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STAAR: a randomised controlled trial of electronic adherence monitoring with reminder alarms and feedback to improve clinical outcomes for children with asthma.

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Abstract

Background

Sub-optimal adherence to inhaled steroids is common in children with asthma and is associated with poor disease control, reduced quality of life and even death. Previous studies using feedback of electronically monitored adherence data have demonstrated improved adherence but have not demonstrated a significant impact on clinical outcomes. The aim of this study was to determine whether introduction of this approach into routine practice would result in improved clinical outcomes.

Methods

Children with asthma aged 6-16 years were randomised to the active intervention consisting of electronic adherence monitoring with daily reminder alarms together with feedback in the clinic regarding their inhaled corticosteroid use or to the usual care arm with adherence monitoring alone. All children had poorly controlled asthma at baseline, taking inhaled corticosteroids (ICS) and long acting beta agonists (LABAs). Subjects were seen in routine clinics every three months for one year. The primary outcome was the Asthma Control Questionnaire (ACQ) score. Secondary outcomes included adherence and markers of asthma morbidity.

Results

77 of 90 children completed the study (39 intervention, 38 control). Adherence in the intervention group was 70%, vs 49% in the control group ($p = <0.001$). There was no significant difference in the change in ACQ but children in the intervention group required significantly fewer courses of oral steroids ($p= 0.008$) and fewer hospital admissions ($p= <0.001$).

Conclusion

The results indicate that electronic adherence monitoring with feedback is likely to be of significant benefit in the routine management of poorly controlled asthmatic subjects.

What is the key question?

Can electronic adherence monitoring with feedback and alarms improve clinical outcomes in children with poorly controlled asthma?

What is the bottom line?

Electronic monitoring with feedback and alarms improved adherence, decreased hospital admissions and courses of oral steroids required.

Why read on?

This study provides the first unequivocal evidence that adherence monitoring with feedback can impact on important clinical outcomes when used in the management of children with poorly controlled asthma.

Twitter

Electronic adherence monitoring with feedback and alarms can be effectively used in the clinical setting to improve clinical outcomes for children with asthma.

INTRODUCTION

Adherence to inhaled steroids is often sub-optimal in children with asthma, resulting in poor disease control¹, increased need for oral steroids² and decreased lung function³. This leads to increased healthcare utilisation, and associated cost⁴. The recent National Review of Asthma Deaths in the UK reported that poor adherence was associated with 34% of deaths due to asthma, emphasising the significance of the problem⁵.

Large adult population studies, and smaller paediatric studies have shown that adherence rates of 75-80% are required to significantly improve asthma control⁶⁻⁸. However, when objectively measured, the average rate of adherence in children with asthma is around 50%, some way below this desired therapeutic level⁹.

Subjective and indirect measurements of adherence have been shown to over-estimate rates due to patients wanting to be looked upon favourably by their clinician- the social desirability bias^{10,11}.

Electronic monitoring devices (EMDs) record adherence rates by logging the exact date and time an inhaler is actuated, with modern devices proven to be highly accurate and reliable in the clinical setting¹². Monitoring allows intentional adherence barriers such as negative illness perceptions or medication beliefs to be identified, and addressed with regular open dialogue¹³⁻¹⁵. Reminder alarms built into the devices can address non-intentional practical barriers such as simply forgetting to take the medication¹⁶.

Due to their cost, the British Thoracic Society (BTS) has questioned the viability of EMDs outside the research setting¹⁷, and The National Institute of Health and Excellence (NICE) has recommended further studies to investigate their use clinically¹⁸.

An Australian study demonstrated that children's adherence increased when electronic data was fed back, although it was under-powered to show any improvement in clinical outcomes¹⁹. A recent study in New Zealand improved self-reported asthma control in the short-term using electronic monitoring, no feedback and reminder alarms²⁰. However, this study involved seeing subjects out of clinic every 2 months, used covert monitoring, and had no effect on objective clinical outcomes.

In order to determine whether clinically relevant benefits could be observed in a routine clinical setting through the use of electronic monitoring we undertook a study in which children with poorly controlled asthma were randomised to the use of reminder alarms and feedback or routine care.

