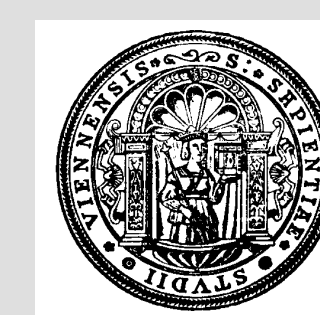


Creep of Carbon Fibres



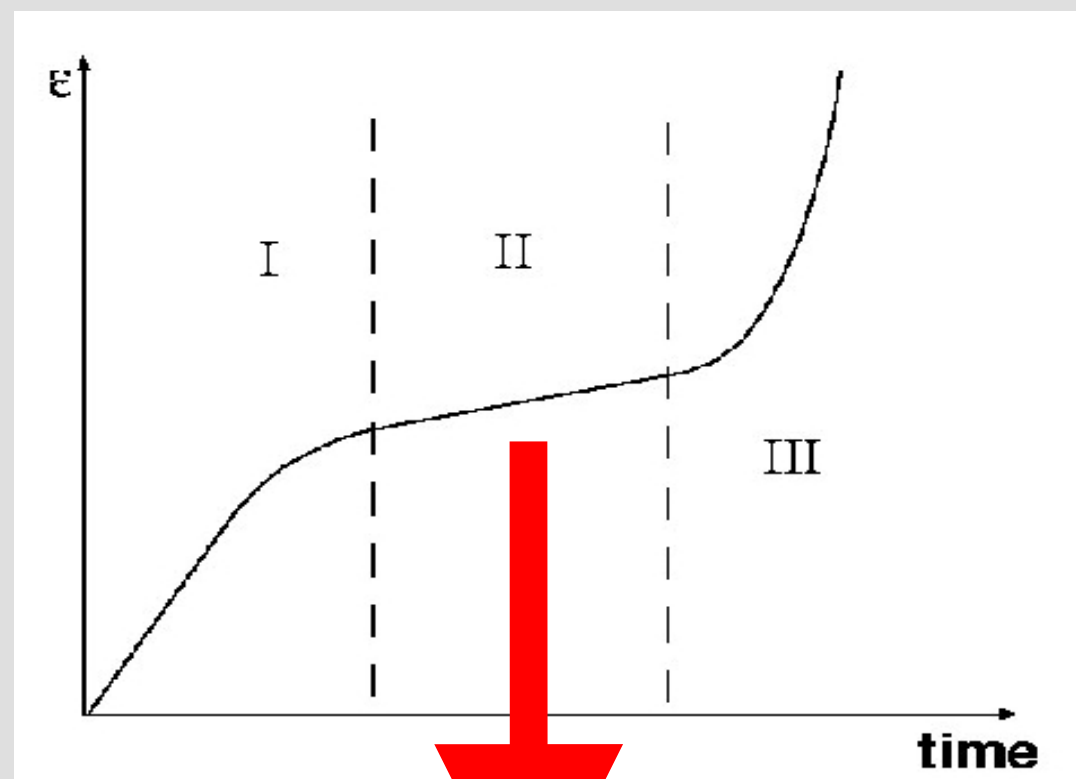
Measurement of mechanical properties and analysis of post-creep structure with X-ray scattering



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What is creep?



Elongation of carbon-fibres under stress at high temperatures because of structural change:

