

# **Overview to Power-to-Gas and methanation as an example**

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#### Institute for Micro Process Engineering



KIT – The Research University of the Helmholtz Association



## Babylonian confusion (First book of Moses, 11,1 - 11,9)

- Power-to-Gas, P2G, PtG
- Power-to-Chemistry, P2C, PtC
- Power-to-Fuel, P2F, PtF
- Power-to-Liquid, P2L, PtL
- Power-to-X, P2X, PtX
- Power-to-Methane, P2M, PtM
- Power-to-Ammonia, P2A, PtA
- Power-to-Methanol, P2M, PtM
- Power-to-Molecules, P2M, PtM
- 0

0

Green hydrogen

In this talk, <u>Power-to-Gas</u> means e-CH<sub>4</sub>







Pieter Bruegel the Elder - The Tower of Babel (Rotterdam) https://commons.wikimedia.org/wiki/File:Pieter\_Bruegel\_the\_Elder\_-<u>\_The\_Tower\_of\_Babel\_(Rotterdam) - Google\_Art\_Project - edited.jpg?</u> <u>uselang=de#filelinks</u>



## Some facts about natural gas in Germany (1/2)

### Natural gas H

- LHV: 9,4 11,8 kWh/m<sup>3</sup>
- HHV: 10,4 13,1 kWh/m<sup>3</sup>

#### Natural gas L

- LHV: 7,6 10,1 kWh/m<sup>3</sup>
- HHV: 8,4 11,2 kWh/m<sup>3</sup>

#### • Pure methane

- LHV: 9.97 kWh/m<sup>3</sup>
- HHV: 11.1 kWh/m<sup>3</sup>

m<sup>3</sup> at 1013 mbar, 0°C

Source: https://www.geothermie.de/bibliothek/lexikon-dergeothermie/b/brennwert.html

Total consumption in 2021 was 91 billion m<sup>3</sup> (bcm, thereof 1 bcm biomethane)

Equals ca. 800 - 900 TWh

• FNR sees **biomethane** potential of 35 bcm by 2030 (residues and energy crops)

Source: https://biogas.fnr.de/ biogas-nutzung/biomethan





# Some facts about natural gas in Germany (2/2)

- considered). This would store ca. **250 TWh** SRG (HHV, about 30% of the current consumption per year).
- Making 250 TWh SRG via PtG needs a lot of green electricity 500 TWh at 50% overall efficiency
  - **312,5 TWh** at 80% overall efficiency
- Current curtailed electrical energy per year in Germany (mostly from wind energy) is ca. 6 TWh
- Current stock market price of natural gas in Germany is 40 50 €/MWh
- Ourrent stock market price of electrical energy in Germany is 80 90 €/MWh
- Current price of CO<sub>2</sub> certificates in Germany is about 90 €/t
- 1 MWh methane converts into 0.2 t of CO<sub>2</sub>, which then gives currently a penalty of 18 €/MWh for fossil methane

To make PtG economically viable, we need a much higher  $CO_2$  certificate price or natural gas price, or a much lower electricity price; Imports from sweet spots and load-flexible plants



• Storage capacity for natural gas in Germany is 22.7 bcm (21 porous rock storages, 29 cavern storages, pipelines not

• Current gross consumption of electrical energy in Germany is 549 TWh (Wind: 125, PV: 60.8, Hydro: 17.5, Biomass: 50.2)



## Stock market gas price in Germany







## Stock market electricity price in Germany





## Where could the CO<sub>2</sub> come from?



Source: Fröhlich et al. (2019), DEHST 2021: Emissionshandelspflichtige Anlagen in Deutschland 2020 (Stand 03.05.2021), eigene Darstellung IZE Umweltbundesamt 2021 UFZ) HELMHOLTZ Zentrum für Umweltforschung DBFZ