We hypothesized that by addressing both the intentional and non-intentional adherence barriers, rates would increase to a degree necessary to improve asthma control and clinical outcomes.

METHODS

The STAAR study (STudy of Asthma Adherence Reminders) was a multi-centre, open label, parallel group randomised controlled trial, with an allocation ratio of 1:1. Written consent was taken from the parents/ carers of all participants, and ethical approval for the study was granted by the South Yorkshire Research Ethics committee, REC reference 13/YH/0289. The protocol was registered with ClinicalTrials.gov, number NCT02451709.

Participants & setting

Children with doctor-diagnosed asthma aged 6-16 years attending clinics in Sheffield or Rotherham were screened for eligibility. Participants had to be taking regular inhaled steroids, with no change in their medication in the last month. At recruitment, participants had to have an Asthma Control Questionnaire (ACQ) score of at least 1.5, indicating they had poorly controlled asthma²¹.

Participants were excluded if they couldn't speak English, or if they had another significant chronic condition. The EMDs available for this trial were only compatible with seretide or symbicort inhalers. Therefore all participants were at BTS level 3 at the start of the trial.

Interventions

Prior to randomisation, all participants had their inhaler technique checked by a qualified asthma nurse, and received a brief asthma education session, emphasizing the importance of taking inhaled steroids regularly. All participants were seen in their standard asthma clinics 3 monthly and all treatment decisions were made by the clinical team. A member of the study team downloaded data from the EMD at each visit.

Intervention Group

Participants in the intervention group had an EMD attached to their regular inhaler. The devices used in this study were "Smartinhalers" and "Smartturbos", manufactured by Adherium, [Auckland, New Zealand] (figure 1). The EMDs are commercially available, have a CE safety mark, and are validated for adherence monitoring in asthma¹². Participants were told the devices monitored the date and time of all actuations. At clinic visits, the adherence data from the previous 3 months was uploaded to the website www.smartinhalerlive.com, which displays the data graphically. These data were reviewed with the patient and parent/ carer (figure 2). Open, non-judgemental discussions were held about the adherence rate, barriers identified and, if necessary, personalised strategies for improvement were devised. Devices were set to play reminder alarms (children's music or character noises), with different times agreed for weekdays and weekends. Alarms sounded for 5 seconds,

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every minute for 15 minutes (or until actuation) if the inhaler hadn't been actuated within the previous 6 hours of the specified time. The devices were locked to prevent tampering. Times were reviewed each study visit and changed if necessary. An example was a child who missed evening doses due to Arabic classes, therefore alarms were changed to later in the evening on weekdays.

Control Group

Control participants had the same EMDs attached to their regular inhaler, they were also told the devices monitored how much the inhalers were taken, but that these data would not be reviewed. Participants were seen in their standard asthma clinic and the data was downloaded, but not reviewed. The alarms were disabled, and the devices locked.

Primary Outcome

The primary outcome for the study was change in the ACQ score at 3, 6, 9 and 12 months. The ACQ is a questionnaire validated for the use in children aged 6-16²². There are 6 clinical questions related to symptoms in the previous week, and the 7th score corresponds to the Forced Expiratory Volume in one second, per cent predicted (FEV1%). A mean of the 7 values is calculated, giving a score of 0-6, with a higher score indicating more poorly controlled asthma.

Secondary Outcomes

Secondary outcomes recorded at baseline and each visit were FEV1%²³, number of unplanned attendances to GP/ED for asthma since last visit (as reported by parents), number of courses of oral steroids required, number of days off school due to asthma, use of β -agonists in the past week and BTS level of asthma therapy. Quality of life was measured using the Mini Paediatric Asthma Quality of Life Questionnaire (mini PAQLQ)²⁴. The adherence rate was calculated for each 3 month period, both morning and afternoon doses, and recorded as a percentage. This was calculated as number of doses actually taken/ number of doses prescribed x 100. The daily adherence was capped at 100%, to avoid falsely increased values due to dose dumping. The overall 3 monthly figure was a mean of each daily %. Parental beliefs about inhaled steroids and perceptions about asthma were recorded at baseline and 12 months with the Beliefs about medicines questionnaire (BMQ) and the Brief Illness Perceptions Questionnaire (IPQ)^{25, 26}.

