Presentation of Case

Dr. Adam B. Cohen (Neurology): A 59-year-old man was admitted to the neurology service of this hospital because of chronic daily headache, fever, and myalgia.

Approximately 6 months earlier, the patient had begun having headache, accompanied by muscle spasms, generalized myalgia, weakness, fatigue, difficulty sleeping, and anxiety. The headache was constant, affected both temporal and frontal regions, and was not affected by posture. The temperature rose daily but remained below 37.8°C. Five months before admission to this hospital, he saw his internist at another hospital. The physical examination was normal. Amitriptyline at bedtime was prescribed, without improvement. The patient took ibuprofen every 4 hours and oxycodone–acetaminophen intermittently, with transient improvement in the headache.

Three months before admission, computed tomography (CT) of the head without contrast material revealed increased density and possible thickening of the dural and subdural spaces along the frontal lobes. Results of liver-function tests were normal; other results of laboratory tests are shown in Table 1.

On follow-up evaluation, the patient reported difficulty sleeping and night sweats, dry mouth at night, polydipsia and polyuria (urinating up to 25 times per day and up to 6 times per night), and pain in the jaws when chewing. His wife noted that he snored at night, but she had not observed choking or apneic episodes. Approximately 30 years earlier, for several years, he had had annual episodes of fever, sweats, and myalgia of 1 week’s duration, each of which had resolved after the use of antibiotics. Twenty-three years earlier, a mediastinal mass, 7.5 cm by 10 cm, was resected, and pathological examination reportedly revealed caseating granulomas within a lymph node. Testing for tuberculosis was negative. The annual febrile episodes had ceased after the surgery.

On physical examination, there were enlarged nasal inferior turbinates and discomfort in the maxillary areas with pressure and percussion. A thoracotomy scar was present. The remainder of the examination was normal. Chest radiography showed calcified right hilar and paratracheal lymph nodes that were slightly more...
prominent than they had been 2 years earlier. There was evidence of a right thoracotomy, and the lungs were clear. Magnetic resonance imaging (MRI) of the brain revealed diffuse thickening and gadolinium enhancement of the dural surfaces, including the falx and tentorium.

The patient was referred by his internist to a neurologist and an infectious-disease specialist at another hospital. Two-and-one-half months before admission, a lumbar puncture was performed (Table 2). A skin test for tuberculosis was negative. Echocardiography revealed normal left ven-

