Insulin-like growth factor and growth hormone secretion in juvenile chronic arthritis

Roger C Allen, Mark Jimenez, Christopher T Cowell

Abstract
Insulin-like growth factor (IGF-1) concentrations were determined in a series of 23 children with juvenile chronic arthritis in conjunction with anthropometric assessment. When standardised z scores were used significant decreases in height and weight were shown in comparison with the normal age/sex matched means. Severe growth disturbance was seen, particularly in those with prolonged disease duration, which was independent of corticosteroid treatment, indicating disease activity itself is a major factor in the growth retardation. Eight children had low IGF-1 z scores—that is, less than −2.00 from age/sex matched mean. Low IGF-1 z scores were associated with low weight z scores but not with low height z scores. Overnight growth hormone secretory profiles were determined in 10 patients, including seven with low IGF-1, and showed generally normal secretion in all but one patient, who subsequently attained normal concentrations coincident with catch up growth. Increased pulse frequency of overnight secretion was commonly seen. Low IGF-1 concentrations probably result from varying factors, particularly nutritional, but do not reflect marked endocrinological abnormalities in most patients.

Poor somatic growth is a feature of juvenile chronic arthritis, particularly in the systemic and polyarticular onset forms of the disease. In Still's description 'arrest in development' was specifically noted, preceding by many years various therapeutic interventions, particularly use of corticosteroids, which have been implicated in contributing to the abnormal growth in juvenile chronic arthritis. Bernstein et al also showed that disease activity itself contributes to growth impairment.

Normal growth is a complex interrelation of factors, including genetic, hormonal, and nutritional requirements. In the presence of normal thyroid function the secretion of pituitary growth hormone and the growth hormone dependent insulin-like growth factors (IGF), particularly IGF-1, form the predominant hormonal axis of postnatal growth. Normal growth hormone secretion has been shown in juvenile chronic arthritis by pharmacological stimulation or by secretory profiles of physiological secretion, yet, increased rate of growth has resulted from exogenous growth hormone treatment. Previous studies of IGF-1 concentrations in juvenile chronic arthritis have reported conflicting results—either reduced or normal values, with or without correlation with corticosteroid treatment. Concomitant growth hormone secretion was not determined in any of these studies. Low IGF-1 secretion may occur in 'normal' short children, related to low growth hormone secretion rates, despite having normal responses to pharmacological stimulation with arginine/insulin or clonidine.

In this study we monitored growth in relation to IGF-1 and physiological growth hormone secretion by 12 hour overnight sampling, in a group of patients with juvenile chronic arthritis, to investigate if an abnormality of the growth hormone/insulin-like growth factor axis, possibly related to altered sleep patterns, may be contributing to poor growth.

Patients and methods
Twenty three children (15 female, eight male) with juvenile chronic arthritis satisfying the EULAR classification criteria were studied. Disease onset subtypes were seven systemic, six polyarticular, and 10 pauciarticular. All children had continuing evidence of active disease as shown by the presence of synovitis in at least one joint requiring anti-inflammatory drug treatment. With the exception of one patient, referred specifically for the study from another hospital, all patients were attending the rheumatology clinic at the Children's Hospital, Sydney. Informed parental consent was given before entry to the study. All patients were over 4 years of age, the lowest age for which IGF-1 data were available. Mean disease duration was 3.2 years (range 0.25–12). At the time of entry into the study all patients were prepubertal except two (patients 1 and 5) who had attained Tanner stage 2 pubertal development, defined by the degree of pubic/axillary hair or breast development. All patients were taking non-steroidal anti-inflammatory drugs. In addition nine were taking slow acting antirheumatic drugs and 10 were receiving oral corticosteroid treatment on alternate days.

Anthropometric measures included height, weight, mid-arm circumference, and skinfold thickness at four sites—subscapular, sublaural, triceps, and biceps. Height, weight, and rate of growth were compared with age/sex matched United States National Center for Health Statistics growth standards. Weight for height for age ratios were graded by reported standards for nutritional sufficiency, the lower limit of normal being 90%. Indices for nutritional insufficiency were 85–90% (mild), 75–85% (moderate), <75% (severe). Standardised deviation scores (z score) for height and weight were...
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calculated by reported methods\textsuperscript{14}—namely, the difference between the patient’s height (or weight) minus the age equivalent mean height (or weight), divided by the ideal height (or weight) standard deviation for age.

