## **GUEST EDITORIAL**

## Increased Knowledge-based Care is Indicated for Patients with Atopic Dermatitis

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Atopic dermatitis (AD) is the most common inflammatory skin disorder, affecting up to 20% of children and at least 3–10% of adults in the developed world (1). AD has a huge impact on quality of life and is the dermatological condition with the largest disease burden from a societal perspective (2).

The disease can lead to a life with interrupted sleep due to intense pruritis, and periodically being unable to attend school, being placed on sick leave, or being hospitalized. It can cause problems with relationships, studies and work. This, in combination with a deficit in healthcare support, and the fact that many patients feel so affected, and sometimes ashamed, that they hide themselves and their eczema, makes it even worse. Mental health issues are frequent in the affected group and the risk of depression and suicide is higher than among people without AD. Systemic treatments of AD can relieve not only the skin symptoms but also signs of depression in parallel, as shown by Lina Evert at al. in this issue of Acta Dermato-Venereologica (3–5). Furthermore, AD is often associated with concomitant diseases, such as asthma and allergy, and secondary cutaneous infections. A recent, large UK study by Silverwood et al. (6), shows that premature death is more common among people with severe AD. The study revealed that mortality was 60% higher among adult patients with severe AD. It is also known that the inflammatory process in AD can lead to diabetes and cardiovascular disease (7).

There is still a significant unmet need to improve overall management of AD in order that patients of all ages can gain control of their symptoms using safe and effective treatments. However, recent advances in understanding of disease mechanisms have led to the discovery of novel potential therapeutic targets and drug candidates. There are already new systemic drugs on the market in the form of biological treatments and small, synthetic molecules (so-called JAK inhibitors). In addition, there are approximately 30 new molecules in various phases of clinical trials (8).

However, new treatments are often compared only with placebo and not with standard treatments. It is important that new treatments are given to adequately selected patients, and in the right way, and are evaluated with regard to side-effects and effects. To achieve this, more knowledge is needed about the disease and available treatments *at all levels* within healthcare and society. We believe that strong patient organizations can help. Other tools are the national quality registries, in which all patients with AD on systemic treatment should be registered and followed by standardized outcome measurements.

Furthermore, national and European treatment guidelines and governance documents for AD are important (9). All of these tools are needed to create equal and knowledge-based care for *all* patients with AD, regardless of where they live.

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