Combination of mometasone furoate and oxymetazoline for the treatment of adenoid hypertrophy concomitant with allergic rhinitis:

A randomized controlled trial

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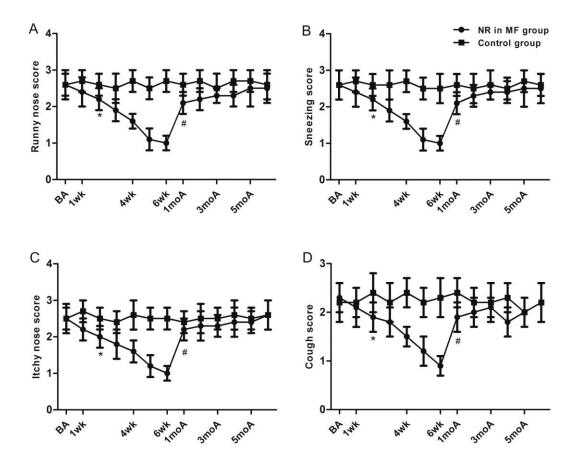


FIG. E1. Individual symptom scores of different time points between non-responders and control during the first treatment stage. A for runny nose score, B for sneezing score, C for itchy nose score and D for cough score. * Compared with baseline score, P < 0.05. * P < 0.05 versus the lowest score during treatment.

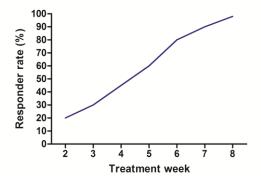


FIG. E2. Responder rate during different treatment weeks.

Table E1 Disposition of subjects and the reasons for discontinuation

	Sta	age 1		Stage 2				
Disposition summary	MF	Placebo	MF+Placebo	MF+OXY	Placebo+OXY	Placebo+Placebo		
Screened subjects, no	120	120	17	17	17	17		
Completed subjects, no	112	109	15	16	16	16		
Discontinued subjects no	8	11	2	1	1	1		
Reason for discontinuation								
Adverse event	1	1	0	0	1	0		
Abnormal test result(s)	0	1	0	0	0	0		
Treatment failure	0	0	0	0	0	0		
Protocol violation	0	3	0	0	0	0		
Noncompliance	2	2	1	0	0	0		
Consent withdrawn	3	2	0	0	0	1		
Lost to follow-up	2	2	1	1	0	0		
Safety population, no. (%)	110 (99)	109 (100)	17 (100)	16 (95)	17 (100)	17 (100)		
ITT population, no. (%)	111 (99)	109 (100)	17 (100)	16 (95)	17 (100)	17 (100)		

ITT, Intent-to-treat.

#Intent-to-treat population includes all subjects who were randomized and had at least 1 postbaseline efficacy observation.

Table E2 Overview of treatment-related adverse events

	Stage 1						Stage 2					
		MF	Placebo		MF+Placebo MF+O		XY Placebo+		+OXY	Y Placebo+Placebo		
Number	110		109		15		16		16		16	
Any adverse event	9		8		2		1		2		2	
Dysgeusia	2		0		0		0		0		1	
Epistaxis	2		2		1		0		0		0	
Headache	1		1		0		1		1		1	
Nasal discomfort		0		1		0		0		1		0
Nausea	1		2		1		0		0		0	
Sneezing		2		1		0		0		0		0
Mucosal erosion		0		1		0		0		0		0
Somnolence		1		0		0		0		0		0

^{*}Safety population includes all randomized subjects who took at least 1 dose of the study drug.