



## Summary of Risk Assessments of MMVFs

The magnitude of any potential carcinogenic risk from occupational exposure to man-made vitreous fibers (MMVFs) has been assessed several times. At present, no disease in humans has resulted from exposure to vitreous fibers. In the absence of positive epidemiological data, the long-term bioassay remains the only method for the *a priori* evaluation of any carcinogenic risks that may be posed by putative carcinogens. For this reason the assessments reported below were based entirely on the results of long-term bioassays.

### Refractory Ceramic Fibers (RCF)

The primary sources of data for a number of risk assessments of RCF were two long-term oncogenicity inhalation studies conducted on rats at the RCC laboratory.

Extrapolating intake from rat to human in terms of fibers per day per kilogram of body weight, Fayerweather et al. (1997) used the linearized multistage model to estimate human risk. The worker activity patterns consisted of an assumed exposure of 4 hours/day, 5 days/week, 50 weeks/year, and 40 years of a 70-year lifespan. Although the authors did not directly compute the excess lifetime risk of developing lung tumors at an exposure of 1 f/cc, it can be inferred from the methods described in their paper to be  $3.8 \times 10^{-5}$  (maximum likelihood estimate).

Moolgavkar et al. (1999) assessed the occupational risk to RCF within the framework of a biologically based model for carcinogenesis, the two stage clonal expansion model (the MVK model), which allows for the explicit incorporation of the concepts of cancer initiation and promotion in the analyses. The model considered the temporal profile of fiber lung burden in the experimental animals and accounted for fiber deposition and clearance in both the human and rat lung. By basing the risk estimates directly on lung fiber burden rather than aerosol exposure, the authors were able to avoid many of the problems associated with interspecies extrapolations. The analysis resulted in a best estimate (maximum likelihood estimate) of the excess probability of lung cancer at age 70 for 30 years of occupational exposure, 8 hours/day starting at age 20, to 1 fiber/cm<sup>3</sup> of  $3.7 \times 10^{-5}$  for a non-smoking population. If the occupational population were assumed to have the smoking habits of a typical workforce, the best estimate of excess risk was computed to be  $1.5 \times 10^{-5}$ .



### **Further extensions of the risk assessment model**

Fiber chemistry is an important determinant of breakage and solubility in tissues, and hence of clearance. Tissue burden and, therefore, fiber carcinogenicity are clearly affected by the composition of the fiber. Moolgavkar and coworkers (2000, 2001) extended the methods used in their assessment of RCF to investigate whether chemical composition of fibers has a role beyond determining biopersistence. Using available data from a number of long-term oncogenicity inhalation experiments, they showed that the results were consistent with the hypothesis that the oncogenic potential of long man-made vitreous fibers is determined mainly by their biopersistence. In other words, the data analyzed were shown to be consistent with the view that “a fiber is a fiber.” The carcinogenic potential is determined by the lung burden of fibers which, in turn, is determined by biopersistence. That is, the authors showed that fiber chemistry influences fiber carcinogenesis primarily through its role in determining biopersistence. A direct mechanistic role, if any, of chemistry in fiber carcinogenesis is of secondary importance. These conclusions allowed the authors to estimate a common potency factor describing the oncogenic potential for all MMVFs.

Turim and Brown (2003) extended these results and considered various means of extrapolating human equivalency concentrations from animal test results. They showed that of all the models considered, including benchmark dose and other statistical models, the weight of evidence argues in favor of the MVK two-stage clonal expansion model for the following reasons:

- The MVK model explicitly takes into account the temporal distribution of the pattern of lung burden. Other models consider only the steady-state level of fibers in the lung.
- It is the only model that explicitly incorporates time-dependent doses.
- By simulating the initiation and promotion activities that are known to underlie cancer induction the model rests on a biologically significant and generally accepted theory of carcinogenesis. Other models rely on purely statistical techniques.
- The MVK model provides a better fit of the observed laboratory data than the other models, taking into account the number of parameters that are used in the model.
- The model is consistent with the results of a number of experiments conducted with synthetic vitreous fibers and was able to detect the effect of overload at high exposure concentrations.
- It is the only model that can be subjected to an external validity check because the parameters estimated in the model must be biologically plausible.



---

1025 Connecticut Avenue, N.W. Suite 1000 c/o Verdant Law Firm Washington, DC 20036 tel 240.343.1738 [www.htiwcoalition.org](http://www.htiwcoalition.org)

## **Risk from exposure to RCF**

Turim and Brown (2003) summarized the results of the previous investigations in estimating cancer risk. They found that the 95% upper bound risk of excess lifetime lung cancer risk to a non-smoking workforce is:

- $3 \times 10^{-5}$  for an exposure of 1 f/cc
- $1.5 \times 10^{-5}$  for an exposure of 0.5 f/cc
- $0.3 \times 10^{-5}$  for an exposure of 0.1 f/cc

For an occupational workforce with typical smoking habits, the corresponding 95% upper bound excess lifetime risks are approximately three times higher.

## **References**

Fayerweather WE, Bender JR, Hadley JG, and Eastes W. (1997). Quantitative risk assessment for a glass fiber. *Regulatory Toxicology and Pharmacology*. 25, 103-120.

Moolgavkar SH, Luebeck EG, Turim J, and Hanna L. (1999). Quantitative assessment of the risk of lung cancer associated with occupational exposure to refractory ceramic fibers. *Risk Analysis*. 4, 138-146.

Moolgavkar SH, Luebeck EG, Turim J, and Brown RC. (2000). Lung cancer risk associated with exposure to man-made fibers. *Drug and Chemical Toxicology*. 23, 223-242.

Moolgavkar SH, Turim J, Brown RC, and Leubeck EG. (2001). Long man-made fibers and lung cancer risk. *Regulatory Toxicology and Pharmacology*. 33, 138-146.

Moolgavkar SH, Brown RC, and Turim J. (2001b). Biopersistence, fiber length, and cancer risk for inhaled particles. *Inhalation Toxicology*. 13, 755-772.

Turim J and Brown RC (2003). A dose-response model for refractory ceramic fibers. *Inhalation Toxicology*, 15, 1103-1118.