

# Efficacy and Safety of Roflumilast Foam 0.3% in Patients With Seborrheic Dermatitis in a Phase 3 Trial: Assessment of Pruritus

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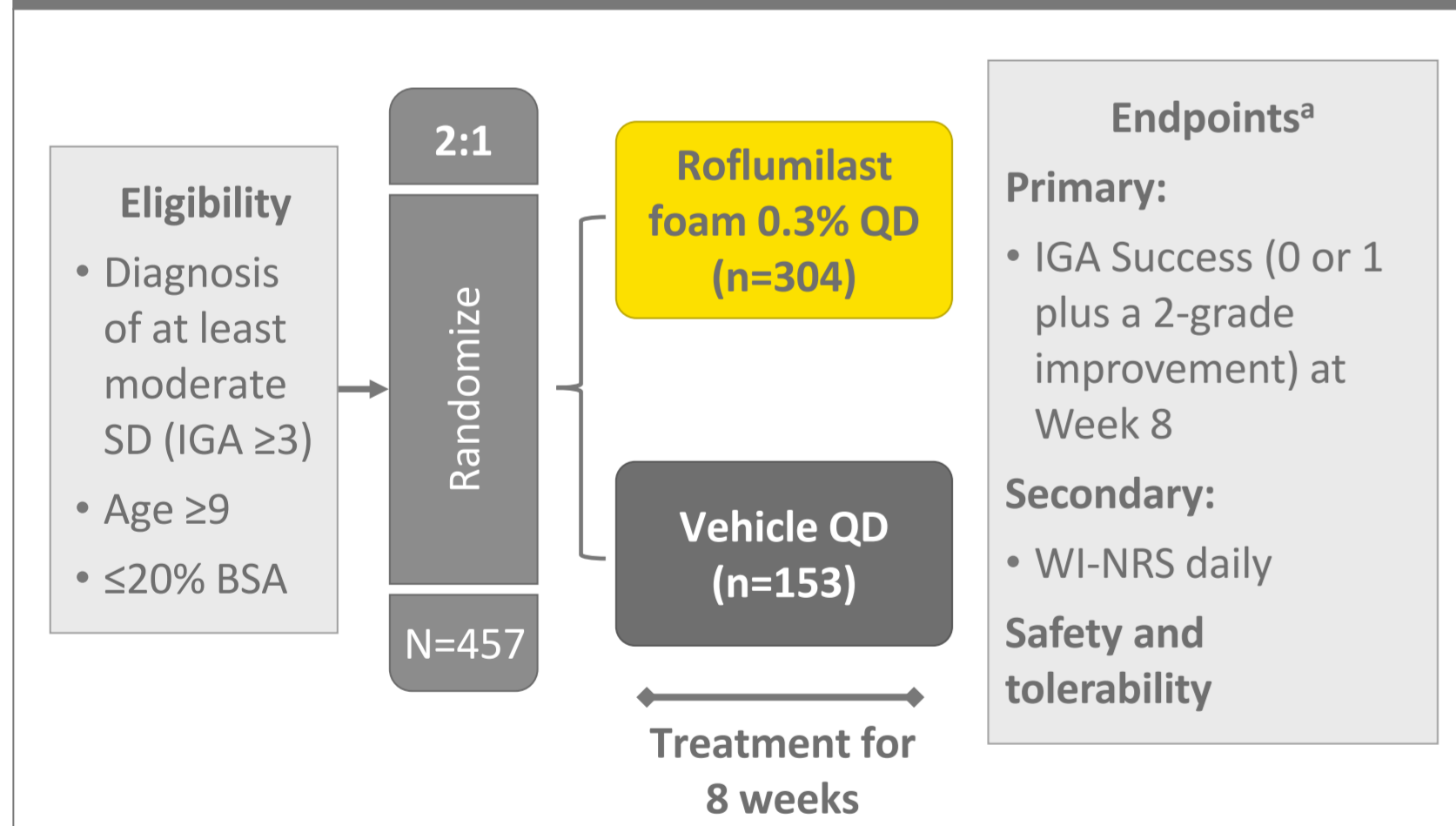
## INTRODUCTION

