#### MAT-XU-2402378 (v1.0) September 2024

## Uncapping Potential in the Diagnosis & Management of Atopic Dermatitis

#### **Atopic Dermatitis: The facts**

Skin is our largest organ, acting as a protective barrier from infections and diseases. However, skin conditions can affect this vital barrier, leading to further impacts on an individual's health and wider society.

Moderate-to-severe atopic dermatitis, also known as atopic dermatitis (AD), is a chronic inflammatory autoimmune condition that causes the skin to become itchy, inflamed, dry and cracked.<sup>1</sup>

- It is estimated that about 10 30% of children, and about 2 10% of adults are living with this condition.<sup>2</sup>
- Atopic dermatitis can increase the risk of bacterial skin infections and viral skin infections, which can become more serious if not treated.<sup>3</sup>
- While treatments can relieve symptoms of atopic dermatitis, there is currently no cure.<sup>1</sup>
- There is widespread variation in waiting times for dermatological services across England, ranging between 7 to 17 weeks across integrated care systems (ICSs).<sup>4</sup>
- One of the significant challenges in dermatology currently is due to workforce shortages with some integrated care systems having as few as 0.6 dermatologists per 100,000 patients.<sup>5</sup>

#### Key recommendations:

- **1.** Integrated care system's must implement clinical pathways for the diagnosis and management of atopic dermatitis in line with NICE guidelines to ensure a consistent approach across the country.
- 2. The Government and NHS England must increase the number of dermatology specialists, with compulsory dermatology training in primary care and greater opportunities for GPs with extended roles in dermatology.
- **3.** Patient education on AD is important so they can make more informed decisions about their own care, improving self-management, outcomes and reducing pressure on primary care as a result. NHS regional teams and dermatology clinical networks must therefore ensure the provision of adequate patient education and supported self-management as a core part of the treatment pathway.
- **4.** ICSs, GPs, dermatology services, mental health teams, and the dermatology patient community should collaboratively establish formal processes to provide mental health support for dermatology patients.







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#### Atopic Dermatitis, The Impact:

Atopic dermatitits can have substantial financial, social, and mental health impact on both patients and society:<sup>6,7</sup>

**78%** of patients said dermatitis had negatively impacted their mental health<sup>(n = 237)</sup>

# 23%

of patients reported taking off more than 15 days of work due to their condition<sup>(n = 92)</sup>

# £6,741 to £14,166

per patient in productivity lost per year for patients with atopic dermatitis People diagnosed with atopic dermatitis should first be offered a topical cream from their GP in primary care.<sup>8</sup> In the event that initial treatment does not improve symptoms, they should then be referred to specialised dermatologists in secondary care.



However, challenges exist in making this a reality, including:

- Public awareness of atopic dermatitis.
- An overburdened and stretched NHS workforce.
- Challenges in equitable access to specialist support.
- Potential tensions between local pathway prescribing and primary care.

## Despite the significant impact of atopic dermatitis on patients, the NHS and the broader economy, improving access to support for patients living with atopic dermatitis is not currently a policy priority at either a national or local level.

Integrated care systems (ICSs) have competing priorities across a range of disease areas, limiting their capacity to sufficiently prioritise atopic dermatitis despite its impact. Specialist dermatology services are often understaffed, oversubscribed, and overloaded with cases across 4,000 skin conditions.

There is also limited ICS bandwidth for prioritising atopic dermatitis, meaning specialist clinicians are important advocates. It is vital that ICSs focus on and implement Dermatology Clinical Networks consistently across the country.

