When can personal best peak flow be determined for asthma action plans?

H K Reddel, G B Marks, C R Jenkins

Background: Written asthma action plans based on personal best peak expiratory flow (PEF) consistently improve health outcomes, whereas those based on predicted PEF do not. Guidelines state that personal best PEF should be assessed over 2–3 weeks during good asthma control, but it is unclear how long to wait after commencing or changing treatment.

METHODS

Electronically recorded spirometric data from 61 subjects with initially poorly controlled asthma from a 72 week budesonide study were analysed. For each week, average morning pre-bronchodilator PEF was calculated and personal best PEF was determined as the highest PEF in the previous 2 weeks. The time to plateau was defined as the week beyond which no further improvement occurred.

RESULTS: At baseline, average morning PEF was 61% predicted and personal best PEF was 87% predicted. Personal best PEF from twice daily monitoring increased to a plateau of 95% predicted (p<0.0001) after only 3 weeks of budesonide treatment. However, average morning PEF continued to improve for 3 months and “as needed” reliever use for 7 months.

Conclusions: Personal best PEF is a useful concept for asthma self-management plans when determined as the highest PEF over the previous 2 weeks. With twice daily monitoring, personal best PEF reaches plateau levels after only a few weeks of corticosteroid treatment.
plateau in personal best PEF based on twice daily PEF measurements was reached after only 3 weeks of treatment when reliever use was still 0.9 occasions/day (IQR 0.3–2.9). The plateau in personal best PEF was delayed to 8 weeks if once daily (morning) PEF values were analysed. Average morning pre-bronchodilator PEF continued to improve to week 13 (467 l/min, 84% predicted, 95% CI 79 to 90) and reliever use to week 30 (0.1 occasions/day, IQR 0.0–0.8); p<0.0001 compared with week 3 for each.

**DISCUSSION**

This study showed that personal best PEF from 2 weeks of twice daily measurements reached a plateau very early after initiation of high dose budesonide in patients with poorly controlled asthma. These findings indicate that the action points for a PEF based action plan can be assigned after only a short period of inhaled corticosteroid treatment, without reliever use to week 30 (0.1 occasions/day, IQR 0.0–0.8); p<0.0001 compared with week 3 for each.

Written asthma action plans are an essential component of asthma self-management education, and their use has been associated with significant improvements in health care utilisation, morbidity, and quality of life. In Australia, although written action plans have been recommended in asthma guidelines for 15 years, the proportion of patients possessing a plan declined by almost half between 1996 and 2001 to only 22%. Confusion for clinicians may have contributed to this decline as, despite general advice to individualise the plan to each patient, asthma guidelines contain few specific instructions about how this should be done. A recent review of the critical components of action plans found additional benefits when they were based on personal best rather than predicted PEF. However, the need to standardise the assessment and recording of personal best PEF has been highlighted by a study of 104 patients presenting with acute asthma. At presentation, only 30 patients knew their personal best PEF and, of these, 10/22 recorded a higher PEF over the next 24 days.

Current guidelines state that personal best PEF should be determined during good asthm control. However, some patients only present when their asthma is uncontrolled and do not return after they have completely recovered, leaving no opportunity to reassess their personal best PEF during good asthma control. This problem could be avoided by basing the action plan on predicted PEF, but such plans show fewer benefits. As patients are less likely to follow medical instructions which are discordant with their own lay beliefs, adherence may be poorer with a plan that specifies action at a PEF which the patient considers to be too high or too low.

The present study shows that personal best PEF can be determined long before symptoms and reliever use have stabilised, when the patient may still be motivated to return for review.

Because the present study was based on the analysis of outlying values, rigorous quality control analysis was undertaken. Within-session reproducibility of PEF, FEV1, and FVC was well within guidelines for supervised spirometry. Data falsification was eliminated by electronic monitoring, and the use of spirometric rather than PEF manoeuvres may have contributed to the low rate of artefactual PEF values. Although long term adherence with monitoring using mechanical PEF meters and paper diaries is notoriously poor, the present recommendations should be applicable to conventional monitoring as adherence is reasonable over shorter periods. The chance of inappropriate changes in medication will be reduced by ensuring correct PEF technique, and by re-establishing personal best every few years or if the PEF meter is changed. This advice will be particularly important in Europe with the imminent introduction of non-linear PEF scales. It is of interest that, despite the known limitations of conventional PEF monitoring, significant health benefits have been found with the use of action plans based on this type of monitoring.

Previous studies have examined the time course of change in average morning PEF but not personal best PEF. It is predictable that maximum PEF would improve more rapidly than average morning PEF, as inhaled corticosteroids reduce PEF variability and improve lung function. The present study was carried out in patients with very poorly controlled asthma at entry who were started on inhaled corticosteroid treatment. The recommendations should be more widely applicable as the time course of improvement in asthma outcomes is even shorter for patients with milder asthma. An earlier plateau in personal best PEF would also be expected with use of long acting β2 agonists, given their more rapid onset of effect than inhaled corticosteroid alone. One limitation of the present study is that patients were excluded if they experienced a viral respiratory infection during the run-in period; however, because of the observed reduction in PEF during viral exacerbations (mean 10 days), it would appear advisable to delay determination of personal best PEF until after clinical resolution of a respiratory infection.

The prolonged improvement in bronchodilator use in the present study is of interest. The time course of improvement for asthma symptoms was similar (data not shown), suggesting that reliever medication was not just being used habitually. This prolonged improvement may instead reflect a
reduction in predisposition to bronchoconstriction, associated with continuing improvement in airway hyperresponsiveness over 18 months despite down-titration of inhaled corticosteroids. Preliminary results from the GOAL study indicate that patients treated with fluticasone/salmeterol combination also have a gradual reduction in bronchodilator use over many months. The findings from these studies suggest that action plans based on reliever use may need revision after several months of inhaled corticosteroid or combination treatment.

The phrase “personal best” is already familiar to the general public through sports reports which commonly refer to an athlete’s “PB”—that is, their “best ever” time/height/distance. The distinction between the sports definition of PB as “best ever” and the guidelines based definition of personal best PEF to help reduce the risk of re-presentation.

References


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