A Pharmacokinetic and Dosing Study of Intravenous Insulin-Like Growth Factor-I and IGF-Binding Protein-3 Complex to Preterm Infants

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ABSTRACT: In preterm infants, low levels of Insulin like growth factor 1 (IGF-I) have been associated with impaired growth and retinopathy of prematurity. Our objective was to study safety and pharmacokinetics of i.v. administered rhIGF-I with its binding protein 3 (rhIGFBP-3) to preterm infants. At 3 d chronological age, an i.v. 3 h infusion of rhIGF-I/rhIGFBP-3 was administered followed by serial measurements of IGF-I and IGFBP-3. Infants were evaluated for physiologic safety measurements. The individual dose of rhIGF-I ranged from 1 to 12 µg/kg. The study was conducted at Queen Silvia Children’s Hospital, Gothenburg, Sweden, between January and November 2007. Five patients (3 F) with mean (range) post menstrual age 27 wk (26–29) and birth weight 1022 g (810–1310) participated. IGF-I and IGFBP-3 levels before infusion were median (range) 18 (25–59) and 838 (754–1182) ng/mL, respectively. Median (range) half-life for IGF-I and IGFBP-3 was 0.79 (0.59–1.42) and 0.87 (0.85–0.94) hours, respectively. Blood glucose, insulin, sodium, potassium, and physiologic safety measures were within normal ranges. The rhIGF-I/rhIGFBP-3 equimolar proportion was effective in increasing serum IGF-I levels and administration under these study conditions was safe and well tolerated. (Pediatr Res 65: 574–579, 2009)

IGF-I plays a central role in regulating fetal growth, particularly in the third trimester as demonstrated by experimental studies (1), which seems to be the case in humans as well, as demonstrated with patients with IGF-I gene defects (2,3). In addition, IGF-I has been shown to play important roles in the intrauterine development of the CNS and the vasculature (2,3). IGF-I promotes proliferation, maturation, and differentiation of neural stem cells and of neuronal and oligodendrocyte precursors (4). IGF-I has also been shown to play an important role in retinal vascularization, both physiologic and pathologic, in both experimental and clinical studies (2,5).

The bioavailability of IGF-I is regulated by several binding proteins. IGF binding protein 3 (IGFBP-3) is the predominant binding protein in the blood circulation. Normally, virtually all the IGF-I/IGFBP-3 is in a ternary complex with their binding to the acid-labile subunit (ALS) (6) and are protected from protease activity. In preterm children (born <32 gestational weeks), we have earlier found very low serum levels of ALS during the first weeks of life (data not published). The dissociation of IGF-I and IGFBP-3 is controlled by proteases, which regulate tissue availability of bioactive free IGF-I, which can then bind to cellular IGF-receptors (6). IGFBP-3 can modulate cellular mechanisms independently of IGF-I, and we have earlier shown that IGFBP-3 promotes vascular regrowth in experimentally induced retinopathy (7).

Serum concentrations of both IGF-I and IGFBP-3 fall rapidly, below in utero levels, after preterm birth (8). IGF-I levels in fetal serum rise in the third trimester and are positively correlated with gestational age and birth weight (9,10). Thus, in general, the more premature the infant, the lower is the level of serum IGF-I at birth.

In preterm infants, low serum levels of IGF-I have been associated with slow weight gain and slow head (brain) growth as well as with the later development of retinopathy of prematurity (5,11,12). The aim of our study was to evaluate the safety and pharmacokinetic parameters of a 3-h infusion of rhIGF-I/rhIGFBP-3-glucose to five very preterm infants and to determine the dose of rhIGF-I required bringing IGF-1 into the physiologic range, defined as the in utero levels for corresponding GA (20–60 µg/L) (13).

MATERIALS AND METHODS

Study subjects. A total of five infants were recruited to the study, which was conducted in the neonatal ward at Queen Silvia Children’s Hospital in...
Gothenburg, Sweden, between January 2007 and November 2007. Inclusion criteria were gestational age (GA) at birth between 26 wk + 0 d and 29 wk + 6 d (GA estimated by ultrasound in pregnancy week 17), a birth weight > −2 SD and < +2 SD based on z score for age and sex and a serum IGF-I concentration measured at postnatal day 2 below 25 μL/L.

