

Risheng Xu, DO

CHI St. Luke's Health-The Vintage Hospital, Houston, TX; Assistant Professor, Department of Medicine, Baylor College of Medicine, Houston, TX

Nicola Abate, MD

Bay Area Metabolic Health, Houston, TX

Nalini Ram, MD

Professor, Department of Medicine, Division of Endocrinology, Baylor College of Medicine, Houston, TX

Kristina Little, MD

Assistant Professor, Department of Medicine, Division of Geriatrics, Baylor College of Medicine, Houston, TX

Most elderly patients with subclinical hypothyroidism do not need to be treated

ABSTRACT

Whether subclinical hypothyroidism should be treated in elderly patients (≥ 65 years) is controversial. The authors argue for a personalized, wait-and-see approach rather than universal treatment, pointing out that randomized clinical trials have not shown that levothyroxine treatment makes any difference in terms of hard clinical end points, quality of life, or hypothyroid symptom relief in elderly patients with this condition.

KEY POINTS

Subclinical hypothyroidism is categorized as either mild (grade 1; thyroid-stimulating hormone [TSH] level 4.0–10.0 mIU/L) or severe (grade 2; TSH > 10 mIU/L).

High TSH levels in patients older than 65 years may be due to aging and do not necessarily require treatment.

The French Endocrine Society proposes using the patient's age divided by 10 as the upper limit of normal for TSH (in mIU/L) when screening and following elderly patients.

Studies have not found levothyroxine replacement therapy to make any significant clinical difference in most cases of mild subclinical hypothyroidism in older patients. However, lack of consistent age-appropriate TSH screening methods limits definitive conclusions.

In most mild cases, TSH can be remeasured 2 to 3 months after diagnosis. Treatment decisions are individualized, and potential risks and benefits must be considered.

doi:10.3949/ccjm.92a.24098

MOST PATIENTS WITH SUBCLINICAL hypothyroidism do not need to be treated for it, and for many, it may be a normal part of aging and can be monitored without active intervention.

The US Preventive Services Task Force defines subclinical hypothyroidism as an elevated serum thyrotropin (thyroid-stimulating hormone, TSH) level (> 4.50 mIU/L), but with a normal free thyroxine (T4) level.¹ Despite the term *subclinical*, symptoms may or may not be present, although they tend to be mild and nonspecific.

See related article page 233

Guidelines for diagnosing and managing overt hypothyroidism (in which the TSH level is elevated and the T4 level is low) enjoy broad consensus.^{2,3} However, whether to treat subclinical hypothyroidism is controversial, especially in people 65 and older. A host of factors, including age, can affect TSH levels. Adding to the challenge, the ideal TSH cutoff point for initiating treatment remains a topic of debate.⁴ Mildly elevated TSH does not necessarily lead to long-term adverse consequences, and overtreatment can increase the risks of fractures, cardiovascular disease, and dysrhythmias.⁵

Here, we review the challenges of managing subclinical hypothyroidism in the elderly (age 65 and older) and argue against routinely treating it with levothyroxine in this age group. We do not cover how to manage it in younger adults, women of childbearing age, or children.

TABLE 1

Advantages and disadvantages of different thyroid function tests

Test	Advantages	Disadvantages	Preferred use
Thyrotropin (thyroid-stimulating hormone, TSH) ⁷	Sensitive to changes in thyroid hormone levels Useful in diagnosing primary and secondary hypothyroidism	Can be influenced by nonthyroid factors (eg, illness, medications)	Primary test for diagnosing hypothyroidism and monitoring thyroid hormone replacement therapy
Free thyroxine (T4) ^{2,8}	More accurate reflection of thyroid hormone levels than total T4, less influenced by changes in binding proteins	Can be affected by protein-binding changes, especially in certain conditions (eg, pregnancy, liver disease)	Essential for assessing thyroid hormone status, especially in conditions affecting thyroid-binding proteins
Free triiodothyronine (T3) ^{7,9}	More accurate reflection of thyroid hormone activity than total T3, less influenced by changes in binding proteins	Lower concentration, weaker protein binding, less precise measurement Often normal in early and mild hypothyroidism due to mostly peripheral conversion, especially in early hypothyroidism	Consider in specific clinical scenarios, such as suspected nonthyroid illness syndrome or hyperthyroidism
Thyroglobulin	Elevated levels may indicate residual or recurrent thyroid cancer	Can be affected by thyroid inflammation and other factors	Used in the management of patients with thyroid cancer
Thyroid peroxidase antibodies	Positive test suggests autoimmune thyroiditis; often associated with Hashimoto thyroiditis	Not specific—can be positive in other autoimmune conditions	Used in the diagnosis and management of autoimmune thyroid diseases

■ PRIMARILY A LABORATORY DIAGNOSIS

The clinical presentation of hypothyroidism often deviates from the classic textbook description. Carlé et al⁶ report that tiredness is the most important symptom of overt hypothyroidism. However, many patients with subclinical hypothyroidism experience no symptoms or only nonspecific symptoms. Consequently, subclinical hypothyroidism is primarily a biochemical or laboratory diagnosis.

TSH is more sensitive than T4

TSH levels have a log-linear inverse relationship with T4 and triiodothyronine (T3) levels, so that a 2-fold decrease in T4 results in a 100-fold increase in TSH.⁷ Therefore, TSH is more sensitive to changes in thyroid function.

Free T4 measurements, complementing TSH, help classify hypothyroidism as overt or subclinical and determine the need for treatment. Free T4 is a better marker of hormone action than total T4, as the latter is 99.97% bound to proteins (thyroid-binding globulin, transthyretin, prealbumin, and albumin), and condi-

tions that alter these binding proteins affect the accuracy of total T4 measurements.⁸ Also, unbound (free) T4 is the biologically active fraction and is widely accepted as a better activity marker.²

T3 is 99.7% protein-bound, and the free T3 level (the other 0.3%) is theorized to be a better marker than total T3. However, free T3 has a lower concentration than free T4 and a weaker affinity for protein carriers, which renders it more susceptible to free fatty acids (which inhibit its binding to its receptor) and drug interactions.⁹ Consequently, the precision and reproducibility of free T3 measurements are less ideal than those of free T4. Also, T3 levels tend to be within reference ranges in cases of suspected hypothyroidism or elevated TSH and so have limited clinical value (Table 1).^{2,7-9} As a result, in the special cases in which T3 measurements are needed, such as euthyroid sick syndrome, many laboratories prefer total T3 assays rather than free T3.

