

Immunological and respiratory changes in coffee workers

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ABSTRACT Immunological status and respiratory function were studied in a group of 45 coffee workers. Skin tests with coffee allergens demonstrated the highest percentage of positive reactions to dust collected during emptying bags (40.0%), followed by dust of green (12%) and then roasted coffee (8.9%). Among 34 skin-tested control workers, 14.7% had positive skin reaction to dust collected during emptying bags, but none had positive skin reaction to green or roasted coffee. Serum levels of total IgE were increased in 24.4% of coffee workers and in 5.9% of control subjects. The prevalence of all chronic respiratory symptoms was significantly higher in coffee workers than in control subjects. Coffee workers with positive skin tests to coffee allergen had a significantly higher prevalence of chronic cough (63.6%) and chronic phlegm (72.7%) than those with negative skin tests (32.4% and 23.5% respectively). There was a significant mean decrease over the Monday work shift in the maximum expiratory flow rate at 50% of vital capacity (MEF₅₀: -7.9%) and at 25% vital capacity (MEF₂₅: -17.8%), suggesting an obstructive effect mostly in smaller airways. Coffee workers with positive skin tests to coffee allergens had larger acute reductions in flow rates than those with negative skin tests but the difference was not statistically significant.

In one of our previous epidemiological studies we have shown that exposure to green or roasted coffee is likely to contribute to the development of chronic respiratory symptoms and lung function changes in exposed workers.¹ Rhinitis, asthma, conjunctivitis, or dyspnoea in subjects exposed to coffee have been reported by several authors.²⁻⁵ However, there have been few studies dealing with the more detailed immunological changes related to coffee dust exposure.⁶⁻¹⁰

In the present epidemiological investigation we have studied the relationship between some immunological tests and lung function changes.

Methods

The study was performed in 45 non-smoking female workers employed in processing roasted or green coffee. Their mean age was 31 years (range: 20 to 56 yr) with a mean duration of employment in the coffee industry of seven years (range: one to 18 yr). In addition, a group of

45 non-smoking female control workers employed in the production of soft drinks was studied.

IMMUNOLOGICAL STUDIES

All coffee workers and 34 control workers were skin-tested with aqueous extracts of occupational and common allergens using the standard intradermal test. Occupational allergens were prepared from three different types of settled dust collected on operating machines in the workroom. These included roasted coffee, green coffee, and dust collected during emptying green coffee from bags. Intradermal skin tests with these allergens were performed using a dilution of 1:500 (0.02 ml of solution). Workers were also tested with skin-test material of house dust, bacteria, *Dermatophagoides pteronyssinus*, moulds, histamine base (0.1 mg/ml) and buffer as control solution. Bacterial antigen consisted of *H influenzae*, *Str pneumoniae*, *Str viridans*, *Str pyogenes*, *Neisseria*, and *Staph aureus* in a concentration of 60×10⁶ in 1 ml. Mould antigen was a mixture of *Alternaria*, *Penicillium*, *Mucor*, *Cladosporium*, *Aspergillus niger*, and *Aspergillus fumigatus* in 0.2% solution.

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The skin reactions were read after 20 minutes. An intradermal skin test was considered positive if the diameter of the observed wheal was larger than 9 mm (corrected for the control reaction).

Serum levels of total IgE antibody were measured in coffee workers and in controls by PRIST (Pharmacia Diagnostics AB Uppsala, Sweden), a direct radioimmunological sandwich technique based on paper discs as a solid phase.¹¹ In addition, in coffee workers concentrations of IgA, IgG, and IgM were determined by single radial diffusion on partigen plates.¹²

Levels of IgE below 125 IU/ml were considered normal. Values of 50–270 IU/ml for IgA, 80–220 IU/ml for IgG, and 60–250 IU/ml for IgM were taken as normal. Normal values of all immunoglobulins were determined according to the Behringwerke AG, Marburg-Lahn.

CHRONIC RESPIRATORY SYMPTOMS

Respiratory symptoms were recorded by using the British Medical Research Council questionnaire¹³ with additional questions on occupational asthma.¹⁴

Chronic cough/phlegm: cough and/or phlegm production on most days for at least three months per year.

Chronic bronchitis: cough and phlegm for a minimum of three months in the year and for not less than two successive years.

Dyspnoea grades: grade 3—shortness of breath when walking with other people at an ordinary pace on the level; grade 4—shortness of breath when walking at own pace on the level.

