CALCIUM METABOLISM IN A CASE OF GARGOYLISM, STUDIED WITH THE AID OF RADIOCALCIUM

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A number of studies have been reported on the fate of radiocalcium (Ca-45) in humans (1-12), but none have included chemical analyses of the concentration of Ca-45 in the skeleton and in such tissues as liver, brain, or intestinal tract. We are now reporting the results of a metabolic study with Ca-45 performed on a 10 year old patient with terminal gargoylism. The analyses of periodic samples of serum, urine, cerebrospinal fluid, and feces obtained during the patient’s illness and of various tissues removed at necropsy 16 days after the administration of Ca-45 have made it possible to formulate a relatively complete picture of the calcium metabolism of this individual, although the lack of comparable information makes it difficult to evaluate whether and how gargoylism modified the calcium metabolism of our patient.

Gargoylism is now well recognized clinically (13), but its pathogenesis is largely unknown. Evidence is accumulating (14-19) to indicate that gargoylism is a storage disease, characterized by the accumulation of abnormal types (20) and quantities of mucopolysaccharides in a large variety of tissues, notably liver and spleen.

Recently, it was proposed that polysaccharides, such as chondroitin sulfate, are involved in the calcification process in bone and cartilage [see Amprino (21) for a review of the evidence]. Although gargoylism appears to be characterized by a defect in structural polysaccharides, it is possible that calcification is also defective in this condition. Dawson (17) has reviewed the histological changes seen in the bones of patients with gargoylism and has concluded that the changes are dystrophic and are probably of direct metabolic origin. However, no metabolic studies have been reported. It seemed of interest, therefore, to study in detail the calcium metabolism of a patient with terminal gargoylism.

MATERIAL AND METHODS

Description of patient. The patient was a male child, 9 years 10 months old, 137 cm. high, who weighed 16.4 Kg. He had been institutionalized since the diagnosis of the disease almost five years earlier. When the study was initiated, the disease had progressed to a point where the boy’s life expectancy appeared short. The patient was almost completely unaware of his surroundings, had lost his ability to communicate except by a bird-like whine, and showed no response to sounds. He appeared incapable of any muscular control, was incontinent of urine and feces, and had to be handled like a 3 months old baby. His arms and legs were flexed, with ankylosis of the joints. The hands were clenched, the fingers showed clubbing, and the skin was taut and shiny. There was general hirsutism.

Design. Although the patient was on the danger list, his condition had remained unchanged for several weeks preceding the study. It was planned to follow the course of the injected Ca-45 intensively during the first five days, then at weekly, and later at monthly intervals. The death of the patient on Day 16 prevented an extended follow-up.

Dietary regimen. Before the initiation of the study and throughout its duration, the patient was spoon-fed with commercial baby foods, baby cereals, and milk. No attempt was made to regulate his intake. During the course of the study, i.e., on days when specimen collection was carried out, food equivalent to that fed to the patient was collected by having the nursing personnel place in a collection jar quantities of food identical to those fed the patient. His calcium intake was analyzed as 3.8 Gm. on Day 1, 5.2 Gm. on Day 2, 4.3 Gm. on Day 3,
The equation predicts the course of the specific activity of the serum with time. $C_i$, equals per cent dose per mg. Ca, $t$ equals hours. Average calcium content of serum: 0.085 mg. per ml.; range, 0.081 to 0.090. Average calcium content of cerebrospinal fluid: 0.049 mg. per ml.; range, 0.044 to 0.059.

1.3 Gm. on Day 4, 1.0 Gm. on Day 5, 0.5 Gm. on Day 8, and 3.2 Gm. on Day 15.\(^4\)

**Calcium-45.** The Ca-45 was diluted in a solution containing 0.8 per cent of NaCl and 0.1 per cent of CaCl\(_2\). A total of 3 ml. of solution was injected intravenously; it contained 80 $\mu$C of Ca-45, 1 mg. of Ca\(^{++}\), 10 mg. of Na\(^{+}\), and 16 mg. of chloride, and had been sterilized by autoclaving.

**Specimen collections.** Blood was obtained by venipuncture at intervals indicated in Figure 1. Samples of cerebrospinal fluid were obtained by spinal puncture (Figure 1). Urine samples were obtained from the catheterized patient at three hour intervals during the first day; thereafter (Days 2 to 5, 8 and 15), 24 hour specimens were collected. Following collection, the urine was acidified by the addition of glacial acetic acid in the proportion of 2 ml. per 100 ml. urine. *Feces* specimens were collected on diapers. On the first day, these were pooled to constitute eight hour specimens; on subsequent days, 24 hour specimens were analyzed.