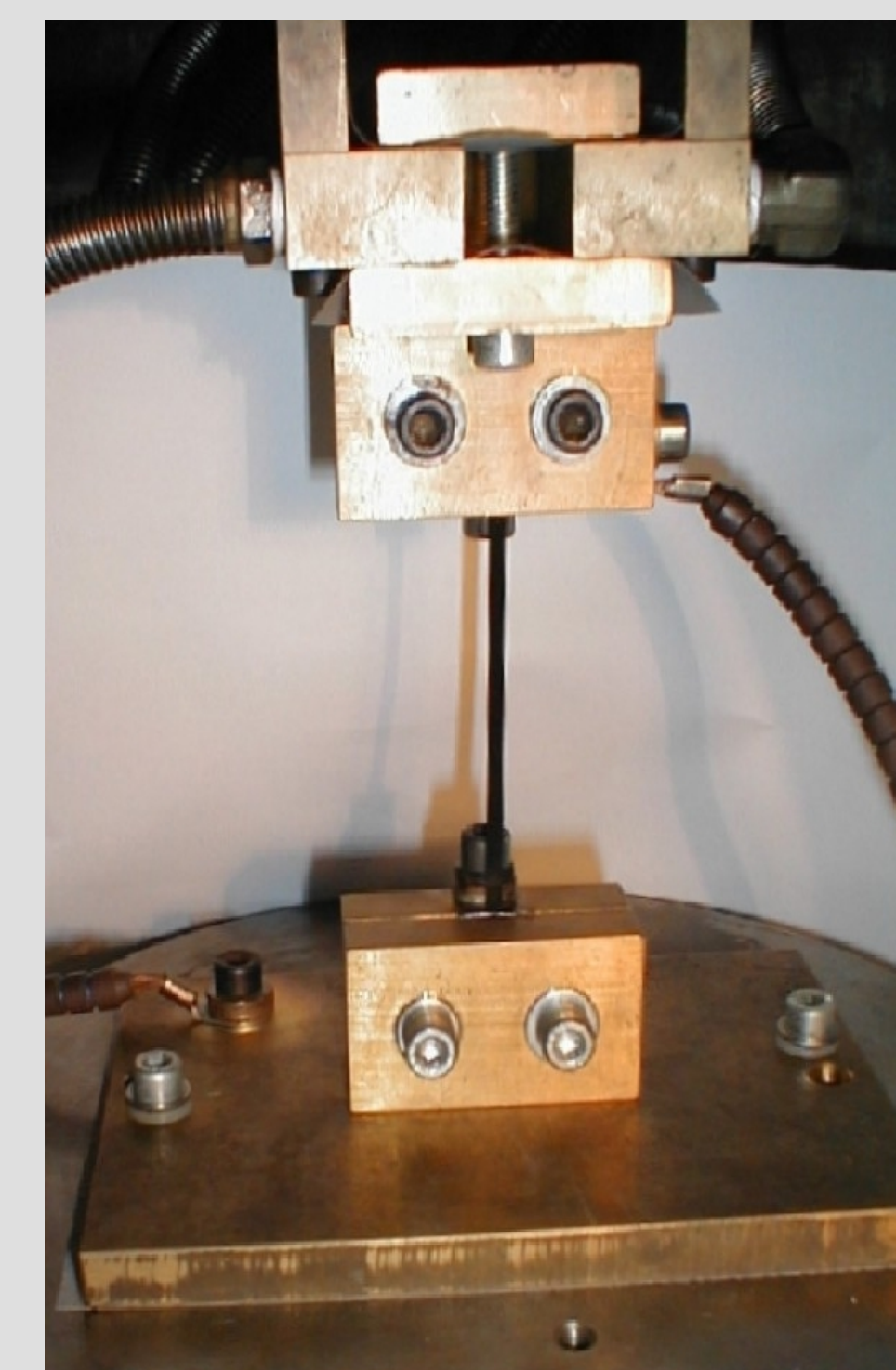
- I... creation of pores
- II... growth of pores
- III... cracks, coalescence of pores