- 25.11.2021
- A. Brosowski, F. Müller-Langer, V. Lenz, J. Horst, R. Dittmeyer, L. Uzor, M. Borchers, D. Thrän, P. Viebahn, U. Zuberbühler, FVEE Jahrestagung 2021

## And there is air, be it ambient or indoor...



Anzahl	t/a] Anza		
28	.694		
40	.603		
57	.546		
21	5.779		
9	.326		
155	.948		



Source: P. Heinzmann, S. Glöser-Chahoud, KIT-IIP, reFuels-Projekt am KIT, 2019 Wuppertal Institut

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Institute for Micro Process Engineering (IMVT)



#### 100 km

Raffinerieanlagen [14] Biomethananlagen [214] Biogasanlagen [10459]



## The PtX landscape in Germany - dena Strategieplattform Power to Gas (2011)

ÜBER DIE DENA



WIR UND DIE ENERGIEWENDE

THEMEN & PROJEKTE V

#### PROJEKT

### Strategieplattform Power to Gas

Seit 2011 leitet die dena die Strategieplattform Power to Gas. Zusammen mit Partnern aus Wirtschaft, Verbänden und Wissenschaft werden die Bedeutung von Power to Gas für die Nutzung erneuerbaren Stroms analysiert und die Rahmenbedingungen für die Nutzbarmachung der Systemlösung für den wirtschaftlichen und großtechnischen Einsatz definiert.

Strategieplattform Power to Gas

DENA.DE > THEMEN & PROJEKTE > PROJEKTE > STRATEGIEPLATTFORM POWER TO GAS

Source: https://www.dena.de/themen-projekte/projekte/projektarchiv/strategieplattform-power-to-gas/



## The PtX landscape in Germany - DVGW / H2020 Project Store&Go



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## Methanation of carbon dioxide

### Main reaction

 $CO_2 + 4H_2 \rightleftharpoons CH_4 + 2H_2O$  $\Delta H_R^0 = -165.1 \ kJ/mol$ 

#### Side reactions

 $\Delta H_{R}^{0} = 41.2 \ kJ/mol$  $CO_2 + H_2 \rightleftharpoons CO + H_2O$  $\Delta H_R^0 = -206.3 \ kJ/mol$  $CO + 3H_2 \rightleftharpoons CH_4 + H_2O$  $\Delta H_R^0 = -172.5 \ kJ/mol$  $2CO \rightleftharpoons CO_2 + C$  $\Delta H_R^0 = 74.9 \ kJ/mol$  $CH_4 \rightleftharpoons C + 2H_2$ 

#### Catalysts

- Typically Ni supported on Al<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub>, ZrO<sub>2</sub>, TiO<sub>2</sub>, MgAl<sub>2</sub>O<sub>4</sub>
- Typically 5-40 wt.-% Ni
- Issues: carbon formation, sintering, stability in transient operation



#### Sabatier reaction

- Reverse water gas shift reaction (rWGS)
- CO methanation
- **Boudouard reaction**
- Methane pyrolysis





## **Detailed reaction mechanism**

Actual surface reaction mechanism is much more complex; 42 elementary steps proposed