The American^{2,3} and European thyroid associations¹⁰ classify subclinical hypothyroidism as either mild (grade 1) or severe (grade 2) based on the TSH

level (Table 2)—about 90% of patients with subclinical hypothyroidism have TSH levels of 10 mIU/L or lower (ie, mild or grade 1).

Before diagnosing subclinical hypothyroidism, one must make sure the thyroid function has been stable for at least several weeks to exclude transient changes caused by nonthyroidal illness, thyroiditis, or medications.² We remeasure TSH 2 to 3 months after the first measurement.

COMMON IN THE ELDERLY

Western countries are getting older. Consequently, the prevalence of subclinical hypothyroidism is also rising, reflecting the growing proportion of the population at risk.¹¹

In the third National Health and Nutrition Examination Survey (NHANES III),¹² in people age 12 years and older in the United States, the prevalence of subclinical hypothyroidism was 4.3% and the prevalence of overt hypothyroidism, characterized by low free T4 levels, was 0.3%. In the Cardiovascular Health Study,¹³ in participants age 65 and older, the prevalence of subclinical hypothyroidism was 15%, and it was higher in women than in men.

MANY FACTORS AFFECT TSH LEVELS

When T4 and T3 levels are low, the pituitary gland releases TSH to increase thyroid hormone production, whereas high T4 and T3 levels inhibit TSH release. However, many other factors can affect the TSH level, making abnormal values difficult to interpret (Table 3).^{14–21} TSH levels are not static but fluctuate within individuals over time,¹⁷ owing to a range of factors:

- **Circadian rhythm.** TSH levels peak in the early morning hours and then decline, reaching a trough in the early afternoon and evening
- **Pulsatile secretion.** TSH is secreted in pulses, contributing to further fluctuations
- **Season.** Levels are generally higher in winter and lower in summer
- **Age.** TSH levels mildly increase with age, and elevations may be normal for some individuals^{12,18}
- **Other factors,** which include inconsistencies in testing methods, thyroid antibodies and thyroid hormone resistance,^{11,19,20} silent or granulomatous thyroiditis, and certain medications.²¹

Genetic differences also affect thyroid hormone levels, so that some patients whose TSH levels are within the normal range can have symptoms while others with slightly elevated TSH might not.²² Also, iodine intake varies in geographic regions.

TABLE 2
Classification of subclinical hypothyroidism

	Thyroid-stimulating hormone range (mIU/L)	Free T4 range (ng/dL)
Grade 1	4.0–10.0	Normal (0.9–1.7)
Grade 2	> 10.0	Normal (0.9–1.7)

Based on information from references 2 and 10.

The complex interplay of factors that influence thyroid hormone regulation makes the distinction between subclinical hypothyroidism and overt hypothyroid disease arbitrary.

IS THE UPPER LIMIT OF NORMAL FOR TSH TOO HIGH?

The true upper limit of normal of TSH to diagnose subclinical hypothyroidism is still debated.¹⁴

Currently, the normal reference range for TSH is extrapolated from population-based data, calculated from Gaussian distribution with 95% confidence intervals of studies of people who do not have thyroid disease, thyroid peroxidase antibodies, or thyroglobulin antibodies.²³ In most studies, the lower limit of TSH (the 2.5th percentile) was between 0.2 and 0.4 mIU/L, but the upper limit (the 97.5th percentile) varied between 2.4 and 4.2 mIU/L.²⁴

Why was this range so wide? Many people in the studies actually had undiagnosed autoimmune thyroid disease (Hashimoto thyroiditis) or other factors that affect TSH, such as medications, nonthyroidal illness, pregnancy, subacute thyroiditis in the recovery phase, heterophilic antibodies, thyroid hormone resistance, TSH-receptor mutations, bio-inactive TSH, and TSH-producing pituitary tumors, or had their blood drawn at a high point in their circadian rhythm.²⁵ This renders the TSH distribution non-Gaussian, with a tail at the upper end.²⁶ If we take out these confounding factors, and a normal Gaussian distribution with a bell-shaped curve is achieved, the normal reference range becomes 0.4 to 2.5 mIU/L.²⁷

In view of these studies, some have proposed lowering the upper limit of normal from 4.5 to 2.5 mIU/L.^{24,27} However, Surks et al²⁸ argue against changing the upper limit, estimating that if we lower the upper limit of normal for TSH to 3.0 mIU/L to screen elderly patients with vague symptoms, it will label 22 to 28 million

TABLE 3

Factors that can affect serum thyroid-stimulating hormone (TSH) levels

Factor	Explanation
Time of day	TSH levels naturally fluctuate throughout the day, peaking in the early morning hours
Season of year	TSH levels may be slightly higher in winter than in summer, potentially due to changes in sunlight exposure or other environmental factors
Stress	Can temporarily suppress TSH production
Illness	Infections or autoimmune diseases can disrupt the hypothalamic-pituitary-thyroid axis, leading to changes in TSH levels, depending on the severity or duration of the illness
Medications	Some medications, particularly those used to treat thyroid disorders or other conditions, can influence TSH levels
Interindividual variation	There can be significant individual differences in TSH patterns, even among healthy individuals; genetic factors and personal characteristics may play a role
Age	TSH levels tend to increase with age, particularly in older adults
Sex	Some studies suggest that there may be sex-specific differences in TSH regulation, so that women tend to have higher TSH levels than men ¹⁴
Autoimmunity	Autoimmune thyroid diseases such as Hashimoto thyroiditis can lead to elevated TSH levels, particularly in the early stages of the disease

Based on information from references 14–21.

more Americans as having subclinical hypothyroidism without evidence-based therapeutic benefit from this diagnosis. Instead, they suggest measuring the TSH level every 1 to 2 years in patients whose TSH levels are between 2.5 and 4.5 mIU/L.

■ WHY DOES TSH RISE WITH AGE?

Why TSH increases with age is not known.²⁹ In NHANES III,¹² TSH levels increased with age, and, interestingly, so did the prevalence of thyroid peroxidase antibodies and thyroglobulin antibodies. Some 14% of people older than 85 years had TSH levels higher than 4.5 mIU/L.

