Presented at **ISPOR 2022** Washington, DC, USA and Virtual • May 15–18, 2022

Examining Treatment Patterns of Topical Therapies Among Adolescent and Adult Patients With Atopic Dermatitis in the United States

Incyte Corporation, Wilmington, DE, USA

*Presenting author

Vijay N. Joish, MS, PhD,* Jennifer H. Lofland, PharmD, PhD, MPH, Samyuktha Darbha, MBA, Ahmad B. Naim, MD

Introduction

- Atopic dermatitis (AD) is a highly pruritic inflammatory skin disease¹
- Topical treatments, such as topical corticosteroids (TCS), topical calcineurin inhibitors (TCI), and the phosphodiesterase-4 inhibitor crisaborole, are commonly used as first-line treatments for patients with AD^{2,3}
- Many patients do not achieve adequate disease control with topical first-line therapies and are escalated to systemic agents, such as oral corticosteroids (OCS) and biologics (eg, dupilumab)^{4,5}
- The exact treatment patterns used by patients with AD in the United States and the extent of systemic treatment use after initial topical treatment have not been thoroughly examined

Objective

 To describe treatment patterns in US individuals with AD during the first 12 months of topical treatment using data from a US claims database

Methods

Study Design

- This was a retrospective observational study using data from the IBM MarketScan Commercial and Medicare supplemental claims database between 2017–2019 (Figure 1)
- Eligible individuals were ≥12 years old with ≥1 medical claim for AD in 2018 and 12 months of continuous enrollment in a healthcare plan before and after the first topical prescription claim for AD (index event)
- Individuals with a history of topical AD prescriptions in the 12 months before the index event were excluded

Figure 1. Study Design

2	2017	2018	2019	
		First topical prescription claim for AD (index event)	Last follow-up date:	
		Individual Identification Period: January 1 to December 31, 2018	December 31, 201	

Assessments

- Demographics and clinical characteristics were summarized at the individual level
- Treatment patterns were summarized based on prescription claim refills during the 12 months after the index event
- Data are reported using descriptive statistics

Results

Population Characteristics

- A total of 47,572 individuals were included in the analysis
- Mean (SD) age was 38.0 (18.8) years, and 63.7% were female

- The most common comorbid conditions included allergic rhinitis (27.2%), asthma (15.7%), and major depressive disorder (11.9%)

- Distribution of baseline demographics and clinical characteristics was similar across treatment groups (Table 1)
- At the index event, the initial topical treatments prescribed were medium-potency TCS (n=20,849; 43.8%), high-potency TCS (n=10,675; 22.4%), low-potency TCS (n=9822; 20.6%), TCI (n=3107; 6.5%), and crisaborole (n=2725; 5.7%)
- A small number of individuals were prescribed dupilumab as initial treatment (n=394; 0.8%)

Table 1. Demographics and Baseline Clinical Characteristics by Initial Treatment Group