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**Table 1. Results of Laboratory Tests.**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Reference Range, Adults†</th>
<th>2–3 Mo before Admission‡</th>
<th>2 Wk before Admission</th>
<th>1 Day before Admission</th>
<th>Second Hospital Day</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hematocrit (%)</td>
<td>41.0–53.0, in men</td>
<td>39.8</td>
<td>34.8</td>
<td>35.6</td>
<td></td>
</tr>
<tr>
<td>Hemoglobin (g/dl)</td>
<td>13.5–17.5, in men</td>
<td>13.1</td>
<td>11.1</td>
<td>11.2</td>
<td></td>
</tr>
<tr>
<td>White cells (per mm³)</td>
<td>4500–11,000</td>
<td>8,100</td>
<td>14,600</td>
<td>11,300</td>
<td></td>
</tr>
<tr>
<td>Differential count (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Neutrophils</td>
<td>40–70</td>
<td>73</td>
<td>76</td>
<td>68</td>
<td></td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>22–44</td>
<td>12</td>
<td>8</td>
<td>12</td>
<td></td>
</tr>
<tr>
<td>Monocytes</td>
<td>4–11</td>
<td>5</td>
<td>4</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>Eosinophils</td>
<td>0–8</td>
<td>9</td>
<td>11</td>
<td>13</td>
<td></td>
</tr>
<tr>
<td>Basophils</td>
<td>0–3</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Platelets (per mm³)</td>
<td>150,000–350,000</td>
<td>433,000</td>
<td>776,000</td>
<td>670,000</td>
<td></td>
</tr>
<tr>
<td>Erythrocyte sedimentation rate (mm/hr)</td>
<td>0–17, in men</td>
<td>31</td>
<td>61</td>
<td>72</td>
<td></td>
</tr>
<tr>
<td>Reticulocytes (%)</td>
<td>0.5–2.5</td>
<td></td>
<td></td>
<td></td>
<td>1.9</td>
</tr>
<tr>
<td>Mean corpuscular volume (μm³)</td>
<td>80–100</td>
<td>78</td>
<td>76</td>
<td>76</td>
<td></td>
</tr>
<tr>
<td>Toxoplasma IgM and IgG antibodies</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coccidioides antibody</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lyme IgM and IgG antibody on Western blotting</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aspergillus antibody</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td></td>
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<tr>
<td>Histoplasma antibody</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
<td></td>
<td></td>
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<tr>
<td>Rheumatoid factor (IU/ml)</td>
<td>&lt;30‡</td>
<td>24.2</td>
<td>89</td>
<td></td>
<td></td>
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<tr>
<td>Angiotensin-converting enzyme (U/liter)</td>
<td>7–46‡</td>
<td>24</td>
<td>21</td>
<td></td>
<td></td>
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<tr>
<td>Direct antinuclear antibody (U/liter)</td>
<td>&lt;100‡</td>
<td>17</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Antinuclear antibody</td>
<td>Negative at 1:40 and 1:160, speckled</td>
<td>Positive at 1:160, speckled</td>
<td>Positive at 1:40, speckled</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C-reactive protein (mg/liter)</td>
<td>&lt;8.0</td>
<td>95.6</td>
<td>127.7</td>
<td>130.4</td>
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</tr>
<tr>
<td>Total thyroxine (μg/dl)</td>
<td>4.5–12.0</td>
<td>6.5</td>
<td></td>
<td></td>
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<tr>
<td>Thyroid-stimulating hormone (μU/ml)</td>
<td>0.40–5.00</td>
<td>3.93</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Vitamin B₁₂ (pg/ml)</td>
<td>&gt;250</td>
<td>215</td>
<td></td>
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</tr>
<tr>
<td>Iron (μg/dl)</td>
<td>45–160</td>
<td>17</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Iron-binding capacity (μg/dl)</td>
<td>228–428</td>
<td>239</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ferritin (ng/ml)</td>
<td>30–300, in men</td>
<td>260</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* To convert the values for vitamin B₁₂ to picomoles per liter, multiply by 0.7378. To convert the values for iron to micromoles per liter, multiply by 0.1791. To convert the values for total thyroxine to nanomoles per liter, multiply by 12.87.

† Reference values are affected by many variables, including the patient population and the laboratory methods used. The ranges used at Massachusetts General Hospital are for adults who are not pregnant and do not have medical conditions that could affect the results. They may therefore not be appropriate for all patients.

‡ At the other hospital, the reference range for rheumatoid factor was 0 to 13.9 IU per milliliter; for angiotensin-converting enzyme, 12 to 68 U per liter; and for direct nuclear antibody, less than 100 units per liter.
tricular ejection fraction and chamber sizes and no pulmonary hypertension. Four weeks before admission, culture of a nasal swab grew *Klebsiella pneumoniae*, *Candida albicans*, and normal flora. Pathological examination of a biopsy specimen of the lip revealed salivary-gland lobules with scattered plasma cells and a single, prominent lymphoid aggregate. Amoxicillin–clavulanate and nystatin were administered.

Two weeks before admission, the patient saw an infectious-disease specialist at this hospital. The patient reported a 2-month history of mild swelling of the lower legs. He had allergic rhinitis, for which he had received immunotherapy injections for 2 years; a septrhinoplasty had been performed for repair of a deviated septum. Hypothyroidism had developed more than 20 years earlier, after an episode of hyperthyroidism. A tonsillectomy and an appendectomy had been performed. He had lived in a rural area of Indiana until 3 years earlier, when he moved to New England. He had traveled to Mexico 2 months earlier and the southwestern United States 20 years earlier. He had never lived on a farm or been exposed to tuberculosis. He had owned a dog in the past. A sister had thyroid disease; a brother of the patient's child were healthy. There was no family history of headache or neurologic or rheumatic disease. He had no allergies to medications. Medications included levothyroxine, aspirin, ibuprofen, oxycodone–acetaminophen, amoxicillin–clavulanate, and nystatin.

