URINARY EXCRETION OF ACID POLYSACCHARIDE IN RHEUMATOID ARTHRITIS AND OTHER DISEASES

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The excretion of acid mucopolysaccharide in the urine was described by Kerby (1954), using a method devised by Astrup (1947) for the estimation of heparin. Scott (1955) introduced the precipitation of acid mucopolysaccharides by quaternary ammonium salts; this method was adapted to urine by Di Ferrante and Rich (1956a, b). Di Ferrante (1957) found that patients with rheumatoid arthritis showed increased excretion of acid mucopolysaccharide when compared with normal controls. Di Ferrante, Robbins, and Rich (1957) described similar excess excretion in cases of disseminated lupus erythematosus. It was concluded (Rich, Di Ferrante, and Archibald, 1957) that “the urinary excretion of acid polysaccharide is elevated in several diseases associated with abnormal metabolism of mucopolysaccharide-rich connective tissues”. It was therefore decided to compare the excretion of acid mucopolysaccharide in patients with rheumatoid arthritis with that in patients suffering from various other maladies, and to define, by isolation of the material, the type of acid polysaccharide in each particular specimen of urine, in order to explore possible differences and to give some idea of the origin of the material. Pooled samples were collected for more extensive analysis.

Materials and Methods

24-hour samples of urine were collected from patients in hospital. Most of the patients with rheumatoid arthritis had clinically active disease and a raised erythrocyte sedimentation rate, and the majority of the adult patients had a positive Rose-Waaler test. The urines were kept, without preservative, at 4° C. 300 ml. urine was used for each estimation.

The method used for the isolation of acid mucopolysaccharide was similar to the modification by Meyer, Grumbach, Linker, and Hoffman (1958) of the method of Di Ferrante and Rich (1956b). Samples with a specific gravity of 1.020 or more were diluted with an equal volume of water. The urine was filtered and acidified to pH 6.0 and a warm aqueous 5 per cent. solution of cetyl trimethylammonium bromide was added (1.7 ml./100 ml. urine). After standing overnight at 4° C., the solution was centrifuged in the cold. The precipitate was washed three times in a large volume of ethanol. The material insoluble in ethanol was dissolved, with agitation and warming to 37° C., in a small volume of 10 per cent. aqueous sodium acetate, and the pH was adjusted to 9.0. Insoluble material was centrifuged down and re-extracted with water. The combined supernatant was acidified to pH 5.0. The crude acid polysaccharide was then precipitated with 1.5 volumes of ethanol and left at 4° C. overnight. This was centrifuged and the precipitate washed with 80 per cent. ethanol. It was extracted with a small volume of water and centrifuged at high speed. To the clear solution, calcium acetate and acetic acid were added to final concentrations of 5 per cent. and 0.5 N, respectively. The solution was fractionally precipitated with alcohol, according to the method of Meyer, Davidson, Linker, and Hoffman (1956). Resulting fractions were washed in increasing concentrations of alcohol and dried in ether, and this was followed by desiccation over phosphorus pentoxide. Each sample was weighed and analysed for uronic acid both by the orcinol method and by Dische’s carbazole method. All methods of analysis were as used by Loewi and Meyer (1958), and described by Meyer and others (1956) and Rapport, Meyer, and Linker (1951).

To obtain purified material for analysis, urine pools of about 10 l. were collected. After isolation by the above procedures, the material was dissolved in water and digested with pepsin for 24 hrs, followed by trypsin for a similar period. The material was reprecipitated and subjected to the procedure of Sevag (1934). After another precipitation, treatment with kaolin and Lloyd’s reagent was carried out, and finally Dowex 50 (H+) was used. The method is essentially that used by Meyer and others (1956).
Results

