meta-analysis, and only reported on outcomes where several studies contributed data. There are likely to be differences in the way the different self-management programmes were implemented. This does not seem to have a major impact on the results since there was no statistical heterogeneity identified in the key results.

#### Conclusion

The provision of individualised written action plans is of benefit to patients with asthma. Effective plans can be based on symptoms or PEF and use two, three or four action points. PEF based plans should use personal best PEF for the action point. The treatment instruction should include both ICS and OCS. With the data available, there was no clear distinction between other variations in the components of asthma action plans such as whether they were based on a traffic light system, the number of action points, or use of ICS alone or OCS alone. For these variations to be fully evaluated they need to be compared directly with each other, not a control group, in a randomised controlled trial setting of individualised written action plans. The observations of this review provide a guide to the types of variations possible with written action plans and strongly support the use of individualised complete written action plans.

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Figures S1-S14 and tables S1-S4 are available on the *Thorax* website at www.thoraxjnl.com/supplemental.

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### LUNG ALERT.....

## No clear benefit of parent initiated oral steroids in preschool children with viral

▲ Oommen A, Lambert PC, Grigg J. Efficacy of a short course of parent-initiated oral prednisolone for viral wheeze in children aged 1–5 years: randomised controlled trial. *Lancet* 2003;**362**:1433–8

sthma in children aged 1–5 years is characterised by recurrent transient episodes of wheeze triggered by viral colds. This is labelled "preschool viral wheeze" and is commonly treated with inhaled bronchodilators and oral corticosteroids. Persistent wheeze is associated with increased systemic eosinophil priming which is thought to be a risk factor for the development of atopic asthma.

In this randomised placebo controlled trial, children aged 1–5 years with known viral wheeze were stratified into either high or low eosinophil priming groups and parents administered 20 mg prednisolone or placebo for 5 days at the start of the next episode of viral wheeze. The primary outcomes were 7 day mean daytime and night time respiratory symptom scores.

217 children were randomised and outcome data were available for 51 and 69 children who received prednisolone or placebo, respectively. There was no improvement in daytime and night time symptom scores and no reduction in salbutamol use and hospital admissions in children treated with prednisolone compared with placebo. There were no differences between high or low eosinophil priming groups.

This study suggests no clear benefit of parent initiated oral prednisolone for preschool viral wheeze.

S Birring

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Fig S1: A comparison of the effects of personal best PEF vs % predicted action points on hospitalisations for asthma

Study or sub-category	VVAP n/N	Usual Care n/N	RR (fixed) 95% Cl	Weight %	RR (fixed) 95% Cl
01 Individualised WAP - PEF %	predicted				
Perneger 2002	4/57	6/58	<del></del>	16.30	0.68 [0.20, 2.28]
Yoon 1993	1/28	7/28	<del></del>	19.19	0.14 [0.02, 1.09]
Heard 1999	2/97	5/94		13.92	0.39 [0.08, 1.95]
Moudgil 2000	10/304	18/289	<del></del>	50.59	0.53 [0.25, 1.12]
Subtotal (95% CI)	486	469	<b>◆</b>	100.00	0.46 [0.26, 0.81]
Total events: 17 (WAP), 36 (U	sual Care)				•
Test for heterogeneity: Chi <sup>2</sup> = 1 Test for overall effect: Z = 2.7 02 Individualised WAP - PEF po	2 (P = 0.007)	,,			
Ignacio-Garcia 1995	0/35	5/35	<del>-</del>	8.22	0.09 [0.01, 1.58]
Lahdensuo 1996	2/56	3/59		4.36	0.70 [0.12, 4.05]
Cote 1997	2/50	2/54		2.87	1.08 [0.16, 7.38]
Cowie 1997	2/46	6/48		8.77	0.35 [0.07, 1.64]
Ghosh 1998	38/140	50/136	=	75.77	0.74 [0.52, 1.05]
Subtotal (95% Cl)	327	332	•	100.00	0.66 [0.48, 0.91]
Total events: 44 (WAP), 66 (U:	sual Care)				
Test for heterogeneity: Chi² = : Test for overall effect: Z = 2.5		%			
			0.01 0.1 1 10	100	
			Favours WAP Favours Usi	ual Care	

