

HYPERURICEMIA RISK IN SUBJECTS WITH MODERATE TO HIGH LEVELS OF PCDD/FS EXPOSURE

Jung-Wei Chang, *Department of Environmental and Occupational Health, College of Medicine, National Cheng Kung University, Tainan, Taiwan*

Hong-Yih Ou, *Division of Endocrinology and Metabolism, Department of Internal Medicine, National Cheng Kung University Hospital, Tainan, Taiwan*

Po-Chi Liao, *Department of Environmental and Occupational Health, College of Medicine, National Cheng Kung University, Tainan, Taiwan*

Ching-Chang Lee, *Department of Environmental and Occupational Health, College of Medicine, National Cheng Kung University, Tainan, Taiwan*

Background and Aim: Hyperuricemia, the predisposing condition for gout, has been associated with hypertriglyceridemia and diabetes mellitus (Berkowitz 1966), and also be a risk factor for the development of coronary artery disease (Abbott et al. 1988). PCDD/Fs have been found to cause renal toxicity in experimental animals and elevate uric acid in epidemiology studies. The potential role of PCDD/Fs exposure as a preventable risk factor has raised attention of public health, especially since it's ubiquitous worldwide. We aimed to investigate and clarify the effect of moderate-to-high exposure to PCDD/Fs on hyperuricemia risk.

Methods: This cross-sectional study recruited 1657 healthy participants near a deserted pentachlorophenol factory (Chang et al. 2010). Renal function related factors were measured to examine associations with serum PCDD/Fs. We also investigated associations between serum PCDD/Fs and hyperuricemia risk. We compared associations of serum PCDD/F levels and estimated glomerular filtration rate (eGFR, renal function index) in both gender under and above 50-years-old.

Results: The eGFR reduces gradually with serum PCDD/F levels in all groups (men < and \geq 50 50-years-old, women < and \geq 50-years-old), especially for a sudden falling observed in men over than 50 years old. Serum PCDD/Fs were significantly increased with uric acid in male participants (Men: $\beta = 0.230$, $p = 0.008$; Women: $\beta = 0.035$, $p = 0.753$; All: $\beta = 0.148$, $p = 0.037$). In addition, we further found that male participants with higher serum PCDD/F levels had a higher hyperuricemia risk than did the reference group (serum PCDD/F levels <11.4 pg WHO₉₈-TEQ_{DF}/g lipid) after adjusting for confounding factors (25 th to <50 th percentile, adjusted Odds Ratio (AOR) = 1.87 [95% CI = 1.13-3.10]; 50th to <75 th percentile, AOR = 2.44 [1.46-4.09]; \geq 75 th percentile, AOR = 3.29 [1.89-5.74]).

Conclusions: We hypothesize that accumulated PCDD/Fs may reduce urate excretion and may heighten the risk of hyperuricemia in the general population.

References:

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