**Autopsy.** Representative samples of mineralized and soft tissues were removed at autopsy and analyzed (see Tables I to III). In the case of certain hard tissues (Table II), the analysis of the whole section was supplemented by analysis of scrapings, obtained with a dental burr at anatomically defined regions (22, 23).

**Analytical procedures**

**Calcium analysis.** Blood calcium content was determined by direct precipitation of the calcium from the serum as the oxalate, which was titrated with perchlorocetic acid (24). Urine calcium was determined by precipitation of the calcium either directly from the urine or from an ash solution of the urine. Calcium contents of stool specimens were determined from fecal ash solutions. The coefficient of variation (standard deviation times 100 divided by the mean) of the calcium determinations was usually below 5 per cent.

**Ca-45 analysis.** Ca-45 was determined by measuring the radioactivity of calcium oxalate samples which contained 4.0 mg. of Ca. The solid samples were counted under a thin end-window (<1.5 mg. per cm.\(^2\)) Geiger-Müller counter, connected to an automatic sample-changing device. Suitable standards were employed in all counting runs and all results are reported as corrected for decay from the day of injection. Counting errors varied,
but their coefficients of variation were generally less than 10 per cent. The term "dose" refers to the quantity of Ca-45 injected.

The analytical techniques employed have been described in detail by Bronner, Harris, Maletskos, and Benda (3).

RESULTS

Serum

When the serum data are plotted on semilogarithmic coordinates (Figure 1), they fall on a complex curve which is approximated by the following equation:

\[ C_t = 0.177e^{-8.11t} + 0.061e^{-0.322t} + 0.035e^{-0.004t} + 0.018e^{-0.006t} \]

where

\[ C_t = \text{Per cent of dose per mg. of Ca in serum} \]

\[ t = \text{Hours.} \]

| Table I | Specific activity of selected soft tissues obtained at autopsy of a patient with gargoylism who had received by intravenous injection 80 \( \mu \text{C} \) of Ca-45Cl2 16 days previously |
|---------------------------------|---------------------------------|-----------------|
| **Tissue**                      | **Ca-45 content**              | **Ca-45 content** |
|                                 | (% dose/Gm. Ca)                | (mg. Ca/Gm.)    |
| Serum                           | 2.26                           | 0.083           |
| Brain                           | 3.06*                          | 0.082           |
| Spleen                          | 2.79                           | 0.089           |
| Kidney                          | 1.55                           | 0.086           |
| Liver                           | 2.52                           | 0.085           |
| Diaphragm                       | 1.67                           | 0.084           |
| Intestinal tract               |                                |                 |
| Duodenum                        | 1.57                           | 0.084           |
| Jejunum                         | 2.90-2.82†                     | 0.084           |
| Ileum                           | 2.42                           | 0.084           |
| Ascending colon                 | 1.51-1.76                      | 0.084           |
| Transverse colon                | 1.25                           | 0.084           |
| Descending colon                | 2.10                           | 0.084           |
| Sigmoid flexure                 | 1.44                           | 0.084           |
| and rectum                      | 1.83                           | 0.084           |
| Bile                            | 2.92                           | 0.084           |
| (Ca content: 0.692 mg./ml.)     |                                |                 |
| Washings                        |                                |                 |
| Small intestine                 | 1.54                           | 0.084           |
| Colon                           | 0.75                           | 0.084           |
| Sigmoid flexure and rectum      | 1.01                           | 0.084           |

* See Figure 2 for location of sampling sites.

This equation is not valid for the first five minutes of the study and therefore its first term has little significance. Its general course resembles that observed in other children (5, 7), but differs from these significantly (Table IV), inasmuch as the exponents of the second and third terms of equation 1) are only about one-tenth of the corresponding terms of the normal boys, and are also smaller than those of the adult. The exponent of the fourth term is as fast as that of the fourth term in the equation of the adult, but much slower than
the corresponding exponent in the equation of the normal boys.

