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Smell and Taste Disorders in Primary Care

Dillon J. Savard, MD, David Grant Medical Center, Travis Air Force Base, California; Uniformed Services University of the Health Sciences, Bethesda, Maryland

Francesca G. Ursua, MD, University of Nebraska Medical Center, Omaha, Nebraska Heidi L. Gaddey, MD, Christie Clinic, Champaign, Illinois; Carle Foundation Hospital Family Medicine Residency Program, Urbana, Illinois

Disorders of smell and taste are reported by approximately one-fifth of people 40 years and older, and one-third of people 80 years and older. These disorders affect quality of life and the ability to identify smoke and toxins. Smell and taste disorders can be early signs of dementia or Parkinson disease and are associated with increased mortality. Dysfunction may

be apparent or may develop insidiously. Screening questionnaires are available, but many patients are unaware of their disorder. Most smell and taste disorders are due to sinonasal disease but also could be caused by smoking, medications, head trauma, neurodegenerative disease, alcohol dependence, or less common conditions. The differential diagnosis should guide the evaluation and include anterior rhinoscopy and an examination of the oral cavity, head, and cranial nerves. Further investigation is often unnecessary, but nasal endoscopy and computed tomography of the sinuses may be helpful. Magnetic resonance imaging of the head with contrast should be performed if there is an abnormal neurologic examination finding or if trauma or a tumor is suspected. Olfactory testing is indicated in refractory cases or for patients with poor quality of life and disease associated with smell or taste dysfunction. Smell and taste disorders



may resolve when reversible causes are treated, but improvement is less likely when they are due to trauma, age, or neurodegenerative disease. Olfactory training is a self-administered mindful exposure therapy that may improve olfactory function. Physicians should encourage patients to ensure that smoke and other alarms are operational and to adhere to food expiration dates. (Am Fam Physician. 2023;108(3):240-248. Copyright © 2023 American Academy of Family Physicians.)

Disorders of smell and taste affect a person's quality of life, health, and safety. The COVID-19 pandemic renewed attention to these disorders, and one of the Healthy People 2030 objectives is to increase the proportion of adults with smell or taste disorders who discuss the problem with a clinician. Olfactory dysfunction is reported by 23% of people in the United States 40 years and older and 39% of those 80 years and older. Altered taste is reported by 19% of people 40 years and older, and 27% of those 80 years and older. Smell and taste disorders can affect food consumption habits, which may lead to chronic disease (*Table 1*4). Depression, poor sleep, dementia, unintentional weight loss, and malnutrition are associated with disordered smell or taste. The ability to detect sources of danger,

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such as smoke or poison, depends on smell or taste. 10,11 When directly tested, 20.3% of a nationally representative sample of adults 70 years and older without known

WHAT'S NEW ON THIS TOPIC

Smell and Taste Disorders

Dysfunction in smell and taste occurs in 40% to 50% of patients diagnosed with COVID-19, often without rhinorrhea or nasal congestion.

A prospective cohort study found that recovery of smell and taste after COVID-19 occurred for most patients by seven days, and more than 80% recovered by three months.

When directly tested, 20.3% of a nationally representative sample of adults 70 years and older without known olfactory dysfunction misidentified the scent of smoke, and 31.3% misidentified the smell of natural gas.

More than 95% of patients with Parkinson disease have smell impairment, and up to 75% have anosmia or hyposmia.

TABLE 1 **Disorders of Taste and Smell** Disorder Description Taste Ageusia Complete loss of ability to taste Dysgeusia Distorted taste perception (triggered by a taste stimulus) Hypogeusia Reduced ability to taste Phantogeusia Gustatory hallucination (occurs without a stimulus) Smell Anosmia Complete loss of ability to smell Hyposmia Reduced ability to smell Parosmia Distorted odor perception (triggered by a stimulus) Phantosmia Olfactory hallucination (occurs without a stimulus)

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disorders in primary care. Am Fam Physician. 2013;88(12):852.

dysfunction misidentified smoke, and 31.3% misidentified natural gas.³

There are many clinical presentations and causes for smell and taste disorders (*Table 2*). ¹⁶⁻¹⁹ Etiologies range from sinonasal disease to less common conditions such as Sjögren syndrome (*Table 3*). ⁴ When only taste disorder is perceived by the patient, it is often due to olfactory dysfunction. ²⁰⁻²⁶ Therefore, without an apparent taste insult, physicians should focus their evaluation on the olfactory system. *Figure 1* provides an approach to evaluating smell and taste disorders in primary care.