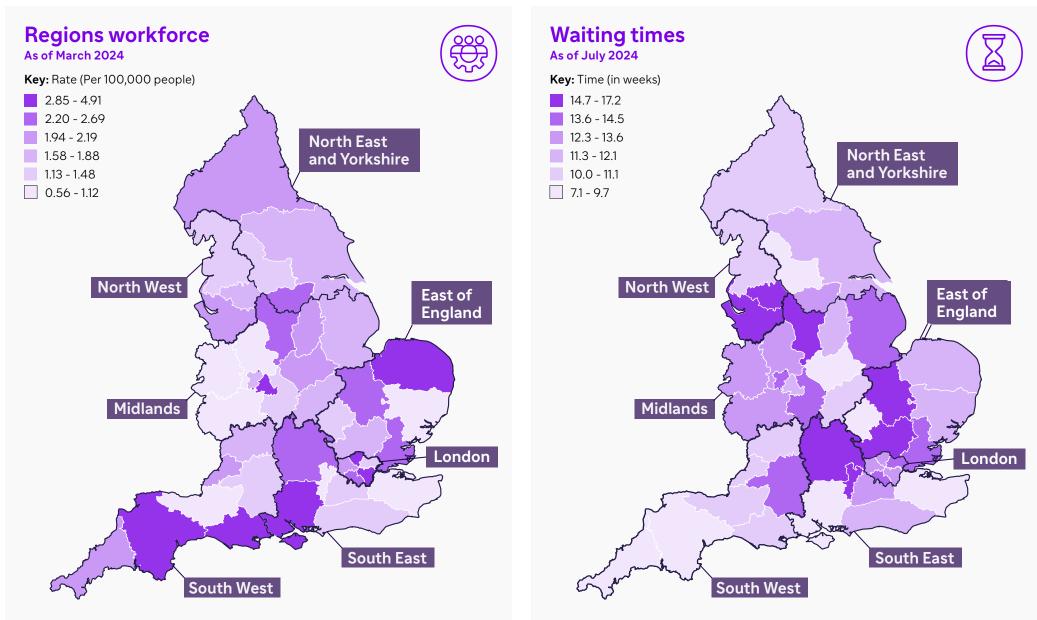
While there is clear NICE guidance on atopic dermatitis,<sup>8</sup> challenges throughout the pathway and their implementation at a local level can impede patient access to treatments in line with this guidance, leading to a postcode lottery in both care and patient outcomes.

It is vital that all people living with atopic dermatitis are equally supported across the whole pathway to better manage their condition.



#### Map of England ICSs

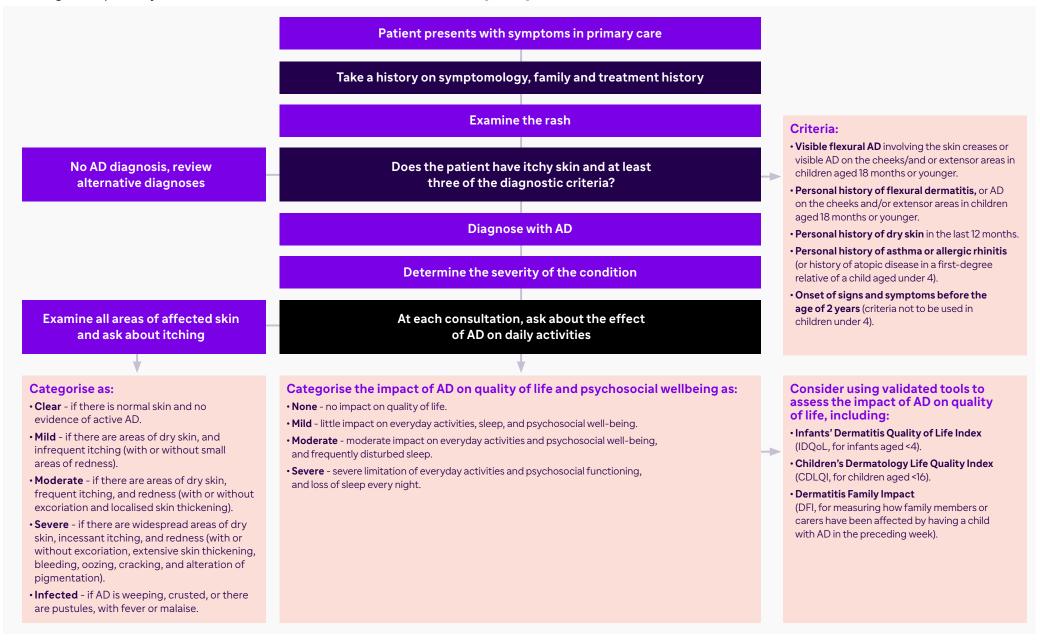
To see a breakdown of data in your specific integrated care system, please click on your respective region below



Sanofi Data on File derived from patient list sizes published by NHS Catalyst and Staff in Post publications

### **Diagnosing Atopic Dermatitis (AD) Dermatitis**

Ideal diagnostic pathway as derived from recommendations in NICE Guideline [CG 57]



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### **Treating Atopic Dermatitis (AD)**

Ideal treatment pathway as derived from recommendations in NICE Guideline [CG 57] and NICE Clinical Knowledge Summaries



