OBSERVATIONS ON THE PROTEOLYTIC ACTIVITY IN VITRO AT
NEUTRAL REACTION OF GASTRIC JUICE FROM
PATIENTS WITH SPRUE

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Previous investigators (1) have demonstrated
the presence in normal human gastric juice of a
protease which is active at neutral reaction. Cer-
tain properties were exhibited which distinguished
the enzyme from pepsin and trypsin. Subsequent
studies (2-4) showed that the protease and the
so-called intrinsic factor of Castle possessed cer-
tain identical properties: both are removed by ad-
sorption with Lloyd’s reagent, both are unable to
penetrate a semi-permeable membrane, they are
not destroyed by Berkefeld filtration or exposure
to alkali, but are destroyed by exposure to heat
and are inhibited by an environment more acid
than pH 3.5. Furthermore, it was found that this
protease activity was absent or greatly diminished
in patients with Addisonian pernicious anemia,
but was normal in patients with pernicious tapeworm
anemia.

The pathogenesis of the macrocytic anemia in
sprue has aroused much speculation. Because it
seemed important to determine whether or not
the intrinsic factor was contained in the gastric
juice of this disease, and because the gastric pro-
tease which is active at neutral reaction may be
identical with Castle’s intrinsic factor, it was de-
cided to measure gastric protease activity at neu-
tral reaction in sprue.

METHODS

The method was similar to that previously described
(1), except for modifications noted below. Gastric juice
was collected free from bile after stimulation with his-
tamine and promptly filtered through gauze. The juice
was brought to pH 10 with sodium hydroxide and kept
for 30 minutes at 40° C, after which it was brought to
pH 7.4. To 50 ml. of such materials were added 50 ml.
of a 1 per cent neutral casein solution, the mixture ad-
justed to pH 7.4 and 2 ml. of toluol added. The mixture
was incubated at 37.5° C for 24 hours, the pH remaining
constant. The procedures used for the formol titration
of amino nitrogen and for the determination of total fil-
trable nitrogen were those previously described (1, 2).
Samples of the digests were taken immediately, at four
hours, and 24 hours for formol titration. At the same
time intervals, samples were removed and analyzed for
the total amount of nitrogen not precipitable by 10 per
cent trichloroacetic acid. The amount of total filtrable
nitrogen and of amino nitrogen produced in 24 hours
was calculated from these determinations. Proteolytic activity
was judged from increases in filtrable nitrogen in milli-
grams per 100 ml. digest. The absence of active trypsin
was shown by the lack of a significant increase in amino
nitrogen.

RESULTS

From a large number of experiments the follow-
ing facts were confirmed:

(1). The effect of incubation of normal human
gastric juice with casein solution at pH 7.4. The
fasting contents from normal subjects were re-
moved and discarded. Histamine phosphate 0.5
mgm. was injected intramuscularly and the gastric
secretion collected during the next hour. Certain
physical changes occurred when such gastric juice
was incubated with an equal quantity of 1 per cent
casein solution at pH 7.4. The digestion mixtures
became chalky white. The trichloroacetic acid
filtrates from these serial samples showed pro-
gressive increases in turbidity. Increases in the
total nitrogen in the trichloroacetic acid filtrates
occurred along with these physical changes.

Peptic activity is not manifest at neutral reac-
tions, and above pH 8 pepsin is destroyed rapidly
by OH ions. Furthermore, it has been shown (5)
that exposure to alkali at pH 10 at 40° C for 30
minutes destroys from 70 to 80 per cent of trypsin
in solution. There was no significant increase in
amino nitrogen in our experiments. This sug-
gests that trypsin and other erepsin-like enzymes
of duodenal origin did not account for increase in

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filtrable nitrogen. The results of ten experiments on normal controls, which agree with previous reports, are shown in Table I.

That infection may have an inhibitory effect is suggested from determinations made on gastric juice obtained from patients at the time of an acute respiratory illness and following convalescence. The results of two such experiments are included in Table I. This would indicate that the gastric protease is inhibited during an acute illness and that the inhibitory factor disappears following recovery.

(2) The effect of incubation of human gastric juice, collected from sprue, with casein solution at pH 7.4. Collections of gastric juice from sprue patients and treatment with alkali before mixing with casein for digestion were performed in a fashion similar to that used on normal controls. The results of 11 of these experiments on sprue are shown in Table II. The gastric juice obtained from cases of sprue in remission showed progressive increases in filtrable nitrogen which compared closely to results in normal controls. On the other hand, the gastric juice from cases of sprue in relapse, with characteristic diarrhea, meteorism, steatorrhea, weight loss, glossitis, and anemia, showed a greatly diminished production of total filtrable nitrogen. Also, it is noteworthy that there was no significant increase in amino nitrogen.