Source: D. Schmider et al., Ind. Eng. Chem. Res. 2021, 60, 5792–5805

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| ole 2. Detailed, Thermodynamically Consiste       | ent Reaction Mechanism fo       | r the Methana | tion of CO and C                           | 0 <sub>2</sub> over Ni <sup>a</sup>        |
|---------------------------------------------------|---------------------------------|---------------|--------------------------------------------|--------------------------------------------|
| reaction                                          | $A_j$ (cm, mol, s) or $S_0$ (*) | $eta_j$       | $E_{\mathrm{a},j}$ (kJ mol <sup>-1</sup> ) | $\varepsilon_{ij}$ (kJ mol <sup>-1</sup> ) |
| $H_2 + 2(s) \rightarrow 2H(s)  (R1)$              | $1.46 \times 10^{-2*}$          | 0             | 0                                          |                                            |
| $2H(s) \rightarrow H_2 + 2(s)  (R2)$              | $4.54 \times 10^{21}$           | -0.138        | 96.1                                       |                                            |
| $CH_4 + (s) \rightarrow CH_4(s)$ (R3)             | $1.06 \times 10^{-2*}$          | 0             | 0                                          |                                            |
| $CH_4(s) \rightarrow CH_4 + (s)$ (R4)             | $2.79 \times 10^{15}$           | 0.085         | 37.0                                       |                                            |
| $H_2O + (s) \rightarrow H_2O(s)$ (R5)             | $1.16 \times 10^{-1*}$          | 0             | 0                                          |                                            |
| $H_2O(s) \rightarrow H_2O + (s)  (R6)$            | $2.04 \times 10^{12}$           | -0.031        | 61.0                                       |                                            |
| $CO_2 + (s) \rightarrow CO_2(s)$ (R7)             | $6.29 \times 10^{-5*}$          | 0             | 0                                          |                                            |
| $CO_2(s) \rightarrow CO_2 + (s)$ (R8)             | $4.99 \times 10^{7}$            | 0.018         | 25.8                                       |                                            |
| $CO + (s) \rightarrow CO(s)$ (R9)                 | $3.74 \times 10^{-1*}$          | 0             | 0                                          |                                            |
| $CO(s) \rightarrow CO + (s)$ (R10)                | $1.14 \times 10^{12}$           | -0.103        | 112.0                                      | $50.0^{\dagger}$                           |
| $CO_2(s) + (s) \rightarrow CO(s) + O(s)$ (R11)    | $1.60 \times 10^{23}$           | -1.001        | 89.3                                       |                                            |
| $CO(s) + O(s) \rightarrow CO_2(s) + (s)$ (R12)    | $5.81 \times 10^{19}$           | 0             | 123.6                                      | $50.0^{\dagger}$                           |
| $CO(s) + (s) \rightarrow C(s) + O(s)$ (R13)       | $2.36 \times 10^{14}$           | 0             | 116.2                                      | $50.0^{\dagger}$                           |
| $C(s) + O(s) \rightarrow CO(s) + (s)$ (R14)       | $2.54 \times 10^{18}$           | 0             | 148.1                                      | 105.0 <sup>‡</sup>                         |
| $CO(s) + H(s) \rightarrow C(s) + OH(s)$ (R15)     | $3.05 \times 10^{18}$           | -0.223        | 105.3                                      | $50.0^{\dagger}$                           |
| $C(s) + OH(s) \rightarrow CO(s) + H(s)$ (R16)     | $2.18 \times 10^{18}$           | 0.128         | 62.8                                       | 105.0 <sup>‡</sup>                         |
| $CO(s) + H(s) \rightarrow HCO(s) + (s)$ (R17)     | $6.82 \times 10^{21}$           | -0.979        | 132.1                                      |                                            |
| $HCO(s) + (s) \rightarrow CO(s) + H(s)$ (R18)     | $2.18 \times 10^{20}$           | -0.021        | 0.2                                        | $-50.0^{\dagger}$                          |
| $HCO(s) + (s) \rightarrow CH(s) + O(s)$ (R19)     | $5.10 \times 10^{15}$           | 0.023         | 81.7                                       |                                            |
