STUDIES ON SERUM GLUTAMIC PYRUVIC TRANSAMINASE IN HEALTH AND IN DISEASES OF THE LIVER *

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It has been reported that serum transaminase activity is found to be altered in certain pathological conditions which are associated with necrosis or other type of cellular damage of cardiac, hepatic or skeletal muscle tissues (5, 6 and 12). It may be that damage to the tissue results in release, from the cells, of the transaminase, which finds its way into the plasma or serum. Wroblewski and La Due (14) studied and compared the activities of glutamic oxaloacetic transaminase (GOT) and glutamic pyruvic transaminase (GPT) in different tissue homogenates. The activity of GPT was found to be relatively greater in liver than in other tissues as compared with the activity of GOT. This would perhaps suggest that the serum glutamic pyruvic transaminase (SGP-T), might possibly be a more specific index of liver cell damage than the serum glutamic oxaloacetic transaminase (SGO-T), because of its selective concentration in liver tissue. Futhermore it might be suggested that since the cardiac tissue activity of GPT is low, myocardial necrosis might not be associated with significant alterations of SGP-T activity. A study was undertaken to find out the levels of SGP-T activity in patients with diseases of the liver, in whom other liver function tests were simultaneously carried out. Normal subjects were also studied for their serum transaminase activity, as a control group for comparison. The results of this study are reported herewith.

MATRIALS AND METHODS

The techinque for analysing the activity of SGP-T is based on the transamination of alanine and α -ketogularate, as shown in the following reaction :---

α -Ketoglutarate + alanine \rightarrow pyruvate+glutamic acid.

Thus the above reaction catalysed by the transaminase results in the formation of a keto-The pyruvate formed is allowed to react with 2-4 dinitrophenylhydrazine. This is then acid. extracted with alkali to give a colour. The optical density of the colour is measured on a spectrophotometer at 505 mµ. (8). The standard calibration curves are prepared for various concentrations within the appropriate range of the standard pyruvate solution. The activity is expressed in terms of SGP-T units/ml. of serum, which is defined as the activity by 1.0 ml. of serum that results in the formation of chromogenic material equivalent to 1 mg. of pyruvate under conditions of the test (8). One hundred and two normal healthy adult male subjects ranging in their age between 30 and 55 years were studied of their serum transminase activity. Each of the estimations was done in duplicate and the average was taken thereof. Similarly fifty male patients with diseases of the liver whose diagnosis has been given separately (Table VII) and ranging, in their age from 39 to 52 years, were similarly investigated for their serum transaminase activity. These were the patients from the S.S.G. Hospital, Baroda and Infectious Diseases Hospital. Other liver function tests, namely, Van den Bergh test and Icteric index test were also performed simultaneously in the samples of sera obtained from these patients (3). A comparision has been made between the results obtained of the SGP-T test on one hand and those of the above two

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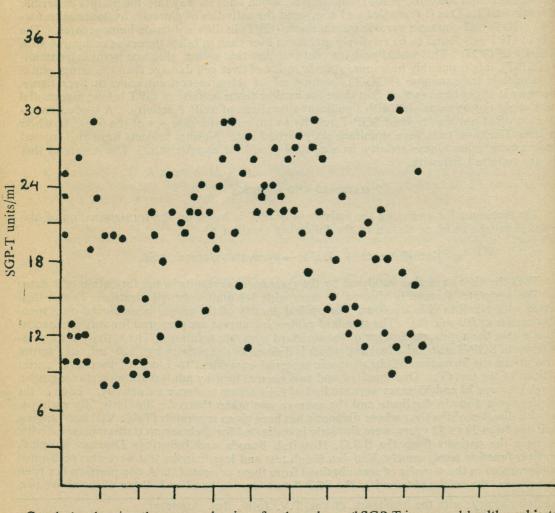
liver function tests performed on the other hand. These detailed results are shown in Tables, VI, VII, VIII, and IX and are represented in graphs.

RESULTS AND DISCUSSION

The values obtained for the serum transaminase (SGP-T) activity in the normal subjects ranged from 8.0 to 34.0 units/ml. with the mean value of 18.9 units/ml. The standard diviation value was found to be 6.4 and the standard error of mean was 0.63 (Table I).

