The effect of asthma and its treatment on growth

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Asthma has little, if any, significant effect on attained adult height. Untreated asthma results in a delay of puberty by approximately 1.3 years, and pubertal delay is likely to explain the majority of apparent growth failure in asthmatics. All currently available inhaled corticosteroids (ICS) result in growth suppression at conventional doses (400 μg/day of beclomethasone dipropionate equivalent), but the growth suppressive effects are relatively short lived, after which growth reverts to pretreatment levels. Younger, prepubertal children, appear more sensitive to the growth suppressive effects of ICS. Asthmatic children receiving conventional doses of ICS (400 μg/day of BDP equivalent) will attain an adult height indistinguishable from their predicted adult height (based on their mid parental height), and no different from non-asthmatics. Adult height could possibly be decreased in severe asthmatics, but this is unlikely to be greater than a 1.2 cm decrement. Recent longitudinal studies offer reassurance that at conventional doses ICS do not have significant long term effects on growth, and that their benefits consistently outweigh their side effects.

THE EFFECT OF ASTHMA ON PUBERTY, GROWTH, AND ATTAINED ADULT HEIGHT

The vast majority of asthmatic children will attain a normal adult height, and most perceived growth failure is due to pubertal delay. In a mixed cross sectional longitudinal cohort study of 331 boys aged between 2 and 20 years of age attending a Belgian residential asthma institute, Hauspie and colleagues noted that, compared to both British and Dutch reference values, boys with asthma were significantly older in attaining each Tanner pubic hair stage. A as a result peak height velocity in the asthmatic boys was delayed by approximately 1.3 years, giving the impression of reduced stature in the asthmatic boys during adolescence, but that growth continued to an older age.

This effect on puberty was reflected in the Melbourne prospective longitudinal asthma study. Children labelled as asthmatic were graded by severity and compared with normal controls. Those with the most severe asthma were significantly shorter at age 14 years, but by 21 years of age their height was no different to mild or moderate asthmatics, or normal controls. Similarly Balfour-Lynn described the growth of a group of asthmatic children as they proceeded through puberty, the majority of who were treated prior to the widespread introduction of ICS. Growth followed a height centile until approximately 10 years of age, but subsequently half the children showed pubertal delay with decreased growth, and height only returned to the baseline centile once puberty had commenced. The mean attained adult height did
not differ significantly from predicted height derived from mid-parental height.

The most comprehensive data to date, provided by countries with military conscription, suggests that adult height in asthmatics is close to normal. A study of 92 310 17-year old Israeli military conscripts reported no significant difference in height between severe asthmatics and non-asthmatic controls, and indeed mild asthmatic males were slightly taller than non-asthmatic controls. A more recent report on the height of 173 034 18-year old Swedish military conscripts (including 8531 asthmatics) showed a small but significant difference in height between asthmatics and normal controls, with a negative correlation between height and asthma severity. The authors were able to analyse four different cohorts between 1983 and 1996, and reassuringly, despite a 40-fold increase in the use of inhaled corticosteroids in Sweden over that time period, the difference in height between asthmatics and controls decreased with each successive age cohort, with the greatest convergence in values observed in severe asthmatics. In the 1996 cohort mean height was 0.7 cm less in the asthmatic conscripts, with the maximum difference being 1.2 cm between the severe asthmatics and non-asthmatics. As many of the subjects in both these studies had not reached their ultimate adult height, it is likely that the observed differences between asthmatics and normal controls are an overestimate.

**THE EFFECT OF TREATMENT OF ASTHMA ON GROWTH**

Regular systemic corticosteroids such as oral prednisolone result in a dose dependent impairment of growth. Hence the advent of inhaled corticosteroids in the early 1970s revolutionised asthma care in children. Early reports concentrated on their therapeutic efficacy, while their side effects appeared negligible compared to oral corticosteroids. The first suggestion that inhaled corticosteroids might have a adverse effect on growth was a report in 1988 of a significant negative inflection in the height standard deviation score (SDS) at the onset of commencing regular beclamethasone dipropionate (BDP). Over the following 25 years all of the currently available ICS have been shown to cause a dose dependent growth suppression. When reviewing the effect of ICS on growth, it is useful to divide the reports into short term studies (less than six months), medium term studies (6–24 months), and longer term studies (more than 24 months and up to adulthood).

