Chronic Spontaneous Urticaria: Backgrounder
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What is chronic spontaneous urticaria (CSU)?
Chronic spontaneous urticaria (CSU) is a distressing skin condition that causes red, swollen, itchy and sometimes painful hives or “wheals” on the skin. At any given time, the worldwide prevalence of CSU is 0.5% to 1%, a similar figure to those living with HIV. Urticaria is defined as chronic when the symptoms spontaneously present and reoccur daily or almost daily for at least six weeks. It is described as spontaneous when symptoms are not triggered by any identified external factor. Women are twice as likely as men to be diagnosed with the disease and most people first develop symptoms between the ages of 20 and 40.

CSU has negative effects on quality of life, which frequently include sleep deprivation and psychological comorbidities such as depression and anxiety. Research indicates this negative impact on quality of life is similar to patients with ischemic heart disease, with patients feeling a similar lack of energy, social isolation and emotional upset. The frequent failure to identify a specific underlying cause of CSU, its unpredictable symptoms and a high disease burden lead to frustration among patients and their doctors.

What are the symptoms of CSU?
Symptoms of CSU include the appearance of flare-type skin reactions and itchy welts, often described as hives or “wheals”.

The itch is often persistent, even at night-time, and is not relieved with an approved dose of anti-histamine. Between 40% and 50% of CSU patients also experience angioedema, a swelling in the deep layers of the skin.

What may trigger a person to develop CSU?
In CSU symptoms appear spontaneously and there is no specific external trigger for the individual hives or swelling that develop. Research into the underlying cause of the disease has uncovered the role of the autoimmune system in many patients, and other aggravating factors may include stress and infection.
How is CSU diagnosed and treated?

CSU is difficult to diagnose and manage because the trigger is unknown or, in the case of factors like the autoimmune system, cannot be easily changed. If a person presents with symptoms, he or she should be evaluated by a healthcare professional who may ask the patient to record any potential triggers e.g., cold, heat, exercise, pressure or other physical triggers, and the area and frequency of the hives for a period of one to two weeks. This aids the diagnosis to exclude obvious underlying causes and/or eliciting triggers for the condition. Further CSU testing may include evaluations of the skin, blood and urine.

Antihistamines are currently the mainstay of therapy for CSU; however, more than 50% of patients on approved doses do not achieve symptom relief. Medical guidelines allow for increased doses of antihistamines, up to four times the approved dose, to increase symptom control in some patients. However, there remains a critical need for new treatment options as up to 40% of CSU patients fail on these increased antihistamine doses.

How is severity measured?

Once CSU is diagnosed, it can be assessed in a variety of ways. Disease severity can be measured by the presence of itch - the Itch Severity Scale (ISS) and rash - the severity of the wheals. Both scales can range from 0 (no symptoms) to 3 (severe symptoms). The two scales are combined in the Urticaria Activity Score (UAS) - with a maximum daily score of 6. The weekly urticaria activity score (UAS7) measures disease activity over the previous 7 days, with a score ranging from 0 to 42. A UAS7 score of 6 or less represents well controlled disease.

Apart from the assessment of physical symptoms, how CSU affects the patients’ quality of life can be measured with the Dermatology Life Quality Index (DLQI), which consists of 10 questions with a score range from 0 to 30 - a higher score represents greater impairment of the patients QoL. Another tool that may be used is the Chronic Urticaria Quality of Life Questionnaire (CU-QoL).

References