Table I shows the results of estimations of acid mucopolysaccharide in the urines of 73 individuals. Since the carbazole values differed somewhat from sample to sample, the amounts of polysaccharide found were adjusted by calculation to a carbazole value of 20 per cent. in each case, so that comparison could be made. The actual readings were nearly all close to this figure. In all cases, rheumatoid and others, acid polysaccharide was precipitated at between 40 and 50 per cent. ethanol concentration, and gave a ratio of carbazole to orcinol close to 1:0, indicating chondroitin sulphate A or C. Though some adult cases of rheumatoid arthritis showed high excretion values, others showed values similar to those found in other diseases and in normal controls. The difference between female rheumatoid arthritics and normal controls approached significance ($P = <0.10, >0.05$). Eight cases showed a daily excretion of more than 19 mg. acid mucopolysaccharide. These included four cases of rheumatoid arthritis, one of which was complicated by pyelo-nephritis, three cases of epilepsy, and one of pneumonia. Excretion of less than 1 mg./24 hrs was seen in two cases of rheumatoid arthritis complicated by amyloidosis. In the children studied, excretion of acid polysaccharide in rheumatoid arthritis was similar to that occurring in other diseases, but significantly lower than that found in normal controls ($P = 0.01$). The figure taken for any particular patient was the first one obtained after admission, though in most cases, several estimations were made subsequently. In two patients, in each of whom daily estimations were done on six consecutive days, the day-to-day variations were found to be relatively small (3.1 mg. with a standard deviation of ±0.65 in one case and 1.24 mg. with S.D. ±2.38 in the other). The majority of the figures were lower than those obtained by Di Ferrante (1957); the reason for this may be sought in differences in the techniques used, since, in the present investigation, it was decided to sacrifice completeness of recovery to removal, as far as possible, of non-acid mucopolysaccharide material. Using the almost identical methods, Meyer and others (1958) obtained figures for normal controls which are similar to those reported here.

Table II shows figures for acid mucopolysaccharide excretion in four patients before and during salicylate administration. In two cases a fall was recorded, while in one (Case 4) a rise in excretion.

**Table I**

<table>
<thead>
<tr>
<th>Condition</th>
<th>Age (yrs)</th>
<th>No. of Cases</th>
<th>Rheumatoid Arthritis</th>
<th>Other Diseases*</th>
<th>Connective Tissue Disorders other than Rheumatoid Arthritis</th>
<th>Epilepsy</th>
<th>Normal</th>
<th>Rheumatoid Arthritis with Amyloidosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Children</td>
<td>10</td>
<td>9</td>
<td>5.81 ± 2.66</td>
<td>6.82 ± 13.92</td>
<td>0.09 to 36.8</td>
<td>0.09 to 6.5</td>
<td>13.55 ± 7.7</td>
<td></td>
</tr>
<tr>
<td>Adult</td>
<td>12</td>
<td>3 to 15</td>
<td>3 to 10</td>
<td>2.96 ± 2.84</td>
<td>26.9 and 31.6</td>
<td>26.9 and 31.6</td>
<td>6.1 to 24</td>
<td></td>
</tr>
<tr>
<td>Females</td>
<td>34-3</td>
<td>16 to 63</td>
<td>3.1 to 39</td>
<td>1.9 to 15.1</td>
<td>19.15</td>
<td>19.15</td>
<td>6.57 ± 2.9</td>
<td>6.57 ± 2.9</td>
</tr>
<tr>
<td>Males</td>
<td>29</td>
<td>17 to 50</td>
<td>6.2 to 10</td>
<td>8.6 and 1.0</td>
<td>14.2</td>
<td>14.2</td>
<td>6.88 ± 5.1</td>
<td>6.88 ± 5.1</td>
</tr>
</tbody>
</table>

**Table II**

<table>
<thead>
<tr>
<th>Case No.</th>
<th>APS (mg./24 hrs)</th>
<th>Change while taking Salicylates</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control Period</td>
<td>Salicylate (4-5 g./day)</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>8.7; 9.0</td>
<td>3.5; 3.8; 2.6; 2.9; 3.0</td>
</tr>
<tr>
<td>2</td>
<td>25.4; 21.3</td>
<td>12.3; 9.4</td>
</tr>
<tr>
<td>3</td>
<td>9.3; 10.2</td>
<td>14.2; 2.0</td>
</tr>
<tr>
<td>4</td>
<td>5.6; 6.3</td>
<td>12.0; 17.9</td>
</tr>
</tbody>
</table>

* Tuberculosis, nephritis, pneumonia, appendicitis.
occurred, although this patient’s temperature became normal and the erythrocyte sedimentation rate was falling during the administration of salicylate. Case 3 showed an initial rise, followed by a fall in acid polysaccharide excretion.