#### Despite this clear pathway, there are a few headline challenges which can prevent best practice care:

- 1. There are few contact points between patients and clinicians given that most treatments for atopic dermatitis are administered in the home, making it difficult for patients to frequently discuss their care. Touchpoints between clinicians and patients are key to supporting the tailoring of treatment options to patient's needs in line with NICE guidelines.<sup>9</sup>
- 2. Insufficient dermatology specialists in primary care. There is a lack of dermatology training among primary care professionals like GPs, pharmacists and nurses working in primary care where AD care is mostly centred. The majority of AD patients will never see a specialist in secondary care, instead pharmacists are becoming more and more commonly the first point of contact for patients.<sup>10</sup>
  - a. Long waits for specialists
  - b. Increasing use of high-cost locum staff
  - c. Considerable variation across the country in care provision and outcomes.
- **3. Corticophobia.** Studies have also shown that concerns around treating flare ups with topical corticosteroids have led to some patients developing phobias, leading to treatment non-compliance and ultimately reduced disease control particularly with patients under the age of 12.<sup>11</sup>
- 4. While the majority of AD patients will never see a dermatology specialist, for the few that need to they face long waits of up to one year (in 27% of cases) with an average wait time of 3 months, as opposed to the two weeks recommended. Notably, individuals from a higher socioeconomic background were also more likely to receive specialist referrals for atopic dermatitis pointing to potential variation in care associated with deprivation.<sup>12</sup>

#### Making the case for a streamlined Atopic Dermatitis patient pathway

## Unequal knowledge of atopic dermatitis across primary care can lead to disparate diagnoses

There is variation in the time taken to diagnose moderate-to-severe atopic dermatitis. Average referral-to-treatment waiting times for dermatology services in England can range from 7 to 17 weeks across England's integrated care systems (ICSs). In some ICS's, only 51% of all patients are being seen within 18 weeks.<sup>4</sup>

Dermatology cases reflect 15 – 20% of GPs workload.<sup>14</sup> Yet, training on dermatology is not compulsory in UK medicine studies, with educational resources for primary care professionals often seen as optional.<sup>13,14</sup> It is vital that GPs and other healthcare professionals receive sufficient training on dermatology to ensure timely and accurate diagnoses.

Furthermore, more than two-thirds of healthcare professionals (n = 30) consulted in our survey think there is a lack of awareness amongst GPs on how to manage and treat atopic dermatitis.<sup>6</sup>

GPs may not be well informed enough to accurately diagnose patients with atopic dermatitis in the first instance or understand the severity of their condition.<sup>14</sup> In turn, this can lead to delays in patient referral to secondary care.<sup>14</sup>

#### ICSs need to implement increased and equitable training on dermatology conditions across primary care to improve patient diagnosis.

Unequal prescribing of full treatment options impacts patient outcomes through the pathway.

#### Immunosuppressants

People with moderate-to-severe atopic dermatitis may be referred to specialist care services in secondary care, where they can be prescribed immunosuppressants or biologics.<sup>9</sup>

Immunosuppressants can help to suppress or control the body's immune system to slow the symptoms of atopic dermatitis.<sup>15</sup> These can be prescribed to children and adults with moderate-to-severe atopic dermatitis for a short time to help stop itching, in turn allowing the skin's protective barrier to heal.<sup>15</sup> Immunosuppressants can have side effects, such as an increased infection risk, the increased risk of certain types of cancer, and an upset stomach or vomiting.<sup>15</sup>

While these treatment options can support patient outcomes, challenges with their prescribing remain.

Immunosuppressants for atopic dermatitis are intended for short-term use, with patients intended to be changed to a topical treatment for their long-term condition management.<sup>15</sup> However, patients can be prescribed multiple immunosuppressant cycles with little respite. Dermatologists must be provided with adequate support and capacity to move patients through the pathway, either onto a topical treatment where appropriate, or on to other treatment classes.

In addition to benefiting patient care and outcomes, supporting patients to move through the pathway can also help to minimise capacity challenges in the dermatology workforce. Patients on immunosuppressants can require careful monitoring,<sup>9</sup> including for example regular blood tests. This monitoring can be time-consuming and labour intensive for an already stretched workforce.