The values obtained in the present series have been found to be well comparable with those given by other workers (Table II).

A similar comparison of the values for SGP-T in patients with liver diseases, has been shown (Table III).



Graph 1-showing the scattered points for the values of SGP-T in normal healthy subjects.

TABLE I

Showing the values for SGP-T in normals and in patients with liver diseases

| No. of normal subjects | SGP-T units/ml | | | | | |
|-------------------------------------|----------------|-----------------|------|------------|--------------|--|
| No. of normal subjects | Range | Mean | S.D. | S.E. of me | an | |
| 102 | 8.0-31.0 | 18.9 | 6.4 | 0. | 63 | |
| No. of patients with liver diseases | | SGP-T units/ml. | | | | |
| not of patients with not diseases | | Range | Mean | S.D. | S.E. of mean | |
| 50 | 1.00 | 42.0-229.0 | 79.4 | 25.4 | 3.59 | |

TABLE II

Showing the comparison of the values for SGP-T in normal healthy subjects as given by different workers

| Authors | No. of | SGP-T units/ml | | | |
|-------------------------------|----------|----------------|------|-------|--|
| Aumors | subjects | | Mean | S.D. | |
| 1. Chinsky (2) | | 3.0-30.0 | | · ··· | |
| 2. Reitman and Frankel (8) | 22 | 5.0-30.0 | | | |
| 3. Wroblewski and La Due (14) | | 5.0-35.0 | 16.0 | | |
| 4. Albaum et al (1) | 26 | | 17.0 | 1.0 | |
| 5. Sacks et al (10) | 20 | 5.0-35.0 | 21.3 | | |
| 6. Present series | 102 | 8.0-31.0 | 18.9 | 6.4 | |

TABLE III

Showing the comparison of the values for SGP-T in patients with liver diseases as given by different workers

| ter at optimitie togeneiter staan op at sel | NE of | SG | SGP-T units/ml | | | |
|---|------------------------|--------|----------------|------|--|--|
| Author | No. of subjects | Range | Mean | S.D. | | |
| 1. Mosley et al (7) | 13 | 26-260 | | | | |
| 2. Wroblewski and La Due (14) | Analysis . differen | 24—224 | | | | |
| 3. Wroblewski (13) | T-DD S. Marken | 26—258 | de las este | | | |
| 4. Rymenant et al (9) | All have A Clifference | 45—350 | | | | |
| 5. Present series | 50 | 42—229 | 79.4 | 25.4 | | |

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These patients were studied for their SGP-T levels during the onset of the disease before any treatment was started and during their stay in the wards of the Hospital, where they were admitted for their complaints and for further investigations. Two other liver function tests, name ly, Van den Bergh reaction and Icteric index, were also simultaneously performed in the samples of their sera. The results of these tests are shown in Tables IV and V.

TABLE IV

Showing values of SGP-T with their range, mean, S.D. and S.E. of mean in patients with liver diseases, investigated with Van den Bergh reaction

| | | Group I | Group I | I Group III |
|-----------------|------------------------------|----------------------|----------------------|----------------------|
| No. of patients | | 18 | 19 | 13 |
| S.GP-T units/ml | Mean S.D. S.E. of mean | 99.6 39.9 7.05 | 73.2 10.8 2.48 | 60.6 13.8 3.83 |
| Grp. I—Biphasic | Grp. II—Delayed | Grp. III—Imme | ediate | |

TABLE V

Showing the values of SGP-T with their range, mean, S.D. and S.E. of mean, in patients with liver diseases, investigated with Icteric index test

| | | Group I | Group II | Group III | Group IV |
|-----------------|------------------------------|----------------------|----------------------|----------------------|------------------------|
| No. of patients | | 8 | 23 | 13 | 6 |
| SGP-T units/ml | Mean S.D. S.E. of mean | 57.1 12.7 4.49 | 70.3 11.7 2.44 | 84.5 17.1 4.74 | 133.3 13.7. 5.59 |
| | Grp I | Icteric index | 12—25 | | |
| | Grp II | ,, ,, | 26—38 | | |
| | Grp III | 33 33 | 39—50 | | |
| | Grp IV | ,, ,, | 51 and al | bove | |

STATISTICAL ANALYSIS OF THE DATA

On comparing the data for SGP-T values obtained in normals with that obtained in patients, it was found that the value for the d.statistic was 14.00. When the data for the SGP-T test in patients was compared with that of the liver function test, namely, the Van den Bergh reaction, the value for the variance ratio F was found to be 46. When the data for the SGP-T levels obtained in the patients was compared with that of the other liver function test, namely, Icteric index test, the corresponding value of F was found to be 42. All these above values of d-statistic and F were found to be very highly statistically significant, even at 0.1% level (p=0.001).