Sample Size

The sample size was calculated using a repeated measures analysis for 4 follow-up visits. The Minimal Clinically Important Difference (MID) for the ACQ is 0.5²². Using a repeat measure analysis

table we calculated for a significance of 5% ($\alpha = 0.05$), and a power of 80%, with a repeated measure correlation of 0.4, that $n = 76$ ²⁷. To allow for a 15% attrition rate we aimed to recruit 90 participants to the study.

Randomisation

Participants were randomised using permuted block randomisation, with an allocation of 1:1, which resulted in equal group sizes of 45, created from a computer generated random number sequence. The allocation of subjects following recruitment involved phoning the independent holder of the randomisation code.

Blinding

Due to the nature of the intervention, neither the participants nor the study team were blinded in this study. In the intervention group adherence data was made available to clinicians if requested, but not in the control group.

Statistical analysis

The ACQ and FEV1% were compared statistically by two approaches. First, by calculating a paired difference between 12-months and baseline for each group separately. The 'difference of the difference' between treatments was compared by an independent t-test. A 95% confidence interval was calculated. Second, an area under the curve (AUC) was calculated for each group and the between-group difference was determined by an independent t-test. The AUC is a mean value weighted by time²⁸. An arbitrary level of 5% statistical difference was assumed (two-tailed). Incident rates were estimated by Poisson regression, and compared by incident rate ratios (IRRs). The assumption of Poisson regression (mean=variance) was confirmed. Continuously distributed data was summarised by the median (quartiles); categorical data by n (%). A sensitivity analysis was carried out on the primary outcome measure (ACQ) using multiple imputation (MI). The approach used is detailed in appendix 1, with charts demonstrating the patterns for missing data. The "Stata" statistical computer package was used for data analysis.

RESULTS

90 children (81 Sheffield, 9 Rotherham) were recruited between October 2013 and August 2014, figure 3 shows the flow of these participants.

Baseline characteristics

The baseline characteristics are shown in table 1.

	Intervention group (n=47)	Control group (n=42)
Age [years]	10.4 (2.9)	10.2 (2.9)
Sex male	28(60%)	22(52%)
ACQ	2.5(0.8)	2.3(0.7)
FEV1 %	87.2 (14.9)	88.0 (13.4)
PQL	4.3(1.5)	4.6(1.2)
ICS dose	697.9(348.6)	664.3(280.1)
BTS	3.5(0.6)	3.4(0.5)
Proportion BTS ≥ 4	51%	43%
GP/ED visits	1.9(2.2)	2.1(2.0)
Beta agonist	2.5(1.3)	2.3(1.3)
School days missed	3.5(4.4)	3.8(5.7)
Oral steroids	1.2(1.8)	1.2(1.3)
Hospital admissions	0.3(0.6)	0.2(0.6)
BMQ score	2.5 (0.5)	2.6 (0.4)
IPQ score	5.6 (1.3)	5.3 (0.9)
Ethnicity WB	30(64%)	24(57%)
BA	3(6%)	6(14%)
BP	11(23%)	11(26%)
BI	0(0%)	1(2%)
AO	1(2%)	0(%)
BC	2(4%)	0(%)
Time from asthma diagnosis (years)	6.0 (3.7)	6.7 (3.7)

Table 1 – Baseline data. Data are mean (SD) or n (%). Notes. Calculations subject to rounding errors. ACQ=asthma control questionnaire. FEV1% = Forced expiratory volume in 1 second, % predicted. PQL=paediatric quality of life questionnaire. ICS dose = beclometasone equivalent. Beta agonist use – score on ACQ question. WB=white British, BA=black African, BP=British Pakistani, BI=British Indian, AO=Asian other, BC=black Caribbean. GP/ED visits, school days missed, oral steroids required and hospital admissions are all parent reported events over the previous 3 months.

Primary Outcome

The ACQ decreased in both groups between the baseline and the 3-month visit. It fell by 1.0 in the control group and 0.9 in the intervention group ($p=0.35$), both significantly exceeding the MID for changes in ACQ scores of 0.5²². This improvement was maintained to 12 months in both treatments arms (figure 4). At 12 months there were no significant differences between the two groups for either the mean change from baseline (table2) or the areas under the curve (table 3). The individual participant trajectories for ACQ across the study visits are shown in appendix 1 (figure A3).

