Hyperhidrosis is excessive sweating that affects patients’ quality of life, resulting in social and work impairment and emotional distress. Primary hyperhidrosis is bilaterally symmetric, focal, excessive sweating of the axillae, palms, soles, or craniofacial region not caused by other underlying conditions. Secondary hyperhidrosis may be focal or generalized, and is caused by an underlying medical condition or medication use. The Hyperhidrosis Disease Severity Scale is a validated survey used to grade the tolerability of sweating and its impact on quality of life. The score can be used to guide treatment. Topical aluminum chloride solution is the initial treatment in most cases of primary focal hyperhidrosis. Topical glycopyrrolate is first-line treatment for craniofacial sweating. Botulinum toxin injection (onabotulinumtoxinA) is considered first- or second-line treatment for axillary, palmar, plantar, or craniofacial hyperhidrosis. Iontophoresis should be considered for treating hyperhidrosis of the palms and soles. Oral anticholinergics are useful adjuncts in severe cases of hyperhidrosis when other treatments fail. Local microwave therapy is a newer treatment option for axillary hyperhidrosis. Local surgery and endoscopic thoracic sympathectomy should be considered in severe cases of hyperhidrosis that have not responded to topical or medical therapies. (Am Fam Physician. 2018;97(11):729-734. Copyright © 2018 American Academy of Family Physicians.)

Etiology and Pathophysiology
The cause of primary hyperhidrosis is not well understood. Eccrine sweat glands—distributed throughout the body, but heavily concentrated on the palms, soles, axillae, and face—are innervated by postganglionic autonomic nerve fibers and stimulated by the neurotransmitter acetylcholine. It is thought that increased or aberrant sympathetic stimulation of the eccrine sweat glands is responsible for the increased sweating rather than an increased number or size of the glands. Persons with primary hyperhidrosis have a higher-than-normal basal level of sweat production and an increased response to normal stimuli, such as emotional or physical stress.

Diagnosis
There are no controlled studies on the sensitivity and specificity of the history, physical
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Examination, or testing to accurately diagnose primary hyperhidrosis or to quantify its severity. Criteria for diagnosis include focal, visible, and excessive sweating for longer than six months without apparent cause, and at least two of the following: bilateral and symmetric sweating, impairment of daily activities, occurrence at least once per week, age of onset younger than 25 years, no occurrence during sleep, and a positive family history. 2 Requiring four criteria increases the discrimination between primary and secondary hyperhidrosis (positive predictive value = 0.99; negative predictive value = 0.85). Laboratory testing is not necessary unless history and physical examination suggest a secondary cause. 2

There are several possible secondary causes for excessive sweating. Conditions and medications that can cause excessive sweating are listed in Table 1 and Table 2, respectively. Hyperhidrosis negatively impacts daily life, especially emotional well-being, self-esteem, interpersonal relationships, and occupational productivity. 6,7 Although several tools have been developed to measure the impact of hyperhidrosis on quality of life, most are too complex to incorporate into office practice. The Hyperhidrosis Disease Severity Scale (HDSS) is a validated single-question survey with four grades of tolerability of sweating and impact on quality of life. 3 This survey can estimate the effect on daily activities and response to treatment.

The HDSS is scored as follows: 1 point for sweating that is not noticeable and does not interfere with daily activities; 2 points for sweating that is tolerable but sometimes interferes with daily activities; 3 points for sweating that is barely tolerable and often interferes with daily activities; 4 points for sweating that is not tolerable and interferes with daily activities.

### Table 1: Causes of Secondary Hyperhidrosis

<table>
<thead>
<tr>
<th>Cause</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alcohol use</td>
<td>Chronic pulmonary disease; acute respiratory failure</td>
</tr>
<tr>
<td>Congestive heart failure</td>
<td>Endocrine/metabolic disorders (e.g., diabetes mellitus, thyrotoxicosis, hypoglycemia, hyperpituitarism)</td>
</tr>
<tr>
<td>Endocrine/metabolic disorders</td>
<td>Febrile illness/infection (e.g., defervescence, tuberculosis)</td>
</tr>
<tr>
<td>Gustatory</td>
<td>Medications (Table 2)</td>
</tr>
<tr>
<td>Gustatory</td>
<td>Neurologic (e.g., Arnold–Chiari malformation, Parkinson disease, spinal cord injury)</td>
</tr>
<tr>
<td>Gustatory</td>
<td>Physiologic (e.g., menopause)</td>
</tr>
<tr>
<td>Gustatory</td>
<td>Psychiatric disease (e.g., generalized anxiety disorder, social anxiety)</td>
</tr>
<tr>
<td>Malignancies</td>
<td>Substance abuse; narcotic withdrawal</td>
</tr>
</tbody>
</table>

Information from references 1 and 2.
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Treatment

Disease severity should be measured using the HDSS. Treatment success is defined as a decrease in the HDSS score. Most treatment recommendations are based on expert consensus, because the evidence is poor (e.g., few study participants, nonrandomized trials, often not based on prospective trials). Table 3 summarizes the treatment options for hyperhidrosis based on location and severity.