$$\dot{\epsilon} = A\sigma^n \cdot e^{-\frac{Q}{RT}}$$

Measurement

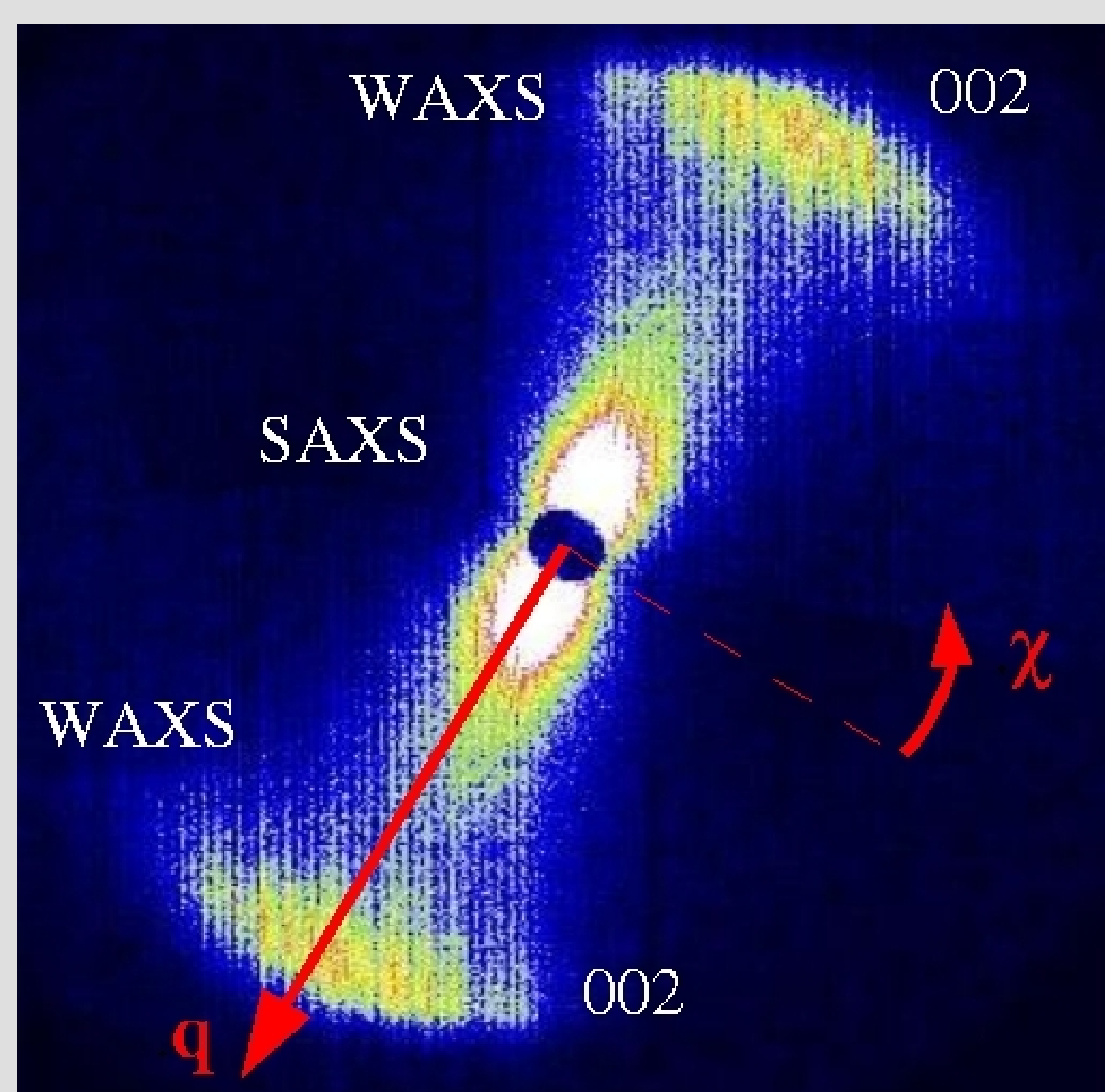
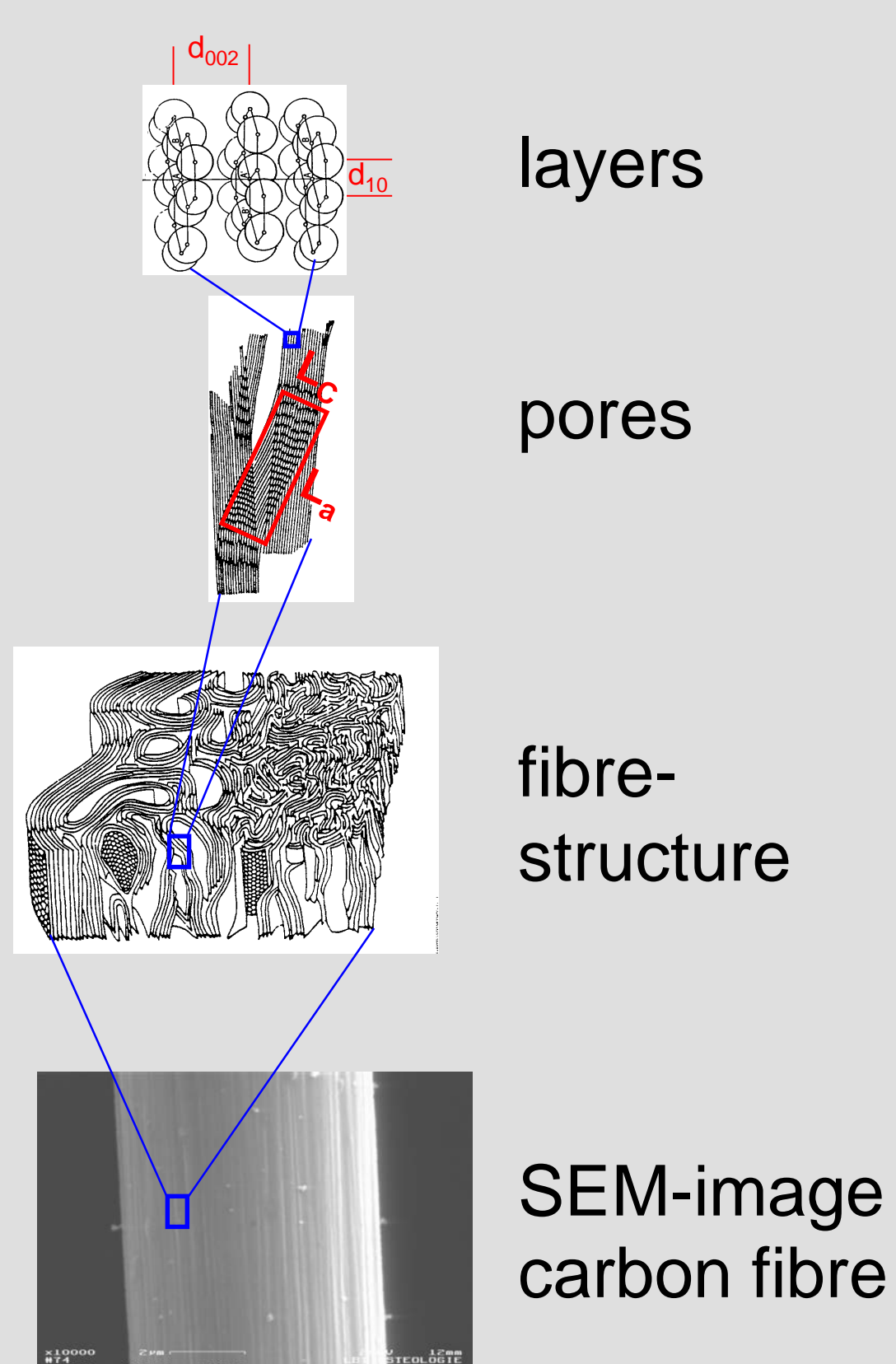
- direct heating (electric current through the bundle of carbon fibres)
- constant load (hydraulic system) in vacuum vessel (pressure <math>< 10^{-3}</math> mbar)
- cooled grips
- (66 +/- 1) mm effective gauge length