On examination at that time by the infectious-disease specialist, the temperature was 37.6°C, the blood pressure 120/76 mm Hg, the pulse 80 beats per minute, and the weight 80.3 kg. There was 1+ pitting edema of the legs to the knees, and the remainder of the examination was normal. Serum levels of glucose and tests of liver and renal function were normal; testing for syphilis and antineutrophil cytoplasmic autoantibodies (ANCAs) was negative. Serum protein electrophoresis revealed a normal pattern, a moderate diffuse increase in IgG, and no M component. A urinalysis was normal, the specific gravity was 1.004, and the culture was sterile; other test results are shown in Table 1.

One day before admission, MRI of the brain with and without gadolinium revealed smooth thickening and diffuse enhancement of the dura. The brain parenchyma, ventricles, sulci, and major intracranial flow voids were normal in appearance. Later that day, the patient saw a neurologist at this hospital. He reported that the headaches were pulsating in quality. He reported nausea, but no vomiting, and severe pain in the jaws on chewing, which caused him to reduce his food intake. There was no positional component to his headache and no double vision, loss of vision, photophobia, dysarthria, dysphagia, seizures, ataxia, weakness, numbness, or weight loss. On examination, the vital signs were normal. The neck was supple. The temporal arteries were tortuous, enlarged, and nontender, with arterial pulsations easily palpated. The pupils were equal in size, round, and reactive to light, and visual acuity was 20/20 in both eyes with correction. Fundus examination revealed sharp optic-disk margins. The rest of the cranial nerves were normal on examination, as were the motor and sensory functions, coordination, and gait; deep-tendon reflexes were 2+, with bilateral flexor plantar responses. Tests for serum antibodies against double-stranded DNA, against ribonucleoprotein, and against anti-Ro, anti-La, and anti-Smith antibodies were negative. Results of additional tests ordered by the infectious-disease consultant (1 day before admission) are shown in Table 1; a specimen of urine was negative for histoplasma antigen.

The next day, the patient was admitted to this hospital. An electrocardiogram was normal. A tuberculin skin test was negative. Aspirin, levothyroxine, and multivitamins were administered orally, and oxycodone was given as needed for pain; dalteparin was given subcutaneously. Levels of glycated hemoglobin, phosphorus, magnesium, creatine kinase, and lactate dehydrogenase were normal, as was the result of coagulation testing; testing was negative for anticardiolipin IgG and IgM antibodies and partial-thromboplastin time–lupus anticoagulant; other results are shown in Table 1.

On the third hospital day, biopsy of the right temporal artery and a lumbar puncture were performed (Table 2). Vitamin B₁₂ and ferrous sulfate were begun. On the fifth hospital day, an 18-hour water-deprivation test was performed. The plasma sodium level rose from 136 to 142 mmol per liter, serum osmolality from 287 to 295 mOsm per kilogram, and urine osmolality from 101 to 182 mOsm per liter, with a urine volume of 200 to 275 ml per hour. Desmopressin acetate (20 μg) was administered nasally, 2.5 hours after which the urine osmolality was 472 mOsm per kilo-
gram. The temperature rose to 37.7°C on several occasions and to 38.1°C on the sixth day. Pathological examination of the biopsy specimen of the temporal artery was reported to show mild intimal hyperplasia and no evidence of giant-cell arteritis.

On the seventh hospital day, MRI of the brain and pituitary with and without gadolinium showed extension of the dural thickening and enhancement to the sella, with nodular thickening and enhancement of the pituitary stalk and posterior pituitary. A diagnostic procedure was performed.

## Differential Diagnosis

Dr. Steven D. Brass: May we see the imaging studies?

Dr. John W. Chen: Three months before admis-
sion, CT of the head without contrast medium, obtained at the other hospital (Fig. 1A), revealed diffuse low-attenuation, extra-axial foci that were of slightly higher density than would be expected for cerebrospinal fluid. Two and one half months before admission, follow-up contrast-enhanced MRI of the brain performed at the other hospital (Fig. 1B) showed diffuse dural thickening and smooth pachymeningeal enhancement involving nearly all the dural surfaces.