FIRST- AND SECOND-LINE THERAPIES

First-line treatment of all primary focal hyperhidrosis, regardless of severity, is topical 20% aluminum chloride (Drysol). This solution is applied nightly to the affected areas for six to eight hours until the HDSS score decreases, at which time the application interval can be lengthened to maintain sweat control. The aluminum salts cause an obstruction of the eccrine sweat glands and destruction of the secretory cells. This solution can result in skin irritation, but it can be diluted to decrease irritation if necessary. Use of over-the-counter “clinical strength” antiperspirants containing aluminum zirconium trichlorohydrate has shown a decrease in excessive sweating (as measured by sweat production, not the HDSS score), with less skin irritation than prescription-strength aluminum chloride solutions.

For craniofacial hyperhidrosis, topical 2% glycopyrrolate (compounded by a pharmacy) may be considered first-line treatment. It has shown a 96% success rate (as measured by gravimetric chemical analysis and non-HDSS quality-of-life surveys) with minimal adverse effects (mild skin irritation), and can be applied once every two to three days.

For palmar and plantar hyperhidrosis, iontophoresis may be effective as first- or second-line treatment. Iontophoresis is the passing of an ionized substance, usually water, through the skin by the application of a direct electrical current. Its mechanism of action is unknown. Tap water is poured into the device tray, and then the hands or feet are submerged while a direct electrical current is applied for a specified time, depending on the current. There are three devices registered with the U.S. Food and Drug Administration: RA Fischer MD-1a, RA Fischer MD-2, and Drionic. The procedure can be easily performed at home, and adverse effects (e.g., erythema, vesiculation, tingling) are typically mild and do not require cessation of the treatments. If tap water alone is not effective, adding a tablespoon of baking soda or one or two crushed tablets of the anticholinergic glycopyrrolate (Robinul) to each pan may be beneficial. A detailed description of a recommended application of the procedure is available in the literature.

Botulinum toxin injection is the most studied hyperhidrosis treatment and demonstrates consistent improvement in HDSS scores and in sweat production as measured in the axillae and...
palms.\textsuperscript{3,4} It may be considered first- or second-line therapy for hyperhidrosis affecting the axillae, palms, soles, or face.\textsuperscript{2,4} Botulinum toxins bind synaptic proteins, blocking the release of acetylcholine from the cholinergic neurons that innervate the eccrine sweat glands.\textsuperscript{17} There are several commercially available botulinum toxin preparations approved by the U.S. Food and Drug Administration that are available to physicians who are trained in this procedure. The most commonly used is onabotulinumtoxinA (Botox).\textsuperscript{17}

OnabotulinumtoxinA is administered intradermally in the affected area. It is packaged as a 100-unit vial that is commonly divided into 50 units total for each side. The toxin is injected intradermally in 0.1 mL aliquots per cm\textsuperscript{2}.\textsuperscript{17} It is important to determine the precise area to treat using the Minor starch-iodine test.\textsuperscript{17,18}

For this test, a 3\% to 5\% iodine solution is first applied to the area to be treated and allowed to dry, and then starch is applied. The sweat turns purple when in contact with the iodine and starch, precisely identifying the areas to inject (see a video example of the Minor starch-iodine test and axillary injection at https://www.youtube.com/watch?v=U08PJhRQD0s).\textsuperscript{17,18} In most cases, treatment results last six to nine months.\textsuperscript{17,19,20} Adverse effects typically include injection-site pain and bruising, decreased grip strength when injected into the palms,\textsuperscript{17,19,20} and frontalis muscle weakness when used on the forehead.\textsuperscript{8}

**ALTERNATIVE THERAPIES AND PROCEDURES**

Canadian guidelines recommend oral anticholinergics for treating primary hyperhidrosis with an HDSS score of 3 or 4 that does not resolve with topical aluminum chloride, onabotulinumtoxinA, or iontophoresis.\textsuperscript{4} The most commonly used oral anticholinergic

