

# Evaluation of a Nurse-run asthma clinic in general practice

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## ABSTRACT

**Background:** Since 1985, nurse-run asthma clinics have been developing and are now widespread in the United Kingdom, having been greatly stimulated by the New Contract for General Practice (1990). To date, there have been few definitive studies evaluating such clinics. This study, in two adjacent general practices in semi-rural Somerset, evaluates the effect of nurse-run clinics on patients' symptoms, pulmonary function, and healthcare utilisation over a four-month period.

**Methods:** One hundred and twenty-nine subjects aged 2-79 years, were recruited from the two practices over a one-year period (1991-1992). All patients identified in either practice who had used preventive therapy in the preceding 12 months were eligible and were included if they had a positive exercise or reversibility (20%) test and had not previously received formal asthma education.

A pre- and post-experimental study design was used. Subjects' symptoms, pulmonary function, and health care in the preceding eight weeks were assessed at intake and at 16 weeks; the nurse education being delivered at intake, one and four weeks.

**Results:** Both mean number of days off work/school and mean number of nights on which waking occurred fell ( $p < 0.001$ ). Peak expiratory flow rate and forced expiratory volumes improved during the study period ( $p < 0.001$  and  $p < 0.01$  respectively). Use of antibiotics, oral steroids, and rescue bronchodilator use all improved significantly ( $p < 0.001$ ). Physician attendances were also reduced.

**Conclusions:** A nurse-run asthma clinic in general practice is an effective way to improve symptoms, pulmonary function, and health care utilisation for asthmatic patients.

## Introduction

Nearly fifteen years ago, Greta Barnes, in a letter to the *Journal of the Royal College of General Practitioners*<sup>1</sup> described nurse-run asthma clinics in general practice. She and Robert Pearson in Stratford UK, developed a protocol for one-on-one patient education by practice nurses.

The New Contract for General Practice 1990<sup>2</sup> which is based on wellness and health, and financially favours health promotion and health maintenance activities in general practice, has also been a stimulus to the development of such clinics.

A 1992 survey of 22 practices by Arthur Hibble in East Anglia has found that practice nurses had almost doubled their working hours since 1989 and in 1990 spent 10% of their time on health promotion activities, 28% of the latter being in asthma clinics<sup>3</sup>.

To date, there have been few studies evaluating nurse-run clinics<sup>4-6</sup>. Hoskins has recently demonstrated patient satisfaction with nurse-run asthma clinics in General Practice. Previously reported studies have not isolated the nurse education component or educational programs in general practice. In 1986 Mitchell in New Zealand assessed the effects of nurse education conducted in the asthmatics' home. Three hundred and sixty-eight children were involved but there was no demonstrable change in the school attendance, symptoms, or frequency of severe attacks<sup>7</sup>.

Usherwood in Sheffield (1988) evaluated a doctor-run clinic comparing 31 children with an equal number from another practice. They found a marginal improvement in absences from school and in the number of home visits and out-of-hours calls. There was no difference in symptoms between the groups but the study group used their inhalers more and had more booked follow-up appointments at the clinic<sup>8</sup>.

In 1989, Beasley described a significant improvement in patient morbidity using a self-management plan but this was based on a hospital clinic and lacked control group<sup>9</sup>. Charlton's 1991 study in East Anglia evaluated the combined effect of a nurse-run clinic and a self-management plan but did not distinguish which part of the package was responsible for the observed improvements in steroid use, nebuliser use, physician consultations, and out-of-hours calls<sup>10</sup>.

In 1995, Kevin Jones reported a study of peak flow based self-management in general practice but his patients saw either physicians or nurses when attending the clinic<sup>1</sup>. In September 1995, Kevin Jones, writing in the *British Journal of General Practice* suggested "a randomised-controlled trial of nurse-run asthma care would now be difficult to conduct, and so it may be necessary to accept nurse-run asthma care without definitive proof of its clinical effectiveness"<sup>2</sup>. However, Greta Barnes has called for "research programs to evaluate asthma management and measure patient outcomes as a result of health professionals receiving training"<sup>3</sup>. The purpose of this study is to provide a measure of the required proof.

