

SUPPLEMENTARY INFORMATION

Switching From Dupilumab to Abrocitinib in Patients With Moderate-to-Severe Atopic Dermatitis: A Post Hoc Analysis of Efficacy After Treatment With Dupilumab in JADE DARE

Jonathan I. Silverberg,¹ Eric L. Simpson,² Andrew E. Pink,^{3,4} Stephan Weidinger,⁵ Gary Chan,⁶ Pinaki Biswas,⁷ Claire Clibborn,^{8a} Erman Güler⁹

¹The George Washington University School of Medicine and Health Sciences, 2300 I St NW, Washington, DC, USA; ²Oregon Health & Science University, 3303 SW Bond Ave, Portland, OR, USA; ³St. John's Institute of Dermatology, King's College London, 2 Lambeth Palace Rd, London, UK; ⁴Guy's and St Thomas' NHS Foundation Trust, Westminster Bridge Road, London, UK; ⁵University Hospital Schleswig-Holstein, Arnold-Heller-Str 3, Kiel, Germany; ⁶Pfizer Inc., 500 Arcola Road, Groton, CT, USA; ⁷Pfizer Inc., New York, 66 Hudson Boulevard, NY, USA; ⁸Pfizer Ltd., Walton Oaks, Dorking Rd, Surrey, UK; ⁹Pfizer Inc., Büyükdere Cd. No:199, 34394 Levent Istanbul, Turkey

^aAffiliation at the time this analysis was conducted.

Table S1. Baseline demographics and clinical characteristics of dupilumab responders and nonresponders by efficacy outcomes at Week 26 of JADE DARE

	IGA 0/1^a (n = 310)		EASI-75 (n = 310)		PP-NRS4 (n = 311)	
	Responders (n = 179)	Nonresponders (n = 131)	Responders (n = 254)	Nonresponder s (n = 56)	Responder s (n = 217)	Nonresponders (n = 94)
Age, mean ± SD, years	36 ± 13	37 ± 14	36 ± 13	38 ± 14	37 ± 14	34 ± 13
Female, n (%)	93 (52)	45 (34)	122 (48)	16 (29)	102 (47)	37 (39)
Race, n (%)						
White	127 (71)	80 (61)	171 (67)	36 (64)	148 (68)	59 (63)
Other ^b	52 (29)	51 (39)	83 (33)	20 (36)	69 (32)	35 (37)
Weight, mean ± SD, kg	76 ± 19	82 ± 23	77 ± 19	88 ± 26	78 ± 20	79 ± 23
IGA score of 4, n (%)	61 (34)	67 (51)	102 (40)	26 (46)	97 (45)	32 (34)

EASI, mean $\pm$ SD	25 $\pm$ 10	29 $\pm$ 13	27 $\pm$ 12	26 $\pm$ 12	28 $\pm$ 13	25 $\pm$ 10
PP-NRS, mean $\pm$ SD	7.1 $\pm$ 1.6 ^c	7.0 $\pm$ 1.6 ^d	7.0 $\pm$ 1.6 ^e	7.1 $\pm$ 1.6 ^f	7.2 $\pm$ 1.5 ^g	6.6 $\pm$ 1.8 ^h

^aIGA response defined as IGA score of 0 (clear) or 1 (almost clear) and ≥ 2 -point improvement from baseline

^bPatients who identified as Black or African American, Asian, multiracial, or did not report their race

^c $n = 177$

^d $n = 127$

^e $n = 250$

^f $n = 54$

^g $n = 216$

^h $n = 89$

EASI Eczema Area and Severity Index, *EASI-75* $\geq 75\%$ improvement from baseline in EASI, *IGA* Investigator's Global

Assessment, *PP-NRS* Peak Pruritus Numerical Rating Scale, *PP-NRS4* ≥ 4 -point improvement from baseline in PP-NRS

Table S2. Proportions of dupilumab responders and nonresponders at Week 26 of JADE DARE, based on various response criteria, who switched to abrocitinib 200 mg in JADE EXTEND^a

Response criterion at Week 26 of JADE DARE	Dupilumab-treated patients in JADE DARE who switched to abrocitinib 200 mg in JADE EXTEND (N = 309) ^b	
	Dupilumab responders	Dupilumab nonresponders
IGA response ^c , n (%)	178 (58)	131 (42)
EASI-50, n (%)	292 (94)	17 (6)
EASI-75, n (%)	254 (82)	55 (18)
EASI-90, n (%)	168 (54)	141 (46)
PP-NRS4, n (%)	217 (70)	92 (30)
PP-NRS 0/1, n (%)	113 (36) ^d	197 (64) ^d

^aData were collected during the end of treatment visit of JADE DARE/screening visit of JADE EXTEND

^bData were missing for some patients who did not have their efficacy responses measured at the end of treatment visit of JADE DARE/screening visit of JADE EXTEND

^cIGA response defined as IGA score of 0 (clear) or 1 (almost clear) and ≥ 2 -point improvement from baseline

^dN = 310

DLQI Dermatology Life Quality Index, *DLQI 0/1* DLQI score of 0 or 1, *EASI* Eczema Area and Severity Index, *EASI-50* $\geq 50\%$ improvement from baseline in EASI, *EASI-75* $\geq 75\%$ improvement from baseline in EASI, *EASI-90* $\geq 90\%$ improvement from baseline in EASI, *IGA* Investigator's Global Assessment, *PP-NRS* Peak Pruritus Numerical Rating Scale, *PP-NRS 0/1* PP-NRS score of 0 or 1, *PP-NRS4* ≥ 4 -point improvement from baseline in PP-NRS