- Seborrheic dermatitis (SD) is a chronic inflammatory skin condition that negatively impacts quality of life, particularly in patients with more severe disease<sup>1</sup>
  - Itch is a major complaint among patients with SD<sup>2</sup>
- Topical treatments include antifungals, steroids, immunomodulators, and dandruff shampoos,<sup>3,4</sup> but efficacious and safe options are needed, especially those that improve itch
- Roflumilast is a selective, potent, phosphodiesterase 4 inhibitor being investigated as a once-daily foam for treatment of SD<sup>5</sup>

## METHODS

- This phase 3, randomized, parallel-group, double-blind, vehicle-controlled trial (NCT04973228) was conducted in patients ≥9 years old with at least moderate SD affecting scalp and/or non-scalp areas
- Eligible patients had clinical diagnosis of SD of ≥3-month duration, Investigator Global Assessment (IGA) score ≥3 (at least moderate severity), and affecting ≤20% of the body surface area (BSA; **Figure 1**)
- Patients were randomized 2:1 to apply once-daily roflumilast foam 0.3% (n=304) or vehicle (n=153) for 8 weeks
- The primary efficacy endpoint was IGA Success (Completely Clear/Almost Clear [score 0–1] plus ≥2-grade improvement) at Week 8
  - Secondary efficacy endpoints included Worst Itch Numeric Rating Scale (WI-NRS), which was completed daily by patients
- Safety and local tolerability were also evaluated

**Figure 1. Study Design**



<sup>a</sup>As this study is a single pivotal trial, the statistical significance of the primary endpoint was assessed at the 1% significance level (2-sided). To control for multiple testing, the 1% alpha was partitioned to 0.0033 for WI-NRS endpoints and 0.0067 for other secondary endpoints. BSA: body surface area; IGA: Investigator Global Assessment; QD: once daily; SD: seborrheic dermatitis; WI-NRS: Worst Itch Numeric Rating Scale.

## RESULTS

- Demographics and baseline characteristics were similar in the treatment groups (**Table 1**)
- Overall, significantly more roflumilast-treated patients than vehicle-treated patients achieved IGA success (79.5% vs 58.0%;  $P<0.0001$ ) and IGA status of Clear (50.6% vs 27.7%;  $P<0.0001$ ) at Week 8 (**Figure 2**)
- Significantly greater percentages of roflumilast- than vehicle-treated patients had ≥4-point improvement on WI-NRS at Weeks 2 (32.7% vs 15.5%;  $P=0.0005$ ), 4 (47.6% vs 29.1%;  $P=0.0003$ ), and 8 (62.8% vs 40.6%;  $P<0.0001$ ; **Figure 3**)