Anatomy and Physiology

Chemicals that cause smells (i.e., odorants) are absorbed through the mucosa of the nose, nasopharynx, and oropharynx. Odorants stimulate olfactory neurons in the neuroepithelium over the cribriform plate. Olfactory neurons project directly to the olfactory bulb, which communicates with the

Causes, Pathology, and History of Olfactory Dysfunction					
Cause	Pathology	History			
Age-related (presbyosmia)	Loss or replacement of olfactory neurons with respiratory epithelium Decreased basal cell proliferation Decrease of interneurons in olfactory bulb with reduced activity in olfactory cortex	Insidious onset; unlikely to see improvement Olfactory dysfunction predicts five-year mortality			
Neurodegenera- tive causes (e.g., Alzheimer disease, Parkinson disease)	Alzheimer disease: neurofibril changes in the olfactory bulb and higher olfactory network Parkinson disease: Lewy bodies deposited in the olfactory tract, olfactory bulb, and anterior olfactory nucleus ¹⁷	Insidious onset; unlikely to see improvement Onset can be early in Alzheimer and Parkinson diseases			
Other medication/ toxin exposure/ iatrogenic	Altered receptor function by binding G-protein coupling or affecting calcium or sodium channel activity	Sudden onset; mostly resolves with ceasing expo- sure; can persist in some cases requiring treatment can have medicolegal implications ¹⁸			
Postviral olfactory dysfunction	Viral common cold pathogens, COVID-19, influenza, respiratory syncytial virus, parvovirus, HIV Long-term inflammation causes neuroepithelial remodeling to respiratory epithelium	Sudden onset with little fluctuation; often associated with parosmia; one in three recovers in the 14 months after presentation for care ¹⁹			
Posttraumatic olfactory dysfunction	Severing of olfactory nerve filament or central nervous system injury	Sudden or delayed onset; fluctuation is uncommon; phantosmia and parosmia are common 10% recover by 13 months, but up to 35% can recover depending on severity ¹⁹			
Sinonasal (e.g., chronic rhinosi- nusitis with polyps)	Obstruction of the olfactory cleft; temporary interference with olfactory receptor binding due to inflammation; eventual neuroepithelial remodeling/central nervous system changes	Gradual onset and fluctuation over time; typically improves with nasal/systemic corticosteroids Not commonly associated with parosmia			

TABLE 3				
Principal Etiologies of Olfactory Dysfunction				
Cause	Leading etiologies*			
Most common Sinonasal conditions	Upper respiratory tract infection (especially viral, including COVID-19), allergic rhinitis, chronic rhinosinusitis, nasal polyps			
More common Head trauma	Damage to cribriform plate, shearing forces, and intracranial damage; facial trauma			
Neurodegenerative disorders	Parkinson disease, parkinsonism, Alzheimer disease, mild cognitive impairment, multiple sclerosis			
Less common Medications	Chemotherapy, angiotensin-converting enzyme inhibitors, angiotensin receptor blockers, dihydropyridine calcium channel blockers, diuretics, intranasal zinc, antimicrobials (macrolides, terbinafine, fluoroquinolones, protease inhibitors, griseofulvin, penicillins, tetracyclines, nitroimidazoles [metronidazole (Flagyl)]), antiarrhythmics, antithyroid agents, antidepressants, anticonvulsants, lipid-lowering agents			
Intoxicants or illicit substances	Alcohol dependence, cocaine use			
Toxins	Ammonia, hairdressing chemicals, gasoline, formaldehyde, paint solvents, welding agents, benzene, sulfuric acids, cad- mium, acrylates, iron, lead, chromium			
Chronic medical conditions	Renal or hepatic failure, complicated type 2 diabetes mellitus, cancer, HIV			
Structural or mechanical conditions	Ischemic stroke, subarachnoid or intracranial hemorrhage, brain or sinonasal tumor			
Nutritional deficiencies	Malnutrition, pernicious anemia or vitamin $\rm B_{12}$ deficiency, deficiencies in vitamins $\rm B_6$ or A, niacin, zinc, or copper			
Postsurgical state	Nasal surgery (septal or sinus)			
Postradiation	Especially to head and neck			
Congenital conditions	Kallmann syndrome, anosmia			
Psychiatric conditions	Anorexia nervosa (not bulimia), major depressive disorder, bipolar disorder, schizophrenia			
Endocrine conditions	Pregnancy, hypothyroidism, Addison disease, Cushing syndrome			
Autoimmune/ inflammatory conditions	Sjögren syndrome, systemic lupus erythematosus, sarcoidosis, herpes encephalitis			
*—Etiologies are listed	in relative order of prevalence.			
Adapted with permission Am Fam Physician. 201	on from Malaty J, Malaty IAC. Smell and taste disorders in primary care 13;88(12):855.			