One day before admission, MRI of the brain with gadolinium (Fig. 1C) again showed diffuse smooth pachymeningeal enhancement involving both cerebral hemispheres, falx cerebri, tentorium cerebelli, and the basilar meninges, consistent with hypertrophic pachymeningitis. The degree of thickening was slightly less than that seen 2.5 months previously. No leptomeningeal enhancement was noted.

Thoracic and abdominal CT with contrast medium performed on the same day (see Fig. 1A and 1B in the Supplementary Appendix, available with the full text of this article at www.nejm.org) revealed large, calcified lymph nodes in the mediastinum and right hilum, parenchymal calcification in the right lung, and punctate calcifications in the spleen and liver, consistent with previous involvement of granulomatous disease. MRI of the brain and pituitary with and without gadolinium performed on the seventh hospital day (Fig. 1D) showed diffuse dural thickening and smooth enhancement extending to the sella, with nodular thickening and enhancement of the pituitary stalk and posterior pituitary.

Dr. Brass: All of the discussants are aware of the diagnosis in this case. This patient presented with chronic daily headache, which is defined as 15 days of headache each month for 3 months. When evaluating a patient with chronic daily headache, a thorough history and physical examination are important, with a focus on red flags (Table 3) that may help differentiate primary headaches — including migraine, tension headache, and cluster headache, which are not related to a structural or systemic illness — from secondary headaches that have an identifiable underlying cause (Table 4).

**CHRONIC DAILY HEADACHE**

There are no evidence-based guidelines for the diagnostic evaluation of patients with chronic daily headache. Routine blood tests have a low yield for the diagnosis of headache but may be helpful when considering an infection, inflammatory disease, or substance abuse. Neuroimaging should be considered, especially for patients with an abnormal neurologic examination, atypical headache that does not fulfill criteria for migraine, or evidence of systemic disease. Lumbar puncture may be helpful, to rule out infectious, neoplastic, and inflammatory causes. Electroencephalography is not routinely done unless there is a seizure or atypical aura.

Numerous red flags were identified in this patient’s evaluation: new onset of headache, with systemic symptoms, in a middle-aged patient. Examination did not disclose nuchal rigidity, temporal-artery tenderness, optic-disk edema, or localizing neurologic signs, but MRI of the brain showed evidence of pachymeningitis, and the lumbar puncture showed an elevated opening pressure with lymphocytosis and an elevated protein level. These features led us to focus on secondary causes for headache that are associated with pachymeningitis, including neoplastic diseases, headache caused by decreased intracranial pressure, idiopathic hypertrophic meningitis, infection, and autoimmune disease.

**NEOPLASM**

We considered dural metastases from cancers that commonly metastasize to the dura — including lung and prostate cancers and melanoma — or dural involvement by primary lymphoma of the central nervous system. Patients may be asymptomatic or may have headache and focal findings such as seizures. This diagnosis was ruled out because our patient had no history of cancer and had smooth dural enhancement as opposed to the focal nodular dural enhancement often seen on T₁-weighted postgadolinium sequences in patients with dural metastases.

**HEADACHE DUE TO LOW CEREBROSPINAL FLUID PRESSURE**

The differential diagnosis of chronic daily headache in a patient who presents with dural enhancement on neuroimaging should include a headache due to low cerebrospinal fluid pressure, which is typically the result of a cerebrospinal fluid leak from a spinal dural tear. The headache characteristically has an orthostatic component, but this feature may disappear over time. Patients may also report photophobia, tinnitus, nausea,
and neck stiffness, which our patient did not have. MRI revealed diffuse dural enhancement on T1 postgadolinium sequences, and there may be lymphocytosis and an elevated protein level in the cerebrospinal fluid, as seen in this patient. The lumbar puncture classically reveals an opening pressure less than 60 mm of water, but the value may be within the normal range.

In this patient, the diagnosis of headache due to low cerebrospinal fluid pressure was unlikely on the basis of clinical history, including the absence of an orthostatic component and presence of systemic symptoms, and was ruled out by the finding of an elevated opening pressure on lumbar puncture.