**Short term studies**

Short term studies have utilised knemometry, an extremely accurate means of measuring lower leg length growth, which allows crossover studies to be performed rapidly. For example, Wolthers and Pedersen showed that both 2.5 and 5 mg of oral prednisolone daily completely interrupted knemometrically measured lower leg growth in mildly asthmatic children. Inhaled corticosteroids do not have such a pronounced effect; the same authors showed no significant effect on lower leg growth for either 200 or 400 μg/day of budesonide (BUD) for two weeks, but a significant 66% decrease in growth while receiving budesonide 800 μg/day. A number of investigators have reported that BUD has no significant effect on short term growth at a dose of 200 μg/day, causes significant growth impairment at 800 μg/day, but results at a dose of 400 μg/day are inconsistent. In contrast BDP has no effect at a dose of 200 μg/day, but consistently results in decreased short term growth at a dose of either 400 or 800 μg/day. Fluticasone propionate (FP) has no effect on growth at a dose of 200 μg/day.

There are however major difficulties over the interpretation of knemometric studies due to the marked intra-subject and inter-subject variability in short term lower leg growth. Even within the same laboratory there are substantial variations between studies in reported lower leg growth while receiving placebo. Furthermore as the growth suppressive effects of ICS are probably short lived (see below), there will be marked differences of effect between subjects who are receiving ICS at onset of the study and those that are not. Thus short term lower leg growth is a very poor correlate of statureal growth, and the short term effects of any given treatment cannot be extrapolated to the longer term.

**Medium term studies**

Medium term studies address growth between 6 and 24 months of follow up. Early reports on the therapeutic effects of ICS were not designed to detect small effects on growth, but in the past 10 years there have been six well designed randomised double blind studies where medium term growth has been addressed: four for BDP (one versus theophylline, one versus placebo, one versus salmeterol, and one versus placebo and theophylline); and two studies comparing FP with either placebo or BDP. All four controlled studies for BDP reported significant growth suppression at a dose of 400 μg/day, and a recent meta-analysis of these four studies calculated that BDP at this dose results in a decrease in linear growth of 1.51 cm/year (95% CI 1.15 to 1.87). Growth in children receiving 200 μg/day of BDP appears unaffected. Younger, prepubertal children, appear more sensitive to the growth suppressive effects of ICS.

A randomised, double blind multicentre study compared FP 100 μg/day or 200 μg/day with placebo via diskhaler in a group of 325 prepubescent children over one year. Mean height increased over 12 months by 6.15 cm in the placebo treated group, 9.94 cm in those receiving 100 μg/day, and by 5.73 cm in those receiving 200 μg/day (p = 0.3 overall). However, a reanalysis of this data established a significant decrease in growth of 0.43 cm/year (95% CI 0.10 to 0.85) in those receiving 200 μg/day of FP compared to placebo, while growth in those receiving 100 μg/day was not significantly different to placebo. In the only comparison of BDP and FP, Rao and colleagues compared FP 200 μg/day with BDP 400 μg/day via metered dose inhaler in a randomised, double blind single centre study in 23 prepubescent children over 20 months. Growth was significantly decreased in those receiving BDP compared to FP (4.94 cm/year compared to 5.75 cm/year; 95% CI 0.45 to 1.16).

A major difficulty with short and medium term studies is that the magnitude of reported growth suppression flies in the face of clinical observations. For example, if the pooled effect of BDP at the conventional dose of 400 μg/day were consistent over a 10 year period, growth decrement would be a not inconsiderable 15 cm. Extrapolation of the of the short term effect of BUD 800 μg/day gives even more inconceivable results. Medium term studies give conflicting results on the duration of the effect, with some authors reporting that the effect is short lived, while others report that the effect persists for the duration of the study.