| $CH(s) + O(s) \rightarrow HCO(s) + (s)$ (R20)     | $3.42 \times 10^{19}$           | -0.023        | 110.2                                      |                                            |
| $H(s) + C(s) \rightarrow CH(s) + (s)  (R21)$      | $1.33 \times 10^{24}$           | -0.456        | 157.7                                      | 105.0 <sup>‡</sup>                         |
| $CH(s) + (s) \rightarrow C(s) + H(s)$ (R22)       | $2.63 \times 10^{22}$           | 0.456         | 22.3                                       |                                            |
| $CH(s) + H(s) \rightarrow CH_2(s) + (s)$ (R23)    | $3.21 \times 10^{25}$           | -0.084        | 81.1                                       |                                            |
| $CH_2(s) + (s) \rightarrow CH(s) + H(s)$ (R24)    | $6.16 \times 10^{24}$           | 0.084         | 95.2                                       |                                            |
| $CH_2(s) + H(s) \to CH_3(s) + (s)$ (R25)          | $7.78 \times 10^{22}$           | -0.048        | 59.5                                       |                                            |
| $CH_3(s) + (s) \to CH_2(s) + H(s)$ (R26)          | $6.16 \times 10^{24}$           | 0.048         | 95.9                                       |                                            |
| $CH_3(s) + H(s) \rightarrow CH_4(s) + (s)$ (R27)  | $3.63 \times 10^{21}$           | -0.048        | 65.7                                       |                                            |
| $CH_4(s) + (s) \rightarrow CH_3(s) + H(s)$ (R28)  | $6.16 \times 10^{21}$           | 0.048         | 53.6                                       |                                            |
| $H(s) + O(s) \rightarrow OH(s) + (s)$ (R29)       | $1.16 \times 10^{24}$           | -0.176        | 104.2                                      |                                            |
| $OH(s) + (s) \rightarrow H(s) + O(s)$ (R30)       | $7.70 \times 10^{19}$           | 0.176         | 29.8                                       |                                            |
| $H(s) + OH(s) \rightarrow H_2O(s) + (s) $ (R31)   | $2.34 \times 10^{20}$           | 0.075         | 44.1                                       |                                            |
| $H_2O(s) + (s) \rightarrow OH(s) + H(s)$ (R32)    | $2.91 \times 10^{21}$           | -0.075        | 90.4                                       |                                            |
| $2OH(s) \rightarrow H_2O(s) + O(s)$ (R33)         | $1.01 \times 10^{20}$           | 0.251         | 95.1                                       |                                            |
| $H_2O(s) + O(s) \rightarrow 2OH(s)$ (R34)         | $1.89 \times 10^{25}$           | -0.251        | 215.8                                      |                                            |
| $H(s) + CO_2(s) \rightarrow COOH(s) + (s)  (R35)$ | $1.29 \times 10^{25}$           | -0.46         | 117.2                                      |                                            |
| $COOH(s) + (s) \rightarrow CO_2(s) + H(s)$ (R36)  | $1.29 \times 10^{20}$           | 0.46          | 33.8                                       |                                            |
| $COOH(s) + (s) \rightarrow CO(s) + OH(s)$ (R37)   | $6.03 \times 10^{23}$           | -0.216        | 54.4                                       |                                            |
| $CO(s) + OH(s) \rightarrow COOH(s) + (s)$ (R38)   | $1.45 \times 10^{21}$           | 0.216         | 97.6                                       | $50.0^{\dagger}$                           |
| $COOH(s) + H(s) \rightarrow HCO(s) + OH(s)$ (R39) | $4.22 \times 10^{23}$           | -1.145        | 104.7                                      |                                            |
| $HCO(s) + OH(s) \rightarrow COOH(s) + H(s)$ (R40) | $3.25 \times 10^{19}$           | 0.245         | 16.1                                       |                                            |
| $2CO(s) \rightarrow CO_2(s) + C(s)$ (R41)         | $6.31 \times 10^{13}$           | 0.5           | 241.7                                      | $100.0^{\dagger}$                          |
| $C(s) + CO_2(s) \rightarrow 2CO(s)$ (R42)         | $1.88 \times 10^{21}$           | -0.5          | 2393                                       | 105.0 <sup>‡</sup>                         |