The children were enrolled during their first day of life. Exclusion criteria were severe sepsis, gross malformation (including cardiac abnormalities other than PDA), known or suspected chromosomal abnormality, genetic disorder or syndrome according to the investigator’s opinion, IGF-I levels ≥ 25 μg/L at study day 2 (to minimize the risk of un-physiologically high IGF-I levels), plasma glucose level < 2.5 mM or > 10 mM at study day 2, and administration of plasma later than 24 h after birth.

The five patients (three females and two males) had a mean gestational age of 27 wk and a mean birth weight of 1022 g. Clinical characteristics of the five infants are given in Table 1. One girl had a serum IGF-I on study day 3 ≥ 25 μL/L, however, as her IGF-I level on study day 2 was 14 μg/L she was included in the study. The study was approved by the Local Ethics Committee in Göteborg and the Swedish Medical Products Agency. All participating parents gave written informed consent before start of study.

rhIGF-I/rhIGFBP—glucose preparation. Individual infusion solutions were prepared at the hospital pharmacy of the Queen Silvia Children’s Hospital in the following manner. The study drug was mescemarin ribafmate (PLEX, Insmed). Insmed is an equimolar preparation of recombinant protein complex of rhIGF-I and rhIGFBP-3. IPLEX (Lots No. DP0506 and DP0609) was diluted with 10% glucose solution in two dilution steps to achieve appropriate doses for each infant. The dilution of Iplex was done based on the individual study drug solution infusion rate, which was 1 mL/kg/h (i.e. 3 mL/kg since infusion duration was standardized to 3 h) in all infants. In both dilution steps, dilution bags (FREKA Mix; Fresenius Kabi AG) were used. Study drug and glucose solution were administered to the bags by different syringes. The solution was transferred to an infusion pump, Alaris Asena CC (Mediagnost GmbH, Tübingen, Germany). The IGF-I samples were diluted 1:50 and the IGFBP-3 samples were diluted 1:300. The intra-assay coefficients of variation (CV) for IGF-I were 18, 11, and 7% at concentrations of 9, 33, and 179 μg/L, respectively. The intra-assay CV for IGFBP-3 was 10, 7, and 6% at concentrations of 716, 1750, and 3929 μg/L, respectively. All samples were analyzed in the same assay. The methods have been described in detail previously (16).

Pharmacokinetic modeling. All serum concentrations of IGF-I and IGFBP-3 at baseline and up to 48 h after the stop of infusion of study drug were analyzed using nonlinear mixed effects modeling in NONMEM VI (17). An underlying endogenous level (baseline concentration) was estimated from the data and the pharmacokinetics of the infused IGF-I and IGFBP-3 were described by compartmental modeling. One- and two-compartment models with linear or nonlinear elimination were evaluated.

The choice of structural model, inclusion of between-subject variability in the parameters and the choice of additive or proportional residual error models were tested for statistical significance using the minimum objective function value (OFV) produced by NONMEM. A drop in OFV of 6.64, which corresponds to a significance level of p < 0.01, was required for adding one extra parameter. Clearance (CL) and volume (V) terms were scaled allometrically by weight (18).

\[ CL = CL_{1\text{kg}} \cdot \text{weight}^{0.75} \]

\[ V = V_{1\text{kg}} \cdot \text{weight} \]

Individual pharmacokinetic parameters of IGF-I and IGFBP-3 were derived using the POSTHOC step in NONMEM. The half-live was computed as \[ t_{1/2} = \ln(2) \times \text{V/C}L \] and the area under the concentration-time curve (AUC) resulting from the mescemarin ribafmate infusion was computed as AUC = dose/CL.

Statistics. All data are expressed as the median and range. Predominantly descriptive statistics are presented. It was not appropriate to perform statistical hypothesis testing due to the small sample size.

