CREATINE AND GUANIDOACETIC ACID METABOLISM IN PITUITARY DISEASE

BY

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Schrire (1937) found that patients with an acidophil tumour of the pituitary excreted creatine in the urine, and this was later confirmed (Cumings, 1944). It was also shown in the latter paper that the fasting blood creatine was raised above the normal in acromegaly, and the results of a creatine tolerance test were given in five cases of pituitary disorders. The cause of the creatinaemia and creatinuria in acromegaly is obscure, but it had been shown previously that the urine excretion of creatine paralleled the height of the blood creatine provided that a level higher than 0.6 mg./100 ml. had been obtained. Guanidoacetic acid has been shown to be a precursor of creatine (Borsook and Dubnoff, 1940; 1941) and its estimation in pituitary disorders does not appear to have been recorded previously. This paper records a large series of patients with pituitary disorders, together with a few cases of thyrotoxicosis, in whom creatine and guanidoacetic acid studies have been made.

Material and Methods

Creatine studies have been made on 39 subjects consisting of 10 normal controls, 11 with acromegaly, nine with chromophobe adenomata or suprapituitary cysts, six with miscellaneous pituitary disorders, and three with thyrotoxicosis. The diagnosis was made in most cases at operation and by histology, or by clinical and radiological evidence which was unmistakable. Guanidoacetic acid was estimated in the urine in 26 out of the 39 subjects, and, in a few patients, more than one examination was made.

All patients were placed on a meat- and fish-free diet and 24-hourly urines were collected over a five- or six-day period. Creatine and creatinine were estimated according to the principles of the Folin and Wu technique (1919). A creatine tolerance test as described by Tierney and Peters (1943) was performed, usually on the third day, and the blood levels of creatine and creatinine estimated according to the technique of Peters (1942) as in my previous paper, but using a photoelectric colorimeter with a filter of 520 μ .

Urinary guanidoacetic acid was estimated as described by Hoberman (1947), following the technique of Dubnoff and Borsook (1941) as modified by Sims (1945),

after each urine had been treated with arginase to remove any arginine present. The arginase was prepared by the technique described by Hoberman (1947).

Results

Examples of results in each group are recorded in Appendix 1 and Appendix 2, and one example in each group is shown in the figures.

Normal Controls.—None of the 10 controls showed any creatinuria, and the amount of urinary creatinine was reasonably constant from day to day. The fasting blood creatinine was 0.5 mg./100 ml. or below in nine cases and in the other case 0.6 mg./100 ml., and this can be regarded as normal since Samuels, Sellers, and McCaulay (1946) obtained figures of up to 0.6 mg./100 ml. in normal controls even though Tierney and Peters (1943) found figures of up to 0.5 mg./100 ml. only.

Guanidoacetic acid was estimated in four of these controls and a normal excretion of 30–50 mg. a day was obtained in all except one, a very large muscular man and a heavy manual worker, who excreted 60–80 mg. a day. The results found in this patient are shown in Fig. 1. It might be mentioned that not infrequently a little creatine is excreted in the first day of the test as a result of the creatine in the diet of the preceding day.

FIG. 1.—Diagram of guanidoacetic acid excretion in a normal control (N. H.).

Chromophobe or Suprapituitary Tumours.

—The nine patients in this group showed results identical with the normal control group as regards urinary creatine and creatinine. The fasting blood creatinine in all cases was within normal limits, but the rise in blood level was somewhat higher after 1 g. of creatine than was to be seen in the control group. Guanidoacetic acid was estimated in all except one case, and raised excretion rates were seen in all the patients except one, the only one who had a suprapituitary cyst. The usual excretion was as high as 80–100 mg. a day. The

FIG. 2.—Diagram of guanidoacetic acid in a case of chromophobe adenoma.

patient with a raised blood creatinine had a severe nephritis from which he died post-operatively. Fig. 2 shows the findings in one case.

Acromegaly.—There was creatinuria in all except two patients, one of whom was examined nine years after removal of the pituitary acidophil tumour, and the other (S.D.) who was clinically normal following hormonal therapy. There was a raised fasting blood creatine in 10 out of the 11 patients, the only normal level of creatine being seen in the clinically normal patient who had received hormone treatment. The creatine tolerance test gave results similar to those previously recorded (Cumings, 1944).