Fig S2: A comparison of the effects of personal best PEF vs % predicted action points on ER visits for asthma

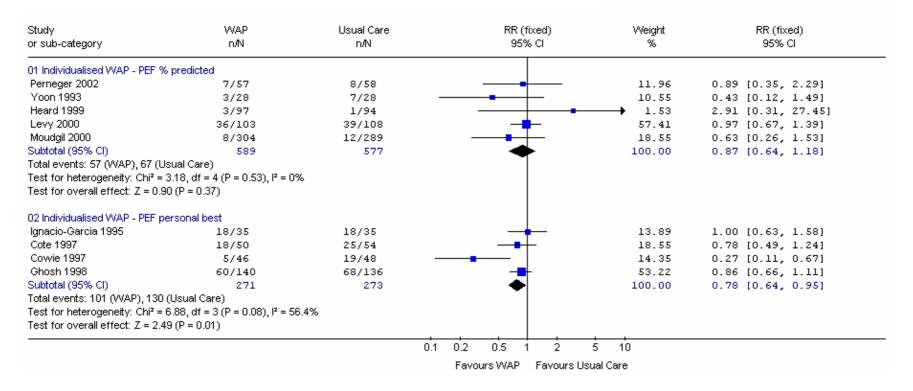


Fig S3: A comparison of the effects of personal best PEF vs % predicted action points on days off work for asthma

Study	WAP	Usual care	RR (fixed)	Weight	RR (fixed)
or sub-category	n/N	n/N	95% CI	<u>~</u>	95% CI
01 Individualised WAP - PEF %	predicted				
Perneger 2002	8/57	11/58	<del></del>	21.19	0.74 [0.32, 1.70]
Yoon 1993	5/28	4/28		7.77	1.25 [0.37, 4.17]
Heard 1999	34/97	36/94	<del>-</del>	71.04	0.92 [0.63, 1.33]
Subtotal (95% Cl)	182	180	•	100.00	0.90 [0.65, 1.26]
Total events: 47 (WAP), 51 (U:	sual care)		٦		
Test for heterogeneity: Chi² = (	$0.50$ , df = $2 (P = 0.78)$ , $I^2 = 0$	%			
Test for overall effect: $Z = 0.6$	0 (P = 0.55)				
02 Individualised WAP - PEF pe	ersonal best				
Ignacio-Garcia 1995	24/35	29/35	<del></del>	54.36	0.83 [0.63, 1.08]
Lahdensuo 1996	13/56	25/59		45.64	0.55 [0.31, 0.96]
Subtotal (95% CI)	91	94	•	100.00	0.70 [0.53, 0.92]
Total events: 37 (WAP), 54 (U:	sual care)		- T		
Test for heterogeneity: Chi² = :	2.21, df = 1 (P = 0.14), $I^2$ = 5	4.7%			
Test for overall effect: $Z = 2.5$	2 (P = 0.01)				
				5 10	
			Favours WAP Favours Us	ual Care	

Fig S4: A comparison of the effects of personal best PEF vs % predicted action points on PEF (mean) for asthma

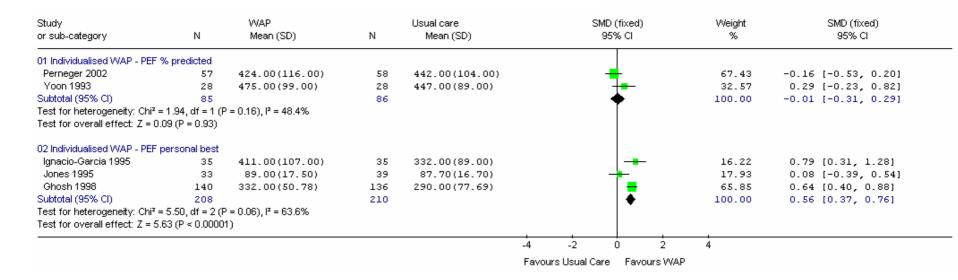


Fig S5: A comparison of the number of action points in written action plans on the outcome of hospitalisations