A comparison of the last term of equation (1), with the coefficients, rate constants, and compartment sizes calculated by Krane, Brownell, Stanbury, and Corrigan (8), shows that the last term, with an intercept of 18.2 per cent dose per Gm. Ca and a slope of 0.138 per day, agrees well with the third term (Days 4 to 9) of equations reported by these authors for euthyroid adults. Similarly, the “calcium pool,” 16.7 Gm., (per cent activity retained at Day 9 divided by the serum specific activity on Day 9) of our subject was comparable to that shown by Krane and collaborators (8) for euthyroid adults. However, on a body weight basis, the “pool” was much greater in our subject than in normal adults, as would be expected from the higher calcification rate seen in children.

Comparison of terms 2 and 3 of equation (1), with the comparable coefficients and rate constants of Krane and co-workers (8), shows (see also Table IV) that the processes which have given rise to these terms in our patient differ markedly from those observed in normal adults or normal children.

**Urine and feces data**

As would be expected, the specific activity data of serum and urine (Figure 1) can be considered to fall essentially on one curve, although the corresponding values for the urine tend, perhaps, to be lower than those for the serum. The very low output of calcium in the urine (Figure 3) may bear on this point.

The endogenous fraction of fecal calcium during the first five days is estimated from the mean serum specific activity during that period, on the assumption that the specific activity of the se-

**TABLE IV**

Comparison of exponents in serum disappearance equations *

<table>
<thead>
<tr>
<th>Boys†</th>
<th>Young adult†</th>
<th>Gargoyle patient</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.281</td>
<td>0.713</td>
<td>0.135</td>
</tr>
<tr>
<td>0.075</td>
<td>0.0112</td>
<td>0.0054</td>
</tr>
<tr>
<td>0.004</td>
<td>0.0004</td>
<td>0.00057</td>
</tr>
<tr>
<td>0.0005</td>
<td>0.00009</td>
<td>0.00010</td>
</tr>
</tbody>
</table>

* Dimension: minute⁻¹.
† Taken from Bronner and Harris (7).
creted intestinal juice always equals that of the serum (5). As the mean specific activity of the serum between 0 and 120 hours was 0.0233 per cent dose per mg. Ca, and as our patient had excreted 9.70 per cent of the dose in his stools during that period, his endogenous calcium output was 416 mg. or 11.0 per cent of the total calcium output in the feces. This value is similar to that observed by us in other instances (7).

In contrast with observations on normal, ambulatory patients (1, 5) or on terminal cancer patients (2, 4), the ratio of total urinary to fecal Ca-45 output was very much below unity (Figure 3). Geissberger (1) has reported ratios of this magnitude for patients with cardiac insufficiency and steatorrhea, as have Laszlo and Spencer [quoted by Comar and Wasserman (9)] for two patients with cancer of the prostate.

**Cerebrospinal fluid**

The specific activity of the spinal fluid approached that of the serum between 5 and 48 hours after injection (Figure 1). Thereafter, its specific activity either was equal to or higher than that of the serum. Postmortem analysis of brain tissue indicated a specific activity of 0.031 per cent dose per mg. Ca, as opposed to the serum specific activity of 0.023 at the time of autopsy. This difference gives support to the possibility that the specific activity of the cerebrospinal fluid may actually have exceeded that of the serum.

**Bile**

Possibly the serum was diluted with calcium released by autolysis during the period which elapsed between death and autopsy (about seven hours); otherwise, the specific activities of bile fluid and serum ought to have been more nearly alike than was found (Table 1). The observed specific activity of the bile fluid was of the same order of magnitude (0.002 per cent per ml.) as reported for adult rats (25).

Singer, Maqsood, Medlen, and Comar (26) have very recently reported on the endogenous and biliary excretion of Ca-45 in dogs. Their data indicate that, within 25 minutes of injection, 21 per cent of the activity found in the contents of the gastrointestinal tract (excluding the stomach) is found in the gall bladder. Our data show that the total activity found in the bile approximated 27 per cent of the activity washed out of the gastrointestinal tract, excluding the stomach (Table 1).

**Soft tissues**

The specific activity of the small intestine increased in a posterior direction. Contrary to expectations, the relative specific activity of the intestine was not highest near the point where the common bile duct enters the duodenum (7 to 8 cm. posteriorly of the pylorus). On the other hand, the low relative specific activity of the washings of the intestinal tract (Table 1) suggests that even at a late stage in the patient’s illness—when his food intake was rather small—labeled body calcium was still being diluted to a fair degree by unlabeled exogenous calcium.