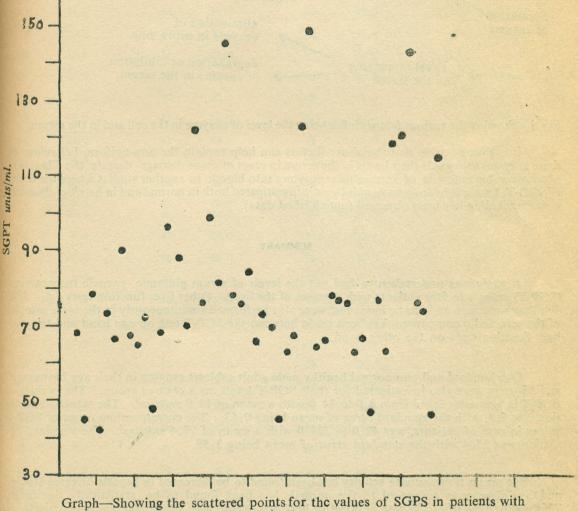
Marked alterations in the serum levels of GO-T and GP-T have been observed during cardiac, hepatic and other muscular diseases (13, 4, and 11). Wrobleski *et al* (14) reported relatively greater activity of GP-T in liver tissue than in other tissues, as compared with the activity of GO-T. The principle of the use of the estimations of the levels of these enzymes in serum, which are generally found to be increased, as diagnostic tests, primarily in the diseases of the damage to the

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tissues of cardiac, liver or skeletal muscle, has been based on the postulated release of these enzymes into the blood stream from the damaged cells of the respective tissues. It is, therefore, usually believed that SGP-T is more sensitive than SGO-T as an index of hepatocellular damage, and also it is further believed to be more specific for hepatic necrosis. But this has not been found to be nniformly true. It has been thus reported that the SGP-T assay, although more specific for hepatic disease as compared to myocardial necrosis, is neither specific for hepatic nectosis nor necessarily more sensitive than SGO-T as an index of hepatic injury (4, 2).

Variations of enzyme activity in serum following the damage to the tissues like liver etc. are not to be interpreted solely as consequences of enzyme leakage from the necrotic tissue cells. These changes should rather be regarded as a more general phenomenon. A commplex reaction of the organism sets in after the damage of the above type, as it does after many other acute diseases. It is this complex reaction that gives rise to a number of symptoms and also that causes



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the changes in enzyme activities. Thus there will be several factors influencing the enzyme level both in the cell and in the serum. The permeability of cell membrane, the enzyme elimination in urine and bile, degradation or inhibition of the enzyme in the serum, may be some of these factors influencing the enzyme level in the serum. These factors have been summarized in the following Fig. (1.)

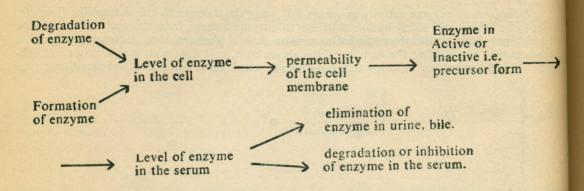


Fig. 1-showing the various factors influencing the level of enzyme in the cell and in the serum.

It is thus possible that the above factors can help explain the non-uniform behaviour of these enzymes as regards their levels in the serum in cases of tissue damage despite the liberation of considerable amounts of intracellular enzymes into blood. In another studies where the levels of SGO-T, Lactic dehydrogenase, aldolase are investigated both in normal and in hepatic diseases, a similar finding has been observed (unpublished data).

