



Dietary intake of vitamin A, lung function and incident asthma in childhood

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A higher intake of preformed vitamin A, but not pro-vitamin β -carotene, in mid-childhood was associated with higher subsequent lung function and lower risk of fixed airflow limitation and incident asthma <https://bit.ly/3d7PUca>

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Abstract

Background Longitudinal epidemiological data are scarce on the relationship between dietary intake of vitamin A and respiratory outcomes in childhood. We investigated whether a higher intake of preformed vitamin A or pro-vitamin β -carotene in mid-childhood is associated with higher lung function and with asthma risk in adolescence.

Methods In the Avon Longitudinal Study of Parents and Children, dietary intakes of preformed vitamin A and β -carotene equivalents were estimated by food frequency questionnaire at 7 years of age. Post-bronchodilator forced expiratory volume in 1 s (FEV₁), forced vital capacity (FVC) and forced expiratory flow at 25–75% of FVC (FEF_{25–75%}) were measured at 15.5 years and transformed to z-scores. Incident asthma was defined by new cases of doctor-diagnosed asthma at age 11 or 14 years.

Results In multivariable adjusted models, a higher intake of preformed vitamin A was associated with higher lung function and a lower risk of incident asthma: comparing top versus bottom quartiles of intake, regression coefficients for FEV₁ and FEF_{25–75%} were 0.21 (95% CI 0.05–0.38; $p_{\text{trend}}=0.008$) and 0.18 (95% CI 0.03–0.32; $p_{\text{trend}}=0.02$), respectively; odds ratios for FEV₁/FVC below the lower limit of normal and incident asthma were 0.49 (95% CI 0.27–0.90; $p_{\text{trend}}=0.04$) and 0.68 (95% CI 0.47–0.99; $p_{\text{trend}}=0.07$), respectively. In contrast, there was no evidence for association with β -carotene. We also found some evidence for modification of the associations between preformed vitamin A intake and lung function by *BCMO1*, *NCOR2* and *SCGB1A1* gene polymorphisms.

Conclusion A higher intake of preformed vitamin A, but not β -carotene, in mid-childhood is associated with higher subsequent lung function and lower risk of fixed airflow limitation and incident asthma.