RESULTS

Dose. Administration of rhIGF-I/rhIGFBP-3 had effects on serum concentrations of IGF-I and IGFBP-3 after the
complex was administered to the infants at the postnatal age of 3 d during a 3-h infusion. The individual percentage change from serum baseline concentrations of IGF-I and IGFBP-3 at the respective sampling time-points are shown in Fig. 1A and B.

Median (range) IGF-I and IGFBP-3 levels immediately before infusion were 18 (12–28) and 771 (651–1047) ng/mL, respectively, Tables 2 and 3. Immediately after completing drug infusion. One (880 g; 6 µg/kg); unfilled boxes and black line, 2 (1,220 g; 24 µg/kg); red filled circle and red line, 3 (760 g; 33 µg/kg); green filled triangles and green line, 4 (1,115 g; 33 µg/kg); dark blue rhomb and dark blue line, 5 (810 g; 59 µg/kg); light blue filled boxes and light blue line. B, Δ% IGFBP-3 from baseline after 3 h infusion of IGF-I/IGFBP-3 in different doses (n = 5) in five study subjects. Sampling time points are immediately before study drug infusion (−3), immediately following completed infusion of drug (0), and at 1, 2, 6, 12, 18, 24, and 48 h post completed drug infusion. One (880 g; 6 µg/kg); unfilled boxes and black line, 2 (1,220 g; 24 µg/kg); red filled circle and red line, 3 (760 g; 33 µg/kg); green filled triangles and green line, 4 (1,115 g; 33 µg/kg); dark blue rhomb and dark blue line, 5 (810 g; 59 µg/kg); light blue filled boxes and light blue line. B, Δ% IGFBP-3 from baseline after 3 h infusion of IGF-I/IGFBP-3 in different doses (n = 5) in five study subjects. Sampling time points are immediately before study drug infusion (−3), immediately following completed infusion of drug (0), and at 1, 2, 6, 12, 18, 24, and 48 h post completed drug infusion. For IGFBP-3, the endogenous concentration, CL1kg and V1kg were estimated to 758 µg/L, 0.0698 L/h/kg, and 0.0906 L/kg, respectively, resulting in a t1/2 of 0.90 h, for a typical child of 1000 g, Table 3. Between-subject variability in the estimated endogenous concentration of IGFBP-3 was 17% and the additive residual error was 60.4 µg/L. The individual model-predicted maximum increase in IGF-I concentration from the estimated endogenous concentration after the mecasermin rinfabate administration was 19.5 (3.4–28.3) ng/mL and AUC was computed to be 61.8 (11.6–123) ng/mL · h.

Table 2. Observed baseline, administered dose (based on weight at day of study drug infusion), maximum IGF-I concentration and estimated half-life of IGF-I in infants after single dose of rhIGF-I/rhIGFBP-3 infusion (rhIGF-I/rhIGFBP-3 [µg/kg/3 h] × 0.21)

<table>
<thead>
<tr>
<th>Characteristics (gender)</th>
<th>Baseline IGF-I (µg/L)</th>
<th>Dose rhIGF-I/rhIGFBP-3 (µg/kg/3h)</th>
<th>Dose rhIGF-I (µg/kg/3h)</th>
<th>Max IGF-I (µg/L)</th>
<th>t1/2 IGF-I (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Girl</td>
<td>18</td>
<td>6</td>
<td>1</td>
<td>25</td>
<td>0.785</td>
</tr>
<tr>
<td>Boy</td>
<td>21</td>
<td>24</td>
<td>5</td>
<td>37</td>
<td>0.939</td>
</tr>
<tr>
<td>Boy</td>
<td>15</td>
<td>33</td>
<td>7</td>
<td>38</td>
<td>0.701</td>
</tr>
<tr>
<td>Girl</td>
<td>28</td>
<td>33</td>
<td>7</td>
<td>59</td>
<td>1.423</td>
</tr>
<tr>
<td>Girl</td>
<td>12</td>
<td>59</td>
<td>12</td>
<td>39</td>
<td>0.593</td>
</tr>
</tbody>
</table>

All pharmacokinetic parameters are based on infant weight on day of study drug administration. Allometric scaling predicts that the increase in distribution volume is directly proportional to body weight whereas clearance is less than proportional to weight, resulting in that the typical half-life increases with body weight. The calculated typical half-life of IGF-I in an infant of 0.5 kg and 1.5 kg was 0.72 h and 0.95 h, respectively, for IGF-1, and 0.76, and 1.00 h, respectively, for IGFBP-3.

