Supplemental Table S4. Adverse Events during the Clinical Trial

Variable	BPS group (n=27)	PCB group (n=25)	P value
Pruritus	1	3	0.34
Urticaria	1	0	1.00
Dyspepsia	1	1	1.00
Heart burn	0	1	0.48
Edema	0	1	0.48
Fatigue	0	1	0.48
Headache	2	0	0.49
Fever	1	0	1.00
Acute pharyngotonsillitis	1	0	1.00
Numbness	1	0	1.00
Total	8	7	1.00

BPS, beraprost sodium; PCB, placebo.