Hashimoto thyroiditis is the most common condition associated with subclinical hypothyroidism in the elderly, and in almost 90% of cases is characterized by antibodies against thyroid peroxidase and thyroglobulin.¹¹ Patients who had these antibodies progressed to having overt hypothyroidism at the rate of 4.3% per year, compared with only 2.6% in those without these antibodies.³⁰

Once these antibodies are present, however, changes in their levels do not add more information while monitoring subclinical hypothyroidism, due to parallel fluctuation of TSH and thyroid peroxidase antibody.³¹ These antibodies are not harmful to the thyroid glands; however, the volume of thyroid glands generally shrinks after 50 years of age, and pathology studies have found lymphocytic infiltration of the gland

and fibrosis.³¹ Thus, aging-associated thyroid cellular damage from cellular and humoral immune mechanisms with possible T lymphocytes was suspected.^{11,32} Changes in iodine absorption and organification have been observed in elderly patients.³³ In addition, the normal nocturnal surge in TSH (possibly related to maintenance and repair mechanisms) is partially or completely lost. Thus, nighttime TSH levels are lower but 24-hour TSH levels are higher, to keep the T4 level normal.¹⁷ Corticosteroid inhibitory function is also compromised in the elderly and leads to decreased hypothalamic-pituitary-thyroid inhibitory function.¹¹

While a direct correlation between aging and mild thyroid failure is debated, there is evidence to suggest that some aspects of aging may resemble the symptoms and physiologic changes associated with mild hypothyroidism.¹¹ Whether this is a protective mechanism in the elderly or rather represents a diseased hypothalamic-pituitary-thyroid axis is debated.³⁴ In a study in southern Italy, Corsonello et al³⁵ found that older people had lower T4 and T3 levels and higher TSH levels—and so did the children and nieces and nephews of centenarians, suggesting that these changes might be associated with longevity.

Also, elderly patients have less circadian variation in TSH and a weaker response to thyrotropin-releasing hormone compared with the young, changes that are believed to conserve energy, particularly in those with reduced physical activity.^{11,18}

■ IS SUBCLINICAL HYPOTHYROIDISM HARMFUL? IS TREATMENT BENEFICIAL?

Findings have been inconsistent regarding the potential harms of untreated subclinical hypothyroidism, and evidence that treatment is beneficial is lacking.

Cardiovascular disease

Thyroid hormones play a crucial role in regulating various bodily functions, glucose metabolism, protein synthesis, and cardiovascular function. However, the impact of thyroid hormones on the cardiovascular system, particularly in older adults with subclinical hypothyroidism, is inconsistent.¹¹

In the Rotterdam Study,³⁶ TSH levels higher than 4.0 mIU/L with normal free T4 were associated with higher risks of myocardial infarction (odds ratio 2.3) and aortic calcification (odds ratio 1.7). When antithyroid antibodies were present, patients with subclinical hypothyroidism had greater progression to more severe atherosclerosis than euthyroid individuals.

Japanese men, average age 58.5 years, with TSH higher than 5.0 mIU/L and normal free T4 had a high risk of ischemic heart disease (odds ratio 4.0).³⁷ In Taiwanese people age 20 and older with subclinical hypothyroidism followed for 10 years, the all-cause mortality rate was 30% higher, and the rate of death due to cardiovascular disease was 68% higher than in euthyroid people.³⁸

In contrast, a study that analyzed data from the Cardiovascular Health Study¹³ found no links between subclinical hypothyroidism and adverse cardiovascular, cerebrovascular, or mortality outcomes. However, the range of TSH values was wide: 4.5 to 20.0 mIU/L.

A meta-analysis³⁹ found significantly greater risks for coronary heart disease morbidity and mortality in those with TSH levels 10 mIU/L and higher, and the risks increased significantly in those age 65 to 79 years, but not at age 80 or above, compared with euthyroid participants. In other meta-analyses that used data from the Thyroid Studies Collaboration cohort,^{40–42} there was no association between subclinical hypothyroidism (with TSH levels in the range of 4.5–19.9 mIU/L) and stroke, atrial fibrillation, or heart failure.¹⁹ Overall, the mortality rate was not higher than in euthyroid patients. However, the subgroup with TSH levels 10 mIU/L or higher had higher risks of heart failure, coronary heart disease events, and death from coronary heart disease than the euthyroid group. In addition, the risks of death from stroke and coronary heart disease were higher in those with TSH levels of 7.0 to 9.9 mIU/L.^{39,40}

In a post hoc analysis of the Prospective Study of Pravastatin in the Elderly at Risk,⁴⁴ patients with sub-

clinical hypothyroidism (age range 70–82 years) had significantly higher rates of heart failure and death, but only those with TSH levels higher than 10 mIU/L.

Razvi et al⁴⁵ performed a meta-analysis and found no difference in ischemic heart disease morbidity or mortality in patients age 65 or older with subclinical hypothyroidism, but did find increased mortality in patients younger than 65 years.

Cognitive dysfunction and depression

Thyroid hormones are essential for brain development; however, whether subclinical hypothyroidism affects cognition in elderly patients is questioned, and studies of this matter are inconclusive.¹¹

Pasqualetti et al⁴⁶ conducted a meta-analysis of prospective and cross-sectional studies and found that subclinical hypothyroidism was associated with cognitive impairment in patients younger than 75, but not older. However, in the 2015 US Preventive Services Task Force study, Rugge et al¹ reviewed trials of treatment vs placebo for subclinical hypothyroidism (TSH 4.5–10 mIU/L or ≥ 10 mIU/L) and found no effects on cognitive skills or performance, speed or capacity of language processing, or psychomotor tests of executive functions. They rated the overall quality of evidence as “poor.”

Dyslipidemia

Rugge et al¹ found only small changes in the average total cholesterol and low-density lipoprotein cholesterol levels with treatment, and no significant differences in high-density lipoprotein cholesterol levels or triglyceride levels between the treated and untreated groups.

Blood pressure

Rugge et al¹ found no evidence that treating subclinical hypothyroidism affected blood pressure in patients with TSH levels higher than 3.6 mIU/L.

Mortality

Some studies¹³ found no significant association between subclinical hypothyroidism and mortality.

Quality of life and symptoms

The only large study that delved into hypothyroid symptoms such as dry skin, poor memory, cognitive slowing, muscle weakness, cold feelings, voice changes, constipation, and puffy eyes was the cross-sectional Colorado Thyroid Disease Prevalence Study.⁴⁷ Interestingly, 25% of individuals with overt hypothyroidism, 20% of those with subclinical hypothyroidism, and 17% of euthyroid individuals reported 4 or more symptoms commonly associated with hypothyroidism.