Safety. There were no changes in blood pressure or heart rate that had clinical importance in any of the five patients, Fig. 2. Blood glucose levels after infusion were all normal.

Median (range) insulin concentrations in the five individuals were stable and normal immediately before, at 1 h or at 24 h after rhIGF-I/rhIGFBP-3 infusion and were 7.8 (6.6–17.0), 6.6 (1.7–19.0), and 8.4 (5.5–14.0) mU/L, respectively.

Serum sodium and potassium levels were within normal ranges at all measuring points. No difference was observed in

serum IGF-I and IGFBP-3 levels were 38 (25–59) and 838 (658–1182) ng/mL, respectively. In all five children, the serum IGF-I levels, immediately after infusion, were within in utero range (13).

**IGF-I and IGFBP-3 pharmacokinetic parameters.** One-compartment models with linear elimination were sufficient to describe the IGF-I data and the IGFBP-3 data. The endogenous serum baseline concentration, clearance (CL1kg), and distribution volume (V1kg) for IGF-I were estimated to 18.8 µg/L, 0.0924 L/h/kg, and 0.114 L/kg, respectively, with a calculated half-life (t1/2) of 0.86 h for a typical child of 1000 g, Table 2. The estimated between-subject variability (CV) in endogenous baseline concentration and CL were 34 and 31%, respectively. The additive residual error was 2.56 µg/L. The individual model-predicted maximum increase in IGF-I concentration from the estimated endogenous concentration after the mecasermin rinfabate administration was 19.5 (3.4–28.3) ng/mL and AUC was computed to be 61.8 (11.6–123) ng/mL · h.

For IGFBP-3, the endogenous concentration, CL1kg and V1kg were estimated to 758 µg/L, 0.0698 L/h/kg, and 0.0906 L/kg, respectively, resulting in a t1/2 of 0.90 h, for a typical child of 1000 g, Table 3. Between-subject variability in the estimated endogenous concentration of IGFBP-3 was 17% and the additive residual error was 60.4 µg/L. The individual model-predicted maximum increase in IGFBP-3 concentration from the estimated endogenous concentration after the mecasermin rinfabate administration was 77.4 (17.4–180.9) ng/mL and AUC was computed to be 351 (61.2–636) ng/mL · h.
after stopping the infusion, although the glucose level was normal. The normoglycemia observed throughout the study possibly reflects the co-infusion of 10% glucose (which is routine for extremely premature infants). Premature infants are also often insulin resistant (19), catabolic and have poor metabolic regulation. Several studies have shown that IGF-I may improve insulin sensitivity (20,21) as well as have positive effects in children who are catabolic from severe burns (22,23). IGF-I is an insulin sensitizer by virtue of its suppression of GH secretion, as well as through direct actions on peripheral tissues (24). Prematurity is a state of partial GH insensitivity that in part can explain the catabolic state of these infants. Elevated GH levels have earlier been shown to be more strongly associated with severe ROP than low levels of IGF-I (25). The complete role of the GH/IGF-I axis for retinopathy development is still not fully understood.

Main clinical indication in humans of recombinant human IGF-I alone (mecasermin) or as an equimolar preparation with rhIGFBP-3 (mecasermin rinfabate) to date, has been growth hormone insensitivity syndrome (GHIS/Laron syndrome) (26) and severe insulin resistance (27).

