Evaluation of Macrocytosis

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Macrocytosis, generally defined as a mean corpuscular volume greater than 100 fL, is frequently encountered when a complete blood count is performed. The most common etiologies are alcoholism, vitamin B₁₂, and folate deficiencies, and medications. History and physical examination, vitamin B₁₂ level, reticulocyte count, and a peripheral smear are helpful in delineating the underlying cause of macrocytosis. When the peripheral smear indicates megaloblastic anemia (demonstrated by macro-ovalocytes and hypersegmented neutrophils), vitamin B₁₂ or folate deficiency is the most likely cause. When the peripheral smear is nonmegaloblastic, the reticulocyte count helps differentiate between drug or alcohol toxicity and hemolysis or hemorrhage. Of other possible etiologies, hypothyroidism, liver disease, and primary bone marrow dysplasias (including myelodysplasia and myeloproliferative disorders) are some of the more common causes. (Am Fam Physician. 2009;79(3):203-208. Copyright © 2009 American Academy of Family Physicians.)

Macrocystosis, defined as a mean corpuscular volume greater than 100 fL, occurs in approximately 3 percent of the general population.¹ Debate persists about the upper limit of normal values of mean corpuscular volume. In a recent study, 7 percent of patients had a mean corpuscular volume greater than 96 fL, and 1.7 percent had a mean corpuscular volume greater than 100 fL.² With the advent of automated complete blood cell counters and the increased use of certain medications, elevations in the mean corpuscular volume may be more commonly encountered. This article describes a strategy for the evaluation of patients with macrocytosis, as well as a brief discussion on treatment of vitamin B₁₂ and folate deficiencies.

Causes of macrocytosis and their prevalence in different populations are shown in Table 1.¹⁻³⁻⁵ Although it is associated with anemia, hypothyroidism is a more common cause in older persons than in other age groups. Results of a study in New York City indicated that medications for treating human immunodeficiency virus (HIV) infection have become a more prominent cause of macrocytosis. Alcoholism is the cause in as many as 80 percent of patients in some populations.⁶ Bone marrow biopsy was only performed in these studies when another cause could not be determined, and resulted in diagnosis in approximately one fourth of these remaining patients.¹ In recent years, an association has been found between Helicobacter pylori infection and vitamin B₁₂ deficiency.⁷

Pathophysiology

The causes of macrocytosis can be broadly classified as megaloblastic and nonmegaloblastic (Table 2).¹⁻³⁻⁵⁻⁷⁻¹⁶ Megaloblastic processes are characterized on the peripheral smear by macro-ovalocytes and hypersegmented neutrophils, which are absent in nonmegaloblastic macrocytic processes (Figure 1). Nonmegaloblastic processes have round macrocytes or macroreticulocytes (Figure 2). Because the mechanisms producing macrocytosis are not completely understood, the separation between megaloblastic and nonmegaloblastic causes is somewhat artificial. However, this concept remains useful for identifying the most predominant etiology for macrocytosis. For example, whereas the effect of alcohol is thought to be primarily a nonmegaloblastic process, in chronic alcoholism there may be concomitant vitamin B₁₂ or folate deficiency.¹⁷,¹⁸

In megaloblastic processes, erythrogenic precursors are larger than mature red blood cells (RBCs) because folate and vitamin B₁₂ deficiencies result in defective RNA and DNA syntheses. Serum elevations in homocysteine and methylmalonic acid result from defective biochemical processes in folate and B₁₂.
deficiencies, and could be used to clarify the cause of megaloblastic anemia, although this is not yet standard clinical practice. The Nonmegaloblastic processes develop from multiple mechanisms and have not been fully outlined. Macrocytosis can occur when there is increased RBC production secondary to peripheral blood cell destruction (i.e., hemolysis) or loss (i.e., hemorrhage), leading to a reticulocytosis. Reticulocytes are incompletely processed RBCs and, therefore, are slightly larger than the average RBC. The mechanisms by which the other diseases listed in Table 1 cause macrocytosis have not been fully explained.