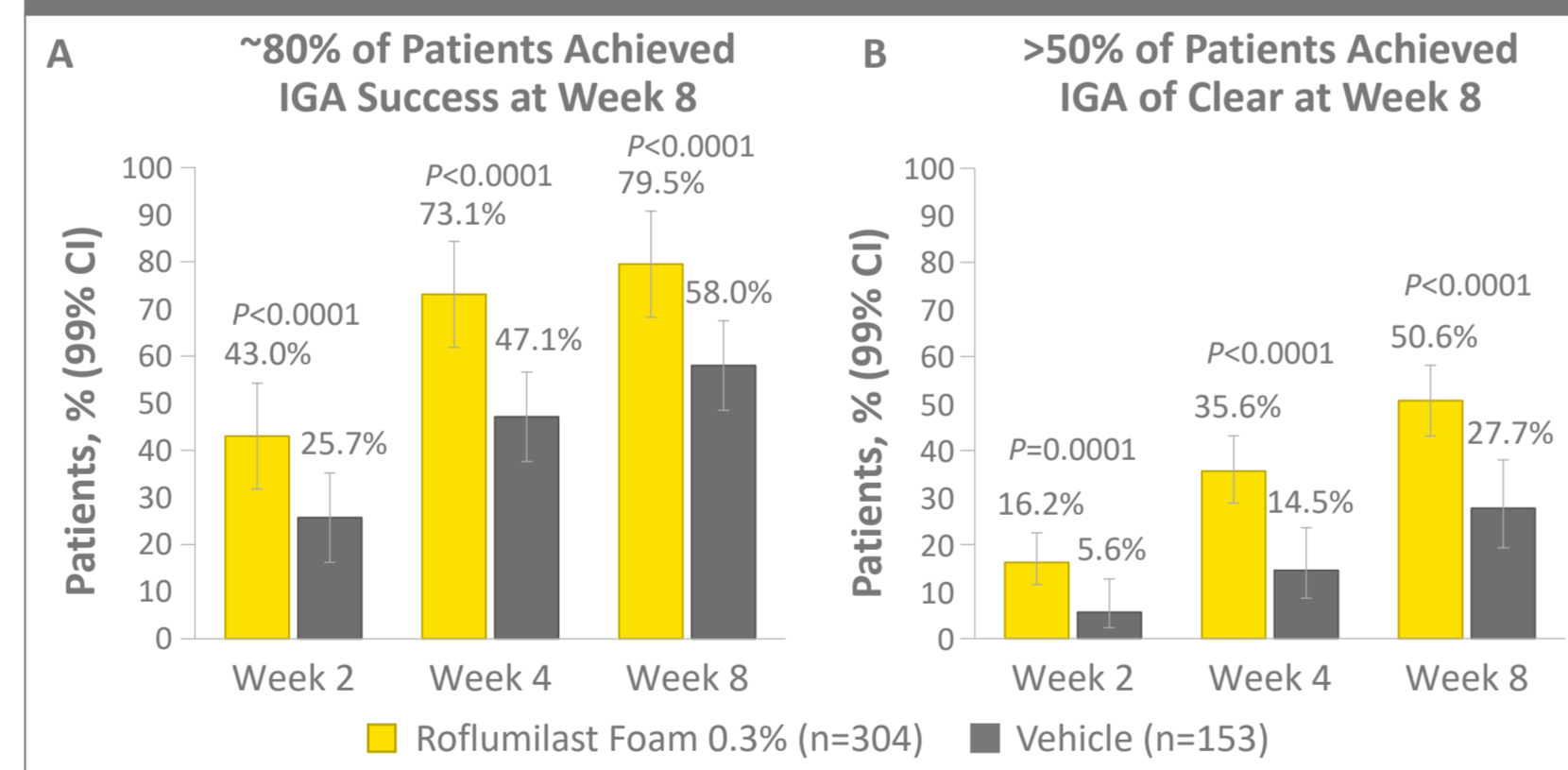
- Greater improvement in itch was observed among roflumilast-treated patients as early as 48 hours after the first application (mean percent change from baseline: -27.87% vs -13.11%; nominal  $P=0.0024$ ; **Figure 4**)
- Changes in SD in patients treated with roflumilast foam 0.3% are shown in **Figure 5**

**Table 1. Baseline Demographics and Disease Characteristics**

	Roflumilast Foam 0.3% (n=304)	Vehicle (n=153)
<b>Age in years, mean (Std Dev)</b>	43.2 (16.8)	41.8 (17.5)
<b>Sex</b>		
Male, n (%)	153 (50.3)	75 (49.0)
Female, n (%)	151 (49.7)	78 (51.0)
<b>Race, n (%)</b>		
American Indian or Alaska Native	4 (1.3)	0
Asian	18 (5.9)	10 (6.5)
Black or African American	36 (11.8)	15 (9.8)
Native Hawaiian or Other Pacific Islander	0	1 (0.7)
White	234 (77.0)	122 (79.7)
More than 1 race	1 (0.3)	1 (0.7)
Other	11 (3.6)	4 (2.6)
<b>Ethnicity</b>		
Hispanic or Latino	69 (22.7)	28 (18.3)
Not Hispanic or Latino	235 (77.3)	125 (81.7)
<b>IGA score, n (%)</b>		
3 (moderate)	287 (94.4)	141 (92.2)
4 (severe)	17 (5.6)	12 (7.8)
<b>WI-NRS, mean score (Std Dev)</b>	5.06 (2.34)	4.74 (2.29)
<b>WI-NRS score ≥4, n (%)</b>	206 (67.8)	98 (64.1)
<b>BSA, mean % (Std Dev)</b>	2.89 (2.03)	2.98 (2.57)