The Thyroid Hormone Replacement for Untreated Older Adults With Subclinical Hypothyroidism (TRUST) trial⁴ included 737 adults 65 or older with TSH levels ranging from 4.6 to 19.99 mIU/L and normal free T4 levels, who were given levothyroxine (median dose 50 µg) or placebo. At 12 months, in the treated group, TSH levels had declined from 6.40 mIU/L to 3.63 mIU/L, but there was no clinical difference in the mean changes of hypothyroid symptoms score or in the tiredness score. These findings raise the question of whether subclinical hypothyroidism is a true disease or just a biochemical cutoff.

Kidney function

In theory, thyroid deficiency could affect kidney function by lowering cardiac output, although evidence does not support an association between subclinical hypothyroidism and kidney dysfunction.⁴⁸ However, patients on hemodialysis who had subclinical hypothyroidism had a higher mortality rate.⁴⁹

Neuromuscular effects

Although there have been suggestions that neuromuscular symptoms are common in patients with subclinical hypothyroidism, more definitive answers are needed for those with TSH 10 mIU/L or below.⁵⁰ In several studies in elderly patients, there was no association between subclinical hypothyroidism and low bone mineral density, fracture risk, or frailty compared with euthyroid patients (reviewed by Biondi et al¹⁹).

HARMS OF OVERTREATMENT

Many patients with hypothyroidism are being overtreated. Biondi and Cooper²⁶ estimate that the overtreatment rate of subclinical hypothyroidism exceeds 20%. This overtreatment can lead to low TSH levels, potentially causing thyrotoxicosis in elderly patients. In 2 studies, 22% to 23% of patients on levothyroxine replacement had suppressed TSH levels.^{47,51} At the Massachusetts General Hospital Thyroid Clinic, 14% of patients on levothyroxine therapy had suppressed TSH levels.⁵²

Subclinical or overt thyrotoxicosis can lead to atrial fibrillation, cardiovascular events, reduced bone density, fractures, and increased mortality. A study in Denmark⁵³ followed a cohort of patients with overt and subclinical hypothyroidism every 6 months for a median of 7.2 years and found that the mortality rates of those who were overtreated with levothyroxine so that their TSH levels were lower than 0.3 mIU/L increased with duration of overtreatment.

GUIDELINES AND RECOMMENDATIONS

Several medical organizations have published guidelines for diagnosing and managing subclinical hypothyroidism. These differ somewhat, as medical research evolves and there is as of yet no universal consensus.

The European Thyroid Association, in its 2013 guideline,¹⁰ recommends classifying subclinical hypothyroidism as mild or severe according to TSH level; if mild, it suggests no treatment but repeating this measurement 2 to 3 months later, along with thyroperoxidase antibodies. If the patient is younger (< 65–70) and has TSH higher than 10 mIU/L, treatment should be started. If a patient in this younger group has TSH lower than 10 mIU/L and normal free T4 but has symptoms, a trial of thyroxine can be considered. If the TSH level is normal after 3 to 4 months of thyroid replacement therapy, then symptoms should be reevaluated and therapy should be stopped if no symptomatic change is observed.

In older people, the European Thyroid Association recommends using age-specific local reference ranges for TSH to diagnose subclinical hypothyroidism (eg, TSH 4–7 mIU/L for those older than 80 years).¹⁰ In patients older than 80 to 85 years with TSH 10 mIU/L or less, a wait-and-see strategy should be prioritized before starting replacement therapy, as deleterious effects of subclinical hypothyroidism on cardiovascular risk vanish in this group. If treatment is to be started in older patients (> 65–70 years), the target TSH range should be the lower half of the reference range, or 0.4 to 2.5 mIU/L. For patients without cardiac disease, a weight-based dose of 1.5 µg/kg/day should be used. In patients with pre-existing cardiac disease, they say to start levothyroxine treatment low at 25 to 50 µg/day and increase it every 2 to 3 weeks to slowly bring TSH down to target.

In patients with persistent subclinical hypothyroidism, thyroid function tests should be checked every 6 months for the first 2 years, and then annually.

The American Thyroid Association and American Association of Clinical Endocrinology 2012 guideline² and their 2014 update³ echo the European classification of mild vs severe subclinical hypothyroidism. In severe subclinical hypothyroidism, thyroxine treatment is generally supported, similar to the European recommendation. However, the American guideline is less specific regarding mild subclinical hypothyroidism. It recommends considering individual factors such as symptoms, thyroid peroxidase antibody status, and evidence of cardiovascular disease, especially in patients younger than 65 years.

The 2014 American Thyroid Association update³ recommends starting low and titrating slowly in elderly

TABLE 4
Guidelines and recommendations on managing subclinical hypothyroidism

Organization	Mild subclinical hypothyroidism (TSH 4.0–10.0 mIU/L)	Severe subclinical hypothyroidism (TSH > 10.0 mIU/L)	Key comments
US Preventive Services Task Force (2015) ¹	Does not recommend routine screening or treatment	Recommends treatment	Insufficient evidence of benefit of treatment in patients without symptoms
European Thyroid Association (2013) ¹⁰	Watchful waiting, unless symptomatic Consider treatment in younger individuals with persistent symptoms	Recommends treatment	Emphasizes a cautious approach, especially in older adults
American Association of Clinical Endocrinology and American Thyroid Association (2012) ²	Does not recommend routine screening or treatment in patients without symptoms Consider treatment in select cases based on individual factors	Recommends treatment	Highlights individualizing treatment decisions and considering factors such as age, comorbidities, and symptom burden
American Thyroid Association (2014 update) ³	Similar to 2012 guidelines, emphasizing individualized approach	Recommends treatment, particularly in younger individuals and those with specific risk factors	Reinforces the importance of considering individual factors and the potential benefits of treatment in certain cases
American College of Physicians (2019) ⁵⁵	Does not recommend routine screening or treatment unless symptoms are present	Recommends treatment	Emphasizes individualizing treatment based on patient factors
American Academy of Family Physicians (2021) ⁵⁴	Does not recommend routine screening or treatment unless symptoms are present	Recommends treatment	Highlights the lack of evidence for routine screening and treatment in patients without symptoms

TSH = thyroid-stimulating hormone

patients with hypothyroidism, and recommends raising the TSH target range to 4 to 6 mIU/L in persons 70 to 80 years old. It also cautions against iatrogenic thyrotoxicosis from overtreatment with levothyroxine, especially avoiding TSH levels lower than 0.1 mIU/L in older and postmenopausal women to avoid cardiac and bone deleterious effects.