a(s) represents an empty surface site.  $\dagger$  denotes coverage dependency on CO(s),  $\ddagger$  on C(s). The mechanism is available in electronic form at www. detchem.com.





## Kinetics for a stoichiometric mixture of $H_2:CO_2 = 4:1$

and a sample of global models.







### Comparison of the predicted conversions for a CO<sub>2</sub> methanation experiment for the detailed kinetic model (blue)

Note that the reaction is fast and highly exothermic, and isothermal conditions are difficult to achieve.

In addition to heat removal from the reactor, heat and mass transport around and inside the catalyst particles may also affect the experimental results. bulk phase





Source: D. Schmider et al., Ind. Eng. Chem. Res. 2021, 60, 5792–5805

## Equilibrium position for a stoichiometric mixture of H<sub>2</sub>:CO<sub>2</sub> = 4:1

Thermodynamics set the boundary for what you may achieve in a practical system



Source: D. Schmider et al., Ind. Eng. Chem. Res. 2021, 60, 5792–5805





- High equilibrium conversion is favoured by low temperature and high pressure
- Carbon formation is expected over a wide temperature range, in particular at low pressure



# How to reach high conversion required for grid injection?

Standard approach in chemical engineering is to use consecutive adiabatic reactor stages with indirect or direct interstage cooling. In the case of methanation, two stages should be sufficient.



Source: D. Schlereth, Dissertation, TU München, 2015







## **Reactors for catalytic methanation**



- Dynamic operation 70-100%
- Footprint 8 x 4 x 15.5 m

Source: <a href="https://bit.ly/3QoRQQi">https://bit.ly/3xqF1MK</a>



Source: R. Bank, J. Dachs, F. Egner, V. Frick, M. Lehr, M. Specht, B. Stürmer, Shell- and-tube reactor for carrying out catalytic gas phase reactions **2012**, WO 2012 035 173 A1







## Strongly exothermic reaction in cooled tubular packed-bed reactors

#### Exponential dependency of the heat generation rate on temperature while cooling rate is linear creates hot spot

Material balance: 
$$\frac{\partial(uc_i)}{\partial z} = \rho_{bed} \sum \nu_{ij} r_j$$
 Heat balance



Source: D. Schlereth, O. Hinrichsen, Chem. Eng. Res. Des. 2014, 92, 702-712.

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### Fluidised bed

Bubbling fluidised bed with built in heat exchanger



Various kinds of contacting of a batch of solids by fluid. Adapted from Kunii & Figure R12.3-2 Levenspiel, Fluidized Engineering (Huntington, NY: Robert E. Krieger Publishing Co., 1977).





Main advantage of fluidised bed over packed bed

Improved heat transport due to solid's movement





## Advanced fluidised bed reactors

Three-phase slurry bubble column



Figure 1: Schematic of a three-phase methanation reactor.

**Source:** M. Götz et al., International Gas Union Research Conference, Copenhagen, **2014**, Paper-Code WP4.6, Abstract ID 210.





Source: M. Held et al., Chem. Ing. Tech. 2020, 92, 595-602.



### Honeycomb vs. slurry bubble column

Mass transfer resistances



Figure 3. Mass transfer phenomena in the honeycomb reactor (a) and the three-phase reaction system (b) based on theoretical considerations.



**Source:** M. Held et al., *Chem. Ing. Tech.* **2020**, *92*, 595-602.





### Honeycomb vs. slurry bubble column

Heat transfer resistances



Figure 4. Heat transfer phenomena in the honeycomb reactor (a) and the three-phase reaction system (b) based on theoretical considerations.



**Source:** M. Held et al., *Chem. Ing. Tech.* **2020**, *92*, 595-602.





### Honeycomb vs. slurry bubble column

Scale-up

Slurry bubble column reactor plant

Honeycomb reactor plant



**Source:** Aerial photograph of the PtX facility at KIT's Energy Lab 2.0. March 2021. Source: MDR WISSEN – Bremst Corona den Verkehr aus? Wie uns ein Virus zum Umlenken zwingt. https://bit.ly/361kR1Y





# **Microreactor technology for CO/CO<sub>2</sub> methanation**

Initiated in 2012 in the frame of KIC InnoEnergy project **MINERVE** ("Management of Intermittent & Nuclear Electricity by highly efficiency lectrochemical Reactor for the Valorization of CO<sub>2</sub> in flexible Energies")