Study or sub-category	VVAP n/N	Usual care n/N	RR (fixed) 95% Cl	VVeight %	RR (fixed) 95% Cl
01 Individualised WAP (4 action	n points)				
Perneger 2002	4/57	6/58		6.76	0.68 [0.20, 2.28]
Cote 1997	2/50	2/54		2.19	1.08 [0.16, 7.38]
Cowie 1997	2/46	6/48		6.67	0.35 [0.07, 1.64]
Ghosh 1998	38/140	50/136	<del></del>	57.64	0.74 [0.52, 1.05]
Heard 1999	2/97	5/94		5.77	0.39 [0.08, 1.95]
Moudgil 2000	10/304	18/289	<del></del>	20.97	0.53 [0.25, 1.12]
Subtotal (95% CI)	694	679	•	100.00	0.65 [0.48, 0.88]
02 Individualised WAP (< 4 act	ion points)				
Yoon 1993	1/28	7/28		45.39	0.14 [0.02, 1.09]
Ignacio-Garcia 1995	0/35	5/35 ←		35.66	0.09 [0.01, 1.58]
Lahdensuo 1996	2/56	3/59		18.95	0.70 [0.12, 4.05]
Subtotal (95% CI)	119	122		100.00	0.23 [0.07, 0.71]
Total events: 3 (WAP), 15 (Ust	ual care)		_		
Test for heterogeneity: $Chi^2 = 2$ Test for overall effect: $Z = 2.53$	2.18, df = 2 (P = 0.34), l² = 8	3.1%			
		0.01	0.1 1 10	100	
			Favours WAP Favours Us	ual Care	

Fig S6: A comparison of the number of action points in written action plans on the outcome of ER visits

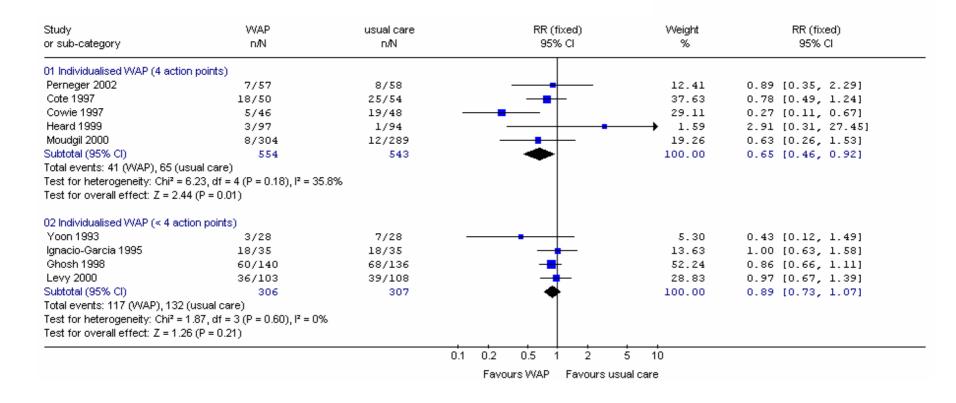


Fig S7: A comparison of the number of action points in written action plans on the outcome of PEF (mean)

Study or sub-category	N	WAP Mean (SD)	N	Usual care Mean (SD)	SMD (fixed) 95% Cl	Weight %	SMD (fixed) 95% Cl
D1 Individualised WAP - 4 action	on points						
Perneger 2002	57	424.00(116.00)	58	442.00(104.00)	<del></del>	25.56	-0.16 [-0.53, 0.20]
Jones 1995	33	89.00(17.50)	39	87.70(16.70)	<del></del>	15.94	0.08 [-0.39, 0.54]
Ghosh 1998	140	332.00(50.78)	136	290.00(77.69)	=	58.51	0.64 [0.40, 0.88]
Subtotal (95% CI)	230		233		♦	100.00	0.34 [0.16, 0.53]
Test for overall effect: Z = 3.6 02 Individualised WAP - < 4 ac		)					
Yoon 1993	28	475.00(99.00)	28	447.00(89.00)	<del> =</del> -	46.15	0.29 [-0.23, 0.82]
	35	411.00(107.00)	35	332.00(89.00)	<del></del> -	53.85	0.79 [0.31, 1.28]
Ignacio-Garcia 1995	35					33.03	
	35 63		63		•	100.00	0.56 [0.20, 0.92]

Fig S8: A comparison of action plan presentations (traffic light vs other) on the outcome hospitalisations

