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Determination of Total Mercury Concentration in Wastewater by Continuous Microflow Analysis with Cold Vapor Atomic Absorption Spectrometry.

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Sub-parts-per-billion levels of mercury compounds in wastewater from the university laboratories and from a factory treating city refuse were determined using "continuous microflow analysis" with a cold vapor atomic absorption technique. The average total mercury content during 1987 for the wastewater of the inlet to the university pond was ca. 0.25 μ g l⁻¹, while that in the outlet was ca. 0.24 μ l l⁻¹. The total mercury concentrations in inlets of the pond depended on the activity of the laboratories.

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High-perforance Liquid Chromatographic Determination of Malonaldehyde using p-Nitrophenylhydrazine as a derivatizing reagent.

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A high-performance liquid chromatographic procedure for the specific determination of free malonaldehyde (MA) is described. MA solution was mixed with 1 M acetate buffer (pH 3.76) and was reacted with p-nitrophenylhydrazine (NPH) hydrochloride solution in ethanol. An aliquot of the reaction mixture was injected into a C_{18} -5 column with a mobile phase consisting of acetonitrile—isopropanol—0.01 M sodium dihydrogenphosphate (30:10:60, v/v/v) and detection at 315 nm. MA reacted readily with NPH in a weakly acidic medium at room temperature, giving 1-(p-nitrophenyl) pyrazole. The method is specific for free MA. The response was linear in the range 36—720 ng/ml of MA and the detection limit was 6ng/ml with a 20 μ l injection.

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Behaviours of Siloxane Polymers Containing Phenyl or Silarylene as Stationary Phases for High-temperature Gas Chromatography.

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Several kinds of mechanisms have been proposed for the thermal degradation of methylphenylsil-oxane and dimethylsiloxane—silarylene copolymer in an inert atmosphere. The termal stability of these stationary phases at temperatures as high as 420°C was examined using a deactivated metal capillary column, which is much more thermally stable than the fused-silica capillary columns. It was found that these phases degraded at 380—400°C by eliminating arylene-containing moieties, which led to cross-linking, and the peak shapes therefore deteriorated. With a column treated at 420°C, elution of solutes was considerably delayed and the elution order became unusual.