The mean half-life of 0.86 h for i.v. administered IGF-I in our study of premature infants is considerably shorter than in both older children and adults with GHIS as well as in healthy subjects in whom a half-life of free IGF-I after a 75 min constant i.v. infusion of rhIGF-I was calculated to be $14 \pm 1$ h (28). Part of the explanation for the shorter half-life of the drug in the present study may be low endogenous levels of IGFBP-3 and ALS (29). The short half-life indicates that administration of IGF-I in preterm infants should be based on continuous infusion of IGF-I to achieve the targeted physiologic concentration.

IGF-I and IGFBP-3 levels both decrease after preterm birth and significant dips in endogenous IGF-I and IGFBP-3 levels at around 2–4 d after birth has recently been recognized (8). This decrease can confound the interpretation of IGFBP-3 levels seen after the 3-h infusion, as the amount of IGFBP-3 infused is not enough to “overcome” the physiologic decrease. This is more apparent with IGFBP-3 since the molar amount of IGFBP-3 infused with IPLEX is less, relative to endogenous IGFBP-3 levels, compared with the infusion of IGF-I. In addition, there is a possibility of degradation of nonglycosylated IGFBP-3 by proteases (nonglycosylated IGFBP-3 is more readily degraded than endogenous glycosylated IGFBP-3), which might be explained by low ALS levels in these infants.

**Figure 2.** Heart rate (bpm) after 3 h infusion of IGF-I/IGFBP-3 in different doses ($n = 5$) in five study subjects. Sampling time points are immediately before study drug infusion (−3), immediately following completed infusion of drug (0) and thereafter at every hour up to 12 h post completed drug infusion. 1 (880 g; 6 μg/kg); unfilled boxes and black line, 2 (1,220 g; 24 μg/kg); red filled circle and red line, 3 (760 g; 33 μg/kg); green filled triangles and green line 4 (1,115 g; 33 μg/kg); dark blue rhomb and dark blue line, 5 (810 g; 59 μg/kg); and light blue filled boxes and light blue line.
In this very limited study on five patients with a narrow range of gestational ages and birth weight SDS, we did not see any correlation between these variables and response. We could only see that birth weight and baseline levels had an impact on the response. Predictions from the developed pharmacokinetic model suggest that in a preterm infant with a birth weight of 1000 g a dose of study drug between 100 and 150 μg/kg/h, i.e. corresponding to 21–32 μg of IGF-I, is required to increase the IGF-1 concentrations by 10 ng/mL to achieve physiologic concentrations of IGF-I during the immediate postnatal period. This is a considerably lower dose than the s.c. dose approved by USFDA for children with severe primary IGF-I deficiency 0.12–2 μg/kg once daily (1 mg Mecasermin rinfabate equates to 0.2 mg of rhIGF-I) (27). The results indicate that the predicted half-life of IGF-I and IGFBP-3 increases with increasing birth weight. This is because clearance is not directly proportional to weight whereas the distribution volume increases proportionally to weight.

The main reason for studying the pharmacokinetics of IGF-I in preterm infants is the implications this growth factor might have for the health of these infants. Postnatal low serum levels of IGF-I in preterm infants have been associated with poor growth, poor head circumference development and retinopathy of prematurity (12,30 –34).

The main limitations of this study is that it would have been very interesting to measure factors like GH and IGFBP-I as this could have given us information about eliminating hypoglycopenotropism and thus demonstrating adequate IGF-I replacement as well as insulin sensitivity. However, in these small infants the amount of blood that can be drawn is very limited and did not allow for these measures in the present study. This study does not provide a rationale for using the equimolar preparation of rhIGF-I/rhIGFBP3 instead of IGF-I alone as an infusion in preterm infants.

In conclusion, we have shown for the first time that infusion of the equimolar preparation of rhIGF-I/rhIGFBP3 elevates IGF-I in extremely preterm infants. The pharmacokinetic results serve as a basis for design of further studies on the efficacy and safety of the systemic use of IGF-I to improve growth, metabolic status as well as for neuro- and vasoprotection in extremely preterm infants.

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