Outcome	Intervention baseline	Intervention 12 months	Paired mean difference	Control baseline	Control 12 months	Paired mean difference	Difference of the difference	95% CI	p
ACQ	2.65 (0.12)	1.58 (0.19)	-1.14 (0.21)	2.47 (0.12)	1.50 (1.07)	-0.95 (0.77)	-0.18 (0.28)	-0.76,0.38	0.51
FEV1%	87.23 (2.77)	91.37 (1.33)	3.00 (1.67)	88.00 (1.07)	88.97 (2.55)	1.54 (2.18)	1.45 (3.68)	-4.00,6.91	0.59

Table 2. Outcome measures for ACQ and FEV1%. Data are means (SD in parentheses). The difference of the difference is an estimative statistic where the SD has limited value.

Intervention	Control	Difference	95% CI	P
1.7 (0.13)	1.6 (0.14)	0.09 (0.18)	(-0.26,0.45)	0.52

Table 3 – Comparison of areas under the curve for ACQ

Secondary Outcomes

Average adherence over the 12 months for the intervention group was 70%, vs. 49% for the control group ($p<0.001$). Higher mean and median adherence rates were maintained for the 12 month period in the intervention group, but declined over time in the control group (figure 5). 20 Participants in the intervention group, and 6 in the control group had a mean adherence rate of >80% for the 12 month period. 4 participants in the intervention group and 11 in the control group had rates of <30%.

Event rates are shown in table 4. The mean (SD) days in study was 351 (117) days for the intervention group, and 358 (101) days for the control group. Figure 5 shows the timing of rescue oral steroids required during the study.

	Event rate (per 100 child days)		P value	IRR	95% CI
	Intervention	Control			
GP/ED visits (n=193)	0.582	0.650	0.316	1.15	0.83,1.63
Days off school due to asthma (n=462)	1.365	1.606	0.1	1.16	0.97,1.39
Courses of oral steroids (n=156)	0.411	0.676	0.008	1.53	1.11, 2.11
Hospital admissions (n=20)	0.0254	0.129	<0.001	4.38	1.46,12.13

Table 4 Incident rates for clinical outcomes. n = total number of events in study as reported by parents. IRR = Incident Rate Ratio. CI = Confidence Interval. Example interpretation: controls are 53% more likely to be prescribed steroids than those in the intervention.

There were no significant differences between the two groups for short acting β -agonist use, change in BTS stage, mini PAQLQ, BMQ or IPQ scores. FEV1% improved in both treatment arms, with no significant difference between treatments at 12-months compared to baseline (table 2). The mean (SD) beclometasone equivalent ICS dose at the end of the study was 673 (303) mcg in the intervention group, and 767 (369) mcg in the control group.

Clinic visits

Patients did not attend (DNA) or cancelled 143 scheduled appointments (73 intervention, 66 control). Appointments were re-scheduled, but where this was not possible, 35 non-clinical study visits (15 intervention, 20 control) were performed. As a result of these missed study visits, only 41% of participants in the intervention group received feedback at all three time points (3,6 & 9 months). Missing data for ACQ and adherence is shown in tabulated form in Appendix 1.

Issues with electronic adherence monitors

Table 5 shows the frequency of devices lost, forgotten or broken. Missed data from forgotten devices was downloaded at a later clinic appointment. Table 5 shows the number of broken, forgotten, and lost devices. Reasons given for being damaged (just reported, or actually broken) included lost/flat battery, dropped on floor, dropped in liquid, alarms not starting, alarms not stopping, screens peeled off.

	Intervention (47 participants)	Control (42 participants)
Device reported as "broken" by child	23(50%)	8 (19%)
Devices damaged beyond repair, (when inspected by study team, requiring replacement device)	17(37%)	2 (5%)
Participant forgot to bring device to clinic	10 (22%)	18 (43%)
Device lost completely	5(11%)	2 (5%)

Table 5: Broken, forgotten and lost devices

DISCUSSION

While use of electronic adherence monitoring with feedback and regular alarms by children with poorly controlled asthma did not significantly improve self-reported asthma control, there was a significant increase in adherence rates that was maintained over the 12 months. This improvement was associated with a significant decrease in the number of exacerbations requiring a course of oral steroids or an admission to hospital. This is the first study to our knowledge that has shown that regular alarms and feedback of electronic adherence data has a significant effect on a number of clinically relevant outcomes. Importantly, the intervention was built into routine clinical care and the benefits were sustained over the 12 month study period.

