**Occupational Burden of Idiopathic Pulmonary Fibrosis (600 word edition)**

Idiopathic pulmonary fibrosis (IPF) is a diagnosis of exclusion. It is made in the presence of a usual interstitial pneumonia (UIP) pattern on high resolution CT scan or biopsy. The diagnosis requires that known causes of interstitial lung disease (such as drug toxicity, connective tissue disease, domestic, and occupational or environmental exposures) be excluded.[1]

There are many review articles of the epidemiology of interstitial lung disease that do not necessarily focus on IPF and only briefly mention occupational factors. We selected review articles that dealt specifically with occupational factors in IPF and cite the case-control studies and case-reports identified.

Turner-Warwick (1998) discusses potential difficulties in establishing attribution and causality in IPF. She observes that there is variation in clinical practice with respect to the standard applied to exclude IPF; some clinicians exclude IPF when exposure to a potential cause is identified, others only when there is clear exposure to an established cause. She explains that diagnosis based on radiologic and clinical findings, and not on lung biopsy or bronchioalveolar lavage, may result in initiating agents for disease being overlooked. Further, that exposures such as asbestos, silica, coal,

graphite, hard metal, and avian proteins, may result in disease that can not be differentiated from IPF.[2]

Reviewing the epidemiology of IPF and case-control studies to date Hubbard (2001) describes the association of IPF with occupational exposures to metal and wood and estimates that 10% of IPF cases may be due to occupational metal exposure and 5% of cases to wood.[3]

Taskar and Coultas (2005) reviews and performs a meta-analysis of six case-control studies investigating environmental and occupational exposures in IPF. They report population attributable risk percentages for agriculture and farming (20.8%), livestock (4.1%), wood dust (5%), metal dust (3.4%), stone/sand/silica (3.5%), and smoking (49.1%).[4]

Gulati and Redlich’s (2015) review of exposures causing usual interstitial pneumonia highlights that asbestosis may appear indistinguishable from IPF and summarises previous case-control studies but did not pool studies to perform a meta-analysis.[5]

We found (as of May 2017) 14 case-control studies looking at occupational exposures in IPF (table 1); the most recent review article covers only eight of them. Associations with metal, wood, silica, and agricultural dust are frequently reported. [6-19]

Two investigators independently reviewed and abstracted data for five exposure categories

common to the identified case-control studies: “vapors, gases, dusts, and/or fumes (VGDF)”, “metal dust”, “wood dust”, “silica dust”, and “agricultural dust”. We calculated PAF as follows: PAF=pc(OR – 1)/OR, where pc is the proportion of cases exposed and OR is the odds ratio.

We calculated pooled OR and pooled PAF for occupational exposures using fixed effects models and random effects models in Stata. When the results of the models differed substantively, we used the results of the fixed effects model, which were more conservative. The pooled PAF relied on the ratio of attributable cases to all cases underlying each risk estimate.

In all, 40 risk estimates from 12 publications (1326 IPF cases in total) were used. Each exposure category was assessed with 5-11 risk estimates. Pooled ORs were significantly elevated for each category; the pooled PAF estimates by category ranged from 3-23% (Table 2).

The observed excess risk could represent disease misclassification of pneumoconiosis or hypersensitivity pneumonitis, but this is unlikely to fully explain the observed effects. Our analysis supports an etiologic role for occupational exposures in IPF, potentially explaining up to 14% of the burden of disease and highlighting a role for workplace exposure reduction in disease prevention.

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