Insulin-like growth factor 1 was measured by a radioimmunoassay,\textsuperscript{15} from which a z score was calculated by the same method as height/weight z scores using normal age and sex data. Seven of eight patients with an IGF-1 z score less than $-2.00$ underwent growth hormone studies. Three additional patients with poor rates of growth were also studied. Overnight sampling every 20 minutes was performed from 2200 to 0800 hours and analysed for the mean growth hormone secretion and the pulsatile pattern of secretion using the pulsar peak identification algorithm program (NICHDB, Bethesda/University of California, Berkeley).\textsuperscript{16} Clonidine stimulation (125 $\mu$g/kg) of growth hormone secretion was performed in the initial six patients studied, but owing to a hypotensive episode in one patient this test was discontinued for the subsequent patients. A normal clonidine stimulation test was defined as a peak in growth hormone of $\geq 20$ mU/l. Growth hormone was measured by radioimmunoassay with second antibody separation using previously reported methods.\textsuperscript{17} Owing to wrist arthritis in many of the patients estimation of bone age was of limited value either because of destructive changes, precluding accurate assessment, or physical advancement of the affected carpus, presumed secondary to local hyperaemia.

The SPIDA statistical computer package (Statistical Laboratory, Macquarie University, Sydney) was used for the analysis, and the rejection level was set at 0.05. Comparison between groups was determined by the two sample $t$ test. Parameters of the secretory patterns were compared by linear regression analysis.

### Results

#### Anthropometric Measurement

Table 1 shows the anthropometric findings. The mean (SD) height z score was $-1.06 (1.24)$ and the mean weight z score $-0.83 (1.14)$, which were both significantly reduced in comparison with age matched normal growth data (height $p=0.0005$, weight $p=0.002$). Patient number 6 had spent a number of years in a transit refugee camp before developing his arthritis and had probably experienced significant preceding nutritional impairment. When this patient was excluded significant differences in height and weight z scores remained between patients values and the normal growth data.

Weight for height for age index was below 90% in five cases, a level indicative of nutritional compromise. An additional six cases had z scores less than $-1.5$ for height or weight, or both, and yet had normal weight for height indexes owing to the proportional reduction in overall growth parameters. Skinfold and mid-arm circumference measurements gave no additional information about nutritional status that than given by the weight z score in this patient group (data not shown).

Significantly lower $z$ scores were found in patients with prolonged disease duration ($>4$ years) for both height ($p=0.001$) and weight ($p=0.027$), but no correlation was found for either height or weight with corticosteroid treatment or disease onset subtype. Of note, however, three patients with pauciarticular onset (Nos 3, 17, and 21) had progressed to a polyarticular course, whereas those with persistent pauciarticular disease did not show such marked somatic disturbance.

#### IGF-1/Growth Hormone

Eight patients (three systemic, two polyarticular, three pauciarticular) had IGF-1 $z$ scores less

### Table 1

<table>
<thead>
<tr>
<th>Case No</th>
<th>Age (years)</th>
<th>Onset type</th>
<th>Disease duration (years)</th>
<th>Corticosteroids</th>
<th>IGF-1* (z)</th>
<th>Height (s)</th>
<th>Weight (s)</th>
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<td>1</td>
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<tr>
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<td>Pa</td>
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<td></td>
<td>$+1.71$</td>
<td>$0.00$</td>
<td>$0.05$</td>
<td>93</td>
</tr>
<tr>
<td>21</td>
<td>4-5</td>
<td>Pa</td>
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<td>$0.57$</td>
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<tr>
<td>23</td>
<td>6-3</td>
<td>Pa</td>
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<td></td>
<td>$+1.12$</td>
<td>$0.22$</td>
<td>$0.08$</td>
<td>91</td>
</tr>
</tbody>
</table>

Mean
- $0.84$
- $-1.06$
- $0.48$

SD
- $1.09$
- $1.24$
- $1.14$

S = systemic; P = polyarticular; Pa = pauciarticular; IGF-1 = insulin like growth factor 1.
than \(-2.00\) from the age/sex matched mean. Only three of the patients with low IGF-1 were receiving corticosteroid treatment. As a corollary, seven further patients receiving corticosteroid treatment had IGF-1 z scores between \(-0.46\) and 2.00.

Seven of the eight patients with low IGF-1 and three additional patients (Nos 8, 9, and 10) who showed poor rate of growth underwent overnight growth hormone estimation (table 2). In comparison with previously published data from our laboratory, the overall mean growth hormone secretion of the patients with juvenile chronic arthritis (5.5 mU/l) was greater than that of children with growth hormone deficiency (1.5 mU/l) and that of short statured children with normal growth hormone response to pharmacological stimulation (4.6 mU/l).18