**Method****The Site**

The study was undertaken in two adjacent general practices in semi-rural Somerset, England. Each practice had a practice nurse trained to diploma level at the National Asthma and Respiratory Training Centre (NARTC)

**The Subject**

The subjects were defined as all patients in the two practices aged from two to 79 years, inclusive, who had used preventive therapy in the preceding 12 months. In this way, patients with mild or seasonal asthma were excluded from the study

Subjects were recruited from physician consultations, from repeat prescription contacts, and from the general asthma registers in the practices. Subjects were recruited over a one-year period at a rate that the practice nurses could accommodate in their schedule of approximately one afternoon per week

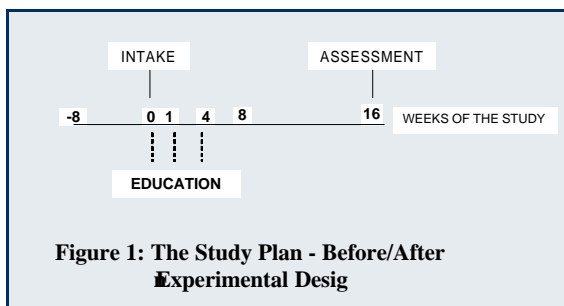
Patients were eligible for inclusion if they had demonstrated 20% reversibility in airflow obstruction measured by peak expiratory flow rate after  $\beta$ -2 agonist inhalation, or if they had a positive exercise test reversed by bronchodilators

Patients were excluded from the study if they regularly used a nebuliser or if they had had contact with the practice nurse about their asthma before the study started. This involved eight patients in one practice and the study did not include the worst or least affected patients.

**The Study Design**

The study design is shown in Figure 1. At the intake visit, an initial assessment of the patient's asthma was made by the nurse. The study was explained, detailed asthmatic history taken, height and weight recorded, and a predicted maximum peak expiratory flow rate calculated.

Diagnosis was confirmed by spirometry before an initial bronchodilator therapy. An exercise provocation



**Figure 1: The Study Plan - Before/After Experimental Design**

test was performed if the history indicated. Details of current medications were discussed and advice given on their proper use. Note, no changes were made to therapy or total daily doses. Inhaler technique was checked with retraining where necessary, and follow-up was arranged for one week's time. Details of asthma related morbidity over the preceding eight weeks were obtained from the patient interview and the patient's chart which, in the U.K., is relatively comprehensive for all physician contacts and hospital visits. The intervention consisted of advice on the us

Analysis of data comparing the pre- and post-intervention data was done by standard statistical methods using the Wilcoxon sign test and two (independent) sample *t*-test. The following data were compared in three groups

**Symptom**

- Number of days off work or school
- Number of nights when waking due to asthma occurred

**Pulmonary Function**

- Spirometry before and after bronchodilator use looking at peak flow rate (PEFR), forced expiratory volume in one second (FEV<sub>1</sub>), and functional vital capacity (FVC)

**Healthcare Utilisation**

- Excess bronchodilator use (number of days on which patient used more than two puffs four times in the day if on regular  $\beta$ -2-agonist or on which relief  $\beta$ -2-agonist was used by patients not on a regular  $\beta$ -2-agonist regimen)
- Courses of oral steroid
- Courses of oral antibiotic
- Days when nebuliser was used
- Physician consultation
- Home visits for asthma

of medication, education in inhaler technique and basic asthma information according to the patient needs and according to the guidelines of the NARTC

Follow-up was arranged for reinforcement of teaching and question answering at one and four weeks. Reassessment took place at 16 weeks. At this point details of asthma morbidity during the preceding eight weeks were again collected; for comparison with the intake data. Note, subjects were then randomised to one of three self-management plans as part of a further study not reported here

**Result**

Table 1 summarises the results; the following table deals with symptoms, pulmonary function and healthcare utilisation separately