Previous studies using this approach have also demonstrated improved adherence through the use of reminders with or without feedback, but they have failed to demonstrate a significant difference in clinical outcomes. These studies have variously involved relatively well controlled and adherent subjects²⁹, relatively mild asthmatic subjects managed in the community³⁰ or have been too small to demonstrate significant difference¹⁹. In this study the subjects all had 'poorly controlled asthma' (ACQ \geq 1.5), and hence there was possibly greater potential to have an impact than those included in previous studies. Our results are consistent with the study of Williams et al., who in a study of 298 adults with asthma found that patients with an adherence in excess of 75%, as assessed by pharmacy records, were significantly less likely to require an ED visit, a course of oral steroids or a hospital admission⁶.

The results from this study suggest that this approach is likely to be beneficial if introduced into routine clinical care, at least in those children with poor asthma control. It is likely that the intervention resulted in an overall reduction in health costs, given that the majority of the direct costs associated with asthma are attributable to hospital admissions, or exacerbations sufficient to require intervention³¹. In this study the hospitalisation rate was five times higher in the control group despite self-reported symptoms being similar. This approximates to the prevention of 12 hospitalisations in one year amongst the 47 intervention subjects giving an approximate figure for the number to treat of 3.25. There would appear to be a cost saving in using this approach, even at current prices (£120 per device). The BTS questioned the utility of this approach due to high device cost, but its introduction into routine practice would reduce unit price, as it would drive competition and innovation.

Adherence in the control group fell progressively over the year, consistent with results from previous studies^{29, 30, 32}. At 49% the median adherence was similar to the value of 53% derived from our

analysis of 18 previous studies using electronic monitors to quantify adherence rates in children⁹. In contrast, the intervention group maintained a significantly higher level of adherence throughout the study and this was associated with the important improvements in clinical outcomes. Whilst both groups required fewer courses of oral steroids during the study compared to baseline rates, significantly more courses were required in the control group throughout. This difference between groups was more marked at nine and twelve months, with rates increasing in the control group in the second six months (figure 6). Similar patterns of increasing control group rates in the second six months were seen for hospital admissions, days off school and GP/ED visits. This would suggest the benefit of this intervention was over a prolonged period of time, maintaining increased adherence rates with regular feedback and discussion.

Which part of the intervention had the greatest impact is unclear. A previous study found reminders alone can have an impact on adherence but this was in a self-selecting group of well controlled adult subjects whose mean adherence was high (74%) in the control group³⁰. It is likely that the feedback in this study played a major role in sustaining improved adherence rates, since other studies using reminders alone have observed declining adherence over time^{20,29}. Burgess et al also reported sustained improvements in adherence over time through feedback though the study only ran for 4 months and subjects were seen every 4 weeks¹⁹. Accurate electronic data that could be discussed with the family appeared to facilitate an open and honest discussion about adherence and the barriers encountered, leading to evolving practical solutions. Social-desirability could be used in a positive way to influence medication taking behaviour. However, for some participants the interventions did not appear to impact on adherence and there were a disproportionate number in this group who reported damaged or lost devices. This approach may not fully address the issue of intentional non-adherence, but it does help to confirm the problem, should the devices remain operational.

In support of the argument that feedback and discussion are important are the results from a recent study which demonstrated that in 220 children, despite improving self-reported asthma control, electronic monitoring and alarms had no effect on objective clinical outcomes²⁰. Subjects were seen out of clinic every 2 months for 6 months, with adherence declining at each time point in both groups, possibly leading to the initial impact on exacerbations disappearing beyond 2 months. Importantly the participants received no direct feedback, preventing the opportunity to initiate an open dialogue. Moreover, covert adherence monitoring was used, an approach that cannot be recommended in routine practice.