Occupational asthma: chest tightness, cough, wheezing, and shortness of breath during exposure to dust at work.

LUNG FUNCTION MEASUREMENT

The acute effect of exposure to coffee dust on ventilatory capacity was studied by recording the maximum expiratory flow-volume (MEFV) curves on Monday before and after work shift. The MEFV curves were recorded on a portable flow-volume spirometer¹⁵ and flow rates at 50% and at 25% of the control vital capacity (MEF₅₀ and MEF₂₅) were read from these curves. The mean of the two highest values on successful MEFV curves was taken as the result of the test.

STATISTICAL ANALYSIS

The results of ventilatory function measurements were analysed by using the *t* test for difference of paired (acute effects) and unpaired (chronic effects) variables. The chi-square test was used for

testing differences in the prevalence of respiratory symptoms, and $p < 0.05$ was considered significant.

Results

IMMUNOLOGICAL STUDIES

Skin reactions to different allergens in coffee workers are presented in the figure. A positive skin reaction to the allergen of roasted coffee was found in 8.9%, to that of green coffee in 12%, and to the allergen prepared from the dust emitted during emptying bags in 40% of the workers tested. Out of four workers with asthma two had positive skin reaction to coffee allergen. Among control workers, 14.7% had positive skin reaction to the allergen prepared from dust collected during emptying bags, but none to green or roasted coffee allergens.

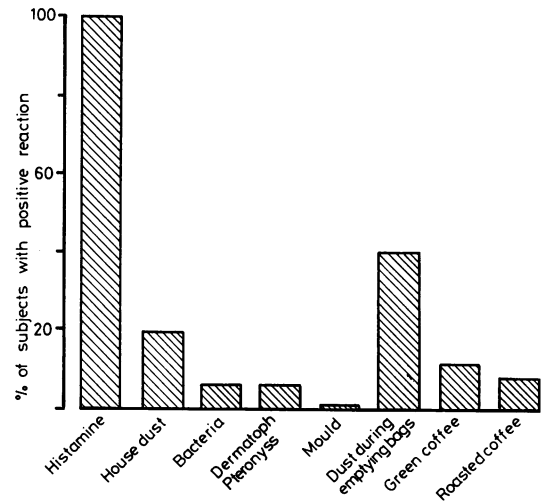


Figure Skin test reactions to occupational allergens (green coffee, roasted coffee, and dust collected during emptying bags) and to different common allergens in coffee workers.

IgE above normal level was found in 24.4% of coffee workers, and 5.9% of control subjects. While IgM in 18.2% of coffee workers was increased, its values in control subjects were normal. Out of 11 coffee workers with increased IgE level, seven (64%) complained of rhinitis and conjunctivitis and also had positive skin reactions to at least one of the coffee allergens. Among four coffee workers with asthma two had increased IgE values (670 and 160 IU/ml). Among those with increased IgM level one worker had a positive skin reaction to coffee allergens. In all those

examined, IgA and IgG serum values were found within normal range.

CHRONIC RESPIRATORY SYMPTOMS

The prevalence of all chronic respiratory symptoms was significantly higher in coffee workers than in control subjects (table 1). A separate analysis of data obtained in workers with positive and negative skin tests respectively, indicated that the former had higher prevalence of respiratory symptoms, the difference being significant for chronic cough ($p < 0.05$) and chronic phlegm ($p < 0.01$). Among four subjects with asthma symptoms, two had a positive skin reaction to coffee allergen. Their durations of employment in the coffee industry were one and 17 years respectively. Thirty-eight per cent of coffee workers reported symptoms of rhinitis or conjunctivitis.

VENTILATORY CAPACITY

Table 2 shows significant acute reductions of ventilatory capacity in coffee workers over the work shift ($p < 0.01$). Acute reductions in MEF_{25} were greater (17.8%) than in MEF_{50} (7.9%).

A comparison of pre-shift values in exposed

and matched control workers (table 2) demonstrated significantly lower MEF_{50} and MEF_{25} in coffee workers than in controls ($p < 0.01$).