The purified acid polysaccharide obtained from large volumes of normal and rheumatoid urine was analysed (Table III). Precipitation by ethanol began at 40 per cent., and was complete at 50 per cent. In the two purified samples cited in Table III, the only amino-sugar found chromatographically was galactosamine, suggesting a chondroitin sulphate. When, however, less highly purified material was subjected to this procedure, a weak glucosamine spot was also obtained. This suggests a relatively small amount of hyaluronic acid or of keratosulphate, but the presence of some glucosamines-containing protein cannot be excluded. The high Dische (carbazole) to orcinol ratio is indicative of chondroitin sulphate A or C; rotational values are suggestive of chondroitin sulphate C and digestion by testicular hyaluronidase (reductimetric method, 24-hr incubation) also suggests chondroitin sulphate A or C rather than B, since the latter is not hydrolysed by this enzyme. That the substance is not chondroitin sulphate B is further shown by the relatively high ethanol solubility, the high ratio of Dische to orcinol value (chondroitin sulphate B gives a ratio of around 0-5) and the low rotation. Precipitation at 40-50 per cent. ethanol and the rotation figures are suggestive of chondroitin sulphate C rather than A. In the absence, however, of an infra-red pattern and of material sufficiently pure for this investigation to be carried out, such differentiation between chondroitin sulphate A and C cannot be made with any degree of assurance.

Discussion

This investigation was undertaken first to determine the exact nature of the acid polysaccharide excreted in rheumatoid arthritis and other diseases, and, secondly, to discover whether the increased excretion, as reported by Di Ferrante (1957), was peculiar to rheumatoid arthritis or was also encountered in unrelated diseases.

The same type of chondroitin sulphate was discovered in all specimens of urine examined in this investigation and the full analysis of pooled material reported in Table III confirmed this. Di Ferrante and Rich (1956a) showed that normal urine contained chondroitin sulphate A or C, and Di Ferrante (1957) indicated that the material obtained from pooled rheumatoid urine was similar, and could be depolymerized by hyaluronidase. Di Ferrante (1957) also reported the presence of some hyaluronic acid in the urinary acid polysaccharide. I have found no evidence of this in the purified material. A trace of glucosamine was found, however, when less highly purified material was examined. Urine from a case of Marfan’s syndrome (personal observation) showed no abnormal constituents qualitatively or quantitatively. Material isolated in the present investigation from patients with epilepsy who were taking hydantoin proved to be, as in the rheumatoid and normal cases, chondroitin sulphate A or C.

With regard to the origin of this urinary constituent, Kerby (1954) showed that acid mucopolysaccharide was present in urine in the renal pelvis and that the urinary content was not raised by prostatic massage. Badin, Schubert, and Vouras (1955) obtained evidence of the presence in increased amount of a similar substance in plasma from cases of rheumatoid arthritis. One may assume, therefore, that urinary chondroitin sulphate is derived from plasma by renal excretion. Meyer (1957), discussing the distribution of acid mucopolysaccharides in various tissues, lists chondroitin sulphate A and/or C as occurring in cartilage and bone. These two tissues must, therefore, be considered as possible sites of origin of the urinary acid mucopolysaccharide, and both cartilage destruction and osteoporosis are indeed found in rheumatoid arthritis. It was noticed, however, that

### Table III

<table>
<thead>
<tr>
<th>Urine Source</th>
<th>Amount (mg.)</th>
<th>N (per cent.)</th>
<th>Hexosamine</th>
<th>Uronic Acid (per cent.)</th>
<th>[α]D</th>
<th>Digestion with Testicular Hyaluronidase</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rheumatoid Arthritis</td>
<td>28·0 (50 per cent. Ethanol)*</td>
<td>3·5</td>
<td>20·8</td>
<td>Galactosamine</td>
<td>18·6</td>
<td>28·6</td>
</tr>
<tr>
<td>Normal</td>
<td>16·5 (50 per cent. Ethanol)*</td>
<td>†</td>
<td>21·0</td>
<td>Galactosamine</td>
<td>18·0</td>
<td>29·1</td>
</tr>
</tbody>
</table>

* Percentage at which precipitate obtained.  † Inadequate material for estimation.
cases of rheumatoid arthritis with extensive osteoporosis and a case of Cushing’s disease did not have an unduly great excretion. Similarly, activity of the disease did not correlate with the amount of acid mucopolysaccharide excreted. It is also not possible to account on such a basis for the relatively high levels of excretion shown by some epileptics.