Out of necessity this was an open label study. The risk of introducing an inadvertent bias was minimised by ensuring both groups had the same number of clinic visits (136 intervention, 124 control) and ensuring clinical management was undertaken by the patients' usual clinicians. The control participants were aware that their adherence was being monitored in a clinical trial, and the increased rates seen at 3 months may be due to this³³. It is possible that true adherence was lower in the intervention group than that recorded as the devices (and all EMDs currently available) simply recorded actuations with no guarantee the medication was inhaled. Participants or family members could have potentially pressed the device randomly, or to silence alarms. Future devices are likely to overcome this problem by detecting inhalation using thermistors or other air flow monitors, an approach used successfully in electronic respiratory rate monitors³⁴.

The lack of difference in ACQ in the two groups despite what appear to be significant differences in adherence rates is consistent with results reported in other studies^{29,30}. Similarly recent community studies have reported a disconnect between self-reported symptoms and other clinical outcomes such as exacerbations^{30,35}. This is possibly because the ACQ score is a subjective measure. Open, honest dialogue facilitated by adherence discussions may have made children more honest about the severity of their symptoms in the intervention group. In contrast, answers from control subjects may have been more guarded, with denial of symptoms, as is the case in standard clinical practice, particularly in adolescents³⁶. Additionally patients who knew their adherence has been poor may have minimised symptom reporting¹³. The ACQ scores and FEV1% may also reflect short term influences such as the recent use of oral steroids, which was significantly more common in the control group. The lack of a significant difference in FEV1% between groups and poor correlation with exacerbations may be due to FEV1% being an inaccurate way of grading asthma severity in children³⁸.

This was designed to be a pragmatic study in which asthmatic children continued with usual care with or without the intervention. There was a high rate of participants cancelling appointments, or simply not attending. This appears to reflect the attitude of many patients and their families who don't necessarily always consider asthma a significant condition. The overall DNA rate in this study was 20% (71/360 appointments, 38 group A, 33 group B), and this compares to a rate of 12% in respiratory clinics in Sheffield. This may demonstrate a reluctance to attend if the participant felt they may be judged or blamed for poor adherence by the study team or parent. Alternatively, due to their improved clinical condition (shown by improved ACQ), participants may have missed appointments because they felt better, a common reason for non-attendance³⁹. Related to this issue were the high levels of devices reported to be 'broken'. Patel and colleagues reported a

Smart inhaler malfunction rate of 1.9%, and 3.5% lost when assessing 2642 monitors in an adult study¹². Our rates of malfunction and loss were much higher (35% and 8% respectively), suggesting some of the devices were deliberately broken or lost. The higher rates in the intervention group (50% broken, 11% lost) suggest that children are more likely to break or lose their device when their adherence is being openly monitored. The next generation of devices provide automated uploading of data via smartphones such that adherence can be monitored in real-time, which offers significant advantages in terms of logistics and opportunity to influence behaviour. These high rates of damage may suggest that this approach was disliked by the participants. However, the devices were popular and positive feedback was received, with the majority of participants asking to keep their device after the study period. A recent qualitative study to investigate young peoples' opinions on electronic monitoring and feedback using these Smart inhaler devices also reported positive opinions, concluding that this is a popular intervention for both adolescents and their parents⁴⁰.

In summary, these data indicate that significant clinical benefits can be derived from using electronic adherence monitoring with feedback and alarms. A sustained improvement in adherence rates was associated with a decreased number of courses of oral steroids required and hospital admissions when used in a population of poorly controlled asthmatic children using combination therapy.

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Conflicts of Interest

None

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Contributorship Statement

RWM wrote the protocol for the study, applied for and secured ethical approval for the study, and applied for and secured funding for the study. He recruited participants, carried out study visits, recorded the results and wrote and edited the manuscript.

HEE helped write the protocol, helped apply for funding, supervised the implementation of the study, and helped write and edit the manuscript.

ASR helped write the protocol, performed all the statistical analysis for the study and helped write and edit the manuscript.

WJD carried out study visits and helped edit the manuscript.

DAK helped apply for ethical approval, helped apply for funding, recruited participants, carried out study visits and helped edit the manuscript.

LJS recorded and analysed the lung function for all the participants in the study and helped edit the manuscript.

MLE devised the concept of the study, wrote the protocol, and helped write and edit the manuscript.