Other organizations^{22,54–56} have also issued recommendations regarding hypothyroidism diagnosis and treatment. Although these recommendations are not specific to subclinical hypothyroidism in the elderly, they generally concur with those above (Table 4).^{1–3,10,54,55}

MANAGEMENT CHALLENGES

There are no uniform national guidelines on screening elderly patients with blood TSH levels for thyroid disease.

Although the association between increasing TSH and aging is well described,^{2,10,47} the practice of using age-adjusted TSH reference ranges to diagnose elderly patients with thyroid diseases is not currently routine.⁵⁷ As a result, the current TSH normal range (0.4–4.5 mIU/L) is inappropriately used to diagnose subclinical hypothyroidism in patients older than 65 years, and this overestimates the prevalence of subclinical hypothyroidism and also leads to overtreatment.

Moreover, Danese et al⁵⁸ estimated the cost of routine screening in women older than 35 years every 5 years at approximately \$9,200 per quality-adjusted year of life (this was in 1996, and the cost would be higher now), raising concerns about the feasibility and potential drawbacks of universal screening.

The concept of using a TSH cutoff value can also be deceiving. The test is based on a sample of healthy volunteers without thyroid disease or thyroid antibodies,

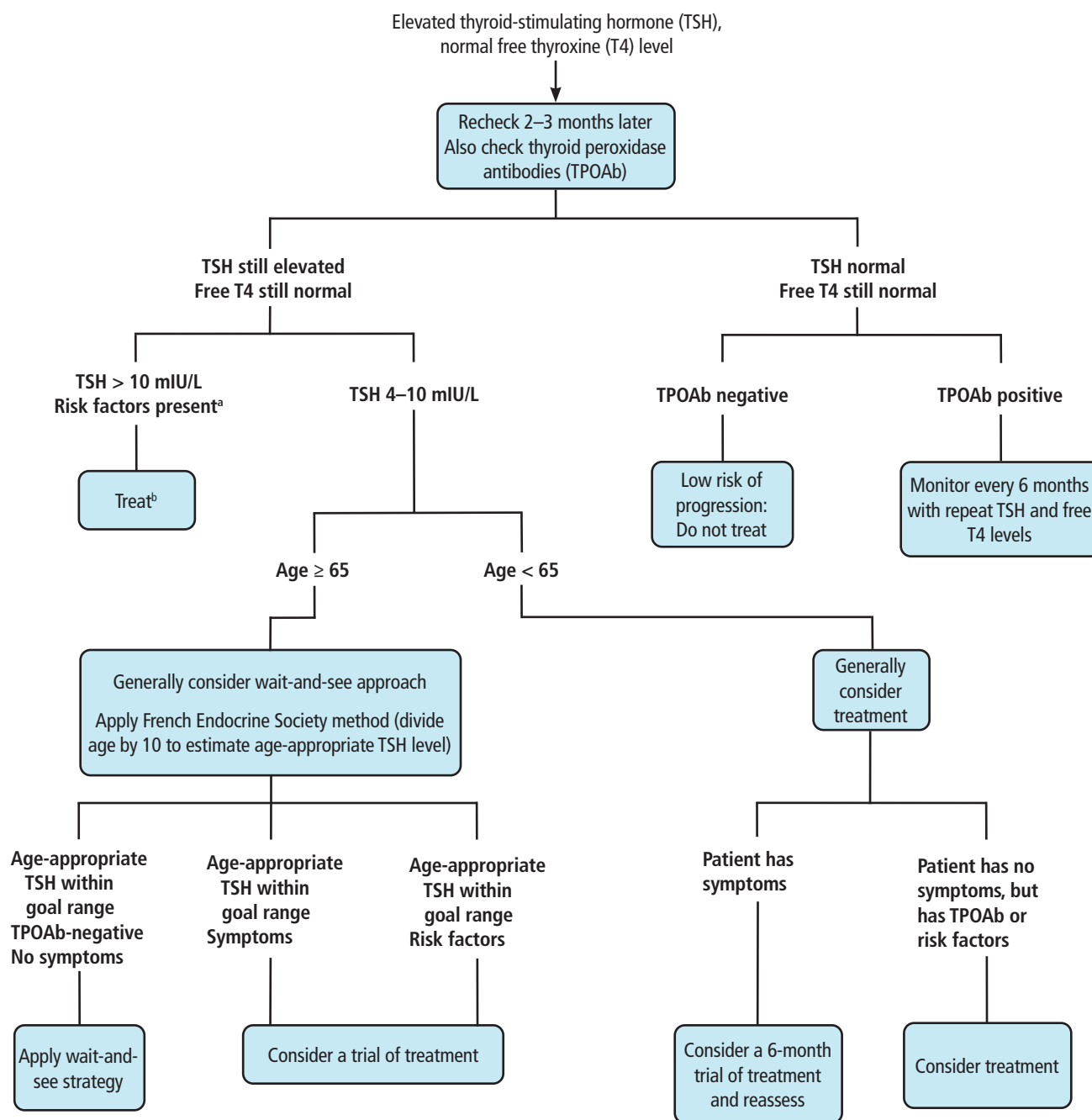


Figure 1. Flow chart for clinical decision-making in subclinical hypothyroidism.

^aRisk factors: TPOAb-positive, goiter, atherosclerotic cardiovascular disease, heart failure, or associated risk factors for these diseases.

^bOral levothyroxine daily is the treatment of choice. For patients with cardiac disease, 1.5 µg/kg/day should be used. For elderly patients with cardiac disease, a dose of 25–50 µg/day is recommended. Increase dose by 12.5 to 25 µg/day every 2 to 3 weeks. Target TSH range is 0.4 to 2.5 mIU/L.

representative of a larger population and calculated with statistical methodology. However, many factors can affect this arbitrary reference range (Table 3).^{17,59} For example, studies in urban populations over age 55 without thyroid peroxidase antibodies or other factors show higher TSH levels in White and Mexican American individuals than in Black individuals.^{12,21} Scobbo et al⁶⁰ demonstrated that TSH fluctuates throughout the day, being significantly higher before breakfast (7:30–9:00 AM) than after breakfast (10:30 AM–12:00 PM), with an average decline of 26.4%.

Another problem with the use of a TSH cutoff to define subclinical hypothyroidism is that TSH changes can be transient and reversible. In a large retrospective longitudinal study,⁶¹ 57.9% of patients with subclinical hypothyroidism reverted to euthyroid TSH levels during a median follow-up of 36 months. Unfortunately, most of the studies dealing with risks and benefits associated with subclinical hypothyroidism included patients who had no repeat TSH assessments to confirm the persistence of TSH elevation.^{21,29} This raises the question of inconsistency in the benefits and risks of treating subclinical hypothyroidism. On top of this, observational studies did not address TSH status while receiving treatment.¹³ All of these uncertainties contribute to the debate about the clinical relevance of subclinical hypothyroidism, especially in grade 1 subclinical hypothyroidism, and in older patients.