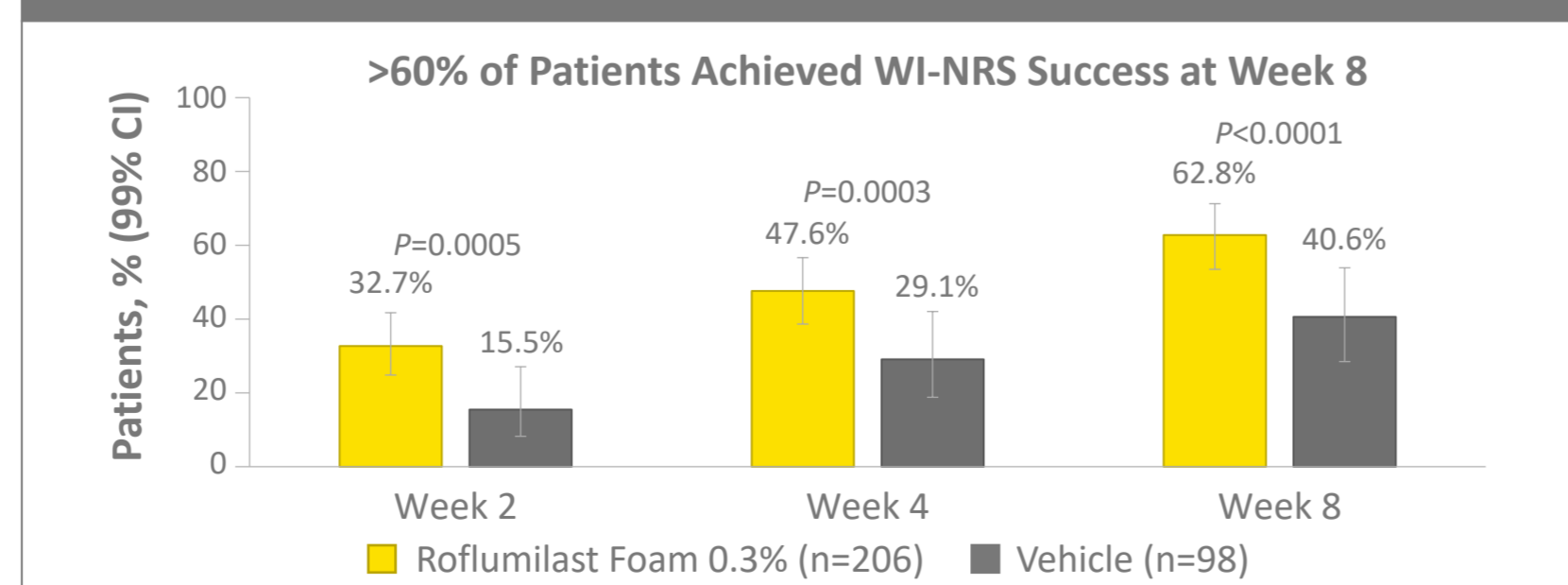
BSA: body surface area; IGA: Investigator Global Assessment; Std Dev: standard deviation; WI-NRS: Worst Itch Numeric Rating Scale.

**Figure 2. Patients Achieving IGA Success and IGA Clear**



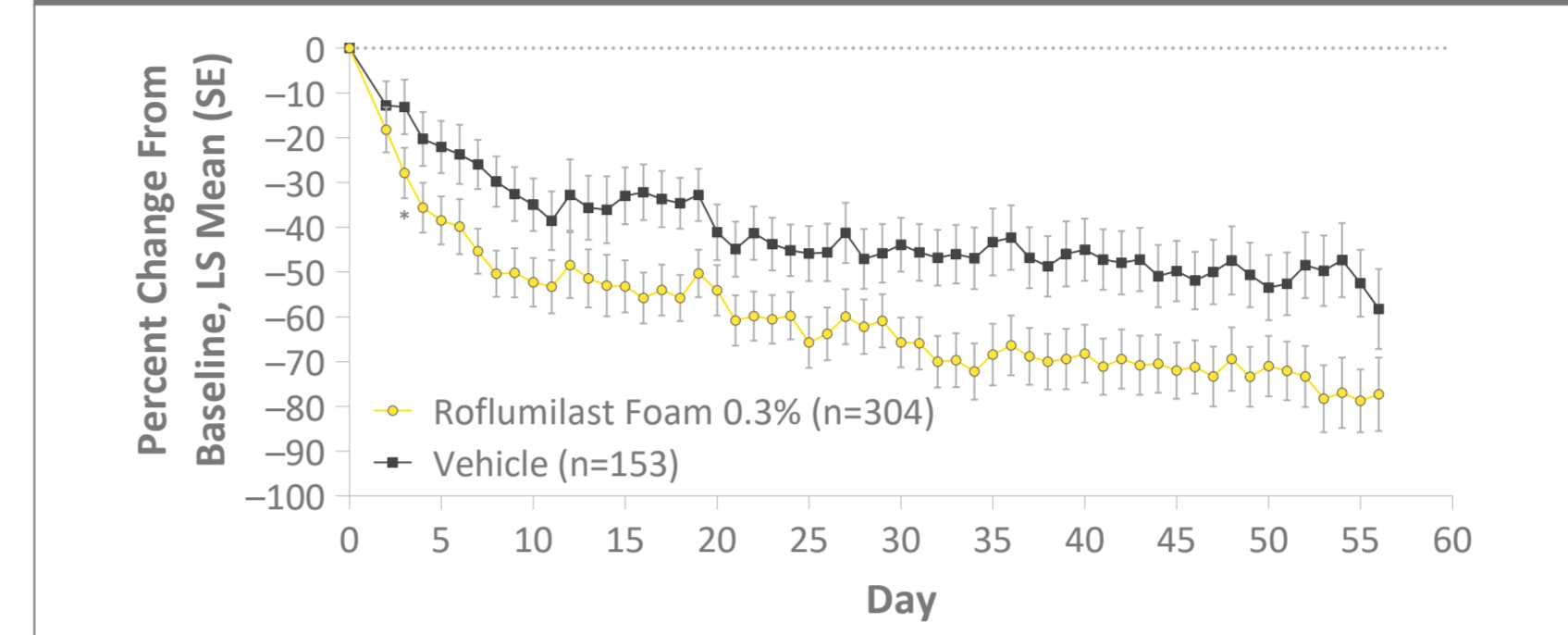
IGA Success = Clear or Almost Clear with at least a 2-grade improvement from baseline. Intent-to-treat population; missing scores imputed using multiple imputations. Error bars represent 99% confidence interval. Statistical significance was concluded at the 1% significance level (2-sided). IGA Clear = IGA score of 0. P-values are not adjusted for multiple testing. CI: confidence interval; IGA: Investigator Global Assessment.

