Effect of cigarette smoking on the specific antibody response in pigeon fanciers

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ABSTRACT Titres of circulating IgG antibodies to pigeon gammaglobulin and end expired carbon monoxide concentrations were measured in 86 pigeon fanciers attending the “Show of the Year.” Antibody levels were significantly higher in non-smokers and in those with end expired carbon monoxide concentrations below 10 parts per million.

Cigarette smoking affects various components of the immune response.1 Morgan et al noted the suppressive effect of cigarette smoking in patients susceptible to extrinsic allergic alveolitis2 and we have reported similar observations, finding circulating IgG specific for pigeon antigen in 55% of non-smoking and 4% of smoking pigeon fanciers.3 We are unaware of any previous studies in which an objective indicator of cigarette smoking has been used to investigate subjects susceptible to the effects of antigen inhalation. Cigarette consumption can be assessed by the measurement of end expired carbon monoxide by means of a portable carbon monoxide meter.4

Methods

During a morning at the “Show of the Year” in Blackpool in January 1987 we assessed 86 volunteer pigeon fanciers. A questionnaire (administered by doctors) concerning antigen exposure and cigarette smoking history was completed and blood obtained for measurement of circulating specific IgG by enzyme linked immunosorbent assay (ELISA), with pigeon gammaglobulin as antigen. End expired carbon monoxide concentrations were determined with a portable hand held meter (ECO-Check, PK Morgan Ltd, England). The subject held his or her breath for 20 seconds and exhaled through a one way valve into the meter, which incorporated an alcohol filter. No information about the purpose of the meter was given before a breath sample was obtained. The carbon monoxide concentration in parts per million (ppm) was displayed digitally on the side of the meter, being freely visible to the subject. For the purpose of this study an index of pigeon antigen exposure was made by combining the fanciers’ own estimates of the numbers of birds kept and the hours spent in the pigeon loft in one week (pigeon exposure index = pigeon number × weekly hours × 100%). Kruskal-Wallis one way analysis of variance was used to assess the significance of the findings.

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Results

The results are shown in figures 1 and 2. Of the 86 subjects studied, 67 were non-smokers and half of these (33) had IgG antibody concentrations of over 8 μg/ml, a level associated with the

![Graph showing specific anti-pigeon gammaglobulin (IgG) antibody and end expired carbon monoxide for smokers and non-smokers (according to smoking history given).]

Fig 1
Effect of cigarette smoking on the specific antibody response in pigeon fanciers

Specific IgG antibody (μg/ml)

Fig 2 Specific anti-pigeon gammaglobulin (IgG) antibody and pigeon exposure index for those with end expired carbon monoxide concentrations of under 10 ppm (●) and 10 ppm or more (△).

The subjects studied were volunteers and they may therefore have been more likely to present if symptoms were present. The finding of circulating antibody in half of the subjects is not unexpected but is likely to reflect the findings in a symptomatic population rather than the prevalence of circulating antibody in the population who attended the pigeon meeting. Previous work suggests that about one third of subjects with higher circulating antibody levels would have symptoms of pigeon breeder’s disease.\(^6\) This study was intended to evaluate a marker of cigarette smoking that might be related to the antibody response, so we have not included an assessment of symptoms.

An estimate of antigen exposure was used to confirm that antigen exposure in non-smokers and smokers was similar. An unexpected finding was the lack of relationship between the antibody response and the estimated degree of antigen exposure. The relationship between antigen exposure and the antibody response is complex. Some workers have reported
that precipitins occur more commonly as the estimated exposure to pigeons increases,1 though antibody levels are often highest in subjects who keep small numbers of pigeons,5 possibly as a result of attempted self regulation in those who are or who have been suffering from symptoms. Pigeon fanciers who develop fulminant disease are more often those who have kept pigeons for less than 10 years.10 Pigeon breeder's disease is rarely encountered in those with more prolonged exposure, which suggests an element of higher innate susceptibility in those most severely affected. There may be other factors that we have not considered in our calculation of exposure. We are unaware of work directly comparing measured antigen exposure and antibody response.

Discrepancies between previous studies and ours are probably explained by the limitations of using this estimated value and a single serum sample to represent a more dynamic reaction within the lung. The impaired antibody response in smoking pigeon fanciers seems not to be due to their having different antigen exposure from non-smokers and may be more dependent on the quantity of exposure to cigarette smoke. An end expired carbon monoxide level of 10 ppm or more was rarely associated with a measurable antibody level in this study. The clinical implication for a regular smoker with undiagnosed lung disease is that, while the long term effects of persistent antigen exposure in cigarette smokers remain unknown, the absence of circulating antibody should not imply absence of exposure to antigen.

We suggest that future studies of the effects of cigarette smoking on the immune response should include measurements of a marker of smoke exposure in order to determine whether the effects are dose related.

References