#### **The NHS Workforce**

The NHS is currently experiencing considerable workforce pressures across the board, but this problem is particularly acute in dermatology, caused in part by long-term restrictions on the number of new dermatology training posts. To manage these pressures, the NHS has often employed the use of highcost locum consultants – reducing the amount invested into training new dermatologists, ultimately contributing to long-term workforce shortages.<sup>10</sup>

In 2021, the Getting It Right First Time (GIRFT) report on Dermatology found that at least 10 NHS Trusts had no dermatologists at all while 1/3<sup>rd</sup> had very severe shortages.<sup>10</sup> These shortages impact on both the timely diagnosis and the efficient treatment of patients with atopic dermatitis, the variation in dermatology staffing also ultimately leads to inconsistencies in care outcomes across the country.<sup>16</sup>

Our recent research found that this considerable workforce variation remains an issue at the integrated care board level. Some integrated care systems had as many as 5.22 dermatologists per 100,000 patients while several integrated care systems had less than one dermatologist per 100,000 patients.<sup>5</sup> Slightly over ½ of all integrated care systems in England have less than two dermatologists per 100,000 patients.<sup>5</sup>

Geographical disparities in the dermatology workforce exist across England. South East London has the highest dermatologists to patient ratio at 4.91 (per 100,000), followed by Norfolk and Waveney at 3.55 (per 100,000) and Birmingham and Solihull at 3.30 (per 100,000).<sup>5</sup> However, Shropshire, Telford and Wrekin has the lowest dermatologist to patient ratio at 0.56 (per 100,000), alongside Kent and Medway at 0.59 (per 100,000) and Frimley at 0.83 (per 100,000).<sup>5</sup>

As a result, dermatology patients face inequity in direct access to an NHS dermatologist which is essential in delivering optimal care for patients and ensuring they are on the best treatment option that works for them. Instead, patients will often seek advice from primary care pharmacists, practice nurses and GPs. This may lead to conflicting guidance and delay diagnosis as some may go through multiple tests and GP visits before seeing a specialist who can diagnose their condition first time.<sup>10</sup>

The disparities also mean that patients in areas with a lower dermatologist to patient ratio may receive a lower standard of care and an even more stretched workforce in cases where patients on immunosuppressants need to be monitored.

This ultimately culminates in a considerable postcode lottery in care for dermatology, a condition under prioritised at a national level despite its considerable primary care burden.

### The Government and NHS England must increase the number of dermatology specialists, with compulsory dermatology training in primary care.

### ICSs should include a focus on local dermatology pathways in their ICS plans to ensure smooth patient pathways.

## Patients experience challenges in being fully informed on atopic dermatitis, often self-managing at home.

People living with atopic dermatitis can face challenges across treatment and condition management – from patient choice and information, to support for mental and emotional wellbeing. Patients can be left to self-manage their condition at home, with limited input from healthcare professionals and insufficient access to information.

#### Patient choice and information on condition management

Patients may not be fully informed of the various condition management options available. To secure greater patient choice and empowerment, it is vital that patients and carers receive improved education on long-term condition management options for atopic dermatitis from healthcare professionals.

**70%** of surveyed patients state that dermatitis has affected their self-esteem or made them self-conscious<sup>6</sup>

#### It is our vision that people living with atopic dermatitis become genuine partners in their own care, empowered to ask for their preferred condition management.

## Impact on patient mental health and access to psychological support

## Patients need to be supported to better manage their condition and their mental health through education and the provision of wider psychological services.

As noted, atopic dermatitis can have negative impacts on an individuals' mental health and daily life – including on everyday activities, social life, sleep, and personal relationships.<sup>6</sup> This can further lead to comorbidities with mental health conditions such as depression and anxiety, as well as suicidal ideation, which are more common amongst people with moderate-to-severe atopic dermatitis.<sup>6,17</sup>

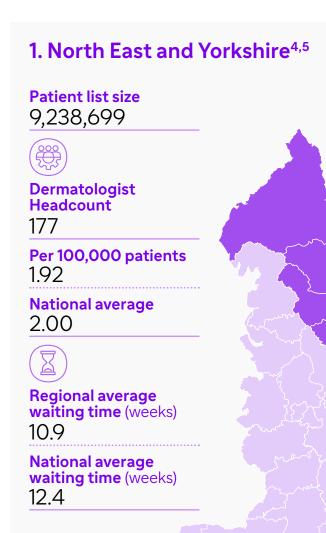
Despite the clear and stark impact of atopic dermatitis on an individuals' mental health, there is insufficient support available for patients. 100% of surveyed healthcare professionals do not think there is adequate mental health support for patients with atopic dermatitis.<sup>6</sup> Patients have reported mixed experiences with advice and support for their mental health. Some patients have received generic advice from healthcare professionals, such as to exercise more but not understanding that this may worsen their condition.<sup>14</sup> However, other patients have received empathetic care from a range of healthcare professionals they engage with.<sup>14</sup>

## ICSs, GPs, dermatology services, mental health teams, and the dermatology patient community should collaboratively establish formal processes to provide mental health support for dermatology patients.

