

## Atopic Dermatitis Polypharmacy and Out-Of-Pocket Healthcare Expenses the United States

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#### **DISCLOSURES**

RC has served as an advisory board member, consultant, and/or investigator for Abbvie, Arcutis, Arena, Dermavant Incyte, National Eczema Association (NEA), Pfizer, Regeneron, and Sanofi-Genzyme, and speaker for Abbvie, Incyte, Regeneron, Revolutionizing Atopic Dermatitis, Sanofi-Genzyme, and UCB.

WSB is an employee of the NEA and has served as an advisory board member and/or investigator for Incyte and Pfizer.

IJT is an employee of the NEA.

JIS served as a advisory board member, consultant, and/or investigator for Abbvie, AFYX Therapeutics, Arena Pharmaceuticals, Asana Biosciences, BiomX Inc. Bodewell, Boehringer Ingelheim, Bluefin Biomedicine, Celgene, Corrona Inc. Dermayant Sciences, Dermira, DS Biopharma, Eli Lilly, Incyte, Galderma, GlaxoSmithKline, Kiniksa Pharmaceuticals, Leo Pharma, MedImmune, Menlo Therapeutics, Ortho Dermatologics, Pfizer, RAPT Therapeutics, Regeneron, Sanofi-Genzyme, TARGET Pharma, and speaker for Eli Lilly, Maui Derm, Pfizer, Regeneron, and Sanofi-Genzyme.

#### BACKGROUND

- Variable severity, symptoms, burden, and course in atopic dermatitis (AD) contribute to to heterogenous treatment patterns, including polypharmacy.1
- Polypharmacy (regular use of ≥5 but anywhere between 2-11 treatments) is associated with increased adverse events, inappropriate medication use, non-adherence, healthcare system burden, medication errors, healthcare visits, and costs,2
- US population-based studies previously showed increased overall OOP expenses among individuals with AD.3-5
- Prescription polypharmacy and its association with out-of-pocket (OOP) expenses in AD is poorly understood.

#### **OBJECTIVE**

To characterize prescription polypharmacy and its association with OOP healthcare expenses among individuals with AD.

#### **METHODS**

- A 25-question voluntary online survey was administered to National Eczema Association members (N=113,502).
- Inclusion criteria (US residents age ≥18 years; self-report of AD or primary caregivers of individuals with AD) was met by 77.3% (1,118/1,447) of respondents.
- Survey areas included sociodemographic factors, AD severity and control measures, categories of OOP expenses, monthly and annual OOP expenses, and household financial impact.
- Chi-square tests were used for comparisons of categorical variables. Kruskal-Wallis one-way analysis of variance was used for comparison of median annual OOP costs

#### **RESULTS**

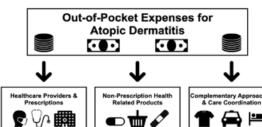
## Polypharmacy was associated with respondent disease burden