Workers with positive skin tests to coffee allergens (table 3) had larger acute reductions in MEF_{50} (13.3%) and in MEF_{25} (21.9%) than those with negative skin tests (MEF_{50} : 6.5%; MEF_{25} : 16.2%), but the difference between these reductions was not statistically significant ($p > 0.05$). Two workers with asthma and with positive skin tests to coffee allergen had demonstrated large acute reductions in MEF_{50} : 31% and 20% respectively, and in MEF_{25} : 18% and 33% respectively.

Pre-shift values of flow rates were found to be significantly lower in all groups of coffee workers than in the corresponding controls, except for MEF_{50} in those with negative skin tests.

Analysis of the acute reductions of ventilatory capacity separately in workers with normal IgE values (MEF_{50} : -7.7%; MEF_{25} : -19.1%) and in those with increased IgE values (MEF_{50} : -9.8%; MEF_{25} : -14.6%) did not reveal significant differences in acute reductions ($p > 0.05$) between these two groups. Workers with increased IgE had sig-

Table 1 Prevalence of chronic respiratory symptoms in coffee workers

Group	Mean age (yr)	Mean exposure (yr)	Chronic cough (%)	Chronic phlegm (%)	Chronic bronchitis (%)	Asthma (%)	Dyspnoea grade 3 or 4 (%)
Coffee workers (n=45)	31	7	40.0	35.6	24.4	8.9	40.0
			< 0.01	< 0.01	< 0.01	< 0.05	< 0.01
Control workers (n=45)	31		6.7	4.4	4.4	0	0
Workers with positive skin tests (n=11)	29	8	63.6	72.7	45.5	18.2	45.5
			< 0.05	< 0.01	NS	NS	NS
Workers with negative skin tests (n=34)	33	7	32.4	23.5	17.6	5.9	38.2

Table 2 Ventilatory capacity in coffee workers and controls

Group	Mean age (yr)	Mean height (cm)	Mean exposure (yr)	MEF_{50}					MEF_{25}				
				Before shift l/s	After shift l/s	Difference l/s	%	p	Before shift l/s	After shift l/s	Difference l/s	%	p
Coffee workers (n=45)	31	163	7	4.80 ±1.29	4.42 ±1.38	-0.38	-7.9	<0.01	2.25 ±0.92	1.85 0.94	-0.40	-17.8	<0.01
Control workers (n=45)	31	163		5.18* ±0.70					2.85* ±0.51				

*Difference between exposed and control workers statistically significant ($p < 0.01$).

Table 3 Ventilatory capacity in workers with positive and negative skin tests to coffee allergens

Group	MEF ₅₀					MEF ₂₅				
	Before shift l/s	After shift l/s	Difference			Before shift l/s	After shift l/s	Difference		
			l/s	%	p			l/s	%	p
Positive skin tests (n=11)	4.35 ±0.80 5.04* ±0.68	3.77 ±0.86	-0.58	13.3	<0.01	1.87 ±0.54 2.90* ±0.34	1.46 ±0.45	-0.41	21.9	<0.01
Negative skin tests (n=34)	4.95 ±1.39 5.20* ±0.79	4.63 ±1.45	-0.32	6.5	<0.01	2.34 ±0.99 2.81* ±0.54	1.96 ±1.03	-0.38	16.2	<0.01

*Control workers.

Difference between exposed and control workers statistically significant except for MEF₅₀ in workers with negative skin tests.

nificantly lower pre-shift values of both MEF₅₀ ($p < 0.05$) and MEF₂₅ ($p < 0.01$) than controls. Workers with normal IgE values had only MEF₂₅ significantly lower than controls ($p < 0.01$).

Discussion

A significantly higher prevalence of all respiratory symptoms was found in coffee workers than in the corresponding control workers. Our results indicate a significantly higher prevalence of chronic phlegm in workers with positive than in those with negative skin tests to coffee allergens, suggesting a higher sensitivity in the former group.

The results of our previous study in two groups of female coffee workers¹ have demonstrated that the mean acute reductions in FVC and FEV₁ were smaller (1.3%–2.8%) than in MEF₅₀ and MEF₂₅ (4.0%–18.5%). This suggests that the inhalation of dust in coffee processing caused significant bronchoconstriction with more pronounced effects in smaller airways. The acute reductions were larger in workers with positive than in those with negative skin tests. The mean pre-shift value of MEF₂₅ in workers with positive skin tests was 64% of the corresponding value in the controls, while in those with negative skin tests it was 83%. There was no significant correlation between increased IgE serum level and acute ventilatory capacity reductions.

