NEW DEVELOPMENTS IN THE DIAGNOSIS AND TREATMENT OF PERNICIOUS ANEMIA*†

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CRITERIA for the diagnosis of pernicious anemia must be reëvaluated in light of the changing pattern of this disease. The profound physical abnormalities and extreme hematologic aberrations associated with classic severe pernicious anemia are not often encountered in modern practice. There are two reasons for this alteration in the character of the disorder. In the first place, patients tend to consult their physicians early, at a time when symptoms may seem inconsequential. The only complaint may be slight weakness or fatigue, or perhaps soreness of the tongue. There may be no detectable pallor at this time, and even if the hemoglobin concentration is determined, anemia may be so mild in degree as to arouse little The diagnosis at this stage is often overlooked. Second, patients concern. with the early symptoms of pernicious anemia are very likely to receive therapeutic preparations which in varying degree are effective in overcoming the manifestations of the disease before the diagnosis has been established. The very nature of the initial symptoms makes it likely that a multivitamin or hematinic preparation will be prescribed, or even procured by the patient without medical advice. Most proprietary vitamin preparations contain folic acid, often in addition to vitamin B12, intrinsic factor and a host of other substances. Administration of these preparations is followed by gratifying clinical improvement. Symptoms subside and anemia disappears. However, the need for adequate therapy for the duration of life is not recognized. Sooner or later relapse occurs, and often neurologic manifestations appear.1

In the past five years, 14 of the new cases of pernicious anemia seen at the Johns Hopkins Hospital have presented with crippling neurologic disease in the absence of an appreciable degree of anemia. In some of these the blood and bone marrow were entirely normal. All of these patients were seriously disabled; several were unable to stand or to control the bladder (figure 1). All have residual neurologic manifestations after prolonged and intensive parenteral therapy with vitamin B12. In most cases it was definitely established that the patient had been taking a vitamin preparation containing folic acid.

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NEW DEVELOPMENTS IN PERNICIOUS ANEMIA

The possibility of pernicious anemia should be considered in any adult patient with an unaccountably subnormal hemoglobin value. Careful examination of the blood will show macrocytosis if anemia is present. However, it is important to remember that red cell abnormalities are slight, and the marrow pattern may not be diagnostic when anemia is mild. If there is a favorable response to treatment with parenterally administered vitamin B₁₂, the diagnosis is reasonably well established. A history of anemia which responded to treatment with a vitamin or hematinic preparation suggests the possibility of pernicious anemia. Appropriate tests should be performed to exclude this disease in any patient who complains of soreness of the



FIG. 1. This 66 year old woman for months had been taking a proprietary multivitamin preparation which contained folic acid. She had never previously had symptoms of pernicious anemia, but while receiving multivitamins rapidly developed extremely severe subacute combined degeneration. She was found to have achlorhydria, and the alimentary absorption of radioactive vitamin B_{12} was markedly impaired. After months of intensive parenteral therapy with vitamin B_{12} she has regained some control of bladder function, but it seems apparent that she will remain a helpless invalid because of permanent damage to the nervous system. A few millionths of a gram of vitamin B_{12} , administered parenterally, could have prevented this incapacitating disorder.

tongue. Neurologic manifestations of subacute combined degeneration should always be considered as probably due to pernicious anemia, even though there are no hematologic abnormalities.

DIAGNOSIS OF PERNICIOUS ANEMIA

An extremely difficult problem is presented by the case in which the diagnosis of pernicious anemia was never definitely proved but in which therapy has been adequate to induce or maintain a remission. The blood and marrow are normal, and the demonstration of gastric achlorhydria makes the diagnosis acceptable but by no means establishes it. If therapy is withheld, relapse may not occur for several years, so that this is an unsatisfactory diagnostic test. A very helpful solution to this problem has become available with the use of tracer tests employing vitamin B₁₂ labeled with radioactive cobalt.²⁻⁸ A small amount of the tagged vitamin is given orally and the fraction which is absorbed from the intestinal tract is measured. In pernicious anemia, absorption is impaired because of the deficiency of intrinsic factor.

