Hydrogen + Fuel Cells 2011: International Conference and Exhibition, Convention Centre in Vancouver, May 15-18, 2011



A DIRECT DME FUEL CELL

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Abstract

A high temperature polymer fuel cell was operated as a vapour fed direct dimethyl ether (DME) fuel cell at ambient pressure and with air as oxidant. A peak power density of 67 mWcm⁻² was measured at 200°C. Conventional polymer based direct DME fuel cells are liquid fed and suffers from low DME solubility in water. When the DME-water mixture is fed as vapour miscibility is no longer a problem.

Dimethyl ether (DME)

DME is:

- A clean colourless gas
- Liquid at 6 bar(a) handled like LPG
- Little or not toxic
- Not a greenhouse gas (Decompose in atmosphere in tens of hours)

As a fuel:

- Excellent diesel engine fuel
- Burns with no particulate matter (soot)
- Cetane rating 55 60
 (45 for petroleum-derived diesel)

Other uses:

- Aerosol propellant
- Cooking gas

Manufacture (like methanol):

- From biomass
- From hydrogen and CO

Direct conversion in a PEMFC?



Methanol

Dimethyl

ether

CH₃OCH₃

18.92

28.62

-24.9

Ethanol

Gasolin

 C_7H_{16}

32.05

43.47

38–204

Ethanol

CH₃CH₂OH

21.09

26.87

Diesel

 $C_{14}H_{30}$

35.66

41.66

125–400

DME

Methane Methanol

CH₃OH

15.82

19.99

64

CH₄

0.0346

47.79

-162

Formula

(kJ cm⁻³)

LHV

LHV

(kJ g⁻¹)

Boil.p (∘C)



The idea is to increase the working temperature of the cell above the boiling point of water and feed the DME-water mixture as vapour. Then **miscibility** is not a problem.

For this a **high temperature polymer fuel cell (HT-PEMFC)** based on a membrane of acid doped polybenzimidazole (PBI) is applied.

A fuel cell system based on phosphoric acid doped polybenzimidazole (PBI) was first presented in 1995 (11).

It has been developed further (*12*) and even entered an early stage of commercialization by companies like BASF Fuel Cells and Danish Power Systems. The working temperature can be 120-200°C







Challenges for direct DME PEMFC

Like in direct methanol fuel cells DME is normally fed as an aqueous solution

Anode reaction: $CH_3OCH_3 + 3H_2O \rightarrow 2CO_2 + 6H^+ + 6e^-$

DME-water ratio 1:3 \Rightarrow 46 %wt solution of DME in water, but DME solubility in water is only ca. 7.6 %wt at 20°C(2).

At 80°C of a traditional polymer fuel cell the solubility is even lower and a saturated DME-solution separates into two phases (*3*). This impedes the process.

Peak power densities reported with conventional polymer fuel cells:

- Ambient pressure, air as oxidant: 20 to 40 mW/cm (3-5).
- Ambient pressure, pure oxygen: 30 to 56 mW/cm (6-10).

For most applications only air is available.

8.

The two-phase problem can in principle be mitigated by pressurizing the system, but this consumes energy and is not an attractive option for direct fuel cell systems which are meant to be simple.

One of the serious drawbacks of direct methanol fuel cells is the methanol crossover. DME crossover is expected to be less pronounced due to the lower solubility in a hydrophilic environment. Polarization curves of a direct DME fuel cell based on acid doped PBI at 200°C. Ambient pressure, air as oxidant.

<u>Anode catalyst:</u> PtRu/C (Johnson-Matthey, 1:1, 60 wt%, 3.7 mg metal pr cm²) <u>Cathode catalyst:</u> Pt/C (in house, 40 wt%, 0.7 mg Pt pr. cm²)

Schematic setup



A similar cell. Comparison of direct fuelling with DME and methanol

Conclusion

- A high temperature polymer fuel cell for direct DME allows for vapour fed operation.
- This solves the problem of phase separation.
- A peak power density of 67 mW/cm² at 200°C with air as oxidant.
- Highest peak power density reported to date for a DME polymer fuel cell at ambient pressure.
- High open circuit voltage indicates that crossover seems less than for direct methanol cells.



Further on DME:

International DME Association DME: 21st Century Energy

http://www.aboutdme.org

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• The Danish Agency for Science, Technology and Innovation (Vedvarende Energiteknologier)

• The ForskEL program (HotMEA)