**Diagnostic Strategy**

Once macrocytosis is identified, the history and physical examination help narrow the differential diagnosis. The presence of anemia, the degree of elevation of the mean corpuscular volume, and the patient’s overall health guide how aggressively the work-up progresses. At least some amount of investigation is warranted if the diagnosis is not readily apparent or if the patient is anemic (defined by the World Health Organization as a hemoglobin level less than 13 g per dL [130 g per L] in men and less than 12 g per dL [120 g per L] in women).

The algorithm outlined in Figure 3 suggests a work-up for macrocytosis. Physicians should begin by ordering a peripheral smear, a reticulocyte count, and a vitamin B₁₂ serum level for all patients with macrocytosis. It may be necessary to specifically order a reticulocyte index in some laboratories, which assesses if there is an adequate bone marrow response. Hemorrhage or hemolysis is the most likely cause if the reticulocyte count is elevated, but anemia recovery also causes an elevation in the

### Table 1. Prevalence of Major Causes of Macrocytosis in Studied Populations

<table>
<thead>
<tr>
<th>Etiology</th>
<th>Hospitalized patients in New York City¹ (%)</th>
<th>Outpatients in Finland¹ (%)</th>
<th>Finnish persons older than 75 years⁴ (%)</th>
<th>Finnish and American patients⁵ (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alcohol</td>
<td>26</td>
<td>65</td>
<td>15</td>
<td>36</td>
</tr>
<tr>
<td>B₁₂ and/or folate deficiency</td>
<td>6</td>
<td>9</td>
<td>28</td>
<td>21</td>
</tr>
<tr>
<td>Medications</td>
<td>37*</td>
<td>3</td>
<td>2</td>
<td>11</td>
</tr>
<tr>
<td>Hypothyroidism</td>
<td>—</td>
<td>1</td>
<td>12</td>
<td>5</td>
</tr>
<tr>
<td>Bone marrow dysplasias</td>
<td>6</td>
<td>1</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>Liver disease (nonalcoholic)</td>
<td>6</td>
<td>—</td>
<td>2</td>
<td>6</td>
</tr>
<tr>
<td>Reticulocytosis</td>
<td>8</td>
<td>—</td>
<td>—</td>
<td>7</td>
</tr>
<tr>
<td>Miscellaneous</td>
<td>3</td>
<td>21</td>
<td>13</td>
<td>7</td>
</tr>
<tr>
<td>Not established</td>
<td>7</td>
<td>—</td>
<td>22</td>
<td>12</td>
</tr>
</tbody>
</table>

**NOTE:** Etiologies listed from most to least common.

*¹—13 percent from zidovudine (Retrovir).

Information from references 1 and 3 through 5.
reticulocyte count. Measures of vitamin B₁₂ are a useful part of the initial work-up, because if vitamin B₁₂ deficiency is present but undiagnosed, folate repletion will correct the megaloblastic anemia, but not the possible neuropathic changes that occur with B₁₂ deficiency.¹⁹

Although uncommon, consider the possibility of spurious macrocytosis. This may be caused by cold agglutinins, hyperglycemia, or leukocytosis. Cold agglutinins cause the RBCs to clump, making them appear larger to the automatic counter.¹⁰ Hyperglycemic blood is more concentrated, and when it is diluted to measure the mean corpuscular volume, the cells swell more than usual, causing a false elevation.¹¹ Increased turbidity of a sample with marked leukocytosis also can cause the machine to overestimate the cell size.¹²

When the history and physical examination, peripheral smear, B₁₂ level, and reticulocyte count have not lead to an obvious diagnosis, consider a comprehensive metabolic panel to look for liver and kidney disease, thyroid-stimulating hormone for thyroid disorders, and methylmalonic acid and homocysteine levels to assess for vitamin B₁₂ deficiency, despite a normal vitamin B₁₂ level. If the cause remains elusive, consider again whether the degree of anemia or the patient’s overall health warrants referral to a hematologist for bone marrow biopsy, or search for rarer causes, keeping in mind that the most extensive work-up will result in a diagnosis in approximately 90 percent of patients.