<table>
<thead>
<tr>
<th>Experiment number</th>
<th>Preparation and reference to clinical observation</th>
<th>Increase in nitrogen in filtrate (mgm. per 100 ml. digest)</th>
<th>Increase in amino nitrogen by formol titration (mgm. per 100 ml. digest)</th>
</tr>
</thead>
<tbody>
<tr>
<td>30</td>
<td>Sprue-relapse gastric juice</td>
<td>12.0 13.6 0.0 0.4</td>
<td></td>
</tr>
<tr>
<td>31a*</td>
<td></td>
<td>7.4 23.2 0.2 0.0</td>
<td></td>
</tr>
<tr>
<td>32</td>
<td>After incubation pH 10</td>
<td>12.0 21.0 0.6 1.6</td>
<td></td>
</tr>
<tr>
<td>33</td>
<td>pH 10 C for 30 minutes</td>
<td>2.7 22.6 0.4 0.8</td>
<td></td>
</tr>
<tr>
<td>34</td>
<td></td>
<td>4.8 14.3 0.2 1.4</td>
<td></td>
</tr>
<tr>
<td>31b*</td>
<td>Sprue-remission sian-gastric juice</td>
<td>40.5 47.6 0.0 0.4</td>
<td></td>
</tr>
<tr>
<td>35</td>
<td></td>
<td>49.7 59.0 0.0 1.4</td>
<td></td>
</tr>
<tr>
<td>36</td>
<td>After incubation</td>
<td>21.4 55.0 0.0 0.7</td>
<td></td>
</tr>
<tr>
<td>37†</td>
<td></td>
<td>18.2 36.7 1.6 1.8</td>
<td></td>
</tr>
<tr>
<td>38</td>
<td>pH 10 40°C C for 30 minutes</td>
<td>32.0 49.0 0.2 0.8</td>
<td></td>
</tr>
<tr>
<td>39</td>
<td></td>
<td>22.5 48.6 0.8 1.2</td>
<td></td>
</tr>
</tbody>
</table>

* Samples of gastric juice 31a and 31b were obtained from the same donor. Sample 31b was obtained five months later during remission.
† Case 37 had a-chlorhydria.

In the majority of sprue patients, gastric secretion was obtained without difficulty in a one-hour period. During the stage of severe relapse there was usually hypochlorhydria and a marked reduction in rate of secretion and in proteolytic activity at neutral reaction. Such patients were observed again and tests repeated six months after treatment had produced remission, and they then showed a return to normal in all these functions. These data suggest that intrinsic factor activity may also be reduced in the gastric secretions of persons with sprue in relapse.

Attempts were made to do a biologic assay of the gastric secretions from these patients in relapse to determine the presence or absence of intrinsic factor. Our efforts to collect sufficient gastric juice for daily feeding experiments failed. However, previous investigators (6) have performed a biologic assay on gastric secretions from a case of sprue. Apparently, they demonstrated the absence of intrinsic factor from the gastric contents of a
case of sprue in relapse by feeding daily collections of gastric juice mixed with beef muscle to a case of Addisonian pernicious anemia.

It has been stated (7) that the deficiency in sprue may come about in a number of ways, among which are (1) a deficiency of extrinsic factor, (2) a defect of gastric digestion resulting from lack of intrinsic factor, and (3) defective absorption from the intestinal tract.

The mal-absorption from the small intestine in patients during relapse can be demonstrated with great frequency and certainly is a major cause of the sprue syndrome. Defective absorption of fats, fat-soluble vitamins, calcium, and carbohydrates can be commonly demonstrated. Patients with severe diarrhea will also show faulty assimilation of protein.

Dietary histories on our sprue patients have not shown any correlation between the ingestion of meat products, or other sources of extrinsic factor, and the incidence of anemia. When fed adequate amounts of beef muscle, the patients in relapse failed to obtain a remission. Subsequently, they made a satisfactory response to injectable liver extract or folic acid.

The data presented in this paper support the concept that a defect in gastric digestion may be important to the occurrence of macrocytic anemia in sprue; though whether such a defect is primary or secondary to gastro-intestinal mal-absorption and chronic debility can not be ascertained. Data reported here show that acute infection can lower the value of the proteolytic activity at neutral reaction to a very low level.

The fact that the anemia of sprue is macrocytic and improved by liver does not mean that a primary deficiency of gastric origin is present. This is well shown by the pernicious tapeworm anemia.

It can only be said that the protease activity of the gastric juice is lessened in sprue in relapse and normal in remission.

**CONCLUSIONS**

1. When equal quantities of normal human gastric juice and 1 per cent casein solution are incubated at 37.5° C and pH 7.4, there is a progressive increase in filtrable nitrogenous substances. The proteolysis was not considered to result from pepsin because the activity was maximal at pH 7.4. Also, since there was no significant increase in amino nitrogen within 24 hours, together with the persistent increase in total filtrable nitrogen after exposure to pH 10, it is considered that the proteolysis was not due to trypsin.

2. The proteolytic activity of the gastric enzyme at neutral reaction, is present to a normal degree in cases of sprue in remission. It is greatly diminished in cases of sprue in relapse. The absence of proteolytic activity in cases of pernicious anemia suggests that the deficiency of intrinsic factor so characteristic of pernicious anemia, may be present in sprue in relapse.

3. In two non-anemic normal subjects, it appears that infection inhibited this type of proteolysis to a considerable degree.

**BIBLIOGRAPHY**


6. Castle, W. B., Heath, C. W., and Strauss, M. B., Observations on the etiologic relationship of achylia gastrica to pernicious anemia. IV. A biologic assay of the gastric secretion of patients with pernicious anemia having free hydrochloric acid and that of patients without anemia, or with hypochromic anemia having no free hydrochloric acid, and of the role of intestinal impermeability to hematopoietic substances in pernicious anemia. Am. J. M. Sc., 1931, 182, 741.