When the quantitative excretion figures were examined, much overlap was found between rheumatoid arthritis, patients with other diseases, and normal controls. Owing to the very wide ranges of readings in each of these categories, no statistically significant differences were found. The reason for these wide variations is apparently not to be sought in the techniques employed, since repeated readings on single patients did not show such variations. Several of the adult patients with rheumatoid arthritis excreted abnormally large amounts of acid polysaccharide, while others excreted only small amounts. The distribution showed no constant relationship to activity of the disease, as assessed by the erythrocyte sedimentation rate. Many of the patients were receiving salicylates and other forms of treatment, which may have caused a decrease in the amount excreted; Di Ferrante (1957) observed such an effect, and a fall occurred in three out of four patients on salicylates, as recorded in Table II. There were, however, some patients who showed a low excretion in the absence of drug therapy. The low level of excretion of acid polysaccharide in children with rheumatoid arthritis may be the result of interference with skeletal growth. Such an explanation is also suggested by the significantly greater excretion in normal children when compared with normal adults (P = <0.02, >0.01). This latter finding was also noted by Rich and others (1957). Relatively high excretion was found in cases of epilepsy receiving hydantoins and in a case of pneumonia (normal controls of comparable age showed much lower excretion). In two cases of rheumatoid arthritis complicated by extensive amyloidosis involving the kidneys, excretion of acid polysaccharide was particularly low. Craddock and Kerby (1955) found decreased excretion in cases of diabetes with renal complications. Patients with other disorders involving connective tissue, such as rheumatic fever and dermatomyositis, did not show high urinary acid polysaccharide excretion. Di Ferrante and others (1957) reported a raised level of excretion in disseminated lupus erythematosus. The failure to demonstrate a high level of acid polysaccharide excretion in many cases of rheumatoid arthritis, although the disease was active, and failure to demonstrate raised levels in some other connective tissue disorders, suggest that the statement by Rich and others (1957) that “the urinary excretion of acid polysaccharide is elevated in several diseases associated with abnormal metabolism of mucopolysaccharide-rich connective tissues” needs qualification. The results of the present investigation do not suggest that urinary acid polysaccharide excretion constantly mirrors any such abnormality in the tissues, such as might occur in the rheumatic diseases. To date, evidence for such a disorder has only been found in gargoyleism, which shows high urinary acid polysaccharide excretion (Meyer and others, 1958).

Summary

Acid mucopolysaccharide has been estimated in the urine of patients with rheumatoid arthritis and with other diseases, and in normal controls. A raised level of excretion was found in some adult patients with rheumatoid arthritis, while others showed normal or low levels. No rise was found in patients with rheumatic fever and dermatomyositis, but a few patients with other diseases showed raised levels of excretion. In children, the excretion was less in rheumatoid arthritis than in normal controls. The acid mucopolysaccharide in all cases was found to be chondroitin sulphate A or C.

REFERENCES


Excrétion urinaire de polysaccharide acide dans l’arthrite rhumatismale et dans d’autres maladies

RÉSUMÉ

On détermine le mucopolysaccharide acide chez des malades atteints d’arthrite rhumatismale et d’autres affections, ainsi que chez des témoins. On en trouva un taux élevé chez quelques adultes atteints d’arthrite rhumatismale, mais chez d’autres ce taux fut normal ou bas. On ne trouva pas de taux augmentés chez des malades atteints de rhumatisme articulaire aigu ou de dermatomyosite, mais on les observa chez quelques autres malades. Chez des enfants rhumatismatiques rhumatisrales il y eut une diminution du taux, absente chez des enfants normaux. Dans tous les cas on identifi...
le mucopolysaccharide acide comme sulfate de chondroitine A ou C.

Excreción urinaria de polisacarido ácido en la artritis reumatoide y otras enfermedades

Sumario

Se determinó el mucopolisacarido ácido en la orina de enfermos con artritis reumatoide, con otras enfermedades y en la de testigos. Se halló una tasa elevada en algunos adultos con artritis reumatoide, pero en otros ésta fue normal o baja. No hubo aumento en enfermos con reumatismo poliarticular agudo o con dermatomiositis, pero se observó una tasa aumentada en algunos otros enfermos. En niños reumatoideo-artríticos la excreción fue disminuida en comparación con la de niños normales. En todos los casos se identificó el mucopolisacarido ácido como sulfato de condroitina A o C.