In summary, factors such as the time of day, ethnicity, and inconsistent TSH levels complicate accurate subclinical hypothyroidism diagnosis. Varied study methodologies and a lack of longitudinal TSH data further obfuscate the issue. The absence of age-appropriate diagnostic guidelines and consensus makes establishing clear evidence-based benefits difficult.

■ PHYSICIAN DECISION-MAKING AND ALTERNATIVES

While some studies have linked untreated subclinical hypothyroidism to cardiovascular issues and heart failure, the US Preventive Services Task Force reviewed multiple studies and found no significant clinical benefit from treatment in grade 1 subclinical hypothyroidism.¹ The 2017 TRUST randomized clinical trial supported these findings.⁴

To personalize the diagnosis and treatment, the French Endocrine Society⁵⁶ proposed a novel approach in their 2019 consensus statement—using the formula patient age divided by 10 to establish the upper limit of normal for TSH in patients older than 60. This approach acknowledges the potential for natu-

ral changes in TSH levels with age. Consequently, for an 80-year-old, the upper limit for TSH would be 8 mIU/L, which differs from the standard cutoff used for younger populations.

Recently, Kuś et al¹⁶ reported a novel approach to personalizing TSH intervals: using a polygenic score system based on 59 genetic variants to calculate TSH reference ranges. This predicted 9.2% to 11.1% of total variance in TSH concentrations, compared with 2.4% to 2.7% with free T4. This reflects the idea that different individuals have different set points in their hypothalamic-pituitary-thyroid axis, and thus can have higher TSH concentrations at the same free T4 levels.⁵⁹

In older patients, Calsolaro et al¹¹ recommend treating severe subclinical hypothyroidism with thyroxine. However, in patients with mild subclinical hypothyroidism, starting levothyroxine can be considered on a case-by-case basis according to symptoms, frailty, cardiovascular risk factors, and personal preferences. If a patient needs treatment, they recommend following the professional guidelines. Alternatively, mild subclinical hypothyroidism can be monitored without therapy by repeating laboratory tests every 3 to 6 months to monitor progression of overt disease and treated if TSH rises to more than 10 mIU/L.^{2,10,58}

We propose an approach that combines the professional society guidelines and the French Endocrine Society 2019 consensus statement⁵⁶ with an age-based TSH cutoff (Figure 1). When the decision becomes complicated, consultation with a team including an endocrinologist and geriatrician is recommended.

■ A PERSONALIZED APPROACH

Subclinical hypothyroidism is common in the elderly and is going to be more so in the coming decades. Its management requires a personalized approach that weights the potential benefits and risks, as current evidence does not show a clear difference in outcomes if mild subclinical hypothyroidism is treated or not. Age-adjusted TSH reference ranges, consideration of individual circumstances, and a wait-and-see approach for mild cases might be more suitable than a universal treatment strategy. ■

Acknowledgments: The authors thank their colleagues in the divisions of Geriatrics and Endocrinology for their insights and recommendations on this topic and their efforts in reviewing and strengthening the content of this manuscript.

■ DISCLOSURES

The authors report no relevant financial relationships which, in the context of their contributions, could be perceived as a potential conflict of interest.