olfactory cerebral cortex. Olfactory receptors regenerate every eight to 10 days, followed by five days for the maturation of cilia.²⁷⁻²⁹ Dysfunction along these pathways can result in a smell disorder.³⁰⁻³²

Odorants and tastants (i.e., particles and liquids that cause the taste sensation) dissolve in saliva and directly contact gustatory receptors in dedicated cells for each of the five tastes (bitterness, saltiness, sourness, sweetness, and umami [savory]). These cells are in taste buds on the tongue, soft palate, pharynx, larynx, epiglottis, and the proximal one-third of the esophagus. Different nerves innervate the anterior two-thirds of the tongue, posterior one-third of the tongue, palate, pharynx, and larynx. The trigeminal nerve also mediates certain characteristics of smell and taste, such as irritation and temperature. Taste is mediated by several different nerves depending on anatomic location; therefore, gustatory loss is less common than olfactory loss. 9,33,34

Clinical Assessment

HISTORY

Validated questionnaires help clinicians screen for subjective olfactory dysfunction (*Table 4*).^{11,35,36} A National Health and Nutrition Examination Survey used three questions that were not formally validated but are a reasonable approach to screen patients for olfactory dysfunction.^{2,3} There are no validated subjective tools to assess taste.¹⁶

Smell influences taste; therefore, it can be challenging to distinguish which sense is disordered and determine the severity of the disorder. The history should include inquiries about medications, supplement use, nose or mouth pain, and symptoms or history of dry mouth, rhinitis, rhinosinusitis, nasal polyps, recent upper respiratory tract infection, or trauma. In older patients, the history should include queries about symptoms of Parkinson disease or dementia.

PHYSICAL EXAMINATION

The physical examination should include anterior rhinoscopy (visual examination of the anterior nasal passages) and examination of the head, oral cavity, and neurologic system. Findings suggestive of a sinonasal etiology include intranasal masses, edematous mucosa, hypertrophic turbinates, purulent mucus, or bleeding. Anterior rhinoscopy is necessary for isolated taste dysfunction because olfactory dysfunction and sinonasal etiologies are commonly found. Nasal endoscopy is much more sensitive (91%) for identifying sinonasal pathology and may be necessary if the initial examination is inconclusive.³⁷ Examination of the oral cavity may find inflammation, plaque, erosions of the tongue or mucosa, or masses. Evidence of head trauma or neurologic dysfunction warrants an additional neurologic examination and specialized testing.

DIAGNOSTIC TESTING

Smell and taste disorders are often due to common conditions readily treated by family physicians without specialized testing. However, because patients often underestimate or do not recognize their dysfunction, many experts recommend having a low threshold for obtaining otolaryngology consultation for objective testing of olfactory function. ^{3,20,38-40} Family physicians should consider olfactory testing for refractory cases and for patients with a poor quality of life who are affected by conditions associated with olfactory or

gustatory dysfunction (e.g., depression, poor sleep, dementia, unintentional weight loss, malnutrition; *Table 5*⁴). Taste function testing is performed in dedicated centers. ^{16,41}

Imaging is not necessary when the initial history and physical examination identify common benign and treatable atraumatic etiologies. If further investigation of the sinuses is warranted, computed tomography without contrast is preferred. Magnetic resonance imaging (MRI) of the head with contrast is warranted when there is a history of trauma, an abnormal neurologic examination finding, or suspicion for a tumor.