	potency TCS	potency TCS	potency TCS	TCI	Crisaborole	Total
haracteristic	(n=9822)	(n=20,849)	(n=10,675)	(n=3107)	(n=2725)	(N=47,572)*
ge, mean (SD), y	36.0 (19.2)	39.9 (18.8)	36.9 (18.7)	36.1 (17.8)	35.5 (17.7)	38.0 (18.8)
12–17 y, n (%)	2424 (24.7)	3300 (15.8)	2304 (21.6)	682 (22.0)	642 (23.6)	9357 (19.7)
≥18 y, n (%)	7398 (75.3)	17,549 (84.2)	8371 (78.4)	2425 (78.1)	2083 (76.4)	38,215 (80.3)
emale, n (%)	6558 (66.8)	12,870 (61.7)	6673 (62.5)	2115 (68.1)	1856 (68.1)	30,294 (63.7)
legion, n (%)						
South	3969 (40.4)	10,312 (49.5)	4874 (45.7)	1246 (40.1)	1320 (48.4)	21,894 (46.0)
Northeast	2642 (26.9)	4329 (20.8)	2408 (22.6)	943 (30.4)	658 (24.2)	11,061 (23.3)
North Central	1810 (18.4)	3413 (16.4)	1720 (16.1)	497 (16.0)	394 (14.5)	7911 (16.6)
West	1354 (13.8)	2734 (13.1)	1643 (15.4)	410 (13.2)	352 (12.9)	6555 (13.8)
Unknown	47 (0.5)	61 (0.3)	30 (0.3)	11 (0.4)	1 (0.04)	151 (0.3)
lan type, n (%)						
PPO	4873 (49.6)	10,719 (51.4)	5333 (50.0)	1765 (56.8)	1531 (56.2)	24,426 (51.4)
CDHP/HDHP	2241 (22.8)	4818 (23.1)	2500 (23.4)	609 (19.6)	697 (25.6)	10,959 (23.0)
HMO	1194 (12.2)	2347 (11.3)	1379 (12.9)	343 (11.0)	168 (6.2)	5464 (11.5)
POS/POS with capitation	767 (7.8)	1492 (7.2)	791 (7.4)	235 (7.6)	157 (5.8)	3475 (7.3)
Comprehensive	415 (4.2)	726 (3.5)	356 (3.3)	66 (2.1)	87 (3.2)	1667 (3.5)
EPO	97 (1.0)	180 (0.9)	93 (0.9)	16 (0.5)	27 (1.0)	418 (0.9)
Unknown/Missing	235 (2.4)	567 (2.7)	223 (2.1)	73 (2.4)	58 (2.1)	1163 (2.4)
comorbidities, n (%)						
Allergic rhinitis	5380 (25.8)	2831 (26.5)	2721 (27.7)	958 (30.8)	934 (34.3)	12,955 (27.2)
Asthma	3030 (14.5)	1774 (16.6)	1542 (15.7)	535 (17.2)	495 (18.2)	7469 (15.7)
MDD	2613 (12.5)	1233 (11.6)	1106 (11.3)	356 (11.5)	296 (10.9)	5660 (11.9)
GAD	1804 (8.7)	917 (8.6)	831 (8.5)	262 (8.4)	228 (8.4)	4071 (8.6)
Urticaria	1314 (6.3)	623 (5.8)	597 (6.1)	187 (6.0)	202 (7.4)	2961 (6.2)
Food allergy	514 (2.5)	377 (3.5)	354 (3.6)	146 (4.7)	119 (4.4)	1537 (3.2)
Nasal polyposis	107 (0.5)	48 (0.5)	41 (0.4)	21 (0.7)	13 (0.5)	236 (0.5)