Specific Causes of Macrocytosis

VITAMIN B₁₂ DEFICIENCY

Vitamin B₁₂ is absorbed by the ileum when it is bound by intrinsic factor, which is produced by the parietal cells of the gastric mucosa. In pernicious anemia, the loss of parietal cells leads to insufficient absorption of vitamin B₁₂, which then leads to vitamin B₁₂ deficiency over time. Pernicious anemia is most commonly caused by autoimmune atrophic gastritis, in which autoantibodies are directed against parietal cells and intrinsic factor. Less commonly, pernicious anemia can be caused by nonautoimmune gastritis secondary to H. pylori infections and Zollinger-Ellison syndrome.

Patients with vitamin B₁₂ deficiency may describe paresthesias related to peripheral neuropathy, poor or strict vegan diet, lack of socioeconomic resources,
bowel-related symptoms (including diarrhea), or a history of bowel surgery for weight loss. Findings on physical examination may include neurologic signs such as ataxia, decreased proprioception, and vibratory sensation. Patients may also have poor dentition or nonspecific oral stomatitis or glossitis.

Because pregnant women take folic acid routinely in prenatal vitamins, macrocytic anemia is much less common during pregnancy. Consider nitrous oxide abuse in at-risk populations, because nitrous oxide inactivates vitamin B\textsubscript{12} through oxidation. Other uncommon causes include *Diphyllobothrium latum* (i.e., fish tapeworm) infection or inherited disorders of cobalamin metabolism, including Imerslund syndrome (a congenital vitamin B\textsubscript{12} malabsorption associated with proteinuria). Only 10 percent of persons with vitamin B\textsubscript{12} deficiency are actually anemic.

The normal range for serum measures of vitamin B\textsubscript{12} varies among laboratories. If the vitamin B\textsubscript{12} level is borderline low (i.e., 100 to 400 pg per mL [74 to 295 pmol per L]), methylmalonic acid and homocysteine levels should be ordered and, if elevated, may provide evidence of B\textsubscript{12} deficiency. The Schilling test (i.e., measuring enteral absorption of vitamin B\textsubscript{12}) is not widely available at this time.

Oral therapy appears to be as effective as intramuscular therapy for the treatment of vitamin B\textsubscript{12} deficiency. Relapse of pernicious anemia occurs at a mean interval

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**Figure 3. Algorithm for the evaluation of macrocytic anemia.** (RBC = red blood cell; MMA = methylmalonic acid.)
of 65 months after cessation of treatment. It is important for patients to adhere to long-term therapy because the deficiency will recur if treatment is stopped, unless a reversible cause is identified.\(^{22}\)

**FOLATE DEFICIENCY**

The history of folate deficiency may mimic the history of vitamin B\(_{12}\) deficiency in regard to poor nutritional intake or absorption. In addition, 35 percent of patients with alcoholism and macrocytic anemia are folate-deficient, which can be caused by poor nutritional intake, malabsorption, hepatobiliary dysfunction, and possibly increased folate catabolism.\(^ {18,23}\) Some medications that are used to treat seizure disorders, cancer, and autoimmune diseases can lead to folate deficiency. For example, methotrexate directly inhibits dihydrofolate reductase, which leads to a functional folate deficiency.\(^ {14}\) Other medications that affect folate metabolism include 5-fluorouracil (Adrucil), hydroxyurea (Hydrea), pyrimethamine (Daraprim), trimethoprim/sulfamethoxazole (Bactrim, Septra), pentamidine (Pentam), and phenytoin (Dilantin).\(^ {24}\) Medications can also affect folate absorption, including metformin (Glucophage) and cholestyramine (Questran). Supplementing with folate may be necessary when treating a patient with such medications.\(^ {15}\)

Serum folate levels are not useful because they fluctuate rapidly with dietary intake and are not cost effective.\(^ {25}\) RBC folate levels more accurately correlate with folate stores and should be performed if folate deficiency is suspected.\(^ {26}\) In differentiating the cause of megaloblastic anemia, a methylmalonic acid level that is within normal range also points toward a diagnosis of folate deficiency, especially if the serum vitamin B\(_{12}\) level is within the normal range. Note that homocysteine levels will be elevated with vitamin B\(_{12}\) and folate deficiencies.