Study or sub-category	WAP n/N	Usual Care n/N	RR (fixed) 95% Cl	Weight %	RR (fixed) 95% Cl
01 Individualised WAP (Traffic	liaht)				
Cote 1997	2/50	2/54		27.47	1.08 [0.16, 7.38]
Heard 1999	2/97	5/94	<del></del>	72.53	0.39 [0.08, 1.95]
Subtotal (95% CI)	147	148		100.00	0.58 [0.17, 1.92]
Total events: 4 (WAP), 7 (Usua	al Care)				, ,
Test for heterogeneity: Chi² = 0	$0.64$ , df = 1 (P = $0.42$ ), $I^2 = 0^\circ$	%			
Test for overall effect: Z = 0.90	0 (P = 0.37)				
02 Individualised WAP (No traft	fic light)				
Yoon 1993	1/28	7/28		7.74	0.14 [0.02, 1.09]
Ignacio-Garcia 1995	0/35	5/35	<del></del>	6.08	0.09 [0.01, 1.58]
Lahdensuo 1996	2/56	3/59		3.23	0.70 [0.12, 4.05]
Cowie 1997	2/46	6/48	<del></del>	6.49	0.35 [0.07, 1.64]
Ghosh 1998	38/140	50/136	-	56.07	0.74 [0.52, 1.05]
Moudgil 2000	10/304	18/289	<del></del>	20.40	0.53 [0.25, 1.12]
Subtotal (95% CI)	609	595	<b>◆</b>	100.00	0.58 [0.43, 0.79]
Total events: 53 (WAP), 89 (Us	sual Care)				
Test for heterogeneity: Chi² = 5	$5.75$ , df = $5$ (P = $0.33$ ), $I^2$ = $13$	3.1%			
Test for overall effect: Z = 3.50	0 (P = 0.0005)				
		0	.01 0.1 1 10	100	
			Favours WAP Favours Usi	ual Care	

Fig S9: A comparison of action plan presentations (traffic light vs other) on the outcome ER visits

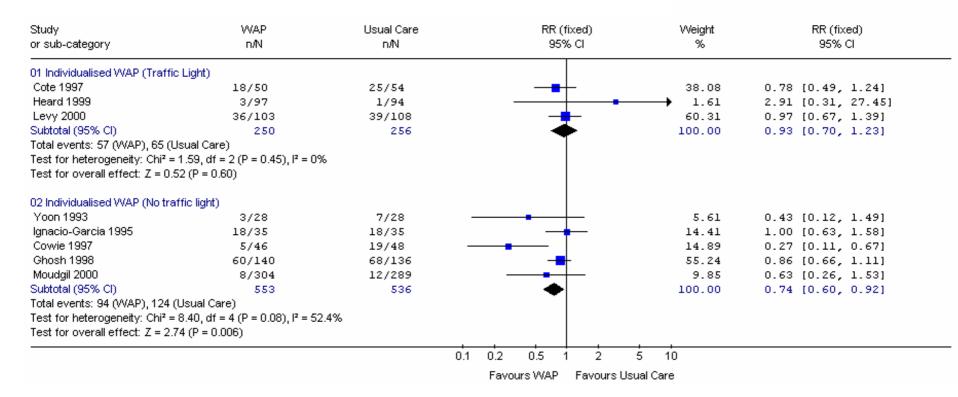


Fig S10: A comparison of action plan presentations (traffic light vs other) on the outcome unscheduled Dr visits

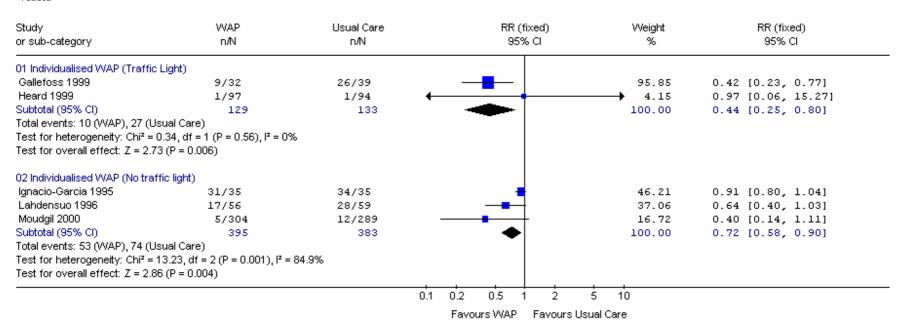


Fig S11: The efficacy of action plan treatments using ICS and OCS based plans on hospitalisations