**Figure 3. Percentage of Patients Achieving WI-NRS Success**



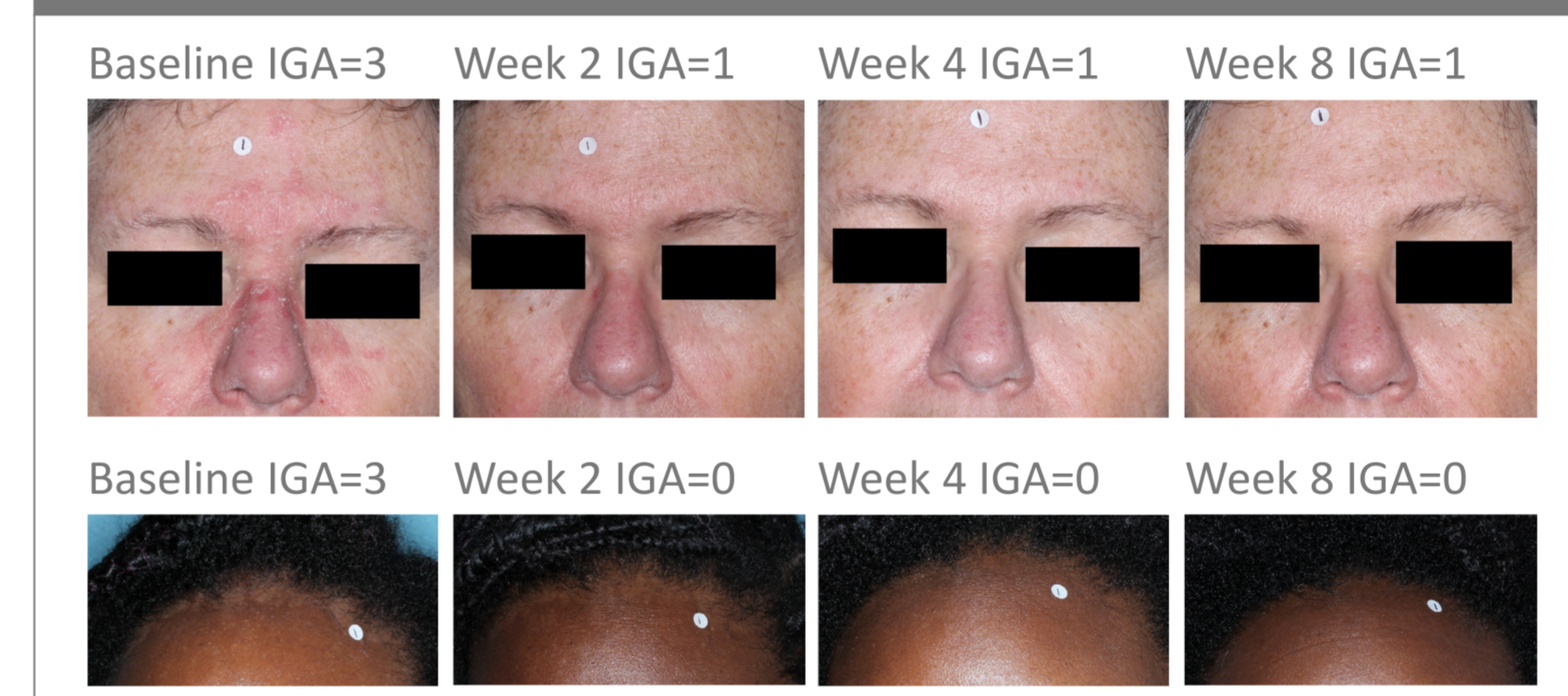
Missing scores imputed using multiple imputations. Error bars represent 99% confidence intervals. WI-NRS Success = ≥4-point improvement in patients with baseline WI-NRS score ≥4, evaluated at n=203. CI: confidence interval; WI-NRS: Worst Itch Numeric Rating Scale.