**Longer term studies**

Longer term studies assess growth for more than 24 months and up to adulthood. There is a single randomised controlled study, and four longitudinal observational studies. The landmark Childhood Asthma Management Program Research Group (CAMP) is unique in its 4–6 year period of follow up, showing that the effects of ICS on growth are relatively short lived. It was designed to measure the effects of regular inhaled anti-inflammatory agents on lung growth, with the primary end point being the change in forced
expiratory volume in 1 second (FEV₁). In a double blind design, 1041 mild to moderate asthmatic children aged 5–12 years were randomised to regular BUD 200 µg twice daily, nedocromil 8 mg twice daily or placebo and followed for 4–6 years. Although both BUD and nedocromil improved control of asthma, with BUD offering greater control than nedocromil, neither BUD nor nedocromil had any significant effects on FEV₁ compared to placebo. Over a mean follow up of 4.3 years, the children randomised to receive BUD grew 1.1 cm less than those who received placebo (22.7 cm versus 23.8 cm, p = 0.005). The difference in growth between those receiving BUD and placebo was confined almost entirely to the first year, and growth in the two groups appeared very similar thereafter. There was no significant difference in growth between those who received nedocromil and placebo. Over the course of the study there were no significant differences in change in bone density or bone age between the three groups.

There are no randomised controlled studies of asthma medication into adulthood, and three small retrospective studies have given slightly contradictory results. Both Balfour-Lynn and Silverstein and colleagues reported no differences in attained adult height between asthmatics that received ICS in childhood and non-asthmatics. In contrast Van Bever and colleagues reported that although there were no significant differences in attained adult height between adult asthmatics who had or had not received ICS in childhood, those who received ICS had significantly lower adult height compared to their predicted adult height based on mid parental height.

The most comprehensive data were reported by Agertoft and Pedersen, who fastidiously recorded the growth of a cohort of over 300 asthmatic children over a 14 year period. In 1986, based on national guidelines, they had elected to change all of their patients with asthma onto regular ICS; virtually all received BUD. A small number of parents were unwilling for their children to commence ICS, and so this group of children and the subject’s healthy siblings acted as controls. The study was not controlled, and the number of asthmatic controls decreased over the years, mostly through commencing ICS. The predicted adult height was calculated on the mid parental height, with adjustment for the subject’s sex.

Growth decreased significantly after commencing BUD, from a mean of 6.1 cm/year (95% CI 5.7 to 6.5) prior to commencing, to 5.1 cm/year in the first year (95% CI 4.7 to 5.5, p<0.001), and 5.2 cm/year in the second year (95% CI 5.1 to 5.9, p = 0.02). Subsequent growth was no different to controls. Over the mean of 9.2 years it took to reach adult height, the subjects in the BUD group received a mean daily dose of 412 µg/day of BUD (range 100–877 µg/day). Compared to their predicted adult height, there was no significant difference in attained adult height between those who received BUD (+0.3 cm; 95% CI −0.6 to 1.2), asthmatic controls (−0.2 cm; 95% CI −2.4 to 2.1), or the sibling controls (+0.1 cm; 95% CI −0.4 to 0.6). There was no significant correlation between the initial reduction in growth after commencing therapy and the difference between predicted and attained adult height, suggesting that the initial growth suppression was transient and had little long consequence.

**REFERENCES**

Pacemaker twiddling

A 10 year old boy with severe learning disabilities had a permanent subpectoral endocardial pacemaker for intermittent heart block. Twenty three months later, he presented with an episode of collapse associated with pacemaker failure. Chest x-ray examination showed that he had twiddled the pacemaker clockwise and pulled the lead out of the right ventricle (radiograph A). A new pacing system was inserted (radiograph B). At revision, surprisingly, there was no fibrosis around the generator or wire. In an attempt to fix the generator more firmly, it was placed in the original subpectoral pocket, but in a dacron pouch (Parsonnet pouch) to promote fibrosis. He presented again, 17 days later, with a further episode of collapse associated with pacemaker failure. Repeat chest x-ray examination showed that he had twiddled the pacemaker anticlockwise and the pacing lead had perforated the right ventricle into the abdomen (radiograph C). This illustrates the difficulties and the potential risk in children with learning disabilities.

Twiddler’s syndrome leading to pacemaker failure, although rare, is recognised in children. Children with learning disabilities are more susceptible because of conscious or subconscious twisting of the pacemaker box and poor comprehension of consequences. Sub-pectoral implantation of the pacing box may prevent this complication, although this was unsuccessful in this child. A chest radiograph should be performed in cases of pacemaker dysfunction to exclude this complication.

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