SUMMARY

A study was undertaken to find out the levels of serum glutamic pyruvic transaminase (SGP-T) activity in fifty patients with diseases of the liver. Other liver function tests e.g. Van den Bergh reaction and icteric index test were also performed simultaneously in the same samples of the sera and a comparison has been made between the SGP-T test on one hand and the two liver function tests on the other hand.

One hundred and two normal healthy male adult subjects ranging in their age between 30 and 55 years were also investigated for their SGP-T levels, as a control group. The values of SGP-T in normals ranged from 8.0 to 31.0 with a mean of 18.9 *units/ml*. The standard deviation was 6.4, with the standard error of mean being 0.63. The corresponding range of SGP-T values in case of patients, was 42.0 to 229.0 with a mean of 79.4 *units/ml*. The standard deviation was 25.4 with the standard error of mean being 3.59.

The serum transaminase activity has been found to be increased in hepatic diseases studied and the difference as compared to the normals has been found to be statistically significant. This was particularly so in case of infective hepatitis, where high serum transaminase activity was found to be present. The significance of the above findings has been discussed in the paper. Volume 10 Number 3

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| | Showing | values . | for SGP-T (units/m | l.) in no | ormal healthy subjec | ts | |
|------------|-----------------|------------|--------------------|------------|----------------------|------------|-----------------|
| Sr. No. | SGP-T units/ml. | Sr. No. | SGPT- units/ml. | Sr. No. | SGP-T units/ml | Sr. No. | SG-PT units/ml. |
| 1 | 25.0 | 31 | 18.0 | 61 | 24.0 | 91 | 12.0 |
| 2 | 10.0 | 32 | 11.0 | 62 | 27.0 | 92 | 18.0 |
| 3 | 23.0 | 33 | 25.0 | 63 | 23.0 | 93 | 31.0 |
| 4 | 20.0 | 34 | 22.0 | 64 | 22.0 | 94 | 9.0 |
| 5 | 12.0 | 35 | 13.0 | 65 | 26.0 | 95 | 11.0 |
| 6 | 13.0 | 36 | 21.0 | 66 | 22.0 | 96 | 30.0 |
| 7 | 10.0 | 37 | 20.0 | 67 | 27.0 | 97 | 17.0 |
| 8 | 12.0 | 38 | 22.0 | 68 | 28.0 | 98 | 10.0 |
| 9 | 26.0 | 39 | 23.0 | 69 | 20.0 | 99 | 12.0 |
| 10 | 12.0 | 40 | 22.0 | 70 | 17.0 | 100 | 16.0 |
| 11 | 10.0 | 41 | 24.0 | 71 | 27.0 | 101 | 25.0 |
| 12 | 19.0 | 42 | 14.0 | 72 | 29.0 | 102 | 11.0 |
| 13 | 29.0 | 43 | 22.0 | 73 | 22.0 | | |
| 14 | 23.0 | 44 | 20.0 | 74 | 26.0 | | |
| 15 16 | 8.0 20.0 | 45 46 | 19.0 24.0 | 75 76 | 14.0 29.0 | | |
| 17 | 9.0 | 47 | 26.0 | 77 | 20.0 | | |
| 18 | 20.0 | 48 | 29.0 | 78 | 15.0 | | |
| 19 | 8.0 | 49 | 29.0 | 79 | 18.0 | | |
| 20 | 14.0 | 50 | 20.0 | 80 | 23.0 | | |
| 21 | 23.0 | 51 | 16.0 | 81 | 14.0 | | |
| 22 | 10.0 | 52 | 27.0 | 82 | 12.0 | | |
| 23 | 9.0 | 53 | 25.0 | 83 | 14.0 | | |
| 24 | 10.0 | 54 | 11.0 | 84 | 13.0 | | |
| 25 | 10.0 | 55 | 28.0 | 85 | 20.0 | | |
| 26 | 15.0 | 56 | 26.0 | 86 | 11.0 | 8 | |
| 27 | 9.0 | 57 | 22.0 | 87 | 21.0 | | |
| 28 | 22.0 | 58 | 25.0 | 88 | 10.0 | | |
| 29 | 20.0 | 59 | 24.0 | 89 | 18.0 | | |
| 30 | 12.0 | 60 | 22.0 | 90 | 22.0 | | |

