

Data Supplement

Supplementary Table 1. Treatment-emergent adverse events reported in >5% of participants in either ozanimod HCl dose group

	Core Period and Blinded Extension	
	Ozanimod HCl 0.5 mg (<i>n</i> =126)	Ozanimod HCl 1 mg (<i>n</i> =123)
Nasopharyngitis	24 (19.0)	21 (17.1)
Upper respiratory tract infection	21 (16.7)	14 (11.4)
Increased alanine aminotransferase	14 (11.1)	16 (13.0)
Increased gamma-glutamyl transferase	13 (10.3)	15 (12.2)
Headache	11 (8.7)	11 (8.9)
Hypertension	10 (7.9)	4 (3.3)
Urinary tract infection	10 (7.9)	5 (4.1)
Back pain	9 (7.1)	9 (7.3)
Pharyngitis	9 (7.1)	6 (4.9)
Depression	8 (6.3)	8 (6.5)
Pain in extremity	8 (6.3)	4 (3.3)

Data are *n* (%).

Supplementary Table 2. Summary of all reported serious treatment-emergent adverse events

	Ozanimod HCl 0.5 mg^a (n=126)	Ozanimod HCl 1 mg^a (n=123)
Acute myocardial infarction	1	0
Cauda equina syndrome	1	0
Clavicle fracture	0	1
Concussion	1	0
Depression	0	1
Headache	0	1
Hepatitis	1	0
Hypertension	1	0
Infectious proctitis	1	0
Injury	1	0
Intracranial aneurysm	1	0
Irritable bowel syndrome	0	1
Lower limb fracture	1	0
Menometrorrhagia	1	0
Pancytopenia	0	1
Optic neuritis	1 ^b	0
Ovarian cyst	1	0
Rheumatoid arthritis	0	1
Somatoform disorder (cardiovascular)	1 ^b	0

Stasis dermatitis	0	1
Urethral stenosis	0	1
Urticaria	1	0
Uterine cervical squamous metaplasia	1 ^b	0
Uterine hemorrhage	0	1

TEAE: treatment-emergent adverse event.

^aOzanimod HCl 0.5 mg, 12 participants experienced a total of 16 serious TEAEs; ozanimod HCl 1.0 mg, 9 participants experienced a total of 9 serious TEAEs.

^bOccurred during the placebo-controlled phase (weeks 0–24).

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