

Patient and Healthcare Provider Perspectives on the Path to Diagnosis of Seborrheic Dermatitis: Results From a National Survey of Adults With Seborrheic Dermatitis in the United States

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Introduction, Methods and Results

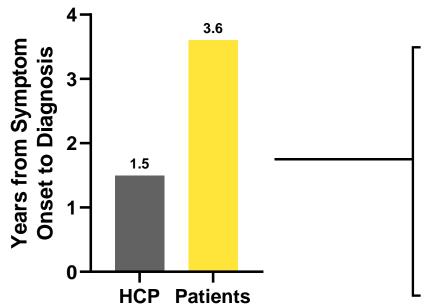
Introduction and Methods

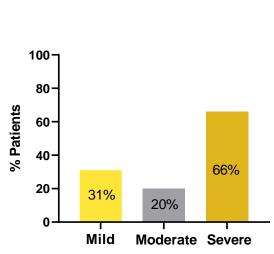
- Seborrheic dermatitis (SD) is a common chronic inflammatory skin disease with a worldwide prevalence of up to 5%¹; however, little is known about patient and provider views and experiences in the path to diagnosis of SD
- Methods and key demographics are described in e-poster 42842
- Here we present the results related to the path to diagnosis of SD

Time from Symptom Onset to Seeking Care

HCP- and Patient-Reported Time From Symptom Onset to Seeking Care







Results

Patient Perceptions of SD prior to diagnosis

"I had not heard of SD prior to my diagnosis"

71% agree

"I mistook my symptoms for a different type of skin condition"

76% agree

"I didn't realize all of my symptoms (e.g., on face, body, scalp, etc.) were due to SD"

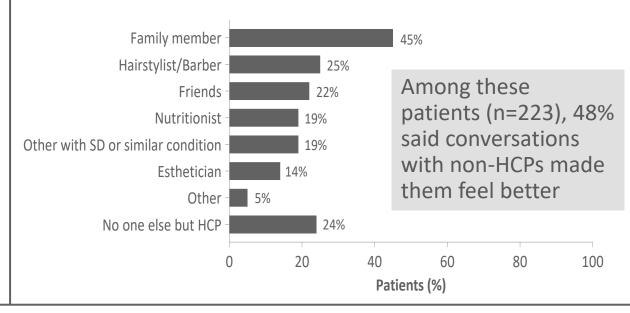
83% agree

"it was hard to find information online about SD before diagnosis"

56% agree

90% of patients wish they had known that there are specific symptoms that can help identify SD

Other People (Non-HCPs) Who Helped Patients Identify their SD **Symptoms**





"I didn't think my symptoms were severe enough to warrant medical attention"



"Most patients did not think that SD symptoms warranted medical attention" **66%** HCPs agree



"I was embarrassed to talk to my family or friends about my **symptoms"** 59% Patients agree



"Most patients were embarrassed to talk to their family or friends about their symptoms"

65% HCPs agree



"I was embarrassed to talk to a HCP about my symptoms"

58% Patients agree

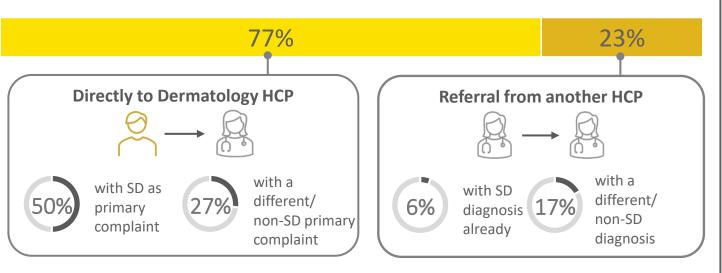


"Most patients were embarrassed to talk to me about their symptoms"

32% HCPs agree

Results

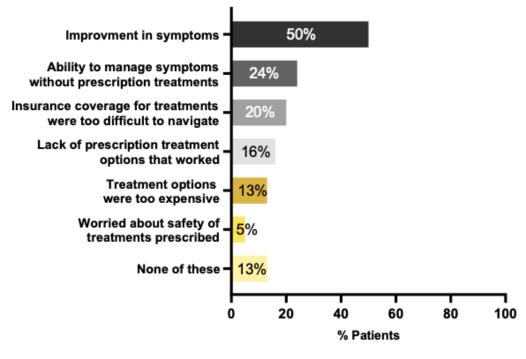
HCP-reported Path to SD Diagnosis



- Most (79%) patients said they prefer a dermatologist for SD management
- Patients reported visiting an average of 2.3 HCPs for SD treatment and **75% have seen more than one HCP**
- Patients* said they visit their primary HCP for SD management an average of **4.6 times per year** (n=289), and **85% said they still** actively meet with their HCP for SD treatment

Reasons Patients Stopped Seeing HCP

Reasons why the 15% of patients stopped seeing their HCP*



^{*}n=43. Small sample size, results are qualitive in nature.



Results and Conclusions

Patient-reported knowledge of SD

- Once diagnosed:
 - Only 20% of patients feel 'very knowledgeable' about what causes SD and 22% about what triggers their SD symptoms
 - Only 30% of patients were 'very satisfied' about the information received about the treatment plan and treatment options they received from their HCP
 - 28% of patients feel very knowledgeable about available treatment options
 - 27% of patients feel knowledgeable about how treatment options work, and how the treatment options are administered

Conclusions

- Prior to diagnosis, most patients said they had not heard of SD and found it difficult to find information online
- HCPs underestimated the time it takes for patients experiencing SD symptoms to reach out, with a 2-year difference between what HCPs thought and patients reported
- These findings reveal major needs in optimization of diagnosis, management, and education for SD
- Future studies are needed to better characterize and address these burdens