Figure legends

Figure 1 – “Smartturbo” and “Smarttrack” electronic adherence monitors. Adherium, New Zealand

Figure 2 – Adherence review graph from <http://www.smartinhalerlive.com>. The number of daily doses taken is the y axis, with the date on the x axis. In this example the participant was prescribed 4 doses on the first date, but only took 2.

Figure 3 – CONSORT 2010 flow diagram showing progress of participants through trial

Figure 4 – Box and whisker plot showing median ACQ scores over time for groups A (intervention) and B (control)

Figure 5 – Box and whisker plot showing median adherence rates over time for groups A (intervention) and B (control)

Figure 6 – Bar chart demonstrating the number of courses of oral steroids required for different time periods during the study for groups A (intervention) and B (control). Group A total 65 courses, Group B total 91 courses

References

1. Jentzsch NS, Camargos P, Sarinho ES, et al. Adherence rate to beclomethasone dipropionate and the level of asthma control. *Respir Med* 2012;106:338-43.
2. Bender B, Zhang L. Negative affect, medication adherence, and asthma control in children. *J Allergy Clin Immunol* 2008;122:490-5.
3. Duncan CL, Hogan MB, Tien KJ, et al. Efficacy of a parent-youth teamwork intervention to promote adherence in pediatric asthma. *J Pediatr Psychol* 2013;38:617-28.
4. McGrady ME, Hommel KA. Medication Adherence and Health Care Utilization in Pediatric Chronic Illness: A Systematic Review. *Pediatrics* 2013;132:730-740.
5. Royal_College_of_Physicians. Why asthma still kills: the National Review of Asthma Deaths (NRAD) Confidential Enquiry Report: London: RCP, 2014.
6. Williams LK, Peterson EL, Wells K, et al. Quantifying the proportion of severe asthma exacerbations attributable to inhaled corticosteroid nonadherence. *J Allergy Clin Immunol* 2011;128:1185-1191 e2.
7. Klok T, Kaptein AA, Duiverman EJ, et al. It's the adherence, stupid (that determines asthma control in preschool children)! *Eur Respir J* 2014;43:783-91.
8. Lasmar L, Camargos P, Champs NS, et al. Adherence rate to inhaled corticosteroids and their impact on asthma control. *Allergy* 2009;64:784-9.
9. Morton RW, Everard ML, Elphick HE. Adherence in childhood asthma: the elephant in the room. *Arch Dis Child* 2014;99:949-953.
10. Bender B, Wamboldt FS, O'Connor SL, et al. Measurement of children's asthma medication adherence by self report, mother report, canister weight, and Doser CT. *Ann Allergy Asthma Immunol* 2000;85:416-21.
11. Jentzsch NS, Camargos PA, Colosimo EA, et al. Monitoring adherence to beclomethasone in asthmatic children and adolescents through four different methods. *Allergy* 2009;64:1458-62.
12. Patel M, Pilcher J, Travers J, et al. Use of metered-dose inhaler electronic monitoring in a real-world asthma randomized controlled trial. *J Allergy Clin Immunol Pract* 2013;1:83-91.
13. Horne R, Weinman J. Self-regulation and Self-management in Asthma: Exploring The Role of Illness Perceptions and Treatment Beliefs in Explaining Non-adherence to Preventer Medication. *Psychology & Health* 2002;17:17-32.
14. Klok T, Kaptein AA, Duiverman EJ, et al. High inhaled corticosteroids adherence in childhood asthma: the role of medication beliefs. *Eur Respir J* 2012;40:1149-55.
15. Everard ML. The Emperor's New Clothes II--time for regulators to wake up and take responsibility for unnecessary asthma morbidity: time for the second aerosol 'transition'. *Thorax* 2013;68:891-3.
16. Burgess SW, Sly PD, Morawska A, et al. Assessing adherence and factors associated with adherence in young children with asthma. *Respirology* 2008;13:559-63.
17. British Thoracic Society. BTS/SIGN Guideline on the management of asthma. Volume 2014, 2012.
18. National_Institute_for_Health_and_Care_Excellence. Asthma: diagnosis and monitoring of asthma in adults, children and young people. NICE guideline, draft for consultation. . Volume 2015, 2015.
19. Burgess SW, Sly PD, Devadason SG. Providing feedback on adherence increases use of preventive medication by asthmatic children. *J Asthma* 2010;47:198-201.
20. Chan AH, Stewart AW, Harrison J, et al. The effect of an electronic monitoring device with audiovisual reminder function on adherence to inhaled corticosteroids and school attendance in children with asthma: a randomised controlled trial. *Lancet Respir Med* 2015.
21. Juniper EF, Bousquet J, Abetz L, et al. Identifying 'well-controlled' and 'not well-controlled' asthma using the Asthma Control Questionnaire. *Respir Med* 2006;100:616-21.