Study or sub-category	VVAP n/N	Usual Care n/N	RR (fixed) 95% Cl	Weight %	RR (fixed) 95% CI
01 Individualised WAP - ICS &	ocs				
Perneger 2002	4/57	6/58	-	5.75	0.68 [0.20, 2.28]
Yoon 1993	1/28	7/28	<del></del>	6.77	0.14 [0.02, 1.09]
Ignacio-Garcia 1995	0/35	5/35	<b>₹</b>	5.32	0.09 [0.01, 1.58]
Lahdensuo 1996	2/56	3/59	A	2.83	0.70 [0.12, 4.05]
Cote 1997	2/50	2/54	111	1.86	1.08 [0.16, 7.38]
Cowie 1997	2/46	6/48	•	5.68	0.35 [0.07, 1.64]
Ghosh 1998	38/140	50/136		49.05	0.74 [0.52, 1.05]
Heard 1999	2/97	5/94	<del>• •</del>	4.91	0.39 [0.08, 1.95]
Moudgil 2000	10/304	18/289	-	17.84	0.53 [0.25, 1.12]
Subtotal (95% CI)	813	801	•	100.00	0.59 [0.44, 0.78]
Total events: 61 (WAP), 102 (	Usual Care)		444000		Control of
Test for heterogeneity: Chi <sup>2</sup> =		%			
Test for overall effect: $Z = 3.6$					
		0	1 0.2 0.5 1 2	5 10	
			Favours WAP Favours Us	ual Care	

Fig S12: The efficacy of action plan treatments using ICS and OCS based plans on ER Visits

Study or sub-category	WAP n/N	Usual care n/N	RR (fixed) 95% Cl	Weight %	RR (fixed) 95% CI
01 Individualised WAP - ICS &	ocs				
Perneger 2002	7/57	8/58	23 <u>2</u> 2 29	4.05	0.89 [0.35, 2.29]
Yoon 1993	3/28	7/28 -		3.57	0.43 [0.12, 1.49]
Ignacio-Garcia 1995	18/35	18/35	(	9.19	1.00 [0.63, 1.58]
Cote 1997	18/50	25/54	-	12.27	0.78 [0.49, 1.24]
Cowie 1997	5/46	19/48 —		9.49	0.27 [0.11, 0.67]
Ghosh 1998	60/140	68/136	17 (17 (17 (17 (17 (17 (17 (17 (17 (17 (	35.21	0.86 [0.66, 1.11]
Heard 1999	3/97	1/94	X	0.52	2.91 [0.31, 27.45]
Levy 2000	36/103	39/108	_	19.43	0.97 [0.67, 1.39]
Moudgil 2000	8/304	12/289		6.28	0.63 [0.26, 1.53]
Subtotal (95% CI)	860	850	•	100.00	0.81 [0.69, 0.96]
Total events: 158 (WAP), 197	(Usual care)		10.40		CONTRACTOR OF THE PROPERTY OF
Test for heterogeneity: Chi² = 1	$10.12$ , df = 8 (P = $0.26$ ), $I^2$ = 1	21.0%			
Test for overall effect: $Z = 2.4$					
	(C)	0.1	0.2 0.5 1 2	5 10	
		0.1			
			Favours VVAP Favours Us	ual Care	

Fig S13: The efficacy of action plan treatments using ICS and OCS based plans on FEV<sub>1</sub> (mean)