REFERENCES

- Rugge JB, Bougatsos C, Chou R. Screening and treatment of thyroid dysfunction: an evidence review for the US Preventive Services Task Force. *Ann Intern Med* 2015; 162(1):35–45. doi:10.7326/M14-1456
- Garber JR, Cobin RH, Gharib H, et al. Clinical practice guidelines for hypothyroidism in adults: cosponsored by the American Association of Clinical Endocrinologists and the American Thyroid Association [published correction appears in *Endocr Pract* 2013; 19(1):175]. *Endocr Pract* 2012; 18(6):988–1028. doi:10.4158/EP12280.GL
- Jonklaas J, Bianco AC, Bauer AJ, et al. Guidelines for the treatment of hypothyroidism: prepared by the American Thyroid Association Task Force on Thyroid Hormone Replacement. *Thyroid* 2014; 24(12):1670–1751. doi:10.1089/thy.2014.0028
- Stott DJ, Rodondi N, Kearney PM, et al. Thyroid hormone therapy for older adults with subclinical hypothyroidism. *N Engl J Med* 2017; 376(26):2534–2544. doi:10.1056/NEJMoa1603825
- Flynn RW, Bonellie SR, Jung RT, MacDonald TM, Morris AD, Leese GP. Serum thyroid-stimulating hormone concentration and morbidity from cardiovascular disease and fractures in patients on long-term thyroxine therapy. *J Clin Endocrinol Metab* 2010; 95(1):186–193. doi:10.1210/jc.2009-1625
- Carlé A, Pedersen IB, Knudsen N, et al. Hypothyroid symptoms and the likelihood of overt thyroid failure: a population-based case-control study. *Eur J Endocrinol* 2014; 171(5):593–602. doi:10.1530/EJE-14-0481
- Hoermann R, Pekker MJ, Midgley JEM, Larisch R, Dietrich JW. Triiodothyronine secretion in early thyroid failure: the adaptive response of central feedforward control. *Eur J Clin Invest* 2020; 50(2):e13192. doi:10.1111/eci.13192
- Oppenheimer JH. Role of plasma proteins in the binding, distribution and metabolism of the thyroid hormones. *N Engl J Med* 1968; 278(21):1153–1162. doi:10.1056/NEJM196805232782107
- Jongejan RMS, Meima ME, Visser WE, et al. Binding characteristics of thyroid hormone distributor proteins to thyroid hormone metabolites. *Thyroid* 2022; 32(8):990–999. doi:10.1089/thy.2021.0588
- Pearce SH, Brabant G, Duntas LH, et al. 2013 ETA guideline: management of subclinical hypothyroidism. *Eur Thyroid J* 2013; 2(4):215–228. doi:10.1159/000356507
- Calsolaro V, Niccolai F, Pasqualetti G, et al. Hypothyroidism in the elderly: who should be treated and how? *J Endocr Soc* 2018; 3(1):146–158. doi:10.1210/js.2018-00207
- Hollowell JG, Staehling NW, Flanders WD, et al. Serum TSH, T(4), and thyroid antibodies in the United States population (1988 to 1994): National Health and Nutrition Examination Survey (NHANES III). *J Clin Endocrinol Metab* 2002; 87(2):489–499. doi:10.1210/jcem.87.2.8182
- Cappola AR, Fried LP, Arnold AM, et al. Thyroid status, cardiovascular risk, and mortality in older adults. *JAMA* 2006; 295(9):1033–1041. doi:10.1001/jama.295.9.1033
- Andersen S, Pedersen KM, Bruun NH, Laurberg P. Narrow individual variations in serum T(4) and T(3) in normal subjects: a clue to the understanding of subclinical thyroid disease. *J Clin Endocrinol Metab* 2002; 87(3):1068–1072. doi:10.1210/jcem.87.3.8165
- Boucai L, Hollowell JG, Surks MI. An approach for development of age-, gender-, and ethnicity-specific thyrotropin reference limits. *Thyroid* 2011; 21(1):5–11. doi:10.1089/thy.2010.0092
- Kuś A, Sterenborg RBTM, Haug EB, et al. Towards personalized TSH reference ranges: a genetic and population-based approach in three independent cohorts. *Thyroid* 2024; 34(8):969–979. doi:10.1089/thy.2024.0045
- van der Spoel E, Roelfsema F, van Heemst D. Within-person variation in serum thyrotropin concentrations: main sources, potential underlying biological mechanisms, and clinical implications. *Front Endocrinol (Lausanne)* 2021; 12:619568. doi:10.3389/fendo.2021.619568
- Bremner AP, Feddema P, Leedman PJ, et al. Age-related changes in thyroid function: a longitudinal study of a community-based cohort. *J Clin Endocrinol Metab* 2012; 97(5):1554–1562. doi:10.1210/jc.2011-3020
- Biondi B, Cappola AR, Cooper DS. Subclinical hypothyroidism: a review. *JAMA* 2019; 322(2):153–160. doi:10.1001/jama.2019.9052
- Fu J, Wang Y, Liu Y, Song Q, Cao J, Peichang W. Reference intervals for thyroid hormones for the elderly population and their influence on the diagnosis of subclinical hypothyroidism. *J Med Biochem* 2023; 42(2):258–264. doi:10.5937/jomb0-39570
- Hennessey JV, Espallat R. Diagnosis and management of subclinical hypothyroidism in elderly adults: a review of the literature. *J Am Geriatr Soc* 2015; 63(8):1663–1673. doi:10.1111/jgs.13532
- Yoo WS, Chung HK. Subclinical hypothyroidism: prevalence, health impact, and treatment landscape. *Endocrinol Metab (Seoul)* 2021; 36(3):500–513. doi:10.3803/EnM.2021.1066
- Xing D, Liu D, Li R, Zhou Q, Xu J. Factors influencing the reference interval of thyroid-stimulating hormone in healthy adults: a systematic review and meta-analysis. *Clin Endocrinol (Oxf)* 2021; 95(3):378–389. doi:10.1111/cen.14454
- Spencer CA, Hollowell JG, Kazarosyan M, Braverman LE. National Health and Nutrition Examination Survey III thyroid-stimulating hormone (TSH)-thyroperoxidase antibody relationships demonstrate that TSH upper reference limits may be skewed by occult thyroid dysfunction. *J Clin Endocrinol Metab* 2007; 92(11):4236–4240. doi:10.1210/jc.2007-0287
- Van Uytanghe K, Ehrenkranz J, Halsall D, et al. Thyroid stimulating hormone and thyroid hormones (triiodothyronine and thyroxine): an American Thyroid Association-commissioned review of current clinical and laboratory status. *Thyroid* 2023; 33(9):1013–1028. doi:10.1089/thy.2023.0169
- Biondi B, Cooper DS. The clinical significance of subclinical thyroid dysfunction. *Endocr Rev* 2008; 29(1):76–131. doi:10.1210/er.2006-0043
- Wartofsky L, Dickey RA. The evidence for a narrower thyrotropin reference range is compelling. *J Clin Endocrinol Metab* 2005; 90(9):5483–5488. doi:10.1210/jc.2005-0455
- Surks MI, Goswami G, Daniels GH. The thyrotropin reference range should remain unchanged. *J Clin Endocrinol Metab* 2005; 90(9):5489–5496. doi:10.1210/jc.2005-0170
- Taylor PN, Lansdown A, Witzak J, et al. Age-related variation in thyroid function—a narrative review highlighting important implications for research and clinical practice [published correction appears in *Thyroid Res* 2023; 16(1):20]. *Thyroid Res* 2023; 16(1):7. doi:10.1186/s13044-023-00149-5
- Huber G, Staub JJ, Meier C, et al. Prospective study of the spontaneous course of subclinical hypothyroidism: prognostic value of thyrotropin, thyroid reserve, and thyroid antibodies. *J Clin Endocrinol Metab* 2002; 87(7):3221–3226. doi:10.1210/jcem.87.7.8678
- Karmisholt J, Andersen S, Laurberg P. Variation in thyroid function tests in patients with stable untreated subclinical hypothyroidism. *Thyroid* 2008; 18(3):303–308. doi:10.1089/thy.2007.0241
- Mariotti S, Caturegli P, Piccolo P, Barbesino G, Pinchera A. Antithyroid peroxidase autoantibodies in thyroid diseases. *J Clin Endocrinol Metab* 1990; 71(3):661–669. doi:10.1210/jcem-71-3-661
- Fiore V, Barucca A, Barraco S, et al. Hypothyroidism in older adults: a narrative review. *Endocr Metab Immune Disord Drug Targets* 2024; 24(8):879–884. doi:10.2174/1871530323666230828110153
- Piers LS, Soares MJ, McCormack LM, O'Dea K. Is there evidence for an age-related reduction in metabolic rate? *J Appl Physiol* (1985) 1998; 85(6):2196–2204. doi:10.1152/jappl.1998.85.6.2196
- Corsonello A, Montesanto A, Berardelli M, et al. A cross-section analysis of FT3 age-related changes in a group of old and oldest-old subjects, including centenarians' relatives, shows that a down-regulated thyroid function has a familial component and is related to longevity. *Age Ageing* 2010; 39(6):723–727. doi:10.1093/ageing/afq116
- Hak AE, Pols HA, Visser TJ, Drexhage HA, Hofman A, Witteman JC. Subclinical hypothyroidism is an independent risk factor for atherosclerosis and myocardial infarction in elderly women: the Rotterdam Study. *Ann Intern Med* 2000; 132(4):270–278. doi:10.7326/0003-4819-132-4-200002150-00004