TABLE VI

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TABLE VII

| | Showing SGP-T units/ml. in patients with hepatic diseases | | | | | | |
|----------------------|---|----------------------------------|----|----------------------|----------|----------------------------|--|
| Sr. No. units/ml. | SGPT- | Disease | | Sr. No. units/ml. | SGP-T | Disease | |
| 1 | 75 | Inf. hepatitis | 13 | 26 | 66 | Obst. jaundice | |
| 2 | 73 | " | | 27 | 73 | Cirrhosis | |
| 3 | 68 | Cirrhosis | | 28 | 70 | Inf. hepatitis | |
| 4 5 | 45 78 | Obst. jaundice Inf. hepatitis | | 29 30 | 45 63 | Obs. jaundice Cirrhosis | |
| 6 | 42 | Obst. jaundice | | 31 | 67 | " | |
| 7 | 73 | Cirrhosis | | 32 | 123 | Inf. hepatitis. | |
| 8 | 66 | Obst. jaundice | | 33 | 229 | Inf. " | |
| 9 | 90 | Inf. hepatitis | | 34 | 64 | Obst. jaundice | |
| 10 | 67 | >> | | 35 | 66 | " | |
| 11 | 65 | Obst. jaundice | | 36 | 78 | ** | |
| 12 | 72 | Cirrhosis | | 37 | 77 | Cirrhosis | |
| 13 | 48 | Obst. juandice | | 38 | 76 | Obst. jaundice; | |
| 14 | 68 | Cirrhosis | | 39 | 63 | Cirrhosis | |
| 15 | 96 | Inf. hepatitis | | 40 | 66 | " | |
| 16 | 88 | Cirrhosis | | 41 | 47 | | |
| 17 | 70 | Inf. hepatitis | | 42 | 77 | ** | |
| 18 | 122 | " 0.0 | | 43. | 63 | Inf. hepatitis | |
| 19 | 76 | Cirrhosis | | 44 | 119 | " | |
| 20 | 99 | ,, | | 45 | 121 | " | |
| 21 | 81 | Obst. jaundice | | 46 | 143 | " | |
| 22 | 145 | Inf. hepatitis | | 47 | 76 | Cirrhosis | |
| 23 | 78 | Cirrhosis | | 48 | 74 | Inf. hepatitis | |
| 24 | 76 | ,, | | 49 | 46 | Obst. jaundice | |
| 25 | 84 | | | 50 | 115 | Inf. hepatitis. | |

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Showing SGP-T units/ml. in patients with hepatic diseases

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| Relationship of SGP-T (units/ml.) with liver function test Van den Bergh reaction | | | | | | | |
|---|------------------|---------|-----------------|-----------|-----------------|--|--|
| Gr | Group I Group II | | ıp II | Group III | | | |
| Big | Biphasic Delayed | | yed | Immediate | | | |
| Sr. No. | SGP-T units/ml. | Sr. No. | SGP-T units/ml. | Sr. No. | SGP-T units/ml. | | |
| 1 | 75 | 1 | 68 | 1 | 45 | | |
| 2 | 73 | 2 | 73 | 2 | 42 | | |
| 3 | 78 | 3 | 66 | 3 | 51 | | |
| 4 | 90 | 4 | 72 | 4 | 48 | | |
| 5 | 76 | 5 | 68 | 5 | 81 | | |
| 6 | 96 | 6 | 88 | 6 | 66 | | |
| 7 | 70 | 7 | 76 | 7 | 45 | | |
| 8 | 122 | 8 | 99 | 8 | 64 | | |
| 9 | 145 | 9 | 78 | 9 | 78 | | |
| 10 | 70 | 10 | 76 | 10 | 76 | | |
| 11 | 123 | 11 | 84 | 11 | 66 | | |
| 12 | 229 | 12 | 73 | 12 | 46 | | |
| 13 | 63 | 13 | 63 | 13 | 66 | | |
| 14 | 119 | 14 | 67 | | | | |
| 15 | 121 | 15 | 77 | | | | |
| 16 | 143 | 16 | 47 | | | | |
| 17 | 74 | 17 | 77 | | | | |
| 18 | 115 | 18 | 63 | | | | |
| | | .19 | 76 | | | | |

TABLE VIII

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