Guanidoacetic acid was estimated in seven patients in this group, which included the patient apparently cured by hormones and another only examined post-operatively. These two patients showed a slightly raised excretion of guanidoacetic acid of up to about 60–80 mg. daily, but one of them excreted very little urine in each 24-hour period. The remaining five typical cases all showed a markedly increased excretory rate, usually of about 100 mg. daily.

A few of the patients were examined after operation and after hormone therapy as well as before these treatments. One patient, five years after operation, showed an improvement in the creatinaemia and creatinuria, and, except for one day during the test period, passed no more guanidoacetic acid in the urine than a normal control. Another patient nine years after operation showed no creatinuria even though the fasting blood creatine was slightly raised. One patient who showed the usual chemical findings seen in acromegaly was given hormone therapy, but after six months there were no improvements in blood or urine chemistry though there had been some clinical improvement. The other patient similarly treated, but for a much longer period, was practically normal when examined clinically and pathologically. An example of one of these cases is shown in Fig. 3.

Miscellaneous Pituitary Disorders.—

This group consisted of two patients with tumours of mixed cell type (both acidophil and chromophobe cells), both of whom were diagnosed in the laboratory as having overactivity of acidophil cells; three patients with gross obesity and other evidence of pituitary dysfunction; and one very small, thin patient who also had a suprapituitary cyst. All these patients showed creatinuria, together with a raised fasting blood creatine. The very thin patient of the 39 patients examined showed the highest rise in blood creatine after ingestion of 1 g. of creatine by mouth. The guanidoacetic acid excretion of this patient one and a half years after operation was normal even though the

FIG. 3.—Diagram of guanidoacetic acid in a case of acidophil adenoma.

creatinuria still persisted. Guanidoacetic acid was also estimated in three other patients before operation. The obese patients so examined both showed a raised urinary output. One patient with a so-called "pituitary abscess" gave a normal rate of excretion of guanidoacetic acid.

FIG. 4.—Diagram of guanidoacetic acid excretion in a case of thyrotoxicosis.

Thyrotoxicosis. — Three patients were examined and all showed creatinuria with a fasting blood creatine, although in no case was this latter figure as high as that seen in the cases of acromegaly. All three cases showed a normal excretion of guanidoacetic acid. The results found in a moderately severe case of thyrotoxicosis with a basal metabolic rate of +75% are shown in Fig. 4. Table I is a summary of the blood creatine curves of the cases recorded in Figs. 1–4.

TABLE I
BLOOD CREATINE CURVES AFTER 1 G. CREATINE IN FOUR CASES RECORDED IN THE DIAGRAMS

Condition	Time	Creatinine (mg./100 ml.)	Creatine (mg./100 ml.)
Normal	Fasting	0.9	0.3
	1 hour after creatine	1.25	0.55
	2½ „ „ „	1.0	0.5
Chromophobe adenoma ..	Fasting	0.7	0.5
	1 hour after creatine	0.9	0.9
	2½ „ „ „	0.8	0.7
Acromegaly	Fasting	0.62	0.88
	1 hour after creatine	0.75	1.75
	2½ „ „ „	0.75	0.85
Thyrotoxicosis	Fasting	0.7	0.9

Discussion

The results obtained in the examination of this large series of pituitary disorders confirm the earlier findings in 1944 as far as creatine metabolism is concerned, and there can be no doubt that in acromegaly with an acidophil tumour of the pituitary there is creatinaemia and creatinuria. It is seen that a large part of the ingested creatine is rapidly excreted in the urine and that the blood soon loses its increased quantity of creatine. Previously it had not been possible to distinguish

biochemically between a normal patient and a patient with a chromophobe adenoma, even though one could distinguish between these two and an acromegalic patient. It was also not possible by creatine metabolic studies to distinguish between thyrotoxicosis and acromegaly. From the results of the guanidoacetic acid studies, however, it is now possible to determine into which of these four groups any one patient falls. Table II shows the distinguishing biochemical features between the normal person and the three other conditions discussed.

TABLE II

DISTINGUISHING BIOCHEMICAL FEATURES OF THE NORMAL AND CERTAIN PITUITARY DISORDERS

Disease	Urine Creatine	Urine Guanidoacetic Acid	Fasting Blood Creatine
Acromegaly	++	++	Over 0.6 mg./100 ml.
Chromophobe tumour	0	+	Less than 0.6 mg./100 ml.
Thyrotoxicosis ..	++	Normal	Over 0.6 mg./100 ml.
Normal	0	„	Less than 0.6 mg./100 ml.