**Table 1: Result**

	n (129)	Mean change	SD
Days off	21	80.8	10.00
Nights Waking	31	90.6	10.00
Pre $\beta$ -2-agonist PEFR**	9	132.5	10.00
Pre $\beta$ -2-agonist FEV <sub>1</sub> ≡	7	90.1	10.00
Pre $\beta$ -2-agonist FVC ≡	7	+0.16	10.00
Post $\beta$ -2-agonist PEF	71	829.2	10.00
Post $\beta$ -2-agonist FEV <sub>1</sub> ≡	5	10.1	10.00
Post $\beta$ -2-agonist FVC ≡	5	40.1	10.00
Antibiotics	12	30.4	10.00
Steroids	51	60.2	10.00
ICS Inhaler*	9	60.9	10.00
Nebuliser	01	-0.15	10.00
Consults	11	31.6	10.00
Visits	41	60.1	10.00

\* Data is incomplete because either the patient or the nurse felt that recall or records were unreliable for that patient

\*\* The number of excess inhaler use subjects are particularly low because of poor recall and because some subjects were on regular  $\beta$ -2-agonist medication four times per day (a standard treatment in 1990)

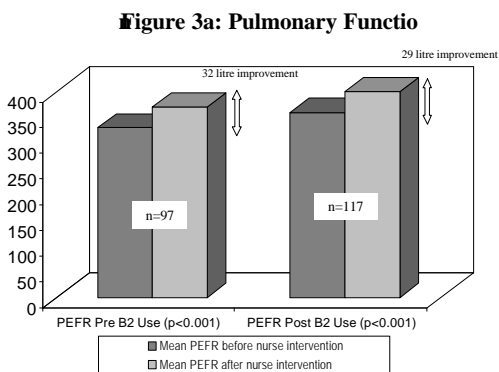
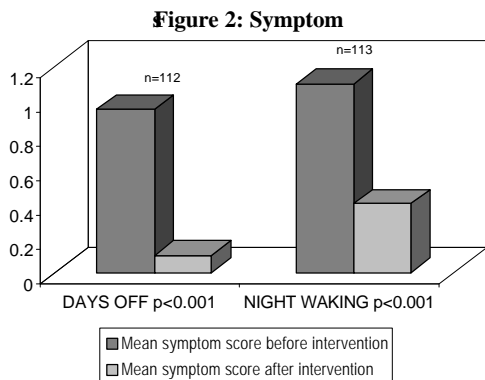
\*\*\* The number of subjects who had PEFR measure before  $\beta$ -2-agonist use is lower (97) than those after (117) because twenty subjects had used the  $\beta$ -2-agonist medication within a few hours prior to their intake interview and their values could not be used

≡ Due to an equipment failure, full spirometry could not be completed at one site for some time during the study and peak flows only are available for some patients

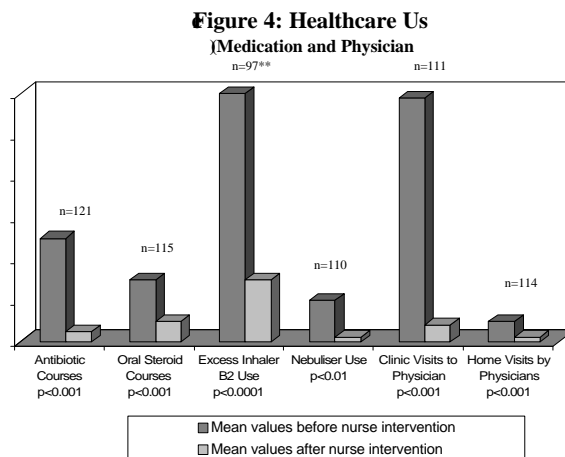
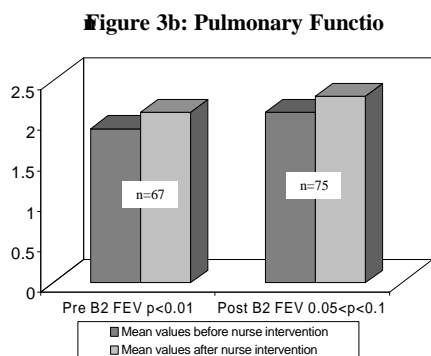
1. Number of Subject  
Table 1 shows that the number of subjects vary for each outcome from the total numbers of 129

2. Symptoms (see Figure 2)  
The mean number of days off work or school fell after the nurse intervention, as did the mean number of nights on which waking occurred. For both, the change is highly significant at  $p < 0.001$