Since 1951 we have used radioactive vitamin B12 in studies of more than 100 individuals. When an appropriate oral dose is administered, the presence of an absorption defect is readily demonstrated. Impaired absorption is regularly found in pernicious anemia and may also be encountered following total gastrectomy, in sprue and in association with lesions of the small intestine. Impaired absorption was not demonstrated in normal subjects of older age groups or in patients with a variety of diseases unrelated to pernicious anemia. Achlorhydria per se was not associated with reduced vitamin B12 uptake. In most of our studies the amount of radioactive material absorbed was determined by measuring the residual radioactivity of the stools. This is a time-consuming and laborious process, requiring total stool collections for not less than six days. The method has the advantage of permitting very accurate measurement. Schilling ^o devised an ingenious technic in which the radioactive material absorbed from the intestine is flushed out into the urine by means of a parenteral injection of a large amount of inert vitamin B12. Radioactive measurements are then made on the urine rather than on the feces. This procedure is less precise, since all of the radioactive vitamin absorbed may not be excreted in the urine. It has the great advantage of simplicity, however, and the test is completed in only 24 hours. The Schilling test has been widely used, with extremely satisfactory results. We have found it to be a reliable clinical test of inestimable value in the diagnosis of pernicious anemia in patients who are in remission as a result of previous therapy. The procedure can easily be carried out in any laboratory in which there are facilities for radioactive measurements.

TREATMENT OF PERNICIOUS ANEMIA

When refined liver extract became available, completely satisfactory treatment for pernicious anemia was at hand. Patients adequately treated with parenteral liver preparations remained in complete remission throughout their lives. No form of treatment can accomplish more, and a number of therapeutic regimens frequently employed in recent years have accomplished much less. In particular, the use of folic acid in the past decade has in many instances permitted neurologic manifestations to develop while the blood remained normal.

The therapeutic effect of liver extract is attributable to the vitamin B12 which it contains. The results of treatment of pernicious anemia with vitamin B12 are no better than those obtained with liver extract. Some authors 10, 11 have reported that vitamin B12 does not provide complete replacement therapy in pernicious anemia, but the experience of other investigators does not support this contention.12 More than 50 patients with pernicious anemia in the Hematology Clinic of the Johns Hopkins Hospital have had no therapy other than vitamin B12 in the past six years and all remain in complete remission. We have not encountered the hypoprothrombinemia * and macrocytosis which have been described by others during treatment with vitamin B12.

Current interest in the treatment of pernicious anemia centers about the use of orally administered preparations. When vitamin B12 is combined with intrinsic factor, absorption of the vitamin is facilitated. Steady progress is being made in the purification of intrinsic factor, and preparations of considerable potency are now available. Several commercial preparations contain a mixture of vitamin B12 and a source of intrinsic factor for oral treatment of pernicious anemia. Preliminary observations indicate that

TABLE 1

Patients with Pernicious Anemia Treated with Orally Administered Vitamin B12

Remission induced by oral therapy	17
Remission induced and subsequently maintained by oral therapy	14
Remission maintained by oral therapy	14
Total	45

these are satisfactory, although careful observation of a large group of patients over a long period of time will be required to establish that these preparations are as reliable as is the parenteral injection of vitamin B12. It can be said with certainty that the amount of vitamin B12 which can be absorbed from the gastrointestinal tract under the most favorable conditions is far less than that which is customarily injected parenterally. Therefore, when intensive therapy is indicated, as in the patient with neurologic manifestations, parenteral therapy should always be used.