measurement at: 173, 217 and 260 MPa
1500, 1600 and 1700°C



Structure

X-ray scattering has been used to investigate the structural change in the carbon fibres caused by creep, dependent on stress and temperature. Information about the pores (small angle X-ray scattering - SAXS) and about the graphite-layers (wide angle X-ray scattering - WAXS) in the fibres is gained.

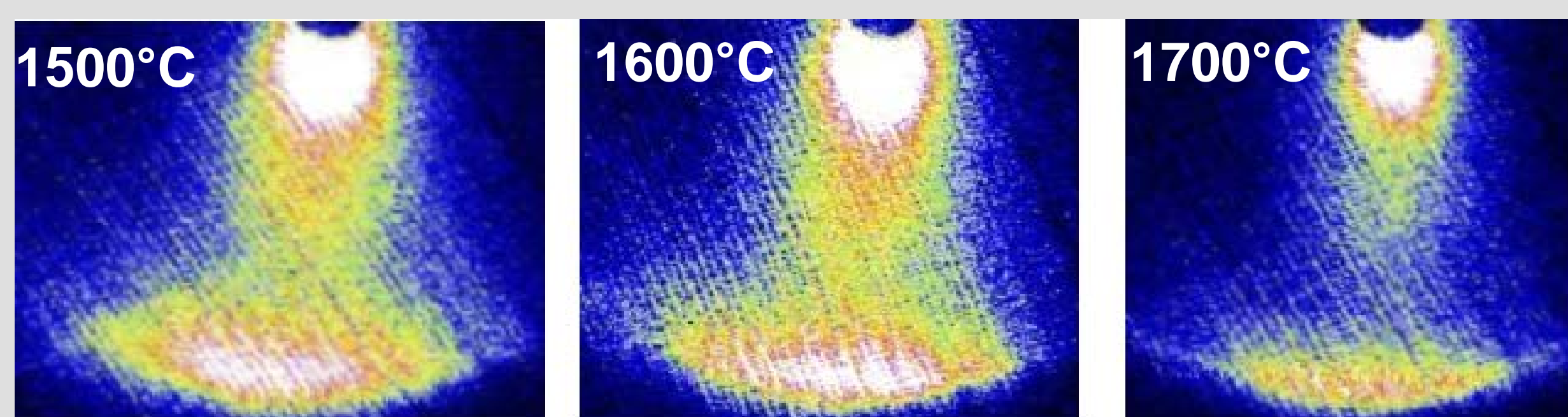


SAXS (information about pores):

- mean diameter 7 Å growing at $T \uparrow$
- mean orientation to fibre-axis 18° declining at $T \uparrow$