Guanidoacetic acid is formed in the kidneys (Borsook and Dubnoff, 1941; Borsook, Dubnoff, Lilly, and Marriott, 1941) by transamidation from the amino-acids glycine and arginine, and is then transformed into creatine by methylation, probably utilizing methionine. This latter process probably takes place in the liver (Borsook and Dubnoff, 1940, 1947; Bloch and Schoenheimer, 1941), although there is some evidence from examinations made in cases of nephritis that this can also take place in the kidneys (Samuels *et al.*, 1946). It would appear, therefore, that the overaction of some of the pituitary cells affects these chains of events and so alters the urinary excretion of these substances.

There is in acromegaly not only an increased production of guanidoacetic acid but also an increased formation of creatine, hence the increase in production of the former substance must be quite considerable. It is different in a patient with a chromophobe adenoma as there is here an excess of guanidoacetic acid but no increase in creatine excretion; it would appear unlikely that any storage of creatine is taking place as this would need to be considerable, and, further, there is no increased excretion of creatinine over the normal, nor is there any creatinaemia. Equally, the excess of guanidoacetic acid must be an excess of production otherwise the blood level of creatine would drop and there would be a lowered excretion of creatinine. Further, because of the normal urinary creatinine and blood creatine it cannot be assumed that the guanidoacetic acid is not being made into creatine. It may be mentioned that in neither acidophil nor chromophobe tumours can any evidence be found biochemically of liver damage, as a number of liver function tests have been carried out in these patients and no abnormality has yet been detected; further, in one patient only was there any evidence of renal disease,

and the results obtained in that case, apart from the level of the blood creatinine, were not significantly different from the others in his group.

Borsook and Dubnoff (1947) showed that the presence of creatine in high concentration may retard the methylation of guanidoacetic acid to such an extent as to give a raised excretion of this substance. This cannot be the mechanism in acromegaly, for then one would expect to see the same process in thyrotoxicosis, but in this condition the guanidoacetic acid excretion is normal. It is therefore unlikely that the pituitary thyrotropic hormone plays any part in acromegaly, and further proof of this is seen in the inability of thiouracil to affect the creatinuria in acromegaly as it does in thyrotoxicosis (Schrire, 1948).

Hoberman, Sims, and Engstrom (1948) showed that methyl testosterone given to normal men produced creatinuria and also an increased urinary excretion of guanidoacetic acid, and they state that this is due to an increase in synthesis of creatine as a result of the effect of this hormone on the reactions involved in the synthesis of guanidoacetic acid. Samuels, Henschel, and Keys (1942) showed that if the methyl testosterone be given for some weeks then there is also creatinaemia and creatinuria occurring usually after 10 days' treatment with methyl testosterone. In the same way testosterone propionate as well as methyl testosterone produced similar results although not giving such a high level of excretion of creatine and guanidoacetic acid in the urine (Hoagland, Shank, and Gilder, 1944). These findings would seem to give a hint as to the mechanism causing the disturbance that is found in pituitary disease. It is not suggested that it is necessarily either of these two hormones that is responsible, but it does seem probable that the acidophil and the chromophobe cells each elaborate hormones that act in the manner suggested by Hoberman *et al.*, and it does seem to be true that the effects of both methyl testosterone and testosterone propionate are those seen in acromegaly. This would seem to be a basis for further study and one on which to build a programme of hormonal therapy for these two types of patients.

Summary

Creatine studies have been made in 10 normal controls, in 26 patients with pituitary disorders, and in three with thyrotoxicosis, and the urinary excretion of guanidoacetic acid followed in 26 of these 39 subjects. It has been shown to be possible biochemically to distinguish between the normal, the thyrotoxic, the acromegalic, and the patient with a chromophobe adenoma of the pituitary.

The possible causes of the biochemical abnormalities have been discussed and the possibilities of abnormal hormonal activity in the two pituitary disorders mentioned.

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APPENDIX 1

In this appendix a representative protocol for one normal control and each type of case discussed is set out.

Normal (E. M.)

