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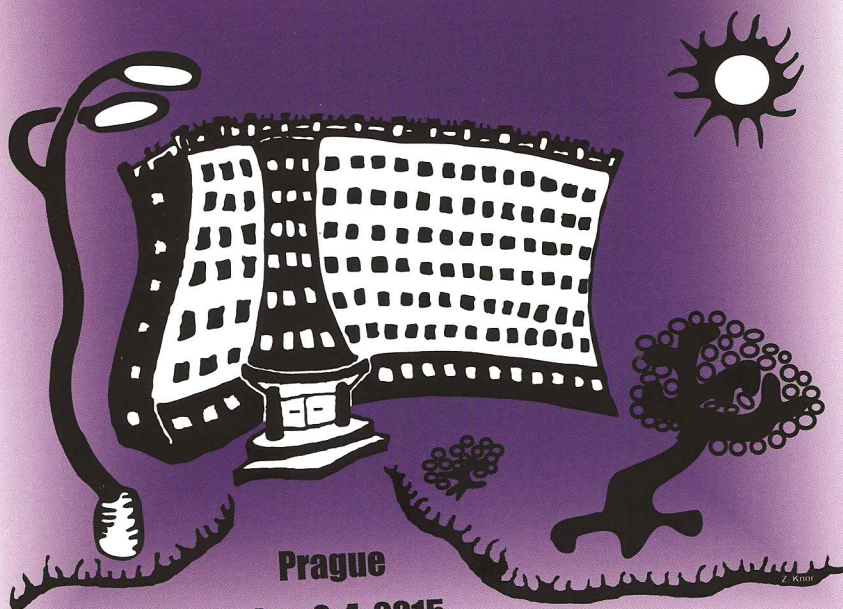
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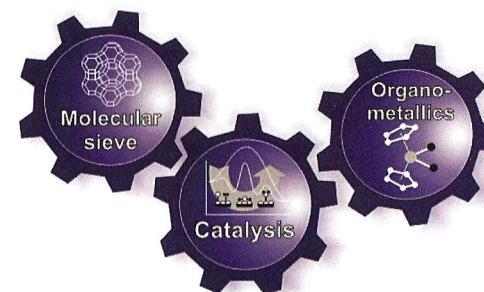
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SYNTHESIS OF Pt/C FUEL CELL ELECTROCATALYSTS: RESIDUAL CONTENT OF CHLORINE AND ACTIVITY IN OXYGEN REDUCTION

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Hydrogen and fuel cells represent a possible solution of modern transportation. Carbon-supported Pt is conventional catalyst in proton-exchange membrane fuel cells (PEMFC). Deposition of Pt over the support is usually based on i) impregnation of the support with a Pt containing solution followed by reduction¹ or on ii) precipitation of Pt out of the support surface and then mixing it with the support. Aqueous solution of H₂PtCl₆ is mostly used as Pt source. However, H₂PtCl₆ is expected to leave a certain amount of residual Cl species on the catalyst surface.² This residual chlorine is believed to decrease the catalytic activity in PEMFC², since Cl⁻ is well-known poison to Pt catalysts and it increases the propensity of PEMFC to corrosion. The aim of this work was to compare Pt/C prepared by impregnation of C followed by reduction and dechlorination (by NaOH(aq)³ or NH₃(g)) with Pt/C prepared by precipitation of Pt (by microwave-assisted polyol method) followed by washing and controlled mixing with C, in terms of residual Cl content, electrochemically active surface area (ESA) and activity towards the oxygen reduction reaction (ORR). It was found that the medium-surface-area C, selected in our previous work³ for the present experiments, resulted in catalysts with improved durability, but somewhat lower surface area and Pt dispersion (for the impregnated catalysts), in comparison to conventional reference catalysts based on high-surface-area C. The microwave-assisted polyol method featured smaller Pt nanoparticles, lower content of residual Cl and higher activities than those prepared by impregnation. On the other hand, the impregnated catalysts showed better durability of the Pt particles. Mass-specific ORR activities of the prepared Pt/Cs were of a similar size as that of the reference catalyst. The novel Pt/C catalysts are thus suitable for high-performance, durable membrane-electrode assemblies (MEAs) for PEMFCs.

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Hydroisomerisation of
industry which improves
freezing point. ^{1,2} Se-
tunstedt zirconia on
Catalysts "A" Pt(111),
ZrO₂ and "C" Pt(111)/
characterized by UV-
analyzed using GC/MS
only C1-C5 products
of C1-C5 gaseous pro-
catalysts "A" or "B"
hydrogenated product
%wt. at 125 °C. The
contained 11.3 ppm of
(35%wt.) and Cl⁻ (Cl)
this case, the hydrogen

Table 1. Tests

Catalyst	Type of test
A	Hydroisomerisation
	Hydroisomerisation
B	Hydroisomerisation
	Hydroisomerisation
C	Hydroisomerisation
	Hydroisomerisation

¹The used catalyst (100-100)

*(%wt. product - test)

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