Of some interest is the observation that several soft tissues (brain, spleen, liver, sections of the ileum and jejunum) had a higher specific activity than serum (Table 1). Bronner (27) has recently shown that in very young rats the specific activity of many tissues is higher than that of serum. Inasmuch as the serum supplies tracer to a multi-

![Fig. 3. Cumulative Output of Ca-45 in Urine and Feces of a Patient with Gargoyleism Following Intravenous Administration of 80 μc. of Ca-45Cl₂](image)
compartment system, it is not surprising that its specific activity drops below that of some compartments.

**Mineralized tissues**

The data in Tables II and III are of interest because they indicate the fate of a single dose of Ca-45 in relation to the architecture of the bones. It is apparent, as has been reported by others (28–36), that the most highly calcified areas contain the smallest quantity of isotope. Thus, the extremities of the ulna showed more labeling than the shaft, undoubtedly because growth was still proceeding at the epiphyses, whereas the center of the shaft had stopped growing and was probably subject to remodelling only. The sternal cartilage of the ribs had a specific activity which was near that of the serum and which was approximately ten times as high as that of the bones [cf., Kulp, Eckelmann, and Schulte (37) for similar observations on strontium deposition].

**DISCUSSION**

The much slower rate of disappearance of Ca-45 from the serum of our patient, as compared with that of normal boys (5, 7), suggests a disturbed calcium metabolism due perhaps to interference with calcium deposition or accretion. It is not surprising, therefore, that evaluation of A ("accretion value") in accordance with the method of Bauer, Carlsson, and Lindquist (38) [see Bronner and Harris (7) for a description and discussion of this approach] reveals an average value of 0.54 Gm. per day. This is about one-fourth of the value calculated for a group of boys, and less than the value calculated for two young men (7).

The average E value, which represents the exchangeable fraction of body calcium (7, 38) of our patient, was 2.8 Gm., or 0.93 per cent of an estimated body calcium content of 300 Gm. This value is comparable to those we calculated for other subjects (7).

Conceivably, the inability to deposit calcium, as indicated by the accretion value, is related to the accumulation of abnormal polysaccharide materials similar to those reportedly found in liver and spleen (16, 18–20). Chondroitin-like materials have recently been reported to occur in bone (40).

Because our patient's urinary output of Ca-45 was depressed, though his output of endogenous calcium in the feces was within the normal range, his absorption of calcium may have been altered, but his death precluded further studies.

If the data of Table II are plotted on semilogarithmic coordinates (Figure 4) they fall along a straight line which is expressed (method of least squares) by the equation

\[
C = 1.18e^{-0.0097A},
\]

where

\[C = \text{Specific activity (per cent of dose per Gm. of Ca), and} \]

\[A = \text{Calcium content of hard tissue (mg. of Ca per Gm. of tissue).} \]

This relationship is significant (p = 0.02) and is a measure of calcification in this individual. Engfeldt, Engström, and Zetterström (31) have hypothesized a similar relationship for the uptake of P-32, although their diagram does not indicate

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**FIG. 4. RELATIONSHIP BETWEEN SPECIFIC ACTIVITY AND CALCIUM CONTENT OF BONES AND TEETH IN A PATIENT WITH GARGOYLLISM 16 DAYS FOLLOWING THE INTRAVENOUS ADMINISTRATION OF 80 μC. OF CA-45Cl₂**

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*Mitchell, Hamilton, Steggerda, and Bean (39) estimate the Ca content of a normal boy weighing 164 Kg. as 185 Gm., and that of a normal 10 year old boy as 396 Gm. We have assumed our patient's body calcium content lay between these limits.*
the exponential nature of the relationship. Plummer, Hansard, Comar, and Beeson (41) have also reported an inverse relation between specific activity of calcium in bones of the bovine fetus and fetal age, but again not as an exponential function. Rogers, Weidmann, and Jones (33) have discussed the difficulty of interpreting these observations from the point of view of mechanisms.

On the whole there is good agreement between the data in Tables II and III, even though the size of the sample was very small in the case of the analyses reported in Table II; this shows that the sampling of bone with a dental burr at well-defined anatomical points can provide both accuracy and definition (23). This method should prove very helpful as a quantitative adjunct to autoradiography.

The findings listed in Tables II and III confirm the observations of Sognnaes, Shaw, and Bogoroch (42) that the teeth do not participate in the general metabolism of calcium. Thus, unerupted teeth showed some labeling, although much less than skeletal structures, while deciduous teeth had almost negligible radioactivity. The two deciduous canine teeth analyzed in toto averaged 168 mg. of calcium. If it is assumed that the calcium content of this patient's body was 300 Gm., then the canine teeth could have contained nearly 0.168 times 100 over 300 or 0.06 per cent of the labeled calcium when, in fact, they contained one-tenth this quantity.

