House dust mite sublingual immunotherapy in patients with receiving subcutaneous immunotherapy maintenance phase: A randomized controlled trial

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from SCIT to SLIT. This study aims to assess the efficacy of switching SCIT to SLIT in patients with house $(\mathbf{I} \mathbf{I} \mathbf{I} \mathbf{I} \mathbf{I}) = 11$





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	SLIT (N = 19)	SCIT (N = 20)	P value
ale (%)	10 (52.6)	14 (70)	0.26
mean±SEM)	39.8±4.4	42.8±2.9	0.57
nitis (%)	19 (100)	20 (100)	>0.99
isease			
thma (%)	4 (21)	4 (20)	>0.99
matitis (%)	4 (21)	1 (5)	0.13
nsitization			
	19 (100)	20 (100)	>0.99
	18 (94.7)	18 (90)	>0.99
ation (%)	4 (21)	10 (50)	0.09
(%)	1 (5.2)	9 (45)	0.008
	1 (5.2)	4 (20)	0.34
)	3 (15.8)	6 (30)	0.45
SCIT, weeks (mean±SEM)	90.9±16.5	148.6±50.9	0.71
sinophil count	176.1±34.9	195.1±25.4	0.28
symptom score	4.8±1.0	3.5±0.7	0.37
ation score	12.8±2.6	14.3±2.9	0.8
edication score	17.8±2.9	17.9±2.8	0.94
gue scale	4.3±0.7	2.6±0.4	0.58



Th2 (CRTH2+CD25-CD4+ cells)



Der p 2- specific lgE/lgG4



randomized controlled study was undertaken in 40 ients with allergic rhinitis with/without asthma who receiving maintenance phase of HDM SCIT TR20200606002). HDM SLIT tablet was given ly for 12 weeks. Switching to SLIT was compared to tinuous SCIT. The principal outcome measure was ptom-medication score (SMS) and asthma control (ACT) score. Immunologic changes in fresh whole od to monitor T cell subsets, including regulatory T ls (Tregs), dysfunctional Tregs, and T helper 2 cells re investigated by the flow cytometry method and p2-specific IgE, Der p2-specific IgG4 and Der p2specific IgE/IgG4 were investigated by ELISA method at bacaling and 19 wasks after awitabing trantment

Results:

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■ SLIT Of 40 patients, 19 patients in the SLIT group and 20 patients in the control group achieved the study. There were no significant differences in SMS and ACT scores between the SLIT group and SCIT group during 12 weeks of treatment. Significantly reduced SMS after 8 weeks compared to baseline (17.6 ± 2.9) to 14 ± 2.4 , p = 0.028) was demonstrated in the patients with SLIT. T cell subsets' frequency, specific IgE, IgG4 and IgE/IgG4 ratio did not change significantly in both groups at the end of the study. No severe adverse drug reactions were reported.

Conclusion:

SCIT can switch to SLIT in the immunotherapy maintenance phase. SLIT was safe and efficacious by reducing the symptoms and medication consumption