**Figure 4. LS Mean Percent Change in Daily WI-NRS Scores**



\* $P=0.0024$  at 48 hours after dosing and  $P<0.05$  for difference from vehicle for all timepoints assessed after. Observed data, intent-to-treat population. LS: least squares; SE: standard error; WI-NRS: Worst Itch Numeric Rating Scale.

**Figure 5. Changes in SD in Patients Treated With Roflumilast Foam 0.3%**



IGA: Investigator Global Assessment; SD: seborrheic dermatitis.

## Safety

- Rates of adverse events (AEs) with roflumilast foam and vehicle foam were low (**Table 2**)
  - Few treatment-related AEs were reported
  - Very few AEs led to study discontinuation, with similar rates of discontinuation between roflumilast and vehicle groups
  - Only 1 patient had a serious AE, and it was considered unrelated to treatment

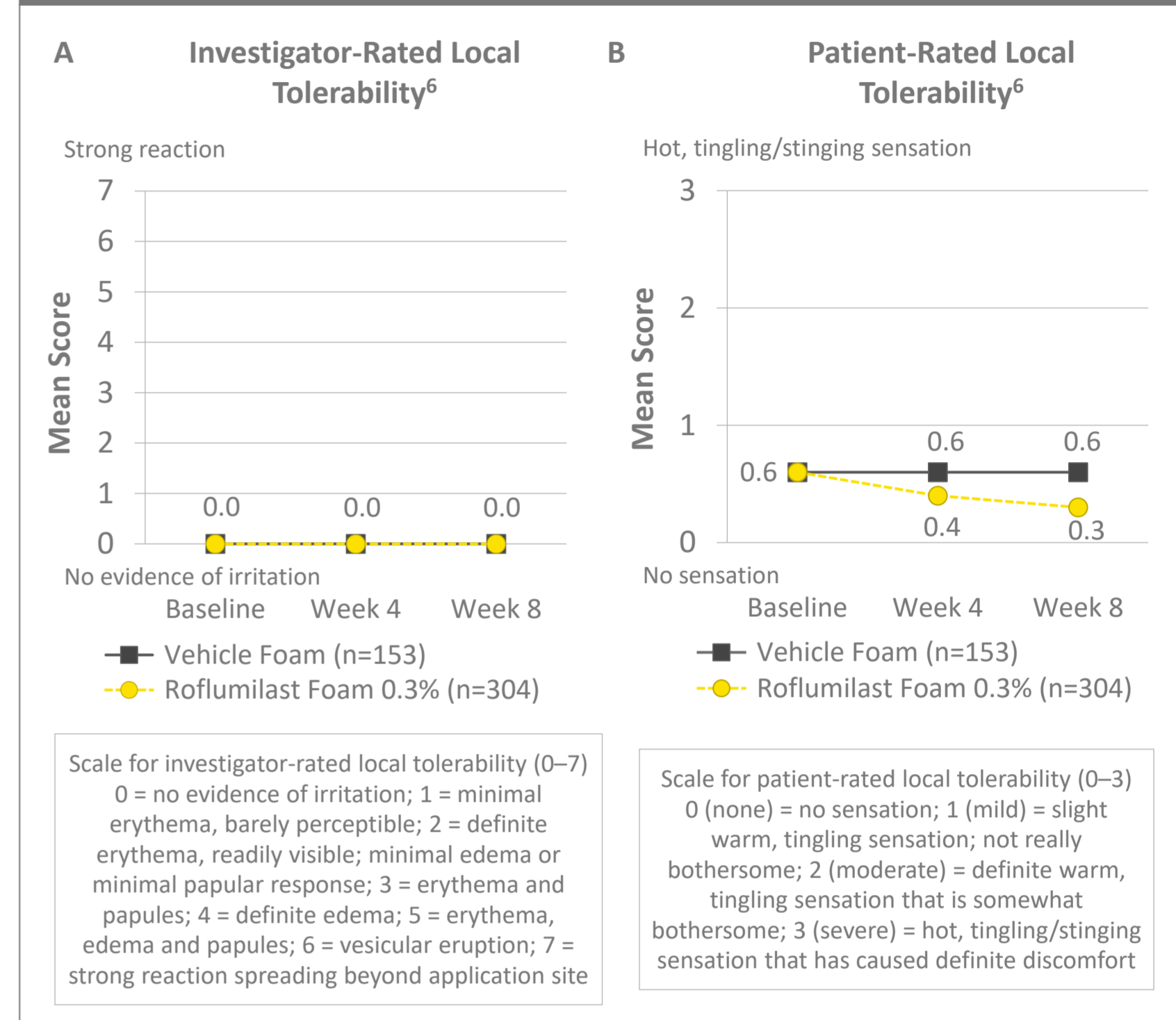
**Table 2. Overall AEs**

n (%)	Roflumilast Foam 0.3% (n=304)	Vehicle (n=153)
<b>Patients with any TEAE</b>	70 (23.0)	33 (21.6)
<b>Patients with any treatment-related TEAE</b>	8 (2.6)	5 (3.3)
<b>Patients with any treatment-emergent SAE<sup>a</sup></b>	1 (0.3)	0
<b>Patients who discontinued study due to AE<sup>b</sup></b>	2 (0.7)	3 (2.0)
<b>Most common TEAE (&gt;1% in any group), preferred term<sup>c</sup></b>		
COVID-19	11 (3.6)	5 (3.3)
Urinary tract infection	4 (1.3)	3 (2.0)
Nausea	5 (1.6)	0
Nasopharyngitis	4 (1.3)	1 (0.7)
Application-site pain	1 (0.3)	3 (2.0)
Sinusitis	0	2 (1.3)

<sup>a</sup>Keratoacanthoma, not in application site, deemed unrelated. <sup>b</sup>Reasons for discontinuation in the roflumilast-treated group include diarrhea/hematochezia/abdominal pain in one patient with a past history of Crohn's and decreased potassium in the second patient. <sup>c</sup>Presented in descending order for overall rates. AE: adverse event; SAE: serious adverse event; TEAE: treatment-emergent adverse event.