Seven of the 10 patients studied, however, had mean growth hormone secretion within the range of the growth hormone deficient group (0.39–5.5 mU/l). The mean number of pulses, 5.5 (range 3–7), was greater than previously seen in our growth clinic (unpublished observation). All six patients with juvenile chronic arthritis studied had normal response to clonidine stimulation. Patient 1, who before the onset of systemic onset juvenile chronic arthritis was growing on the 97th centile, improved in rate of growth, IGF-1 concentration, and mean growth hormone secretion as her disease state went into remission, with catch up growth being seen (figs 1 and 2). She continued to receive steroids throughout this time. Pubertal development also occurred, reflected in increased oestradiol from 44 to 181 pmol/l over this period. Overall, the IGF z scores correlated with both the height and weight z scores (IGF v height \(r=0.43, p=0.042\); IGF v weight \(r=0.46, p=0.028\)). The mean weight z score of those patients with an IGF-1 z score less than \(-2.00\) was significantly less than those greater than \(-2.00\) (\(-1.79\) compared with \(-0.41, p=0.007\)). The height z scores, however, failed to reach a significant difference (\(-1.87\) compared with \(-0.70, p=0.065\)). No significant association was shown between IGF-1 and rate of growth or between mean growth hormone secretion and rate of growth or height/weight z scores (data not shown). The mean growth hormone secretion correlated with the amplitude of the secretory peaks, both maximum (\(r=0.94, p<0.001\)) and mean (\(r=0.87, p<0.001\)), and with the length of the peaks (\(r=0.62, p<0.05\)). No correlation was found between the IGF-1 z score and mean growth hormone secretion (\(r=0.34, p=0.32\)).

**Discussion**

Growth disturbance in juvenile chronic arthritis is of considerable importance as it may be a persisting legacy of the disease after the active inflammatory process has apparently abated.
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Considerable concern has evolved about the limitation of linear growth secondary to corticosteroid treatment, somewhat overshadowing the potential effect of the disease process itself. With the recognition of growth hormone secretion and subsequent IGF-1 production as the predominant hormonal stimulus to growth, studies have investigated possible abnormalities in this axis in a variety of chronic disease states—for example, Crohn’s disease, chronic renal failure, thalassaemia major—though no consistent abnormality has been shown.

Growth hormone secretion has been stimulated by drugs, but no abnormality has been shown in juvenile chronic arthritis irrespective of the corticosteroid regimen used—daily, alternate day, etc. This was confirmed in our study with those children receiving treatment on alternate days. Butenandt, however, showed improvement in rate of growth in patients with juvenile chronic arthritis during exogenous growth hormone treatment. Growth hormone is secreted in a pulsatile nature, especially during sleep; abnormalities in this physiological pattern of secretion, particularly in the amplitude of the pulses, are reported in short children who have responded normally to insulin/arginine provocation of growth hormone secretion. Furthermore, low IGF-1 concentrations are also reported in such children. As low IGF-1 concentrations have been noted in systemic onset juvenile chronic arthritis this study aimed at determining physiological growth hormone patterns in children with low IGF-1, the hypothesis being that poor sleeping patterns due to disease may be blunting the physiological response.

We confirmed that IGF-1 may be low in patients with all disease onset subtypes and not only in those with systemic onset. No consistent growth hormone secretory abnormality was shown, however, though some patients did have low concentrations, falling within the range of growth hormone deficient patients. The increased pulse frequency that we noted may reflect disturbances of hypothalamic control of growth hormone secretion either because of abnormal sleep patterns or poor nutrition. Our sample size limited further analysis of the interaction between nutrition and growth hormone secretion. Variability of physiological growth hormone secretion with overlap between normal and growth hormone deficient children has been described. Furthermore, variability between consecutive periods of sampling shows the possible limitations in the investigation of physiological secretion. Chipman et al also studied physiological secretion in patients with polyarticular juvenile chronic arthritis, concluding that no abnormality was present as all patients had at least one peak exceeding 10 ng/ml during sleep. Considerable variability of 24 hour growth hormone secretion was also shown as in our study.

One patient was found to have a dramatic improvement in rate of growth concomitant with increases in both IGF-1 and mean growth hormone secretion. Although pubertal development was a likely contributor, as confirmed biochemically, the improvement coincided with disease remission, and rate of growth seemed more exaggerated than would be expected for puberty alone. This suggests that catch up growth, in disease remission, may have an endocrinological component in some cases.

Previous reports have used reduction in weight for height for age index to indicate nutritional impairment whereas we also used weight z scores to indicate possible nutritional compromise. This may be more meaningful than calculating the index alone as the index does not detect patients with proportional reduction of both height and weight. Low IGF-1 and normal/raised growth hormone concentrations are found in malnutrition, which, as suggested by Bennett et al, may be occurring in patients with juvenile chronic arthritis. Insulin-like growth factor 1 was in closer association with weight z score than height z score, further reflecting a possible link with nutritional status. The weak correlation of IGF-1 with growth hormone secretion, who showed catch up growth associated with increases in both IGF-1 and mean growth hormone secretion. Studies of a potentially treatable situation, such as poor nutrition, are the basis for further study.

This study does not exclude the possibility that exogenous growth hormone treatment may be of benefit in individual patients with persistently poor rate of growth, but such an approach should only be considered after careful consideration of nutritional factors and maximum control of the disease process.

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