Study or sub-category	N	WAP Mean (SD)	N	Usual Care Mean (SD)		SMD (fixed) 95% Cl	Weight %	SMD (fixed) 95% Cl
01 Individualised WAP - ICS	& OCS							
Yoon 1993	28	2.79(0.81)	28	2.81(0.77)		<del>-</del>	29.02	-0.02 [-0.55, 0.50]
Ignacio-Garcia 1995	35	80.45(19.52)	35	65.48(22.01)		<del></del>	33.99	0.71 [0.23, 1.20]
Jones 1995	33	83.20(18.00)	39	81.20(18.30)		<del></del>	36.99	0.11 [-0.36, 0.57]
Subtotal (95% CI)	96		102			•	100.00	0.27 [-0.01, 0.56]
Test for heterogeneity: Chi <sup>2</sup> : Test for overall effect: Z = 1		= 0.09), I <sup>2</sup> = 59.0%						
03 Individualised WAP - OCS	,							
Zeiger 1991	92	92.90(22.60)	92	88.70(21.90)		<del> -</del>	26.43	0.19 [-0.10, 0.48]
Grampian 1994	250	74.60(27.80)	260	75.40(27.70)		<u>=</u>	73.57	-0.03 [-0.20, 0.14]
Subtotal (95% CI)	342		352				100.00	0.03 [-0.12, 0.18]
Test for heterogeneity: Chi2:	= 1.58, df = 1 (P	= 0.21), I <sup>2</sup> = 36.8%				ſ		·
Test for overall effect: Z = 0	.38 (P = 0.71)							
					-4 -2	0 2	4	
					Favours Usu	ual Care Favours VVA	.P	

Fig S14: The efficacy of action plan treatments using ICS and OCS based plans on PEF (mean)

Study or sub-category	N	VVAP Mean (SD)	N	Usual Care Mean (SD)	SMD (fixed) 95% CI	Weight %	SMD (fixed) 95% Cl
01 Individualised WAP - ICS & 0	ocs						
Perneger 2002	57	424.00(116.00)	58	442.00(104.00)	<del>-</del>	20.16	-0.16 [-0.53, 0.20]
Yoon 1993	28	475.00(99.00)	28	447.00(89.00)	+-	9.74	0.29 [-0.23, 0.82]
Ignacio-Garcia 1995	35	411.00(107.00)	35	332.00(89.00)	-	11.37	0.79 [0.31, 1.28]
Jones 1995	33	89.00(17.50)	39	87.70(16.70)	<del></del>	12.57	0.08 [-0.39, 0.54]
Ghosh 1998	140	332.00(50.78)	136	290.00(77.69)	-	46.16	0.64 [0.40, 0.88]
Subtotal (95% CI)	293		296		♦	100.00	0.39 [0.23, 0.56]
Test for overall effect: Z = 4.66	6 (P < 0.0000	1)					
03 Individualised WAP - OCS o	nly						
Zeiger 1991	88	109.20(36.50)	87	104.30(32.20)	<del> -</del>	25.52	0.14 [-0.16, 0.44]
Grampian 1994	250	335.00(120.00)	260	345.00(130.00)	<u> </u>	74.48	-0.08 [-0.25, 0.09]
Subtotal (95% CI)	338		347		•	100.00	-0.02 [-0.17, 0.13]
Test for heterogeneity: Chi² = 1	1.59, df = 1 (F	P = 0.21), I <sup>2</sup> = 37.3%					
Test for overall effect: $Z = 0.30$	O(P = 0.76)						
				-4	-2 0	2 4	
				•	ours Usual Care - Favoi		

 $\textbf{Table S1} \ \textbf{Outcome comparisons for action points based on \% predicted or personal}$ 

best PEF

Outcome	% Predicted PEF	No. of studies	Personal best PEF	No. of studies
Hospitalisations*	$0.46 (0.26 \text{ to } 0.81)^+$	4[18,23,24,27]	$0.66 (0.48 \text{ to } 0.91)^+$	5[8,9,15,19,2]1
ER visits*	0.87 (0.64 to 1.18)	5[18,22,23,24,27]	$0.78 (0.64 \text{ to} 0.95)^+$	4[8,9,15,19]
Unscheduled Dr	0.44 (0.17 to 1.14)	2[18,23]	$0.69 (0.56 \text{ to } 0.85)^+$	3[19,21,10-14]
visits*				
Days off work*	0.90 (0.65 to 1.26)	3[18,24,27]	$0.70 (0.53 \text{ to } 0.92)^+$	2[19,21]
Nocturnal asthma*	0.65 (0.38 to 1.11)	2[18,24]	1.0 (0.92 to 1.08)	1[19]
$FEV_1^{\ \#}$	-0.02 (-0.55 to 0.50)	1[27]	$0.40~(0.06~\text{to}~0.73)^{+}$	2[19,20]
PEF <sup>#</sup>	-0.01 (-0.31 to 0.29)	2[24,27]	$0.56 (0.37 \text{ to } 0.76)^+$	3[15,19,20]

<sup>\*</sup>Relative risk (95% CI).