IDIOPATHIC HYPERTROPHIC PACHYMENINGITIS

Idiopathic hypertrophic pachymeningitis may cause chronic daily headache, characteristically in middle-aged men, such as this patient. Patients have variable neurologic symptoms and signs, such as cranial-nerve palsies, optic-disk...
edema, hemiparesis, loss of vision, blindness, and cerebellar symptoms. MRI of the brain shows dural enhancement on T1 postgadolinium sequences. On lumbar puncture, there may be an elevated opening pressure, elevated cerebrospinal fluid protein level, and cerebrospinal fluid lymphocytosis on sterile cerebrospinal fluid culture, as in this patient. Idiopathic hypertrophic pachymeningitis is a diagnosis of exclusion, and the recurrent low-grade fever and myalgia made it necessary to rule out other conditions before labeling this condition idiopathic.

Our greatest concern was infection or an autoimmune disorder, and therefore we consulted colleagues in Infectious Diseases and Rheumatology.

**PACHYMENINGITIS DUE TO INFECTION**

*Dr. Marlene L. Durand:* This patient was referred to me to exclude infectious causes of pachymeningitis. Most reports of infectious pachymeningitis are in the older literature and describe spinal pachymeningitis; cases were attributed to syphilis, tuberculosis, or molds. However, documentation of these or other infectious diseases as causes of pachymeningitis is very rare in the modern era, especially in cases of cranial pachymeningitis.

By the time I saw this patient, he already had had an extensive infectious-disease evaluation by his local physicians. Our major concern was histoplasmosis, since he had moved from an area in which the disease is endemic 3 years earlier, had had granulomatous mediastinal lymphadenitis 23 years earlier, and had hepatic and splenic calcifications characteristic of previous histoplasmosis infection. However, he was not immunocompromised, had been well for 22.5 years after excision of the mediastinal node, and had a negative test for urinary histoplasma antigen; thus, progressive disseminated histoplasmosis seemed unlikely. Although isolated histoplasmosis of the central nervous system has been described in immunocompetent hosts, the absence of histoplasma antigen or antibody in the cerebrospinal fluid and the presence of negative fungal cultures made this diagnosis unlikely.

**RHEUMATOLOGIC DISEASES AND PACHYMENINGITIS**

*Dr. John H. Stone:* Several rheumatic diseases can cause pachymeningitis, particularly those that are associated with granulomatous inflammation. Pachymeningitis associated with any of these could cause diabetes insipidus through involvement of the infundibular area of the brain or through associated hypophysitis, but the diagnosis must also account for the patient’s headache, myalgia, weakness, and pain on chewing.

Wegener’s granulomatosis is the rheumatic disease most likely to cause hypertrophic pachymeningitis, but this manifestation is relatively uncommon as compared with other features. Pachymeningitis in patients with Wegener’s granulomatosis can manifest precisely as it did in this patient, with chronic, unrelenting headaches. Two points argue strongly against Wegener’s granulomatosis in our patient. First, he had no features of involvement of other organ systems, such as the upper airway, and second, ANCA assays were negative. A total of 10 to 15% of patients with Wegener’s granulomatosis may be ANCA-negative, including patients with pachymeningitis. However, in the absence of both ANCA and any classic disease features, the consideration of other diagnoses is imperative.
Table 4. Causes of Secondary Headache.

<table>
<thead>
<tr>
<th>Cause</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>Head or neck trauma</td>
<td>Post-traumatic headache</td>
</tr>
<tr>
<td>Cranial or cervical vascular disorder</td>
<td>Arterial dissection</td>
</tr>
<tr>
<td>Nonvascular intracranial disorder</td>
<td>Noninfectious inflammatory disease</td>
</tr>
<tr>
<td></td>
<td>Intracranial neoplasm</td>
</tr>
<tr>
<td></td>
<td>High and low cerebrospinal fluid pressure headache</td>
</tr>
<tr>
<td>Substance use or its withdrawal</td>
<td>Medication-overuse headache</td>
</tr>
<tr>
<td>Infection</td>
<td>Intracranial infection: meningitis</td>
</tr>
<tr>
<td>Disorder of cranium, neck, face, eyes, ears, nose, sinus, teeth, mouth, or other facial structure</td>
<td>Acute glaucoma</td>
</tr>
<tr>
<td>Disorder of homeostasis</td>
<td>Hypoxia or hypercarbia: sleep apnea</td>
</tr>
<tr>
<td></td>
<td>Hypothyroidism</td>
</tr>
<tr>
<td></td>
<td>Fasting</td>
</tr>
<tr>
<td>Psychiatric disorder</td>
<td>Somatization disorder</td>
</tr>
<tr>
<td>Cranial neuralgia</td>
<td>Occipital neuralgia</td>
</tr>
</tbody>
</table>