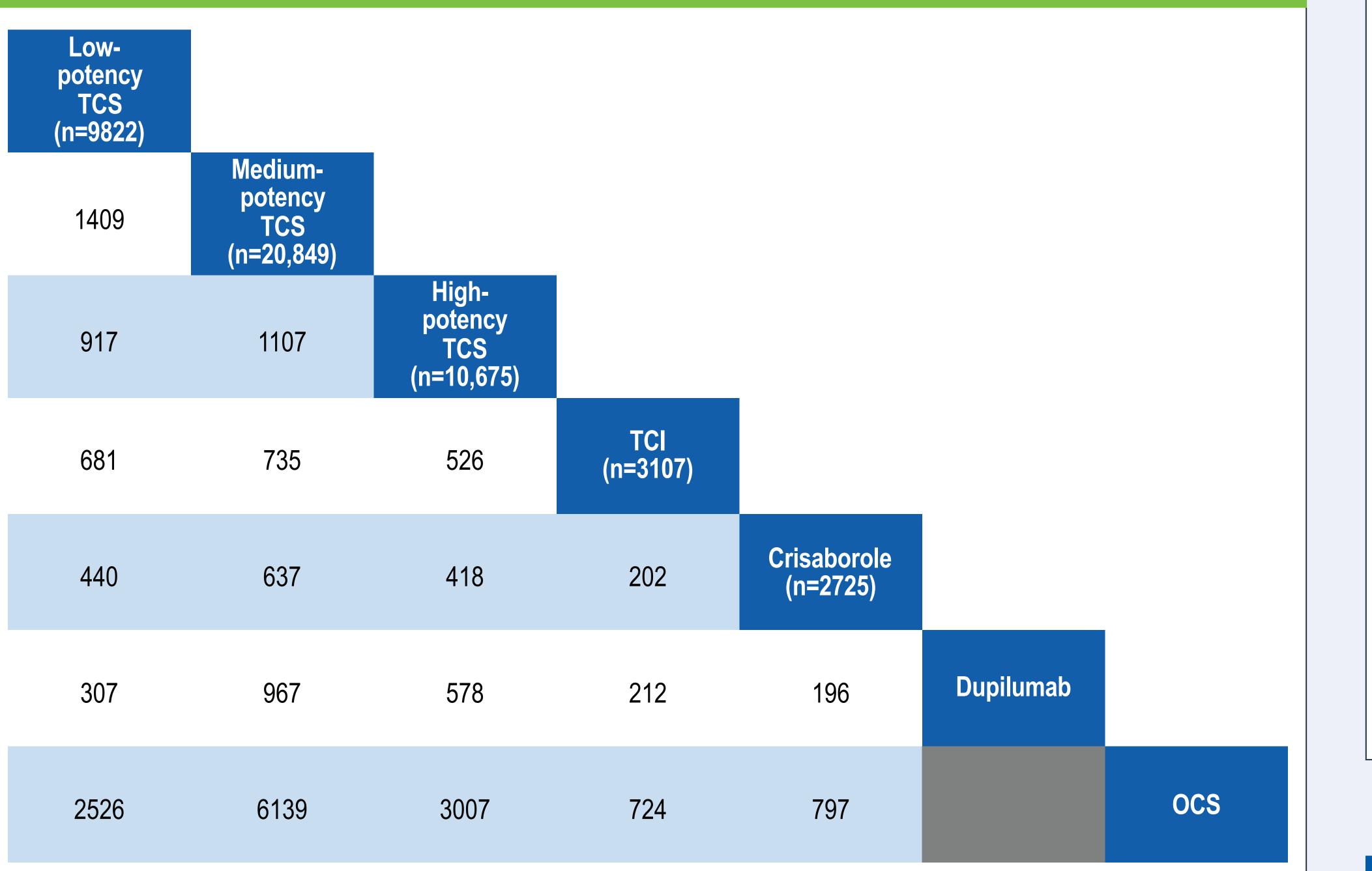
CDHP, consumer-driven health plan; EPO, exclusive provider organization; GAD, generalized anxiety disorder; HDHP, high-deductible health plan; HMO, health maintenance organization; MDD, major depressive disorder; POS, point-of-service; PPO, preferred provider organization; TCI, topical calcineurin inhibitor; TCS, topical corticosteroid.

* Includes data for 394 individuals who were initially treated with dupilumab (subgroup not shown).

