Permeability and p_c-S_w relationship for gas diffusion layers of a **PEMFC**



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Motivation

During two decades the polymer electrolyte membrane fuel cell (PEMFC) as one promising power source has been investigated in detailed experiments as well as in modelling studies. Most of the transport models were based on pore network models or on model types like the dusty gas model. Acosta et al. [1] used a Darcyflow based approach to model the processes in the electrodes of a PEMFC.



Figure 1a: Sketch of a PEMFC

The electrodes (cf. fig. 1a/b) consist of a flow field, a catalyst and a hydrophobic thin material, which is called gas diffusion layer (GDL).



Figure 1b: Cross section with profile of water content

Figure 2: GDL 200x, SEM micrograph

As a crucial point for modelling the counter-current transport processes of gas, water and electrical current occurring in the GDL of a PEMFC were identified. The knowledge of constitutive relationships as well as permeabilities as intrinsic parameters for the model of GDL is essential.

- The properties and tasks of the gas diffusion layer are:
- 200-500 µm thickness
- Enabling diffusion of gas to the catalytic layer • Formation of an electrical connection between
- bipolar plate/flow field and catalyst layer
- Carrying out of the produced water
- Providing a protective layer over the catalyst layer
- It is sometimes covered with micro porous layer
- (MPL) to enhance performance

While assembling the fuel cell stack the different layers are compressed due to sealing purposes, which leads to pore deformation and has a strong influence on capillary pressure - saturation relationship.

In the following sections experimental methods for the direct determination of permeabilities and pc-Sw relationships under well-defined compression levels for thin hydrophobic layers, such as GDLs, are presented.

Experimental setup

Capillary Pressure - Saturation Relationship

Capillary pressure on macro-scale (Darcy-flow based REV-models):

$$p_c = p_n - p_w = f(S_w)$$
 with $S_w = \frac{\Psi_a}{\Phi}$

where p_n denotes the pressure of the non-wetting phase, p_w for the wetting phase. For hydrophobic GDLs p_n is the pressure of the water phase, p_w the pressure of the gas phase.



Operation mode:

- Purging of lower part with water to remove trapped air
- Assembling stack consisting of membranes and sample (cf. figure 3)
- Sealing of cell and adjusting desired compression level with stamp
- Continuous imbibition and drainage cycles till pressure response is repeatable in the range of -30.000 Pa up to 30.000 Pa
- Starting stepwise injection of small portions of water via a syringe pump
- Pausing to enable relaxation of pressure (cf. figure 5/6)
- Withdrawing gradually small amounts of water followed by relaxation periods to gain the drainage curve





Permeability Measurement

The permeability as one key parameter for modelling the processes in the electrodes of a PEMFC was determined for several materials. In-plane (IP) and through-plane (TP) permeabilities (cf. figure 7) were measured with the following framework:

- Accurate flow of air ensured by mass flow controller (MKS 1179) and accounting for actual temperature by a thermocouple (type K)
- Differential pressure manometer (Furness FCO 12 micro manometer, range of 0-2 mbar or 0-20 mbar, accuracy ~0.001 mbar)

In-Plane measurement $R \cdot T$





15 20 25 Compression [%] Figure 11a: In-Plane (IP) permeabilities Figure 11b: Through-Plane (TP) permeabilities

utlook

- \rightarrow Examination of hysteresis behaviour of p_c - S_w function in detail
- → Combination of capillary pressure saturation relationship with permeability measurements within a new device, where:
 - capillary pressure (i. e. amount of water inside sample) is kept constant, while

20

Compression [%]

- · low-flow permeability measurements or/and
- counter-current diffusion experiments will be conducted

to obtain relative permeability - saturation relationships in axial and radial direction for gas diffusion layers.

References:

- [2]
- (ES: osta, M.; Merten, C.; Eigenberger, G.; Class, H.; Helmig, R.; Thoben, B. and Müller-Steinhagen, H.: odeling non-isothermal two-phase multicomponent flow in the cathode of PEM fuel cells. urnal of Power Sources 159 (2006), p. 1123-1141 Stick, J. T.; Joannidis, M. A.; Fowler, M. W. and Pritzker, M. D. : Direct measurement of the capillary pressure character flusion layer systems for PEM fuel cells. Electrochemistry Communications 10 (2008), p. 1520-1523 inveether, J. D.; Cheung, P.; SF-Perre, J. and Schwartz, D. T. : A microfluidic approach for measuring capillary pressure yers. Electrochemistry Communications, 2007, 9, p. 2340-2345 [3] ng capillary pressure in PEMFC gas diffusion layers. Electro





 $Q \cdot \eta \cdot d$

 $\Delta p \cdot A$

the fiber orientation (cf.

fig. 2)

K =