\*Standardised mean difference (95% CI).

+ p<0.05.

 Table S2 Outcome comparisons for number of action points

Outcome	4 Action points	No. of studies	< 4 Action points	No. of studies
Hospitalisations*	$0.65~(0.48~{ m to}~0.88)^+$	6[8,9,15,18,23,24]	$0.23 (0.07 \text{ to } 0.71)^+$	3[19,21,27]
ER visits*	$0.65 (0.46 \text{ to } 0.92)^+$	5[8,9,18,23,24]	0.89 (0.73 to 1.07)	4[19,21,22,27]
Unscheduled Dr	0.44 (0.17 to 1.14)	2[18,23]	$0.79~(0.64~\text{to}~0.97)^+$	2[19,21]
visits*				
Days off work*	0.87 (0.62 to 1.23)	2[18,24]	$0.74 (0.56 \text{ to } 0.97)^+$	3[19,21,27]
Nocturnal	0.65 (0.38 to 1.11)	2[18,24]	1.0 (0.92 to 1.08)	1[19]
asthma*				
$FEV_1^{\#}$	0.11 (-0.36 to 0.57)	1[20]	$0.37~(0.02~\text{to}~0.73)^+$	2[19,27]
PEF <sup>#</sup>	$0.34 (0.16 \text{ to } 0.53)^+$	3[15,20,24]	$0.56 (0.20 \text{ to } 0.92)^+$	2[19,27]

<sup>\*</sup>Relative risk (95% CI).

\*Standardised mean difference (95% CI).

† p<0.05.

 Table S3 Outcome comparisons for 'traffic light' configuration of action plans

Outcome	Traffic light	No. of studies	No traffic light	No. of studies
Hospitalisations*	0.58 ( (0.17 to 1.92)	2[8,18]	$0.58 (0.43 \text{ to } 0.79)^+$	6[8,15,19,21,23,27]
ER visits*	0.93 (0.70 to 1.23)	3[8,18,22]	$0.74 (0.60 \text{ to } 0.92)^+$	5[9,15,19,23,27]
Unscheduled Dr	$0.44~(0.25~\text{to}~0.80)^{+}$	2[10-14,18]	$0.72 (0.58 \text{ to } 0.90)^+$	3[19,21,23]
visits*				
Days off work*	0.92 (0.63 to 1.33)	1[18]	$0.74~(0.56~\text{to}~0.97)^{+}$	3[19,21,27]
Nocturnal asthma*	$0.36 (0.16 \text{ to } 0.81)^+$	1[18]	1.0 (0.92 to 1.08)	1[19]

<sup>\*</sup>Relative risk (95% CI).
+ p<0.05.

 Table S4 Outcome comparisons for corticosteroids

Outcome	ICS & OCS	No. of studies	OCS only	No. of studies
Hospitalisations*	0.59 (0.44 to 0.78) <sup>+</sup>	9[8,9,15,18,19,21,23,24,27]	0.43 (0.08 to 2.18)	1[28]
ER visits*	$0.81~(0.69~\text{to}~0.96)^{+}$	9[8,9,15,18,19,22,23,24,27]	$0.67~(0.46~\text{to}~0.97)^{+}$	1[28]
Days off work <sup>#</sup>	$-0.35 (-0.49 \text{ to } -0.21)^+$	6[8,9,10-14,18,19,21]	-0.13 (-0.34 to 0.08)	3[17,26,28]
Nocturnal asthma*	0.84 (0.67 to 1.05)	3[18,19,24]	$0.30 (0.16 \text{ to } 0.55)^+$	1[28]
FEV1 <sup>#</sup>	0.27 (-0.01 to 0.56)	3[19,20,27]	0.03 (-0.12 to 0.18)	2[16,28]
PEF#	$0.39 (0.23 \text{ to } 0.56)^+$	5[15,19,20,24,27]	-0.02 (-0.17 to 0.13)	2[16,28]

ICS: inhaled corticosteroid. OCS: oral corticosteroid.

<sup>\*</sup>Relative risk (95% CI).

\*Standardised mean difference (95% CI).

<sup>&</sup>lt;sup>+</sup> p<0.05.