Urine					
Day	Volume	Creatinine	Creatine	Guanidoacetic Acid	
1	2,300 ml.	1.20 g.	0	0.035 g.	
2	1,700 "	1.14 "	0	0.052 "	
3*	1,920 "	1.12 "	0.31 g.	0.051 "	
4	1,450 "	0.96 "	0	0.044 "	
5	2,450 "	1.04 "	0	0.056 "	

Creatine Tolerance Test

Fasting	0.5 mg./100 ml.	0.4 mg./100 ml.
1 hour after creatine	0.75 " " "	0.55 " " "
2½ hours " " " " "	0.62 " " "	0.48 " " "

Chromophobe Adenoma.—A single woman of 25 complained of failing vision for seven years, and headache and hirsuties for six years. Menstruation was normal. Examination showed a coarse skin, which was greasy, and there was a definite deposit of hair over the face. There was a bitemporal lower quadrant defect with rather pale

*Creatine tolerance test (1 g. by mouth) on this day.

discs in the eyes. No other abnormality was found clinically or radiologically. Keto-steroid excretion, blood sugar curve, and C.S.F. were normal. At a later admission there was an additional hysterical factor also present. There was insufficient evidence of progression of the lesion causing the eye signs to warrant operative interference.

Urine				
Day	Volume	Creatinine	Creatine	Guanidoacetic Acid
1	910 ml.	0.96 g.	0	0.075 g.
2	945 "	0.95 "	0	0.080 "
3*	1,250 "	0.94 "	0.13 g.	0.085 "
4	1,070 "	1.02 "	0	0.108 "
5	440 "	0.33 "	0	0.029 "

Creatine Tolerance Test

Fasting	0.9 mg./100 ml.	0.5 mg./100 ml.
1 hour after creatine	1.4 " " "	1.6 " " "
2½ hours " "	0.9 " " "	1.5 " " "

Acromegaly.—A woman of 53 with five children noticed enlargement of and pain in the hands for 17 years before admission. The feet enlarged four years later. A diagnosis of acromegaly was made eight years ago. Menstruation ceased five years ago. On examination there were typical acromegalic features. The eye fields were full, but there was a central scotoma in the right eye. The blood pressure was slightly raised, but no other abnormal physical signs were found. Radiologically the jaw was typically acromegalic and the pituitary fossa a little deep. Hormonal therapy was adopted because symptoms had become static.

Urine				
Day	Volume	Creatinine	Creatine	Guanidoacetic Acid
1	2,200 ml.	1.58 g.	0.31 g.	0.075 g.
2	1,050 "	0.90 "	0.08 "	0.078 "
3*	3,208 "	2.16 "	0.12 "	0.107 "
4	2,010 "	1.41 "	0.20 "	0.097 "
5	2,050 "	1.44 "	0.21 "	0.109 "

Creatine Tolerance Test

Fasting	1.4 mg./100 ml.	1.35 mg./100 ml.
1 hour after creatine	2.0 " " "	1.5 " " "
2½ hours " "	1.5 " " "	1.75 " " "

Mixed "Dermoid" Tumour.—A man of 58 complained of failure of vision for four years and of impotence for two years. No headache or vomiting. Physical examination showed optic atrophy and perception of hand movements only on the right side, and on the left side a temporal hemianopia. The skin was smooth and of

* Creatine tolerance test (1 g. by mouth) on this day.

fine texture, and no axillary hair was present. The testes were small. Radiologically there was gross enlargement of the sella. Operation revealed a cystic mass in the pituitary, containing sebaceous-looking material. Histology showed this to be a "dermoid" tumour.

Urine				
Day	Volume	Creatinine	Creatine	Guanidoacetic Acid
1	905 ml.	0.65 g.	0.08 g.	0.053 g.
2	910 "	0.78 "	0.07 "	0.065 "
3*	502 "	0.43 "	0.24 "	0.032 "
4	435 "	0.49 "	0.32 "	0.035 "
5	725 "	0.68 "	0.20 "	0.042 "

Creatine Tolerance Test

Fasting	0.9 mg/100 ml.	0.7 mg./100 ml.
1 hour after creatine	1.25 " " "	1.75 " " "
2½ hours " "	1.12 " " "	1.75 " " "

Obesity with Amenorrhoea.—A married woman of 35, whose third and last child was 8 years old, gave one year's history of increasing obesity, having gained 4 stone. Menstruation was almost absent. She was somewhat drowsy and there was increased sweating. There was no headache or visual symptoms. There was some temporal pallor of both optic discs, but no other abnormal physical signs were found. The B.M.R., blood count, C.S.F., and urinary ketosteroids were all normal. The blood sugar curve was normal; radiological examinations were also negative.