There is also limited healthcare professional knowledge or referral to psychological support groups. NHS psychodermatology services are reliant on specialists with an understanding of the relation between mental health and dermatology conditions – a limited pool of healthcare professionals.<sup>14</sup> Additionally, 93% of surveyed patients themselves were not aware that psychological support services even existed.<sup>6</sup>

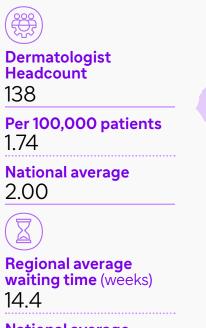
## Healthcare professionals and patient groups to collaboratively increase awareness of psychological support for dermatology patients.

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## 2. North West<sup>4,5</sup>





National average waiting time (weeks) 12.4

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### 3. Midlands<sup>4,5</sup>

#### **Patient list size** 11,968,840



Dermatologist Headcount 232

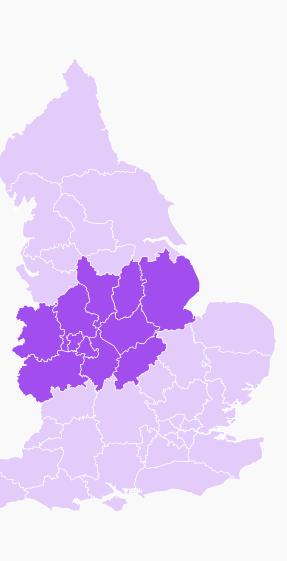
**Per 100,000 patients** 1.94

National average 2.00



Regional average waiting time (weeks) 12.6

National average waiting time (weeks) 12.4



## 4. East of England<sup>4,5</sup>

**Patient list size** 7,305,931



Dermatologist Headcount 147

**Per 100,000 patients** 2.01

National average 2.00



Regional average waiting time (weeks) 13.1

National average waiting time (weeks) 12.4

## 5. South East<sup>4,5</sup>

**Patient list size** 9,834,438



Dermatologist Headcount 161

**Per 100,000 patients** 1.64

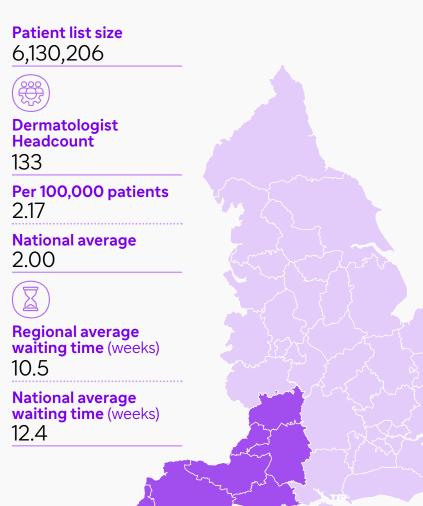
National average 2.00



Regional average waiting time (weeks) 12.4

National average waiting time (weeks) 12.4

## 6. South West<sup>4,5</sup>



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### 7. London<sup>4,5</sup>

#### **Patient list size** 11,049,835



Dermatologist Headcount 290

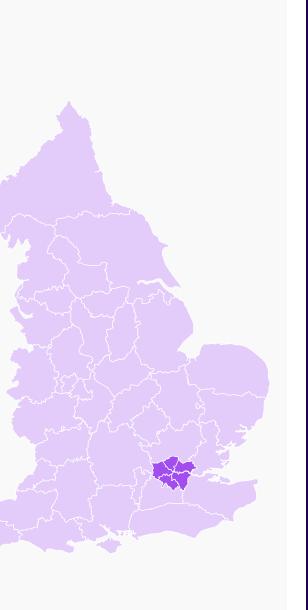
**Per 100,000 patients** 2.62

National average 2.00



Regional average waiting time (weeks) 12.9

National average waiting time (weeks) 12.4



## About Sanofi

Sanofi is a global life sciences company with a strong heritage and presence in the UK. From prevention to treatment, Sanofi transforms scientific innovation into healthcare solutions.

For further information about our work in dermatology, please contact Chris Ward, Specialty Care Public Affairs and Advocacy Manager at **christopher.ward@sanofi.com** 

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