### **Partners**:

- GDF Suez, CRIGEN, Paris
- CEA, Grenoble
- KIT, Karlsruhe
- AGH, Cracow
- Solvay (Rhodia), Lyon

#### **Rationale:**

- Highly efficient solid oxide co-electrolyzer (> 80%)
- CAPEX reduction due to the double function of the co-electrolyzer (steam and CO<sub>2</sub> reduction to produce synthesis gas)
- Higher global efficiency due to utilization of the reaction heat of methanation by steam generation (feed for electrolysis)





## **Reactor for lab-scale testing**



For details on the reactor, see: Myrstad et al., Catal. Today 147 (2009) 301-304



#### **Basic concept:**



## **Microreactor technology for CO/CO<sub>2</sub> methanation**

### Key project outcomes

- Reduced heating requirement in co-electrolysis (slightly exothermic mode) through steam supply at high temperature, i.e. only 2 % of total energy input is required for heating
- "Raw" methane possesses good composition:
  - CH<sub>4</sub> 95.389%
  - CO<sub>2</sub> 0.019%
  - CO 0.000%
  - H<sub>2</sub>O 0.461%
  - H<sub>2</sub> 4.131%
- Total efficiency: 83.5 %
  - AC/DC conversion of SOEC unit excluded
  - Efficiency is defined as: HHV ( $CH_4 + H_2$ ) / [E(SOEC) + Q(aux)]
- $CO_2$  Utilization > 99.8 %







## **Microreactor technology for CO/CO<sub>2</sub> methanation**

#### **Concept: Polytropic reactor with hot spot / falling temperature profile**



#### Thermodynamics







- Optimum operation regime :  $250 < T < 450^{\circ}C$
- Preferential methanation of CO
- High temperature allowed at inlet, low temperature required at reactor exit



# Scale-Up: Conceptual design of the first prototype

#### Geometry 0

- 2 Slits, width 5 cm, length 10 cm, hight 0.2 cm

## Catalyst

- 5 g of a commercial Ni-catalyst; particle size 200 600  $\mu$ m, diluted with SiC
- m<sub>Cat</sub> 10 times over design

## Cooling / Heating

- Co-current, air / steam / pressurized water, 70 channels 500  $\mu$ m x 500  $\mu$ m cross section
- 5 Heating cartridges for pre-heating (330°C ignition temperature)



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## **Key experimental results**

#### **Operation with air and steam cooling**



Reproducible and stable T-profiles up to 1.4 m<sup>3</sup>/h syngas feed (H/C = 4); exit temperature >  $350^{\circ}$ C

330 to 530 W of the generated heat transferred to air (ca. 40-50 W heat loss) 0

- Peak temperature  $< 500^{\circ}$ C in catalyst bed even when applying heat transfer to air  $\bigcirc$
- However, challenges with establishing the desired temperature profile without heating when applying evaporation cooling



- Feed composition: 10% CO, 7% CO<sub>2</sub>, 72%  $H_2$  and  $N_2$  (internal standard)
- Feed temperature: 300°C; coolant inlet temperature: 150°C
- Reduced in-situ in  $N_2/H_2 = 1:1$  (4h at 450°C)

## **Revised prototype for evaporation cooling**

Coolant in / out



see also: Sarvenaz Farsi, PhD-Thesis, KIT, 2021

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### **Start up behaviour**



Top cover plate Blank foil Redirecting foil 1/1 Redirecting foil 2 Redirecting foil 1/2 Redirecting foil 3 Catalyst foil Heating cartridge plate