Urine				
Day	Volume	Creatinine	Creatine	Guanidoacetic Acid
1	1,510 ml.	1.12 g.	0.24 g.	0.108 g.
2	1,660 "	1.24 "	0.35 "	0.136 "
3*	1,224 "	1.40 "	0.55 "	0.105 "
4	1,240 "	1.07 "	0.30 "	0.098 "
5	1,870 "	1.20 "	0.11 "	0.104 "

Creatine Tolerance Test

Fasting	0.8 mg./100 ml.	1.3 mg./100 ml.
1 hour after creatine	1.0 " " "	1.7 " " "
2½ hours " "	0.8 " " "	1.4 " " "

Thyrotoxicosis.—A single woman of 28 complained of mental and bodily fatigue for three years, and scanty periods for one year. There was also a complaint of proneness to ocular fatigue. Examination revealed no evidence of myasthenia or other

* Creatine tolerance test (1 g. by mouth) on this day.

neurological disorder. There was evidence of vasomotor instability. The pulse rate was between 80 and 90; the B.M.R. was slightly raised (+20%). Clinically she was considered to have a mild thyrotoxicosis, and under treatment she improved in hospital and after discharge.

Urine				
Day	Volume	Creatinine	Creatine	Guanidoacetic Acid
1	1,900 ml.	1.42 g.	0.48 g.	0.047 g.
2	2,250 "	1.47 "	0.40 "	0.041 "
3*	2,530 "	1.73 "	0.21 "	0.055 "
4	1,190 "	0.95 "	0.42 "	0.061 "
5	1,700 "	1.32 "	0.23 "	0.065 "

Creatine Tolerance Test

Fasting	0.85 mg./100 ml.	0.65 mg./100 ml.
1 hour after creatine	1.25 " " "	0.77 " " "
2½ hours " "	1.3 " " "	0.65 " " "

* Creatine tolerance test (1 g. by mouth) on this day.

APPENDIX 2

This appendix contains a brief clinical abstract of the cases recorded in Figs. 2, 3, and 4.

Fig. 2 (13106/49. N.H.).—A woman of 32 with a history of amenorrhoea and headaches for 17 years, and poor vision for nine months.

Examination showed a well-built, normal-looking woman. There was primary optic atrophy and a complete temporal field loss on the right side only. There was also an early loss in the upper and outer part of the left field. The uterus and ovaries were small, but there were no other abnormal physical signs. A radiograph of the skull showed considerable enlargement of the pituitary fossa. Operation revealed a typical chromophobe adenoma (verified histologically). The tumour was removed and there was uneventful recovery.

Fig. 3 (7371/48. N.H.).—A woman of 42 whose sole complaint was of failing vision for 3½ months.

Examination showed a middle-aged, alert woman with unmistakable acromegalic features. There was a large head, massive maxillae, and slightly prognathous jaw. There was little soft tissue hypertrophy. Hands and feet were all large and broad. The skin was smooth and of a fine texture. In the eyes both fundi showed definite pallor of the discs, and some bitemporal scotomata were present. The blood pressure was raised, but no other abnormal physical signs were found. Radiologically the pituitary fossa was enlarged.

At operation a pituitary acidophil adenoma (confirmed histologically) was found and partially removed.

Fig. 4 (31707/49. W.H.).—A woman of 50 with a six months' history of tremor. There was said to be no loss in weight.

Examination revealed a middle-aged, rather thin woman with bilateral exophthalmos, with tremor of the outstretched hands, but no obvious sweating. The thyroid showed a diffuse soft enlargement. The pulse rate varied from 90 to 100 a minute, but the blood pressure was normal. B.M.R. +75% ; blood count normal ; blood cholesterol 197 mg. per 100 ml. Thiouracil produced considerable improvement in the clinical condition apart from the exophthalmos. There has been a gain in weight of 4 lb. to 8 st. 10½ lb.