Causes of Smell and Taste Disorders

SINONASAL CONDITIONS

Most olfactory dysfunction is caused by sinonasal conditions, such as nasal polyps, allergic rhinitis, chronic

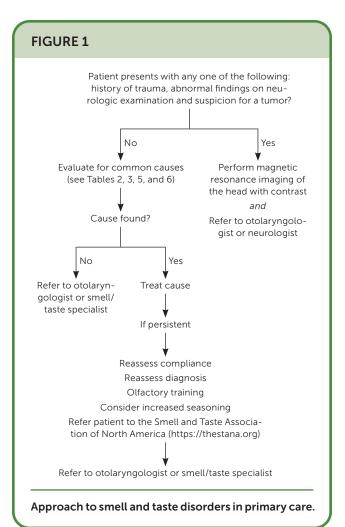


TABLE 4

Tools Available for Primary Care Office—Based Screening for Olfactory Dysfunction

National Health and Nutrition Examination Survey questionnaire:

- 1. During the past 12 months, have you had a problem with your ability to smell, such as not being able to smell things or things not smelling the way they are supposed to? (Yes or No)
- 2. How would you rate your ability to smell now compared with when you were 25 years old (could replace with younger)? (Better, Worse, or No Change)
- 3. Do you sometimes smell an unpleasant, bad, or burning odor when nothing is there? (Yes or No)

Self-reported Mini Olfactory Questionnaire (five items)	https://onlinelibrary.wiley.com/doi/10.1002/ lary.28419#lary28419
Brief Questionnaire of Olfactory Disorders–negative statements (seven items)	https://onlinelibrary.wiley.com/doi/ abs/10.1002/alr.22800
Questionnaire of Olfactory Disorders—negative statements (17 items)	https://onlinelibrary-wiley-com.usu01.idm. oclc.org/doi/epdf/10.1002/lary.23349

Information from references 11, 35, and 36

rhinosinusitis, and upper respiratory tract infection. Olfactory dysfunction associated with sinonasal disease usually resolves with successful treatment of the related condition. Oral and nasal corticosteroids are often used to treat some of these conditions. Corticosteroids may improve the transmission of odorants to the olfactory nerve epithelium

Categories of Gustatory Dysfunction					
Category*	Leading etiologies				
Poor oral hygiene, infection/ inflammation	Gingivitis, oral abscess, oral candidiasis, periodontal disease, viral upper respiratory tract infection				
Oral appliances	Dentures, prosthetics (hypothesized to be related to oral inflammation if poor hygiene with appliances)				
Postsurgical state	Middle ear surgery affecting chorda tympani (usually transient), oral or dental surgery (especially third molar extraction)				
Radiation	Head and neck irradiation with oral mucositis, xerostomia				
Nutritional	Copper supplementation; deficiency of copper, vitamin B_3 (niacin), vitamin $B_{12'}$ or zinc (rare); general malnutrition				
Medications	Intranasal zinc, chlorhexidine, chemotherapy, angiotensin-converting enzyme inhibitors, angiotensin receptor blockers, dihydropyridine calcium channel blockers, diuretics, antimicrobials (macrolides, terbinafine, fluoroquinolones, protease inhibitors, griseofulvin, penicillins, tetracyclines, nitroimidazoles [metronidazole (Flagyl)]), antiarrhythmics, antithyroid agents, antidepressants, anticonvulsants, lipid-lowering agents				
Head trauma	Head or facial trauma				
Toxins	Pepper gas (oleoresin capsicum), weed killer with trifluralin, ammonia, benzene, sulfuric acids, cadmium, acrylates, iron, lead, chromium				
Chronic medi- cal conditions	Renal or hepatic failure, cancer, HIV, complicated type 2 diabetes mellitus				
In a retrospective problems, only 10 (ven causes of gustatory dysfunction are limited review of 1,176 patients with subjective taste (0.85%) had significant gustatory loss that coulc are not many large studies available for review.				
*—Categories are li	isted in relative order of prevalence.				
	nission from Malaty J, Malaty IAC. Smell and rimary care. Am Fam Physician. 2013;88(12):857.				

by decreasing local swelling and inflammation.⁴² Multiple systematic reviews evaluated possible pharmacotherapies for postviral olfactory dysfunction and found relatively low-quality evidence. Nasal corticosteroids showed a small benefit and a minimal risk. Studies on zinc did not show consistent benefits.⁴³ One industry-sponsored, placebo-controlled, double-blind, randomized clinical trial of 323 patients with chronic rhinosinusitis and nasal polyps with anosmia or hyposmia found that fluticasone propionate, delivered via an exhalation delivery system, yielded a clinically significant improvement in patient-reported sense of smell.44 Oral corticosteroids may improve symptoms but, due to potentially severe adverse effects, their use should be limited (three to seven days) and reserved for severe symptoms refractory to other medical therapies. 45 A prospective cohort study found improved olfactory function and olfactory bulb volume six months after endoscopic sinus surgery for chronic rhinosinusitis with nasal polyps. 46