Different catalysts



## A more detailed view into the results

#### Varying the cooling water split in evaporation cooling



see also: Sarvenaz Farsi, PhD-Thesis, KIT, 2021





## Modulation of the feed flow

#### Feed flow reduction from 21.1 to 15.8 l/min

15.8 to 21.1 l/min



 $H_2 / CO_2 = 4$ ,  $p_w = 10$  bar

see also: Sarvenaz Farsi, PhD-Thesis, KIT, 2021



# Feed flow increase from

**CO<sub>2</sub> Conversion** 

## Validation and scale-up

- Scale-up to 100  $m_N^3/d$  together with INERATEC
- Factor 5 larger than 2<sup>nd</sup> prototype, single stacking scheme, 0 two reactor stages
- Successful start-up of power-to-gas pilot plant by gasNatural 0 fenosa at a waste water treatment plant close to Barcelona, Spain (press release from May 31, 2018 at GNF website)

http://www.prensa.gasnaturalfenosa.com/en/gas-natural-fenosalaunches-pilot-project-to-produce-renewable-gas-in-catalonia/

- Testing at 10  $m_N^3/h$  scale in the Energy Lab 2.0
- Factor 6 larger than gasNatural fenosa system, double stacking scheme, two parallel modules **KOPERNIKUS**





Assembled power-to-gas pilot plant at INERATEC site in Karlsruhe before shipping









Synthetic Fuels – Combustibles Sintètics (CoSin), Grant No. COMRDI15-1-0037



Generalitat de Catalunya











## 100 kW Prototype - first operation campaign (1/3)



- Module 1 operated in normal condition
- Module 2 did not ignite (~ 380°C)



- 5 bar
- 5 m<sup>3</sup>/h CO<sub>2</sub>
- 20 m<sup>3</sup>/h H<sub>2</sub>
- 3 m<sup>3</sup>/h N<sub>2</sub>





## 100 kW Prototype - first operation campaign (2/3)



Control of Module 1 within very short time possible; similar ignition behavior like original reactor



- 5 bar
- 5 m<sup>3</sup>/h CO<sub>2</sub>
- 20 m<sup>3</sup>/h H<sub>2</sub>
- 3 m<sup>3</sup>/h N<sub>2</sub>





# 100 kW Prototype - first operation campaign (3/3)



Module 1 contributed mainly to conversion; parameters still to be optimized

- Module 2 did not ignite (~ 380°C) and behaved like a bypass
  - most reasonably due to failed reduction or oil droplets from slurry bubble
  - column (problems with aerosol formation in product line)



2.0 1.8 1.6 % 1.4 1.2 Selectivity 1.0 0.8 0.6 0.4 0.2

- 5 bar
- 5 m<sup>3</sup>/h CO<sub>2</sub>
- 20 m<sup>3</sup>/h H<sub>2</sub>
- 3 m<sup>3</sup>/h N<sub>2</sub>





## 100 kW Prototype - Second operation campaign - reactivation of module 2





Both modules ignited (~ 400°C), partly approval of re-reduction process Although different absolute temperature good control over temperature



# 100 kW Prototype - Third operation campaign - flow modulation



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Successful approval of flow modulation





## 100 kW Prototype - Fourth operation campaign - water cooling only on 2nd inlet



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Control almost possible, however, long adaptation times and final

Further validation and/or design improvements needed



# **Summary / Outlook**

## **Status of PtG**

- Technology is more or less ready for commercial use, but economics are difficult and still have to be improved
- PtG, like all types of PtX, only makes sense (from a CO<sub>2</sub> emission reduction point of view) if CO<sub>2</sub>-free electrical power is used
- Current R&D addresses catalyst improvements, advanced reactor designs, and process integration (heat, material flows) with a focus on higher efficiency, lower cost, capability of transient operation, increased longterm stability all targeting lower production cost and higher operational flexibility

## Uncertainties regarding the commercial implementation of PtG

- Future role of gas in domestic and industry heating ?
- Future role of gas in electric power generation (cold dark doldrums, security of supply)?
- Future of the gas grid: methane or hydrogen, both, or (partial) deconstruction ?
- Role of CRG/LRG in transport ?
- Business model for PtG ?