22. Juniper EF, Gruffydd-Jones K, Ward S, et al. Asthma Control Questionnaire in children: validation, measurement properties, interpretation. *Eur Respir J* 2010;36:1410-6.
23. Quanjer PH, Stanojevic S, Cole TJ, et al. Multi-ethnic reference values for spirometry for the 3-95-yr age range: the global lung function 2012 equations. *Eur Respir J* 2012;40:1324-43.
24. Wing A, Upton J, Svensson K, et al. The standardized and mini versions of the PAQLQ are valid, reliable, and responsive measurement tools. *J Clin Epidemiol* 2012;65:643-50.
25. Broadbent E, Petrie KJ, Main J, et al. The brief illness perception questionnaire. *J Psychosom Res* 2006;60:631-7.
26. Horne R, Weinman J, Hankins M. The beliefs about medicines questionnaire: The development and evaluation of a new method for assessing the cognitive representation of medication. *Psychology & Health* 1999;14:1-24.
27. Frison L, Pocock SJ. Repeated measures in clinical trials: analysis using mean summary statistics and its implications for design. *Stat Med* 1992;11:1685-704.
28. Matthews JN, Altman DG, Campbell MJ, et al. Analysis of serial measurements in medical research. *BMJ* 1990;300:230-5.
29. Charles T, Quinn D, Weatherall M, et al. An audiovisual reminder function improves adherence with inhaled corticosteroid therapy in asthma. *J Allergy Clin Immunol* 2007;119:811-6.
30. Foster JM, Usherwood T, Smith L, et al. Inhaler reminders improve adherence with controller treatment in primary care patients with asthma. *J Allergy Clin Immunol* 2014.
31. Bahadori K, Doyle-Waters MM, Marra C, et al. Economic burden of asthma: a systematic review. *BMC Pulm Med* 2009;9:24.
32. Nikander K, Turpeinen M, Pelkonen AS, et al. True adherence with the Turbuhaler in young children with asthma. *Arch Dis Child* 2011;96:168-73.
33. McCarney R, Warner J, Iliffe S, et al. The Hawthorne Effect: a randomised, controlled trial. *BMC Med Res Methodol* 2007;7:30.
34. Al-Khalidi FQ, Saatchi R, Burke D, et al. Respiration rate monitoring methods: a review. *Pediatr Pulmonol* 2011;46:523-9.
35. Turpeinen M, Pelkonen AS, Selroos O, et al. Continuous versus intermittent inhaled corticosteroid (budesonide) for mild persistent asthma in children--not too much, not too little. *Thorax* 2012;67:100-2.
36. Osman LM. Psychological factors in asthma control and attack risk. *Thorax* 2002;57:190-1.
37. McNicholl DM, Stevenson M, McGarvey LP, et al. The utility of fractional exhaled nitric oxide suppression in the identification of nonadherence in difficult asthma. *Am J Respir Crit Care Med* 2012;186:1102-8.
38. Paull K, Covar R, Jain N, et al. Do NHLBI lung function criteria apply to children? A cross-sectional evaluation of childhood asthma at National Jewish Medical and Research Center, 1999-2002. *Pediatr Pulmonol* 2005;39:311-7.
39. Murdock A, Rodgers C, Lindsay H, et al. Why do patients not keep their appointments? Prospective study in a gastroenterology outpatient clinic. *J R Soc Med* 2002;95:284-6.
40. Howard S LA, Yule C. Exploring the attitudes of adolescents with asthma towards monitoring and sharing of data on their inhaler use. *European Respiratory Journal* 2015;46 (supplement 59).