- Roflumilast foam was well-tolerated (**Figure 6**)
  - ≥98.9% of roflumilast-treated and 100% of vehicle-treated patients had no evidence of irritation on the investigator-rating of local tolerability
  - ≥92.4% of patients had reported "no sensation" or "slight warm, tingling sensation; not really bothersome" on the patient-rated assessment of local tolerability

**Figure 6. Local Tolerability Assessments (Safety Population)**



## CONCLUSIONS

- Once-daily, non-steroidal roflumilast foam 0.3% provided improvement across multiple efficacy endpoints, including rapid itch improvement, versus vehicle in patients with SD in a phase 3 trial
  - 80% of patients achieved IGA Success and >50% achieved complete clearance by Week 8
  - >60% of patients achieved an itch response at Week 8, with significant improvements at the 2- and 4-week assessments
  - Greater improvement in daily itch scores was observed among roflumilast-treated patients as early as 48 hours after the first dose
- Local tolerability was highly favorable on investigator- and patient-rated assessments and was consistent with safety profiles in prior trials

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## DISCLOSURES

AB, ZDD, MG, EL, AYM, KAP, and MZ are investigators and/or consultants for Arcutis Biotherapeutics, Inc. and received grants/research funding and/or honoraria; DK, PB, DRB, and DHC are employees of Arcutis Biotherapeutics, Inc. Additional disclosures provided on request.