COVID-19

Smell and taste dysfunction occurs in 40% to 50% of patients diagnosed with COVID-19.47 The dysfunction tends to improve or resolve more quickly than other etiologies and, with some variants of the virus, it may occur without rhinorrhea or nasal congestion. 48 A prospective cohort study (n = 129) conducted over 18 months found recovery occurred for most patients by seven days, and more than 80% recovered by three months. The severity of the initial dysfunction and degree of improvement in the first seven days predicted earlier resolution of smell and taste dysfunction after COVID. 49 Most studies examining intranasal corticosteroid treatment for post-COVID olfactory dysfunction did not show a benefit. A Cochrane systematic review of interventions for preventing persistent post-COVID olfactory dysfunction found "very limited evidence available."50

HEAD TRAUMA

Olfactory dysfunction after head trauma is common, and the evaluation should include imaging and specialty consultation. The anatomic location and nerve fibers that extend from the olfactory bulb near the cribriform plate are vulnerable. Besides direct trauma, shearing of olfactory fibers can also occur with coup-contrecoup injury. Posttraumatic olfactory disorders often have MRI findings of reduced olfactory bulb volume that correlate with the severity of the disorder. Postoperative olfactory or gustatory dysfunction may occur if the surgical location is near relevant anatomic structures.8,16 Recovery of olfactory function after trauma ranges from 10% to 35% but is less common than after infection and depends on the severity or location of the trauma.

SMOKING

After smoking cessation, olfactory function takes 15 years to return to levels consistent with nonsmokers. The mechanisms by which smoking impairs olfaction have not been definitively identified. Vascular mechanisms may play a role because the risk of coronary heart disease and vascular death in former smokers takes a similar amount of time to resolve to nonsmoker levels (15 and 20 years, respectively).⁵¹

The effect of secondhand smoke exposure on olfactory function is inconsistent. A prospective, case-control study comparing olfactory function in adults younger than 65 years found that olfactory function was similarly worse in people who smoke and those passively exposed to smoke, compared with nonsmokers without passive exposure.⁵² A cross-sectional analysis of 201 children six to 11 years of age showed no difference in olfaction among those with or without smoking parents.⁵³

NEURODEGENERATIVE DISEASES

Family physicians should consider evaluating patients for neurodegenerative diseases when they present with olfactory or gustatory dysfunction without an apparent etiology (Table 6).10 Changes in olfaction can serve as an early warning sign of idiopathic Parkinson disease, mild cognitive impairment, and Alzheimer dementia and are associated with increased mortality in older people. 4,10,54-56 Disease severity in Parkinson and Alzheimer disease correlates with the severity of smell and taste loss.^{57,58} More than 95% of patients with Parkinson disease have smell impairment, and up to 75% have anosmia or hyposmia. 10 Olfactory changes can precede the motor symptoms of Parkinson disease by four to six years. Olfactory dysfunction and rapid eye movement sleep behavior disorder are also associated and can be prodromes of Parkinson disease. 10 In mild cognitive impairment and Alzheimer dementia, more dysfunction is noted on objective testing than in subjective reports from patients.4 Deterioration of smell and taste often precedes cognitive deficits. 59,60

MEDICATIONS

Loss of taste and smell is an adverse effect of many medications. ^{18,61} Loss of taste and smell attributed to medication accounts for 12% of olfactory dysfunction, and a greater proportion of gustatory dysfunction may be related to medication. ³⁷ Xerostomia can impair the transmission of tastants to taste buds. Medication-induced loss of smell or taste may be reversible if the medication is discontinued. ^{37,62}

Olfactory and Taste Training

Olfactory training should be offered to patients with persistently poor quality of life due to olfactory dysfunction. 63-65

TABLE 6

Neurodegenerative Diseases Associated With Olfactory Dysfunction

Idiopathic Parkinson disease

Alzheimer dementia

Lewy body dementia

Familial Parkinson disease

Multisystemic atrophy

Huntington disease

Wilson disease

Friedreich ataxia

Spinocerebellar ataxia (types 2 and 3)

Creutzfeldt-Jakob disease

Note: Diseases are listed from most severely affected to least severely affected.

Information from reference 10.

TABLE 7

Sample Instructions for Olfactory Training

Perform the following twice daily (ideally before a meal) for at least four to six months:

Use essential oils soaked in cotton, or liquid (around 50 mL) in an amber sniffing jar.