* Data are based on information from the International Headache Society and Silberstein et al.

Neurosarcoidosis can occur without other organ-system involvement, most commonly with leptomeningeal disease involving the basilar meninges. In the absence of involvement of other organs, the diagnosis of neurosarcoidosis is challenging. This patient had a history of granulomas in mediastinal lymph nodes, but they were caseating and thus not typical of sarcoidosis, and he did not have chest radiographic findings to suggest sarcoidosis. Bronchoscopic biopsies of the lung may be useful, even in patients with a normal chest radiograph. Pachymeningitis has been reported in the Churg–Strauss syndrome. Patients with the Churg–Strauss syndrome are less likely to be ANCA-positive than are patients with Wegener’s granulomatosis. This patient had allergic rhinitis, a frequent manifestation of the Churg–Strauss syndrome. In the absence of a more clinically significant peripheral eosinophilia, however, this diagnosis is unlikely. Meningeal inflammation is a rare but reported extra-articular manifestation of rheumatoid arthritis. Although our patient was positive for rheumatoid factor, the absence of arthritis effectively rules out this diagnosis.

The final form of granulomatous inflammation that must be considered is giant-cell arteritis. Chronic headache is a cardinal feature of this disease. Headaches vary considerably from patient to patient in their location, quality, and intensity. For any individual patient, the chief distinguishing feature of headache is that it differs from all other types of headache the patient has had before. The unrelenting headache reported by this patient is consistent with giant-cell arteritis. The reported myalgia and weakness are consistent with polymyalgia rheumatica, which occurs in one third to one half of patients with giant-cell arteritis. The pain in the jaw on chewing — jaw claudication — is viewed as the most specific symptom suggesting giant-cell arteritis, although it occurs in only about one third of patients with biopsy-proven giant-cell arteritis. A temporal-artery biopsy is imperative in any patient over 50 years of age with chronic headache and jaw claudication. If the diagnosis is strongly suspected, treatment should begin even before the temporal-artery biopsy, with performance of bilateral biopsies as soon as possible (within a few days). Unilateral biopsy leads to underdiagnosis in 20 to 40% of cases.

Sjögren’s syndrome was considered by the patient’s physicians, and a lip biopsy was performed. The dry mouth and eye irritation suggested xerostomia and keratoconjunctivitis sicca, respectively. The patient is rheumatoid-factor positive, as are most patients with extraglandular Sjögren’s syndrome. However, tests for antibodies to the Ro and La antigens — markers strongly associated with extraglandular disease — were negative. Sjögren’s syndrome cannot explain the presence of pachymeningitis, diabetes insipidus, headaches, and jaw claudication.

In summary, Wegener’s granulomatosis would be most likely to cause this kind of presentation, but several features — the headache, symptoms of polymyalgia rheumatica, and jaw claudication — make it essential to rule out giant-cell arteritis. The only way of differentiating among these disorders is through histopathology. A meningeal biopsy would have a strong likelihood of distinguishing among the disorders discussed above. In addition, biopsy of the contralateral temporal artery is also justified.

Dr. Nancy Lee Harris (Pathology): Dr. Friday, would you give us your impressions when you saw this patient?