3. Pulmonary Function (Figure 3)  
The improvement in peak expiratory flow rate after intervention was highly significant ( $p < 0.001$ ), whether measured before using a  $\beta$ -2 agonist bronchodilator or after use (Figure 3). The improvement in FE<sub>1</sub> measured prior to use of a  $\beta$ -2-agonist bronchodilator was significant at a lower p value possibly due to the smaller number of subjects compared because of spirometry



\*\*\*Note: 20 patients had used regular  $\beta$ 2-agonist just prior to the initial assessment so their data was not included



failure at one of the clinics, and the exclusion of patients who had recently ( $< 4$  hours) used their  $\beta$ -2 agonist inhaler. The numbers of post  $\beta$ -2-agonist subjects is slightly greater because no patients were excluded but there is less margin for improvement in bronchodilated patient. This is reflected in the smaller p values which are much less significant, if at all, due to the reduction in power ( $> 0.05$  for FE<sub>1</sub>) (Table 1)

4. Healthcare Use (Medication)  
Figure 4 shows the effect of the nurse intervention on drug use. The mean numbers of courses of antibiotic and oral steroids were small before the intervention but the reduction after the intervention was still significant for both drugs at  $p < 0.001$ . Nebuliser use was very small, regular users being excluded from the study, therefore, the reduction in use, even though significant ( $p < 0.01$ ), was less so than for the other medications

From this table, the most obvious improvement was the reduction in days on which extra use of  $\beta$ -2 agonist inhaler (as required use) occurred ( $p < 0.001$ ). This is important as it suggests better disease control as a result of the intervention on top of the improved drug use suggested by the reductions in the other medications

5. Healthcare Use (Physician Services)  
Figure 4 shows a highly significant ( $p < 0.001$ ) reduction in consultations for asthma, with the physician both in the clinic and in the patient's home even though the total number of home visits to the subjects was small. Hospital visits, though recorded were too few for analysis

Discussion  
The effects of the educational intervention on asthma outcomes have not previously been isolated from the effects of an asthma clinic generally. The current study attempts to isolate the educational component without medication changes or the introduction of management plans. Therefore, the period under observation after the educational intervention is short each subject in fact went on to be randomised into one of several management plans after the sixteen-week assessment. We have demonstrated a highly significant effect of nurse education on symptoms

pulmonary function, and healthcare utilisation

The nurses who provided the education also collected data, however every attempt was made to be objective in the assessments. This involved not including data judged at the time of collection to be weak such as patient uncertainty over numbers of rescue Ventoli inhalations. All patients had their data included but not all patients had complete data sets. Difficulties arise with the interpretation of results when the data is not complete for each subject for each variable but we feel justified in the way in which we have handled the data analysis, preferring to include all the reliable data we obtained and all the subjects, rather than excluding all the patients with incomplete data sets. We believe this represents a more valid picture of the expected effect in the general practice setting

The findings are limited in their generalisability partly because of the particular environment in which it was conducted in the UK and partly because of the short length of follow-up. This last was unavoidable. A longer period would have raised ethical issues of withholding a management plan, now often considered the standard of care. There is an intrinsic weakness in using the before and after design rather than a parallel control group. Patients were often recruited while attending about their asthma and this may have been during a bad period for their asthma with a natural tendency to improve anyway. Repeat studies in other practice settings would be desirable for support of findings but, as Jones has observed<sup>2</sup> a randomised trial against a control group is probably now not acceptable in the UK, even of education alone. In view of Usherwood's and Beasley's limited success in their clinics<sup>9</sup>, we would like to see a comparative trial of practice nurse education against outpatient education or general practitioner education to compare outcomes and effectiveness. The difficulties associated with general practitioners giving asthma education are described elsewhere<sup>6,4-1</sup>. It will be noted from Figure 4 that physician visits, in the surgery or at home, were reduced but this was at the expense of three visits per subject to the nurse. In the current climate of the primary healthcare team, this may or may not be considered the preferred mode of delivery for asthma education

### Conclusion

This before and after experimental study has demonstrated that education in a nurse-run asthma clinic in general practice can have a highly significant effect on the symptoms, pulmonary function, and healthcare utilisation of patients of all ages with moderate/severe asthma. ■

### Acknowledgement

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