Use the following recommended scents initially: rose, eucalyptus, lemon, and clove.

Smell the first scent for 15 to 20 seconds; while smelling, focus on what you remember it smelling like.

Rest for 10 seconds.

Repeat as above for each scent separately.

Every one to three months consider changing, or adding, scents, such as menthol, thyme, tangerine, jasmine, green tea, bergamot, rosemary, gardenia; any scent of your choosing; consider scents you wish to smell again that you cannot smell now.

Information from reference 69

Two systematic reviews that included 3,173 patients with postviral olfactory dysfunction found that olfactory training yields objectively measured, clinically significant improvement in olfaction. 43,63 Two other meta-analyses showed objective improvement in nonviral olfactory dysfunction after olfactory training. 64,65 A systematic review and meta-analysis of 13 articles examining olfactory training found that 36% of patients with posttraumatic olfactory dysfunction had clinically significant improvement in measured

Clinical recommendation	Evidence rating	Comments
Patients should be referred to otolaryngology for objective testing of olfactory or gustatory function in refractory cases or if they have depression, poor sleep, dementia, unintentional weight loss, or malnutrition. 3,20,38-40	В	Case series of 602 patients, multiple observational studie
Olfactory training should be offered to patients with persistently poor quality of life due to olfactory dysfunction. ⁶³⁻⁶⁵	A	Consistent evidence from multiple systematic reviews ar meta-analyses, including randomized controlled trials ar prospective cohort studies, showing clinically significant improvement after viruses and other etiologies
Physicians should counsel patients with olfactory dysfunction about the risks of occupational chemical exposure, adherence to food expiration dates, signs of food spoilage, and home fire and explosive gas monitoring. ^{3,40}	С	Expert opinion and a cross-sectional study

olfaction. ⁶⁶ One randomized controlled trial (n = 138) found that adding budesonide nasal irrigation to olfactory training in patients without a visible sign of sinonasal inflammation yielded clinically significant improvement in olfaction (odds ratio = 3.93; 95% CI, 1.20 to 12.88). ⁶⁷ Another randomized controlled trial (n = 42) found taste sensitivity improved after three days of taste training. ⁶⁸ *Table 7* provides olfactory training instructions. ⁶⁹

Patient Education

Physicians should address safety and occupational issues with their patients because hazardous events occur two to three times more often in people with olfactory dysfunction. Patients who work with chemicals may have a reduced ability to identify toxic exposure and should consider mitigation strategies. All patients should take extra care to adhere to expiration dates on food, and to ensure smoke or fire and natural gas or propane alarms are operational. Some patients who are dysosmic may benefit from loved ones' assistance in identifying personal hygiene gaps. Patients and family members may benefit from a support network, such as the Smell and Taste Association of North America (https://thestana.org).

This article updates previous articles on this topic by Malaty and Malaty, 4 and Bromley. 71

Data Sources: A PubMed search was completed in Clinical Queries and the MeSH database using the key terms ageusia, anosmia, assessment, dysgeusia, olfactory disorders, parosmia, smell, taste, taste disorder, test, questionnaire, smoke, survey, and COVID-19. The search included meta-analyses, randomized

controlled trials, clinical trials, books, and systematic reviews. The Cochrane database and Essential Evidence Plus were also searched. Search dates: July 31, 2022; October 5, 2022; October 20, 2022; and July 2, 2023.

The opinions and assertions contained herein are the private views of the authors and are not to be construed as official or as reflecting the views of the U.S. Air Force, the U.S. Department of Defense, or the U.S. government.

The Authors

DILLON J. SAVARD, MD, FAAFP, is director of medical education at David Grant Medical Center, Travis Air Force Base, Calif., and an assistant professor in the Department of Family Medicine at the Uniformed Services University of the Health Sciences, Bethesda, Md.

FRANCESCA G. URSUA, MD, is a resident in the Department of Family Medicine at the University of Nebraska Medical Center, Omaha.

HEIDI L. GADDEY, MD, is a family physician in the Department of Family Medicine at Christie Clinic, Champaign, Ill., and faculty at the Carle Foundation Hospital Family Medicine Residency Program, Urbana, Ill.

Address correspondence to Dillon J. Savard, MD, FAAFP, 101 Bodin Circle, Travis AFB, CA 94535 (dillon.savard@gmail.com). Reprints are not available from the authors.

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