Dr. Robert P. Friday (Rheumatology): At the time the rheumatology service was consulted, an exhaustive evaluation had been completed. We agreed.
with our neurology colleagues that a meningeal biopsy was necessary to establish a diagnosis, with Wegener's granulomatosis being the most likely inflammatory cause of this patient's pachymeningitis, despite negative ANCA testing. However, we also requested a left temporal-artery biopsy, since jaw claudication has high specificity for the diagnosis of giant-cell arteritis, which is a “do not miss” diagnosis associated with a risk of sudden, irreversible vision loss. The initial approach to treatment for giant-cell arteritis would be different from that for Wegener's granulomatosis or another rheumatic disease.

CLINICAL DIAGNOSIS

Pachymeningitis due to either Wegener’s granulomatosis or giant-cell arteritis.

PATHOLOGICAL DISCUSSION

Dr. James R. Stone: Review of the biopsy specimen of the lip revealed a minor salivary gland with a lymphoid aggregate and a mild diffuse lymphoplasmacytic infiltrate. These findings suggest Sjögren's syndrome in the appropriate clinical setting but are not diagnostic. The biopsy specimen of the right temporal artery showed mild intimal hyperplasia, with no inflammation. Intimal hyperplasia in atherosclerosis-resistant medium-sized muscular arteries is a nonspecific finding, most often associated with increased age, smoking, or repetitive trauma.

The diagnostic procedures were biopsies of the left temporal artery and of the dura. The left temporal artery showed granulomatous inflammation involving primarily a branch of the main superficial temporal artery (Fig. 2A), extending into adjacent small muscular arteries. There was only focal necrosis and no evidence of compact sarcoid-type granulomas or small-vessel leukocytoclastic vasculitis. The dural biopsy contained a lymphocyte-rich inflammatory infiltrate involving the small arteries (Fig. 2B). Fungal stains were negative.

The pathological features are characteristic of a primary vasculitis. Of the primary vasculitides, the two to consider are giant-cell arteritis and Wegener's granulomatosis. Although Wegener's granulomatosis can involve both temporal arteries and the dura, the lack of substantial necrosis, extravascular granulomatous inflammation, and small-vessel leukocytoclastic vasculitis make it unlikely. The histologic features of the temporal arteries are characteristic of giant-cell arteritis, which may also involve small muscular arteries, as in this patient. Although intracranial involvement by giant-cell arteritis is unusual, a small number of autopsy reports have described the involvement of medium-sized intracranial arteries or the dura in giant-cell arteritis.

Dr. Harris: Dr. Friday, would you tell us about the care of the patient?

Dr. Friday: Corticosteroids are the mainstay of therapy for patients with giant-cell arteritis, typically beginning with 0.5 to 1 mg per kilogram of body weight per day, with adjustments directed by the clinical response and the levels of serum inflammatory markers. A recent report suggests that intravenous pulse corticosteroid dosing at the time of diagnosis may reduce the cumulative corticosteroid dose at 1 year in patients with giant-cell arteritis, but this observation requires further validation. Because of the complex clinical presentation, and since infection continued to be considered, corticosteroid therapy was withheld until the diagnosis was confirmed. Prednisone (60 mg daily) was then begun.

Additional therapies were aimed at reducing treatment-related complications. For prevention of glucocorticoid-induced osteoporosis, we administered calcium, vitamin D, and oral bisphosphonate, according to recommendations from the American College of Rheumatology. Patients who are receiving either antiplatelet therapy (low-dose aspirin or clopidogrel) or anticoagulant therapy (warfarin) for other conditions at the time of a diagnosis of giant-cell arteritis reportedly have fewer ischemic complications than patients not receiving antiplatelet or anticoagulant therapy. The patient was already taking 81 mg of aspirin daily at the time of presentation, and this was continued. Proton-pump–inhibitor therapy was also recommended. Trimethoprim–sulfamethoxazole was prescribed as prophylaxis against Pneumocystis jiroveci pneumonia, since we anticipated extended treatment with moderate-dose or high-dose corticosteroids.

The patient's headache improved overnight, after a single dose of prednisone. After the second dose, jaw claudication ceased, and within a week, myalgias and fatigue subsided. The erythrocyte sedimentation rate and serum C-reactive protein level plummeted. With attempts to re-
duce his prednisone dose during the subsequent 3 months, headache and myalgias recurred, with slight increases in the erythrocyte sedimentation rate and serum C-reactive protein level, but responded well to increases in dose. After approximately 5 months, the prednisone dose was tapered, without recurrence of symptoms, to a daily dose of 5 mg at 1 year. The excellent response to corticosteroid monotherapy is typical for patients with giant-cell arteritis.

