

Pulmonary Epithelial Integrity in Children -Relationship to Ambient Ozone Exposure and Swimming Pool Attendance

Birgitta Json Lagerkvist, Alfred Bernard, Anders Blomberg, Erik Bergstrom, Bertil Forsberg, Karin Holmstrom, Kjell Karp, Nils-Goran Lundstrom, Bo Segerstedt, Mona Svensson, and Gunnar Nordberg doi:10.1289/ehp.7027 (available at http://dx.doi.org/) Online 13 September 2004



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Pulmonary Epithelial Integrity in Children -

Relationship to Ambient Ozone Exposure and Swimming Pool Attendance.

Birgitta Json Lagerkvist¹, Alfred Bernard⁵, Anders Blomberg², Erik Bergstrom³, Bertil Forsberg¹, Karin Holmstrom⁴, Kjell Karp⁴, Nils-Goran Lundstrom¹, Bo Segerstedt¹, Mona Svensson¹, and Gunnar Nordberg¹.

¹Environmental & Occupational Med., ²Respiratory Med. & Allergy, Dep. of Public Health and Clinical Medicine, ³Paediatrics, Dep. of Clinical Sciences, and ⁴Clinical Physiology, Dep. of Surgical and Peri-operative sciences, Umea University Umea, Sweden. ⁵Unit of Ind. Toxicology, Catholic Univ. of Louvain, Brussels, Belgium.

The study was performed at Environmental Medicine, Department of Public Health and Clinical Medicine, Umea University, Umea, Sweden.

Address correspondence to B Json Lagerkvist, Environmental Medicine, Department of Public Health and Clinical Medicine, Umea University, S-901 87 Umea, Sweden. Telephone +46 90-7851343. Fax: +46 90-779630. E-mail: Birgitta.Lagerkvist@envmed.umu.se **Key words:** Clara cell protein-CC16, airway irritants, ozone, nitrogen trichloride, swimming pool, children

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Abbreviations used:

BMI, Body mass index, kg/m²

EPA, Environmental Protection Agency

FORMAS, Forskningsradet for Miljo, Areella Naringar och Samhalle

O₃, ozone

O₂, oxygen

NO₂, nitrogen dioxide

NCl₃, nitrogen trichloride, trichloramine or chlorine azide

FVC, forced vital capacity, litre

FEV₁, forced expiratory volume in one second, litre/s

FEV₁% predicted, percent of the predicted normal value of FEV₁ calculated from age, sex,

height, and weight of the subject

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Abstract

Airway irritants such as ozone (O₃) are known to impair lung function and induce airway inflammation. Clara cell protein (CC16) is a small anti-inflammatory protein secreted by the non-ciliated bronchiolar Clara cells. CC16 in serum has been proposed as a non-invasive and sensitive marker of lung epithelial injury. In this study, we used lung function and serum CC16 concentration to examine the pulmonary responses to ambient ozone exposure and swimming pool attendance. The measurements were made on 57 children 10-11 years old before and after outdoor exercise for two hours. Individual ozone exposure was estimated as the total exposure dose between 7 AM until the second blood sample was obtained, (mean O_3 concentration/m³ x hours). The maximal one-hour value was 118 μ g/m³ (59 ppb), and the individual exposure dose ranged between 352-914 μ g/m³h. These ozone levels did not cause any significant changes in mean serum CC16 concentrations before or after outdoor exercise. Nor was any decrease in lung function detected. However, children who regularly visited chlorinated indoor swimming pools had significantly lower CC16 levels in serum than non-swimming children both before and after exercise, 5.7±2.4 and $5.3\pm1.7 \mu g/l$ versus 8.2 ± 2.8 and $8.0\pm2.6 \mu g/l$, p<0.002. These results indicate that repeated exposure to chlorination by-products in the air of indoor swimming pools have adverse effects on the Clara cell function in children. A possible relation between such damage to Clara cells and pulmonary morbidity (e.g. asthma) should be further investigated.