In the case of one complete metacarpal bone removed at autopsy, a similar calculation would have led to an expected Ca-45 content of 0.01 per cent. This value was found experimentally and indicates high metabolic activity. If ossification was delayed in this as compared to other bones (ossification is often delayed in the metacarpal bone centers of patients with gargoylism (13)], one might expect a relatively higher calcification rate leading to greater concentration of the isotope. In addition, the calcium content of the metacarpal bone as a whole was only 75.6 mg. of Ca per Gm. as contrasted with the higher Ca content of the bone scrapings (cf., Figure 4).

The analysis of the calvarium (Tables II and III) indicates that calcification was not primarily the cause of its immense thickening. The level of the specific activity of the exterior of the calvarium was low, approximating that of the teeth; at the interior, the calvarium had a specific activity typical of most bony tissues. Its rate of calcification was similar to that of the other hard tissues (Figure 4). Anderson, Emery, McAlister, and Osborn (10) have recently reported that the Ca-45 content of the skull of a myelomatous patient was 0.0096 μc. per Gm. or 0.00062 per cent dose per Gm. 60 days after administration of the isotope. Our patient's skull contained 0.0313 per cent dose per Gm. 15 days after administration of the Ca-45. Even if his level had dropped by half at the end of 60 days, our patient's skull would still have contained over 25 times more isotope than reported for the skull of the myelomatous adult.

Because Anderson and collaborators (10) reported that the Ca-45 tended to concentrate in the myelomatous regions, it is difficult to know whether the higher isotope level of the skull of our patient with gargoylism was the result of abnormal metabolic activity or whether the lower level in the myelomatous skull was due to preferential deposition in the myelomatous regions.

**SUMMARY AND CONCLUSIONS**

1. The rate of disappearance of Ca-45 from the serum of a 10 year old patient with terminal gargoylism is described by the following equation:

\[
C_t = 0.177e^{-0.11t} + 0.061e^{-0.322t} + 0.035e^{-0.041} + 0.018e^{-0.061},
\]

where

\[
C_t = \text{Specific activity of the serum (per cent of dose per mg. of Ca)}, \text{ and}
\]

\[
t = \text{Time (hours)}.
\]

The exponents in the second and third terms of this equation, valid for 15 days, indicate an appreciably slower rate of disappearance than observed heretofore in normal boys and one young adult. The exponent in the fourth term indicates a rate comparable to that of normal adults, but slower than seen in normal boys.

2. The urinary calcium output of this patient was severely depressed, so that the urinary-fecal partition ratio of Ca-45 was 0.24, as contrasted with a more normal range of 1 to 2. The specific activities of serum and urine were found to be alike.
3. Fecal output of calcium and Ca-45 was in the normal range. The endogenous fecal output of this patient was calculated to equal 83 mg. per day, or 11.0 per cent of his average total excretion of calcium in the feces. This value is similar to that observed in other humans.

4. The depressed rate of disappearance of the injected Ca-45 from the blood is consistent with the possibility of decreased calcium deposition. The accretion value, 0.54 Gm. of Ca per day, was much lower than that calculated for normal boys. The exchangeable fraction of body calcium, E, was 0.93 per cent of his estimated body calcium content of 300 Gm. This value is comparable to the E value of more normal subjects.

5. Data on the level of tracer in the soft tissues (brain, liver, spleen, gastrointestinal tract) obtained at necropsy 16 days after the start of the study were consistent with similar data in animals.

6. The level of Ca-45 in the mineralized tissues varied inversely and exponentially with the level of calcium in bone and tooth samples obtained with a dental burr at anatomically defined loci. This indicates that the labeling of calcified tissues corresponded to the rate of calcium deposition. Analyses of gross bone specimens supported this conclusion.

7. In the absence of comparable data on normal humans, it is difficult to know whether and how gargoylism caused the abnormal calcium metabolism of our patient.

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We are indebted to Dr. R. F. Sognnaes of the Harvard School of Dental Medicine for many helpful suggestions in planning the analyses of the bone samples and for preparing and identifying the location of the bone and tooth samples. Mrs. Jean R. Moor and Mrs. Gloria Romano rendered skillful technical assistance. We are happy to express our appreciation to the staff of the Walter E. Fernald State School who nursed this patient with skill, self-sacrifice and patience.

REFERENCES


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