Recently Karr *et al*¹⁰ demonstrated positive skin reactivity to green coffee bean and serum IgE antibodies specific to green coffee in six coffee workers with occupational allergic disease (asthma, rhinitis, conjunctivitis, and urticaria). Total serum IgE levels were within normal values in these workers. In our present study, out of 11 workers with increased total serum IgE antibody, eight complained of symptoms such as cough, con-

junctivitis, rhinitis, and headache shortly after exposure to coffee dust.

Layton *et al*⁷ suggested that primary allergy to green coffee does not involve chlorogenic acid and that coffee allergy is an example of atopic hypersensitivity to proteins. Karr *et al*¹⁰ demonstrated that chlorogenic acid produced no RAST inhibition for green coffee bean.

Our previous data¹ as well as the present results indicate that exposure to dust in coffee processing might cause the development of chronic respiratory symptoms and changes in lung function particularly in subjects with positive skin tests to coffee allergens.

The increasing frequency of positive skin reactions to allergens prepared from roasted coffee, green coffee, and the dust released during the emptying of bags, respectively, as well as the stronger bronchoconstricting potency of green than roasted coffee found in our previous epidemiological study led us to the hypothesis that (1) some of the chemical or microbial foreign components present in contents of bags of green coffee may be biologically active, (2) that the potency of these components may be decreased by heat during roasting, and (3) the potency of biologically active components of green coffee may decrease on heating. Some of our preliminary provocation tests seem to have confirmed all three assumptions but the results are not conclusive. The investigation is being continued into the comparative immunological activity heat treated and untreated dust components.

References

- 1 Žuškin E, Valić F, Skurić Z. Respiratory function in coffee workers. *Br J Ind Med* 1979; **36**:117–22.
- 2 Bruun E. Allergy to coffee. An occupational disease. *Acta Allergol* 1957; **11**:150–4.

- 3 Kaye M, Freedman SO. Allergy to raw coffee: an occupational disease. *Can Med Assoc J* 1961; **84**:469-71.
- 4 Turula M, Aho J, Taipale S, Förström L. Raw coffee allergy among coffee roastery workers. *Proceedings of the XV International Congress of Occupational Health*. Vienna: Egerman, 1966: 845-8.
- 5 Somazzi S, Wüthrich B. Asthme professionnel à la poussière de café vert. *Med Hyg* 1975; **33**:677-83.
- 6 Pepys J, Longbottom JL, Jenkins PA. Vegetable dust pneumoconioses. Immunological responses to vegetable dusts and their flora. *Am Rev Respir Dis* 1964; **89**:842-58.
- 7 Layton LL, Panzani R, Greene FC, Corse JW. Atopic hypersensitivity to a protein of the green coffee bean and absence of allergic reactions to chlorogenic acid, low-molecular-weight components of green coffee, or to roasted coffee. *Int Arch Allergy* 1965; **28**:116-27.
- 8 Layton LL, Greene FC, Panzani R. Allergy to green coffee. *J Allergy* 1965; **36**:84-91.
- 9 Van Toorn DW. Coffee worker's lung. A new example of extrinsic allergic alveolitis. *Thorax* 1970; **25**:399-405.
- 10 Karr RM, Lehrer SB, Butcher BT, Salvaggio JE. Coffee workers' asthma: a clinical appraisal using the radioallergosorbent test. *J Allergy Clin Immunol* 1978; **62**:143-8.
- 11 Wide L, Porath J. Radioimmunoassay of proteins with the use of Sephadexcoupled antibodies. *Biochim Biophys Acta* 1966; **130**:257-60.
- 12 Mancini G, Carbonara AO, Heremans JF. Immunochemical quantitation of antigens by single radial immunodiffusion. *Immunochemistry* 1965; **2**:235-54.
- 13 Medical Research Council Committee on the Aetiology of Chronic Bronchitis. Standardised questionnaire on respiratory symptoms. *Br Med J* 1960; **2**:1665.
- 14 Murphy RLH. Industrial diseases with asthma. In: Weiss EB, Segal MS (eds). *Bronchial asthma: mechanisms and therapeutics*. Little Brown, Boston, Mass: Little Brown, 1976:517-36.
- 15 Peters JM, Mead J, Van Ganse WF. A simple flow-volume device for measuring ventilatory function in the field. *Am Rev Respir Dis* 1969; **99**:617-22.