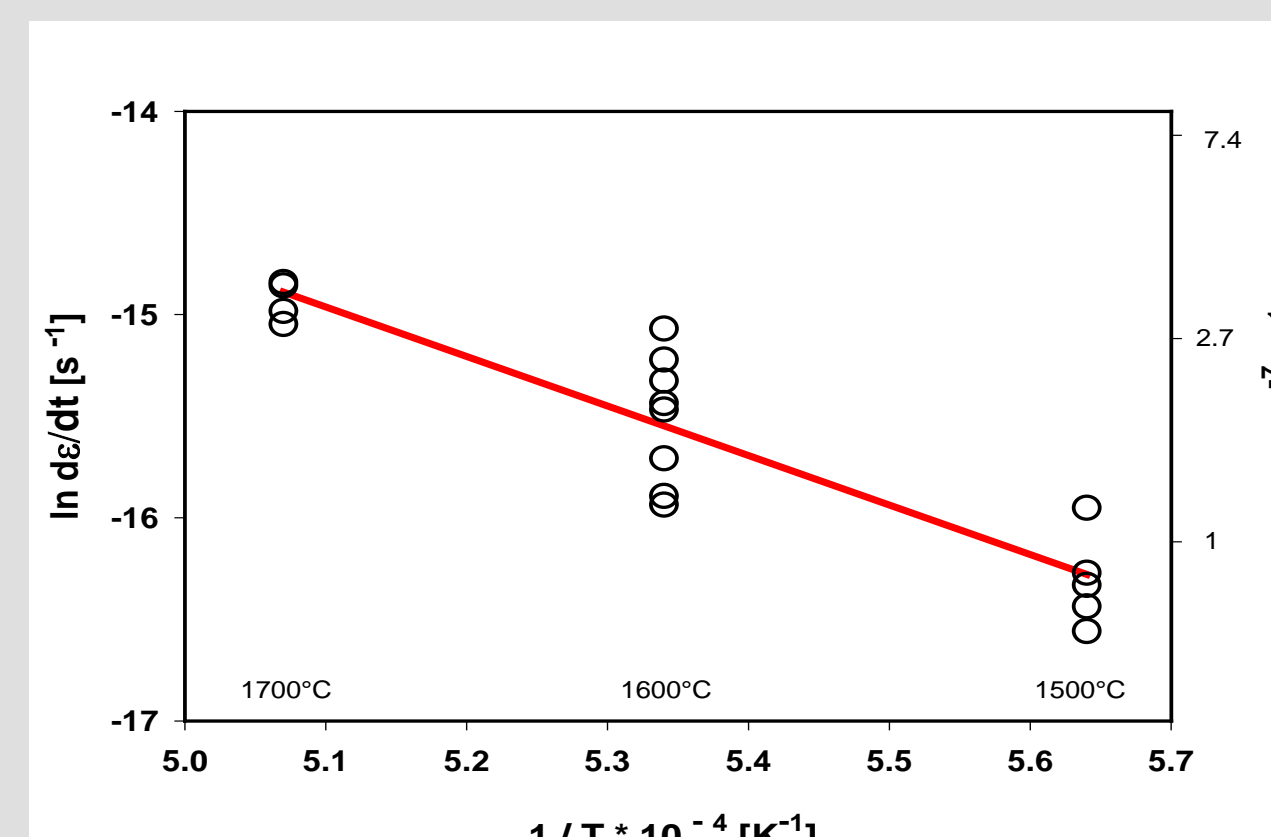
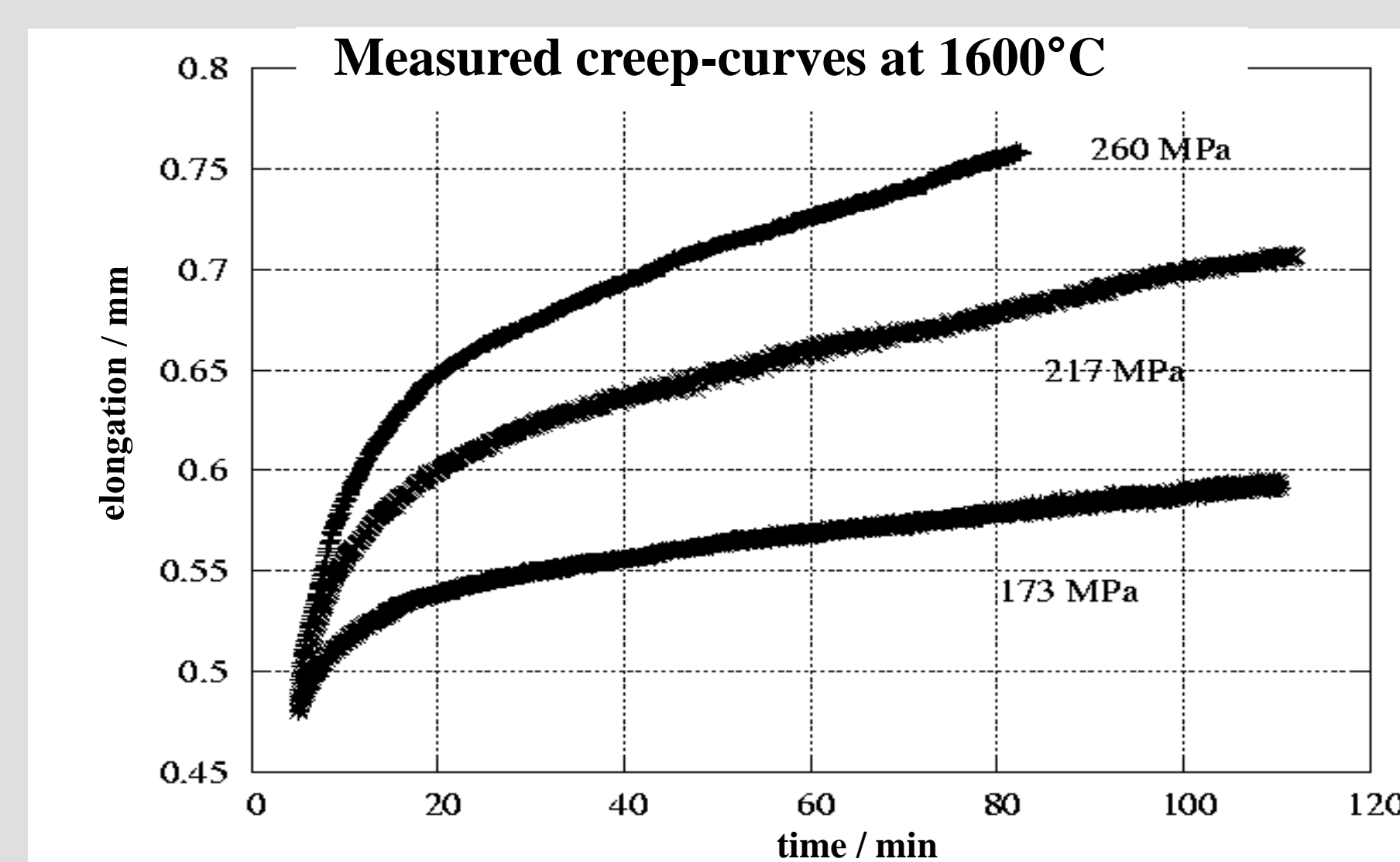
WAXS (information about graphite-layers):