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### TABLE 3

**Treatment Recommendations for Primary Hyperhidrosis**

<table>
<thead>
<tr>
<th>Location</th>
<th>Mild (HDSS = 2)</th>
<th>Severe (HDSS = 3 or 4)</th>
<th>All severities</th>
</tr>
</thead>
<tbody>
<tr>
<td>Axillary</td>
<td>Step 1: Topical 20% aluminum chloride (Drysol)</td>
<td>Step 1: Topical 20% aluminum chloride or onabotulinumtoxinA injection</td>
<td>Step 3: Consider oral anticholinergics if above treatments fail, alone or in combination with above</td>
</tr>
<tr>
<td></td>
<td>Step 2: If topical treatment fails, onabotulinumtoxinA injection</td>
<td>Step 2: If both treatments fail, consider aluminum chloride and onabotulinumtoxinA in combination</td>
<td>Step 4: Consider microwave therapy</td>
</tr>
<tr>
<td>Craniofacial</td>
<td>Step 1: Topical 20% aluminum chloride or topical 2% glycopyrrolate</td>
<td>Step 1: Topical 20% aluminum chloride plus onabotulinumtoxinA injection or iontophoresis; all three are considered first-line treatment</td>
<td>Step 6 (last resort): Sympathetic denervation (i.e., endoscopic thoracic sympathectomy)</td>
</tr>
<tr>
<td>Palmar</td>
<td>Mild (HDSS = 2)</td>
<td>Severe (HDSS = 3 or 4)</td>
<td>All severities</td>
</tr>
<tr>
<td></td>
<td>Step 1: Topical 20% aluminum chloride</td>
<td>Step 1: Topical 20% aluminum chloride plus onabotulinumtoxinA injection or iontophoresis; all three are considered first-line treatment</td>
<td>Step 2 or 3: Consider oral anticholinergics alone or in combination with above</td>
</tr>
<tr>
<td></td>
<td>Step 2: If topical treatment fails, consider onabotulinumtoxinA or iontophoresis</td>
<td></td>
<td>Step 4 (last resort): Sympathetic denervation</td>
</tr>
<tr>
<td>Plantar</td>
<td>Mild (HDSS = 2)</td>
<td>Severe (HDSS = 3 or 4)</td>
<td>All severities</td>
</tr>
<tr>
<td></td>
<td>Step 1: Topical 20% aluminum chloride</td>
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<td>Step 2 or 3: Consider oral anticholinergics alone or in combination with above</td>
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<tr>
<td></td>
<td>Step 2: If topical treatment fails, onabotulinumtoxinA injection or iontophoresis</td>
<td></td>
<td>Step 3 or 4 (last resort): Sympathetic denervation</td>
</tr>
<tr>
<td></td>
<td>Severe (HDSS = 3 or 4)</td>
<td></td>
<td></td>
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<tr>
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<td>Step 1: Topical 20% aluminum chloride plus onabotulinumtoxinA injection or iontophoresis; all three are considered first-line treatment</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Step 2 or 3: Consider oral anticholinergics alone or in combination with above</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

HDSS = Hyperhidrosis Disease Severity Scale.

Information from references 4, and 8 through 10.
medications are oxybutynin and glycopyrrolate.21 One study showed that oxybutynin, 2.5 to 10 mg per day, decreased excessive sweating and improved HDSS and non-HDSS quality-of-life scores (median = 76% of patients; range = 57% to 100%); however, 75% also experienced dry mouth.21 Although dry mouth is the most common adverse effect, patients may also experience abdominal symptoms, constipation, urinary retention, tachycardia, drowsiness, and blurred vision. On average, 10% of patients stop taking oxybutynin because of adverse effects.21 There is no evidence to quantify the benefit of glycopyrrolate; however, it also has a high prevalence (38%) of dry mouth.21

A newer, noninvasive local treatment of axillary hyperhidrosis uses microwave technology.22 The application of microwave energy destroys eccrine sweat glands by creating local heat, resulting in cellular thermolysis.22 This outpatient procedure is applied with a handheld transducer after mapping the axillae using the Minor starch-iodine test. Local anesthesia is required.22 This treatment results in a decrease in the HDSS score of at least one point in 94% of patients and at least two points in 55% of patients.23

Another emerging treatment in axillary hyperhidrosis is fractionated microneedle radiofrequency.24 During this procedure, microneedles are placed 2 to 3 mm under the skin, and radiofrequency energy is applied. This therapy results in a decrease in the HDSS score of at least one point in nearly 80% of patients.24,25

Local surgical therapy has been used to treat axillary hyperhidrosis. Techniques include radical surgical excision (rarely used because of high complication and recurrence rates), limited skin excision, liposuction, curettage, and liposuction-curettage.26,27 Although these techniques can initially reduce measured axillary sweating, they have high relapse rates several months after the procedure.26,27

Because hyperhidrosis is thought to be secondary to excessive sympathetic stimulation, endoscopic thoracic sympathectomy has been used to treat severe cases of hyperhidrosis.28 This procedure, which has evolved from an open procedure to an endoscopic one, involves cutting or clipping sympathetic nerves.20 Referral for endoscopic thoracic sympathectomy may be indicated when less invasive therapies are ineffective.4,9,28

Although the procedure decreases or eliminates sweating in the original problem area, a common late complication is compensatory sweating in other areas, usually in the abdomen, back, gluteal region, and legs.29 This article updates a previous article on this topic by Thomas, et al.30

Data Sources: The authors searched the Agency for Healthcare Research and Quality Evidence Reports, Cochrane Database of Systematic Reviews, National Guideline Clearinghouse, Essential Evidence Plus, and PubMed clinical queries using the following keywords: hyperhidrosis, excessive sweating, primary hyperhidrosis, secondary hyperhidrosis. Search date: September 24, 2017.

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References

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