When very large amounts of crystalline vitamin B12 are given orally in the absence of intrinsic factor, satisfactory therapeutic responses may be obtained. The doses required for these effects are measured in milligrams rather than in micrograms.13, 14, 15 In 1950 we initiated an experimental study to determine whether orally administered vitamin B12 alone is adequate treatment for pernicious anemia.12 Forty-five patients have now been treated in this way † (table 1). Thirty-one were in hematologic

* Plasma prothrombin measured by the two-stage technic in 15 of our patients maintained for years on vitamin B₁₂ alone was well within the normal range. The lowest values obtained were higher than values obtained in some normal subjects. †The vitamin B₁₂ used in these studies was generously provided by Merck and Co.,

Inc., Rahway, New Jersey.

relapse when first treated. Most of these patients were hospitalized and placed on a diet deficient in vitamin B₁₂. After completion of the initial studies, a single oral dose of vitamin B₁₂, ranging from 3,000 to 10,000 μ g, was given in the morning with the patient fasting. After this single dose, additional therapy was withheld until it was clear that no further improvement was taking place. A number of patients were observed for more than a month before additional treatment was given. The clinical and hematologic responses to these large oral doses were in most instances entirely comparable to those seen after parenteral injection of 30 or more micrograms of the vitamin (figure 2). Soreness of the tongue and gastro-



FIG. 2. Hematologic response of a patient with pernicious anemia to a single oral dose of 5,000 μg of vitamin B₁₂.

intestinal symptoms subsided. In several cases neurologic manifestations improved. Several patients developed a complete hematologic remission after a single oral dose. Three patients responded suboptimally to oral therapy. Parenteral injection of the vitamin in each of these three cases failed to accelerate the hematologic improvement. In two the retarded response was associated with infection. The third patient was subsequently allowed to go into relapse and on a second trial responded well to orally administered vitamin B_{12} . In addition to the cases of classic pernicious anemia, one patient with megaloblastic anemia following total gastrectomy and one with a pernicious anemia-like syndrome associated with multiple

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Patient	Months -	Hematocrit	
		Initial	Present
C. J. 55 CF	56	14.0	51.0
E. T. 44 WF	39	13.7	42.0
W M 84 WM	30	22.0	45.8
T. L. 54 WM	27	26.0	45.1
E. W. 68 CF	27	24.0	43.0

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Patients Initially Treated and Maint	tained in Remission with Orally	Administered Vitamin B12*

* These patients were brought out of relapse by oral therapy. The initial dose ranged between 3,000 and 10,000 μ g of vitamin B₁₂. The maintenance dose in each case was 1,000 μ g given as a single tablet once a week.

diverticula of the jejunum also responded well to single oral doses of the vitamin.

Information currently at hand makes it seem probable that extremely high concentrations of vitamin B_{12} in the intestine are required if adequate amounts are to be absorbed regularly in the absence of intrinsic factor. A single dose of less than 1,000 µg appears to be suboptimal, and some patients have failed to respond to daily doses of as much as 250 µg.^{16, 17, 18} In attempting oral maintenance therapy, therefore, we decided to use a single large dose once a week, rather than smaller doses at daily intervals.

Of the 31 patients brought into remission by orally administered vitamin B_{12} , 14 have continued on oral therapy for periods ranging for from four months to almost five years. All 14 remain in complete remission, and eight have now been under continuous maintenance therapy for more than two years (table 2). The maintenance dose has been 1,000 µg of vitamin B_{12} , given as a single tablet once a week.

Patient	Months -	Hematocrit	
		Initial	Present
M. D. 63 WF	39	43.8	45.0
B. S. 73 CF	39	45.0	43.4
R. F. 49 WF	39	44.8	43.7
T. S. 50 WF	39	41.9	40.7
V. F. 48 CF	39	41.5	40.3
A. M. 56 WF	38	44.8	42.2
M. S. 67 WF	38	46.2	47.2
M. G. 70 CF	36	41.9	42.7

TABLE 3 Patients Maintained in Remission for More Than Three Years by Orally Administered Vitamin B₁₂*

* These patients, previously maintained on parenteral therapy, were in remission at the onset of oral therapy. All have remained in complete remission.

Fourteen additional patients had previously been maintained on parenteral therapy and were in remission at the time oral therapy was instituted. These patients have been receiving 1,000 μ g of vitamin B₁₂ orally once a week for 15 months or longer, and all remain in remission. Hematocrit values before and after oral maintenance therapy in those of this group treated for more than three years are shown in table 3.