**Dr. Chen:** Follow-up MRI of the brain and pituitary after 3 months of corticosteroid therapy (Fig. 1E) revealed marked improvement in the pachymeningeal enhancement and thickening, with some mild residual thickening and enhancement posteriorly. The nodular enhancement of the pituitary stalk and posterior pituitary had nearly resolved.

**Dr. Harris:** Dr. Utz, would you comment on the patient’s endocrine evaluation?

**Dr. Andrea L. Utz** (Endocrinology): I initially saw this patient approximately 8 weeks after his discharge, for management of diabetes insipidus. His inpatient fluid-deprivation testing was consistent with, but not diagnostic of, diabetes insipidus, since the testing was terminated slightly prematurely, before development of an elevated serum sodium level or osmolality. An intranasal dose of 20 μg of desmopressin acetate produced an increase of more than 50% in urine osmolality, consistent with central diabetes insipidus. The patient’s symptoms resolved after administration of daily inhaled desmopressin acetate (10 μg), and the serum sodium level remained normal. I therefore opted to continue treatment with desmopressin acetate without repeating the fluid-deprivation test. I stressed that the patient should take desmopressin acetate only as needed for polyuria and to drink fluids as needed for thirst. Since glucocorticoid treatment could lead to resolution of the diabetes insipidus, as-needed dosing of desmopressin acetate would decrease the risk of hyponatremia.

I also tested for other pituitary hormone abnormalities. The patient’s thyroid-stimulating hormone and free thyroxine levels were consistent with those seen with treated primary hypothyroidism. He had two morning testosterone levels that were moderately low, which could be due to pituitary-stalk dysfunction, use of high-dose glucocorticoids, or chronic illness. I initiated testosterone replacement for its anabolic effect on bone and muscle, particularly in the context of therapy with high-dose glucocorticoids. The patient’s insulin-like growth factor I level was in the middle of the normal range, which suggested that he did not have severe growth hormone deficiency. His prolactin level was normal. At 18 months, when his glucocorticoid dose was weaned below a standard replacement dose, co-syntropin stimulation testing was performed to ensure adrenal sufficiency. He continues to take daily desmopressin acetate and testosterone, 2 years after the diagnosis.
Dr. Harris: Dr. Brass, I believe you have some additional follow-up information.

Dr. Brass: Fourteen months after discharge, the patient described a new pattern of headache: intermittent mild morning headache lasting 1 to 2 hours, nocturnal awakenings, snoring, excessive daytime somnolence, feeling unrefreshed after a night of sleep, a 30-pound weight gain, and depressive symptoms. Morning headache is a common symptom of obstructive sleep apnea.5,57 On the basis of the constellation of symptoms and his normal neurologic examination, I obtained a polysomnogram, which revealed a respiratory disturbance index (the number of episodes of obstructive apnea, hypopnea, and respiratory-effort–related arousal per hour of sleep) of 41, which was also associated with oxygen desaturations down to 82% during rapid-eye-movement sleep. A respiratory disturbance index greater than 30 with symptoms of excessive daytime sleepiness is considered diagnostic of severe obstructive sleep apnea.58

Our patient’s internist originally suspected obstructive sleep apnea as the cause of the chronic daily headache. This outcome illustrates the importance of looking for other conditions associated with headache and of reevaluating patients presenting with a new pattern of headache.

ANATOMICAL DIAGNOSIS

Giant-cell arteritis, involving the superficial temporal artery and small dural arteries.

Dr. Brass reports receiving consulting fees from Teva Neuroscience, EMD Serono, Biogen, and Bayer; Dr. John Stone, consulting fees from Merck and Zymogenetics; and Dr. James Stone, consulting fees from Merck, Muscle Tech, GlaxoSmithKline, and Proteon Therapeutics. No other potential conflict of interest relevant to this article was reported.

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The New England Journal of Medicine
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