Treatment Patterns

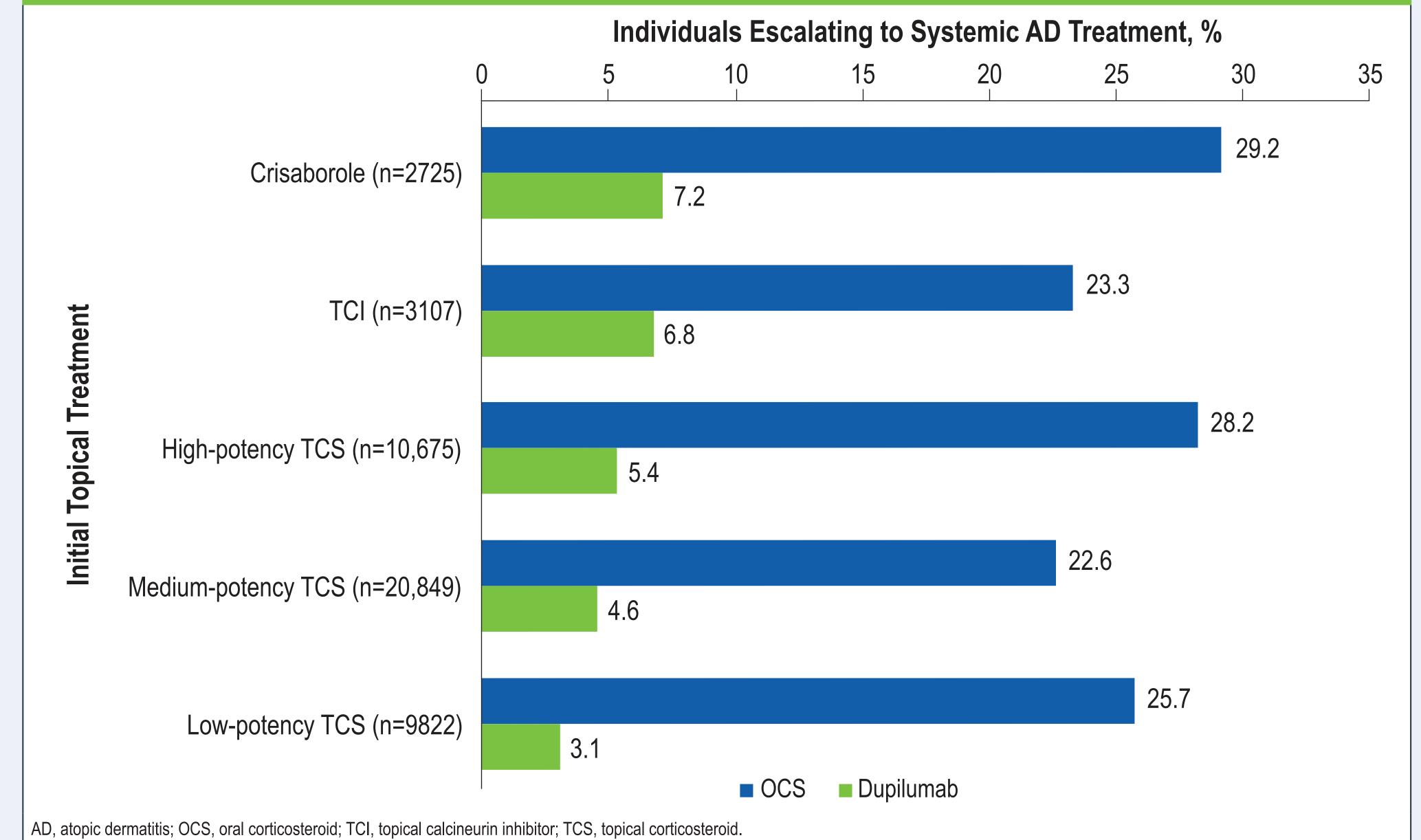
 Across initial topical treatments, 32.8% of individuals were prescribed a systemic treatment (Figure 2)

Figure 2. Number of Individuals Escalating Treatment After Initial Topical Treatment