The results of this and of comparable studies ¹⁰ indicate that patients with pernicious anemia can be satisfactorily treated by the oral administration of large amounts of vitamin B₁₂ in the absence of intrinsic factor or other adjuvants. Therapeutic effects appear to be as good as those obtained with oral preparations containing concentrates of intrinsic factor. The



FIG. 3. The concentrations of vitamin B_{12} in the serum of seven patients maintained for prolonged periods on 1,000 μ g of vitamin B_{12} orally once a week are compared with those of normal subjects and of patients with untreated and parenterally treated pernicious anemia. Blood for these determinations was drawn just prior to administration of a regularly scheduled dose. All of the treated patients were in clinical remission.

advantage of crystalline material is that it can be assayed by weight, whereas the potency of intrinsic factor preparations must necessarily be measured in terms of the response produced when administered to patients with pernicious anemia.

While these studies were in progress, Lear and his associates ²⁰ were determining the serum vitamin B_{12} concentrations of a large group of patients at the Boston City Hospital. We were fortunate to have their collaboration in measuring the vitamin B_{12} levels in the serum of seven of our patients who had been maintained for prolonged periods of time on 1,000 μg of B_{12} orally once a week. The results can be compared directly with those obtained in other patients on various parenteral dosage schedules

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(figure 3). It is clear that the amount of vitamin B_{12} absorbed from the intestine, even with these large doses, is not sufficient to restore the serum concentration to normal. Two patients had extremely low serum levels even though they appeared to be in complete clinical and hematologic remission. These data suggest that the oral dosage schedule employed, 1000 μ g once a week, is suboptimal, and that larger amounts would be required to restore tissue saturation. Evidence has been provided by others ^{14, 19, 21} that the concentration of vitamin B₁₂ in the serum of patients with pernicious anemia can be restored to normal levels if adequate oral doses are given.

It is important to emphasize that relatively little vitamin B_{12} can be absorbed from any orally administered preparation in contrast to the large amount which can be injected parenterally. Furthermore, the ability of various patients to absorb the vitamin differs. The complete effectiveness of parenteral therapy, which requires injections no more often than once a month, has been well established. At the present time it would seem wise for most patients with pernicious anemia to continue to receive parenteral therapy. However, if oral therapy is to be used, crystalline vitamin B_{12} alone in milligram doses appears to be as effective as smaller amounts of the vitamin combined with intrinsic factor.

SUMMARIO IN INTERLINGUA

Patientes monstrante le precoce manifestationes de anemia perniciose es frequentemente tractate con preparatos multivitaminic o hematinic que contine acido folic, mesmo ante le diagnose es definitemente establite. Per consequente, le curso clinic del morbo es alterate, e manifestationes neurologic pote resultar in le absentia de anemia. Le diagnose es frequentemente difficile a establir in patientes tractate in iste maniera. Le uso de tests a etiquettage con un forma radioactive de vitamina B_{12} es de specific valor diagnostic in tal casos.

Therapia parenteral con vitamina B_{12} remane le tractamento de selection, sed certe preparatos que es administrate per via oral pare therapeuticamente efficace.

In le presente studio 45 patientes con anemia perniciose esseva tractate con vitamina B_{12} in forma crystallin, administrate oralmente sin factor intrinsec. Quando 3 a 10 milles μ g de vitamina B_{12} esseva administrate per via oral e in un sol dose a patientes in stato de recidiva, le responsas clinic e hematologic esseva simile al responsas resultante del therapia parenteral. Il esseva possibile mantener le patientes in stato de remission complete per le administration de un sol tabletta con 1000 μ g B_{12} un vice per septimana. Per iste regime 11 patientes ha essite mantenite de maniera satisfactori durante periodos de plus que tres annos. Totevia, recente mesurationes del concentration seral de vitamina B_{12} in 7 de iste patientes indica que le nivellos seral que resulta de iste dosage es probabilemente suboptimal e que plus grande quantitates es requirite pro restaurar le saturation del texitos.

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