- mean interlayer-spacing 3.46 Å constant
- mean orientation to fibre-axis 18° declining at $T \uparrow$



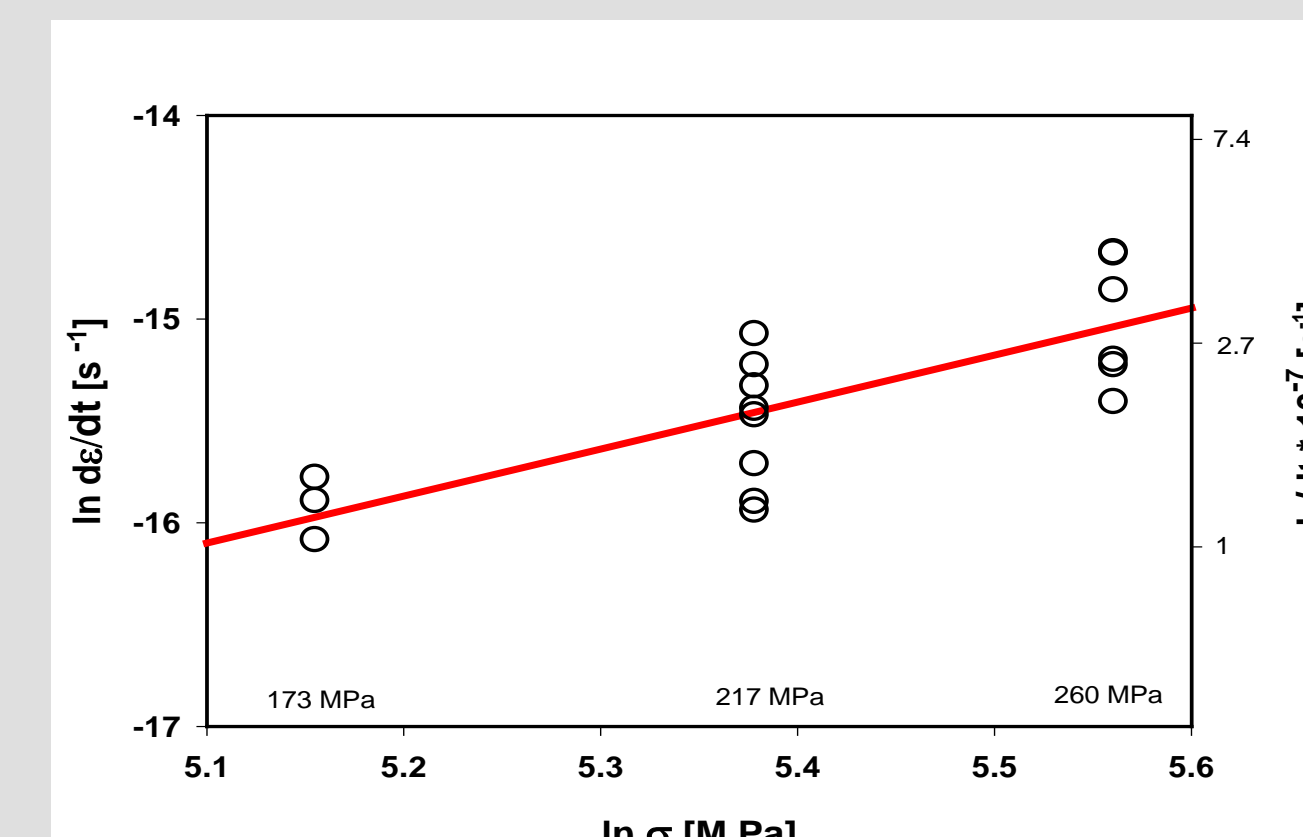
Scattering-images at 217 MPa: the distribution-width is smaller after creep-experiments at higher temperature.

Mechanical Properties



Activation Energy

$$Q = (167 \pm 26) \text{ kJ/Mol}$$



Creep-Exponent

$$n = 2.3 \pm 0.5$$

Conclusion

Low Activation Energie Q



No Diffusion of C-Atoms
[Kanter57]

+ $n \approx 2$
[Cannon83, Cannon88]

The creep-mechanism proposed for these experiments is the glide of grain-boundaries:

Rachinger-sliding

Loss of nitrogen during the heating possibly facilitates this process. Further experiments in argon-atmosphere will be performed for clarification.