37. Imaizumi M, Akahoshi M, Ichimaru S, et al. Risk for ischemic heart disease and all-cause mortality in subclinical hypothyroidism. *J Clin Endocrinol Metab* 2004; 89(7):3365–3370. doi:10.1210/jc.2003-031089
38. Tseng FY, Lin WY, Lin CC, et al. Subclinical hypothyroidism is associated with increased risk for all-cause and cardiovascular mortality in adults. *J Am Coll Cardiol* 2012; 60(8):730–737. doi:10.1016/j.jacc.2012.03.047
39. Rodondi N, den Elzen WP, Bauer DC, et al. Subclinical hypothyroidism and the risk of coronary heart disease and mortality. *JAMA* 2010; 304(12):1365–1374. doi:10.1001/jama.2010.1361
40. Chaker L, Baumgartner C, den Elzen WP, et al. Subclinical hypothyroidism and the risk of stroke events and fatal stroke: an individual participant data analysis. *J Clin Endocrinol Metab* 2015; 100(6):2181–2191. doi:10.1210/jc.2015-1438
41. Baumgartner C, da Costa BR, Collet TH, et al. Thyroid function within the normal range, subclinical hypothyroidism, and the risk of atrial fibrillation. *Circulation* 2017; 136(22):2100–2116. doi:10.1161/CIRCULATIONAHA.117.028753
42. Gencer B, Collet TH, Virgini V, et al. Subclinical thyroid dysfunction and the risk of heart failure events: an individual participant data analysis from 6 prospective cohorts. *Circulation* 2012; 126(9):1040–1049. doi:10.1161/CIRCULATIONAHA.112.096024
43. Rodondi N, den Elzen WP, Bauer DC, et al. Subclinical hypothyroidism and the risk of coronary heart disease and mortality. *JAMA* 2010; 304(12):1365–1374. doi:10.1001/jama.2010.1361
44. Nanchen D, Gussekloo J, Westendorp RG, et al. Subclinical thyroid dysfunction and the risk of heart failure in older persons at high cardiovascular risk. *J Clin Endocrinol Metab* 2012; 97(3):852–861. doi:10.1210/jc.2011-1978
45. Razvi S, Shakoor A, Vanderpump M, Weaver JU, Pearce SH. The influence of age on the relationship between subclinical hypothyroidism and ischemic heart disease: a metaanalysis. *J Clin Endocrinol Metab* 2008; 93(8):2998–3007. doi:10.1210/jc.2008-0167
46. Pasqualetti G, Pagano G, Rengo G, Ferrara N, Monzani F. Subclinical hypothyroidism and cognitive impairment: systematic review and meta-analysis. *J Clin Endocrinol Metab* 2015; 100(11):4240–4248. doi:10.1210/jc.2015-2046
47. Canaris GJ, Manowitz NR, Mayor G, Ridgway EC. The Colorado Thyroid Disease Prevalence Study. *Arch Intern Med* 2000; 160(4):526–534. doi:10.1001/archinte.160.4.526
48. Meuwese CL, van Diepen M, Cappola AR, et al. Low thyroid function is not associated with an accelerated deterioration in renal function. *Nephrol Dial Transplant* 2019; 34(4):650–659. doi:10.1093/ndt/gfy071
49. Rhee CM, Kim S, Gillen DL, et al. Association of thyroid functional disease with mortality in a national cohort of incident hemodialysis patients. *J Clin Endocrinol Metab*. 2015;100(4):1386–1395. doi:10.1210/jc.2014-4311
50. Fatourech V. Subclinical hypothyroidism: an update for primary care physicians. *Mayo Clin Proc* 2009; 84(1):65–71. doi:10.4065/84.1.65
51. De Whalley P. Do abnormal thyroid-stimulating hormone level values result in treatment changes? A study of patients on thyroxine in one general practice. *Br J Gen Pract* 1995; 45(391):93–95. PMID: 7702890
52. Ross DS, Daniels GH, Gouveia D. The use and limitations of a chemiluminescent thyrotropin assay as a single thyroid function test in an out-patient endocrine clinic. *J Clin Endocrinol Metab* 1990; 71(3):764–769. doi:10.1210/jcem-71-3-764
53. Lillevang-Johansen M, Abrahamsen B, Jørgensen HL, Brix TH, Hegedüs L. Over- and under-treatment of hypothyroidism is associated with excess mortality: a register-based cohort study. *Thyroid* 2018; 28(5):566–574. doi:10.1089/thy.2017.0517
54. Wilson SA, Stem LA, Bruehlman RD. Hypothyroidism: diagnosis and treatment. *Am Fam Physician* 2021; 103(10):605–613. PMID:33983002
55. McDermott MT. Hypothyroidism. *Ann Intern Med* 2020; 173(1):ITC1–ITC16. doi:10.7326/AITC202007070
56. Goichot B, Raverot V, Klein M, et al. Management of thyroid dysfunctions in the elderly. French Endocrine Society consensus statement 2019. Long version. *Ann Endocrinol (Paris)* 2020; 81(2–3):89–100. doi:10.1016/j.ando.2020.04.010
57. Ross DS. Treating hypothyroidism is not always easy: when to treat subclinical hypothyroidism, TSH goals in the elderly, and alternatives to levothyroxine monotherapy. *J Intern Med* 2022; 291(2):128–140. doi:10.1111/joim.13410
58. Danese MD, Powe NR, Sawin CT, Ladenson PW. Screening for mild thyroid failure at the periodic health examination: a decision and cost-effectiveness analysis. *JAMA* 1996; 276(4):285–292. PMID:8656540
59. Jonklaas J. TSH reference intervals: their importance and complexity. *Thyroid* 2024; 34(8):957–959. doi:10.1089/thy.2024.0380
60. Scobbo RR, vonDohlen TW, Hassan M, Islam S. Serum TSH variability in normal individuals: the influence of time of sample collection. *W V Med J* 2004; 100(4):138–142. PMID: 15471172
61. Kim TH, Kim KW, Ahn HY, et al. Effect of seasonal changes on the transition between subclinical hypothyroid and euthyroid status. *J Clin Endocrinol Metab* 2013; 98(8):3420–3429. doi:10.1210/jc.2013-1607

Address: Risheng Xu, DO, Department of Medicine, Baylor College of Medicine, 20171 Chasewood Park Drive, Houston, TX 77070; risheng.xu@bcm.edu