Wardahla dana MO	Overall (n=1,118)	Total Number of Prescriptions				
		0	1-2	3-4	5+	P-value
Variable – freq (%)		(n=112)	(n=344)	(n=311)	(n=306)	
Current AD severity						
Clear	29 (2.6%)	5 (19.2%)	10 (38.5%)	7 (26.9%)	4 (15.4%)	< 0.0001
Mild	238 (21.3%)	39 (13.0%)	103 (34.8%)	58 (34.8%)	31 (17.4%)	
Moderate	531 (47.5%)	53 (10.4%)	150 (29.5%)	169 (33.3%)	136 (26.8%)	
Severe	296 (26.5%)	12 (4.2%)	73 (25.6%)	69 (24.2%)	131 (46.0%)	
Current AD control						
Minimally controlled	259 (23.2%)	21 (8.5%)	81 (32.7%)	54 (21.8%)	92 (37.1%)	0.0005
Somewhat controlled	448 (40.1%)	29 (9.5%)	99 (29.2%)	89 (34.1%)	72 (27.2%)	
Moderately well controlled	300 (26.8%)	41 (10.0%)	126 (34.3%)	147 (30.8%)	117 (24.9%)	
Very well controlled	102 (9.1%)	19 (19.8%)	34 (35.4%)	21 (21.9%)	22 (22.9%)	
Number of flare days in past 30 days						
0	44 (4.0%)	7 (17.1%)	17 (41.5%)	7 (17.1%)	10 (24.4%)	< 0.0001
1-3	271 (24.4%)	33 (12.7%)	93 (35.9%)	84 (32.4%)	49 (18.9%)	
4-7	206 (18.6%)	19 (9.4%)	69 (34.2%)	62 (30.7%)	52 (25.7%)	
8-10	151 (13.6%)	11 (7.4%)	47 (31.5%)	55 (36.9%)	38 (24.2%)	
≥11	437 (39.4%)	42 (10.0%)	118 (28.0%)	103 (24.5%)	158 (37.5%)	
Comorbidities						
Asthma	382 (34.5%)	32 (8.6%)	105 (28.3%)	107 (28.8%)	127 (34.2%)	0.013
Allergic rhinitis	557 (50.4%)	48 (8.9%)	158 (29.2%)	159 (29.4%)	176 (32.5%)	0.009
Food allergy	426 (38.5%)	30 (7.3%)	104 (25.2%)	126 (30.6%)	152 (36.9%)	< 0.0001
Frequent/persistent skin infections	210 (19.0%)	10 (4.9%)	31 (15.1%)	54 (26.3%)	110 (54.7%)	< 0.0001
Anxiety and/or depression	404 (36.5%)	33 (8.4%)	122 (30.9%)	114 (28.9%)	126 (31.9%)	0.15
HCP visits in past year						
0	113 (10.6%)	56 (49.6%)	44 (35.4%)	9 (9.7%)	3 (5.3%)	< 0.0001
1-2	435 (40.6%)	40 (10.1%)	219 (50.3%)	61 (26.7%)	23 (12.9%)	
3.4	284 (26.5%)	11 (3.2%)	116 (21.5%)	118 (41.6%)	65 (33.8%)	
≥5	239 (22.4%)	6 (1.3%)	56 (9.6%)	96 (27.2%)	148 (61.9%)	

#### Polypharmacy (≥5 Rx treatments) was associated with:

- increased AD severity
- increased HCP visits
- poorer control increased flares
- allergic and infectious comorbidities

### Polypharmacy was associated with multiple categories of OOP expenditures



- Office visit deductibles Moisturizers Prescription co-pays Anti-itch medications Allergy medications Prescriptions not covere
- Outpatient phototherapy · Lab testing Mental health services
- Hospitalizations Emergency room visit
- Hygiene product
- Adjunctive therapy · Clothing & bedding
- Sleep medication

Figure adapted from Smith Begolka W. Chovativa R. Thibau IJ. Silverberg Jl. Dermatitis, 2021 Oct 1:32(1S):S62-S70

#### Polypharmacy was associated with all queried categories of OOP expenses

- healthcare providers Rx products
- complementary approaches · care coordination
- non-Rx products

### RESULTS

#### Polypharmacy was associated with increased monthly and yearly OOP expenditures

# Monthly CCP expenses for HCP visits \$1-\$100 \$101-\$200 Monthly OOP expenses for Rs not covered by

■ 0 Rx ■ 1-2 Rx ■ 3-4 Rx ■ 5+ Rx

## ■ 0 Rx ■ 1-2 Rx ■ 3-4 Rx ■ 5+ Rx Monthly OOP expenses for Rx covered by insurance

#### Polypharmacy was associated with systemic and topical Rx use

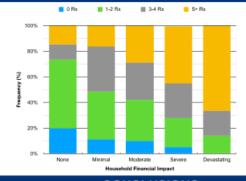
biologics (dupilumab)

\*P.0.0001 for all figures

- oral immunosuppressants (AZA, CsA, MTX, SCS)
- topical calcineurin inhibitor (P≤0.005 for all)

oral antimicrobials

### Polypharmacy was associated increased financial impact



#### CONCLUSIONS

- Individuals with AD report considerable polypharmacy which is associated with increased OOP expenses and significant household financial impact.
- Strategies are needed to reduce polypharmacy, minimize OOP costs, and optimize clinical outcomes

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