**HIV MEDICATIONS**

Treatment of HIV with reverse transcriptase inhibitors (e.g., stavudine [Zerit], lamivudine [Epivir], zidovudine [Retrovir]) will cause macrocytosis because they interfere with DNA production, which may lead to megaloblastic changes. Most patients with HIV who are being treated with reverse transcriptase inhibitors will display macrocytosis without anemia. This indicates medication compliance by the patient, and no treatment is necessary.\(^ {27}\) Table 3 lists other medications that may cause macrocytosis.\(^ {28}\)

**RETICULOCYTOsis (HEMOLYSIS OR HEMORRHAGE)**

For the patient’s history, questions that are pertinent to evidence of blood loss, whether chronic or acute, can point to reticulocytosis. As part of the family history, physicians should ask about the presence of certain hemato-logic syndromes, including sickle cell disease, hereditary spherocytosis, and glucose-6-phosphate dehydrogenase deficiency. In persons who do distance running, hemolysis from constant foot pounding has infrequently been shown to cause macrocytosis.\(^ {29}\) The physical examination may reveal hepatosplenomegaly in hemolysis or other physical manifestations of blood loss (e.g., conjunctiva, mucosal pallor). In further diagnostic testing, the peripheral smear will reveal evidence of hemolyzed RBCs (e.g., bite cells, helmet cells). Macrocytosis results from the marrow’s response to increased cell destruction or blood loss, with release of reticulocytes into the peripheral circulation. Treatment should be focused on the underlying cause of hemolysis or hemorrhage.

**ALCOHOL ABUSE**

The Michigan Alcoholism Screening test and obtaining γ-glutamyltransferase levels were found to be the two most sensitive tests for detecting alcohol abuse in patients with macrocytosis.\(^ {4}\) Physical findings consistent with alcoholism include gynecomastia, caput medusae, and jaundice. Alcohol use more commonly causes macrocytosis through its toxic effect than through folate deficiency secondary to alcoholism. The mean corpuscular volume is generally less than 110 fl with chronic alcohol use. Abstinence from alcohol rapidly corrects the elevated mean corpuscular volume.\(^ {18}\)

**BONE MARROW DYSFUNCTION**

As noted above, myeloproliferative disorders (sometimes called refractory anemia) are a more common cause of

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**Table 3. Medications That May Cause Macrocytosis**

<table>
<thead>
<tr>
<th>Medications</th>
<th>Treatments for human immunodeficiency virus: reverse transcriptase inhibitors (e.g., stavudine [Zerit], lamivudine [Epivir], zidovudine [Retrovir])</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Anticonvulsants (e.g., valproic acid [Depakote], phenytoin [Dilantin])</td>
</tr>
<tr>
<td></td>
<td>Folate antagonists (e.g., methotrexate)</td>
</tr>
<tr>
<td></td>
<td>Chemotherapeutics (e.g., alkylating agents, pyrimidine, purine inhibitors)</td>
</tr>
<tr>
<td></td>
<td>Trimethoprim/sulfamethoxazole (Bactrim, Septra)</td>
</tr>
<tr>
<td></td>
<td>Biguanides (e.g., metformin [Glucophage]), cholestyramine (Questran)</td>
</tr>
</tbody>
</table>

*Information from reference 28.*
Macrocytosis and anemia among older persons than in younger populations. Although the peripheral smear may be suggestive, bone marrow biopsy is required to establish this diagnosis. Referral will likely be necessary for the work-up and management.

OTHER ETIOLOGIES

Medications, hypothyroidism, liver and renal disease, and chronic obstructive pulmonary disease are associated with less dramatic elevations in the mean corpuscular volume. Nonalcoholic liver disease and hypothyroidism account for a substantial portion of macrocytosis. Patient history and physical examination should include a history of these medical diseases. Splenectomy may cause macrocytosis because cells are not processed as thoroughly when the spleen is absent. Down syndrome also may be associated with reticulocytosis or myeloproliferative disorders. Treatment, if needed, can be directed at the underlying disorder.

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Author disclosure: Nothing to disclose.

REFERENCES