

**Figure S1** Progress of the CO conversion rate measured at 210°C, 220°C and 240°C. Apparent CO conversion rate is measured with the same set of process parameters except the temperature, which was always 40°C for any experiment. ( $V_{syngas,210°C} = 2.5 I_{STP}(g_{cat}h)^{-1}$ ;  $V_{syngas,220°C} = 5 I_{STP} (g_{cat} h)^{-1}$ ;  $V_{syngas,240°C} = 30 I_{STP} (g_{cat} h)^{-1}$ ;  $p_{total} = 15$  bar;  $H_2/CO = 2/1$ ).



**Figure S2** Progress of product distribution calculated from measured chain growth probability values using Anderson-Schulz-Flory equation during the initial time of Fischer-Tropsch synthesis.



**Figure S3** Decline of pore filling degree over drainage time for 210°C ( $V_{HGL,total} = 45 I_{STP}/h$ ;  $p_{H2} = 1$  bar; ptotal = 15 bar).



**Figure S4** Comparison of FTS activities before and after pore draining using Hydrogenolysis at 210°C, 220°C and 240°C. FTS run A is measured prior to pore drainage, whereas FTS run B is measured directly after the pore drainage (full drainage, corresponds to last measuring point of the 16 h experiment visible in Figure 4 and Figure S3).



**Figure S5** Typical distribution of products of hydrogenolysis at 210°C and 220°C after various Fischer-Tropsch synthesis time. Selectivity of methane and kerosene fraction  $(C_9-C_{17})$  at different temperatures and pore filling degrees.