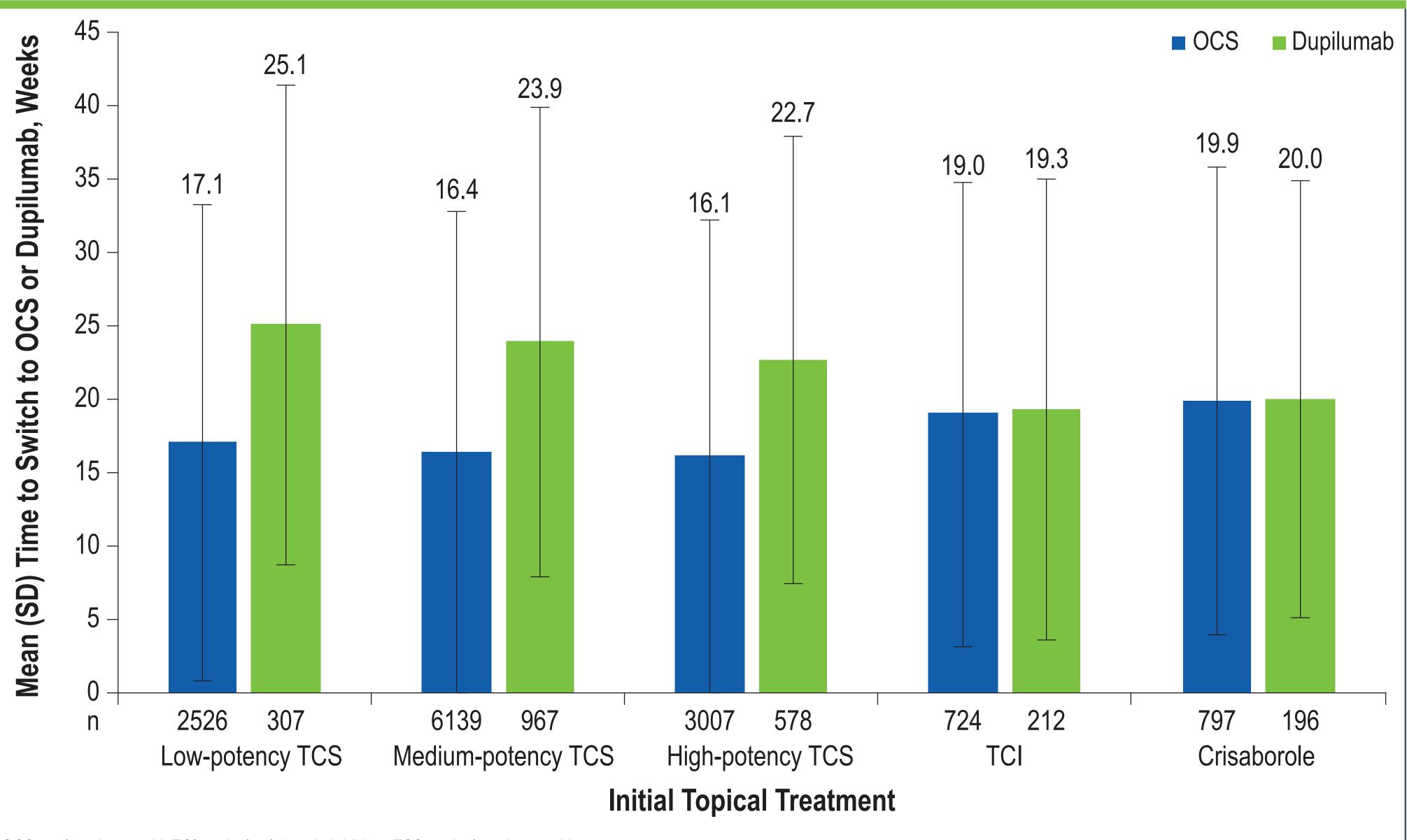
dicated by the dark blue square at the end of each corresponding row.

• 23%–29% of individuals escalated to OCS, and 3%–7% escalated to dupilumab (Figure 3) Figure 3. Proportion of Individuals Escalating to Systemic AD Treatment



• The mean time for adding OCS or dupilumab was 16–20 and 19–25 weeks, respectively, across initial treatments (Figure 4)

Figure 4. Mean Time From Topical Treatment to Use of OCS or Dupilumab



OCS, oral corticosteroid; TCI, topical calcineurin inhibitor; TCS, topical corticosteroid

Some individuals (13.4%) were on ≥2 topical treatments during the study period

Limitations

Limitations to the study are those consistent with retrospective, administrative claims analyses, including the potential for sampling bias, incomplete records, improper data entry, and differences in actual vs physician-directed medication use

Conclusions

- Many US individuals (32.8%) needed a systemic treatment to manage their AD
- A considerable proportion of US individuals with AD (13.4%) needed ≥2 topical treatments in their first year after treatment initiation
- These results suggest that topical treatments (available in 2018) are not sufficient for long-term disease control
- Escalation to systemic treatments occurred after approximately 4–6 months of topical therapy
- These treatment patterns may reflect the real-world effectiveness and treatment success of topical regimens during the analysis period

Disclosures

VNJ, JHL, and ABN are employees and shareholders of Incyte Corporation. SD is a consultant to Incyte Corporation.

Acknowledgments

This study was funded by Incyte Corporation (Wilmington, DE, USA). Writing assistance was provided by Vicky Kanta, PhD, an employee of ICON (Blue Bell, PA, USA) and was funded by Incyte Corporation.

References



of this poster, scan code

1. Langan SM, et al. Lancet. 2020;396(10247):345-360. 2. Eichenfield LF, et al. J Am Acad Dermatol. 2014;71(1):116-132. 3. Papier A, Strowd LC. Drugs Context. 2018;7:212521. 4. Anderson P. et al. Dermatol Ther (Heidelb). 2021;11(5):1571-1585 **5.** Wollenberg A, et al. *J Eur Acad Dermatol Venereol*. 2020;34(12):2717-2744.