

The interactions between lignocellulosics and water at a micro-level

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The ISOBIO project

Title: Development and Demonstration of Highly Insulating, Construction Materials from Bio-derived Aggregates

- Duration: 4 years (February 2015 – January 2019)
- Coordinated by TWI Ltd (Cambridge, UK)
- Budget of budget of 6,3M€
- Funded by the Horizon 2020 programme under a specific call to improve energy performance and reduced embodied energy across the whole life cycle of a building (EeB-01 - Materials for building envelope).

Objectives

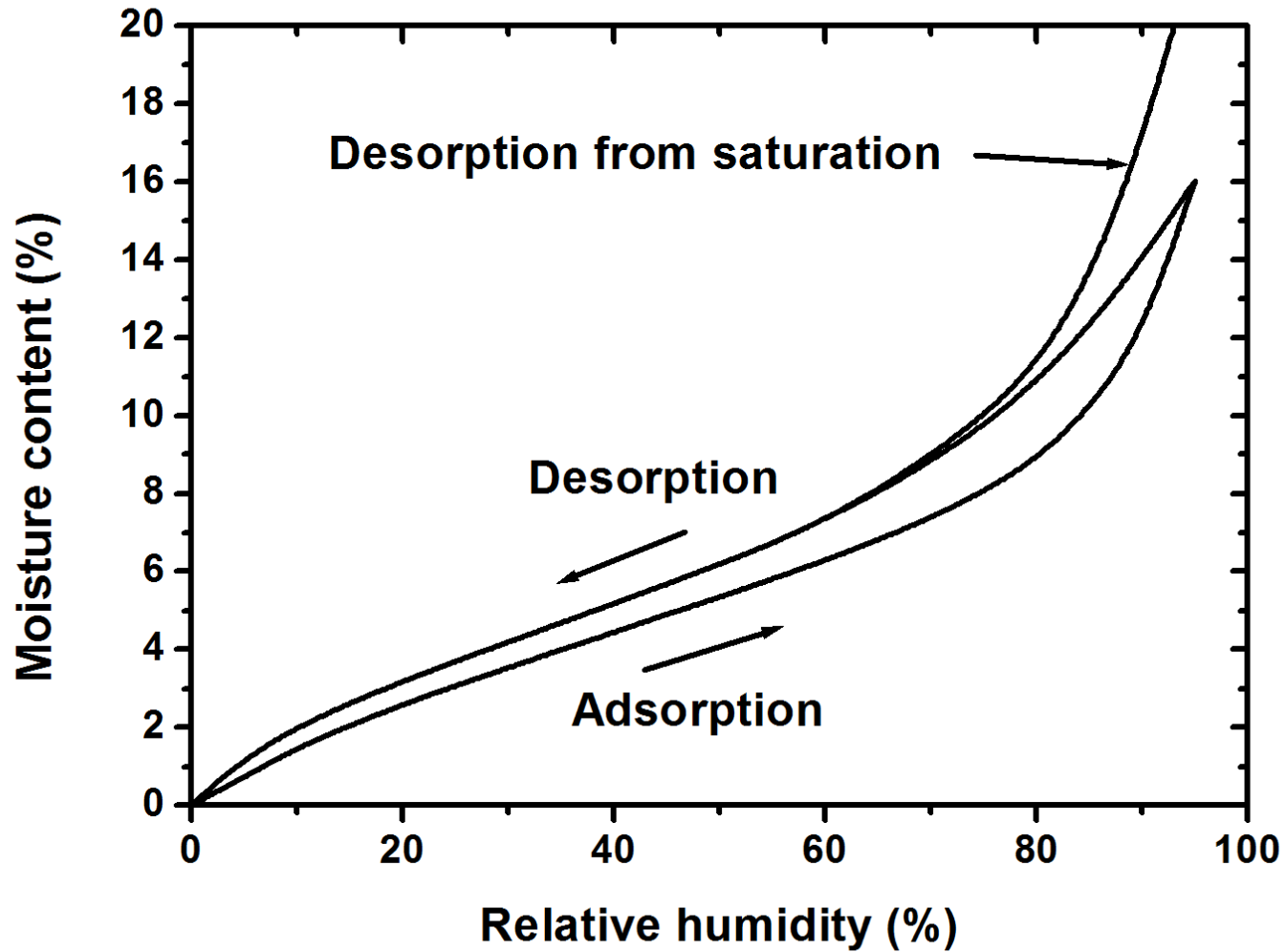
- To develop and bring new bio-based insulation panels and renders into mainstream in order to improve the energy efficiency of buildings;
- To assess and advance the state-of-the-art in natural insulation materials, and hygrothermally (heat and moisture) efficient buffering materials, binders, sol-gels and resins;
- To deliver products with at least:
 - **50% reduction in embodied energy and CO₂ emissions** at component level;
 - **20% better insulation** properties than conventional materials;
 - **15%, lower construction costs**; and
 - **5% reduction in the total energy spent** over the lifetime of a building.

The consortium

Multidisciplinary consortium of 11 partners from 6 different European countries.

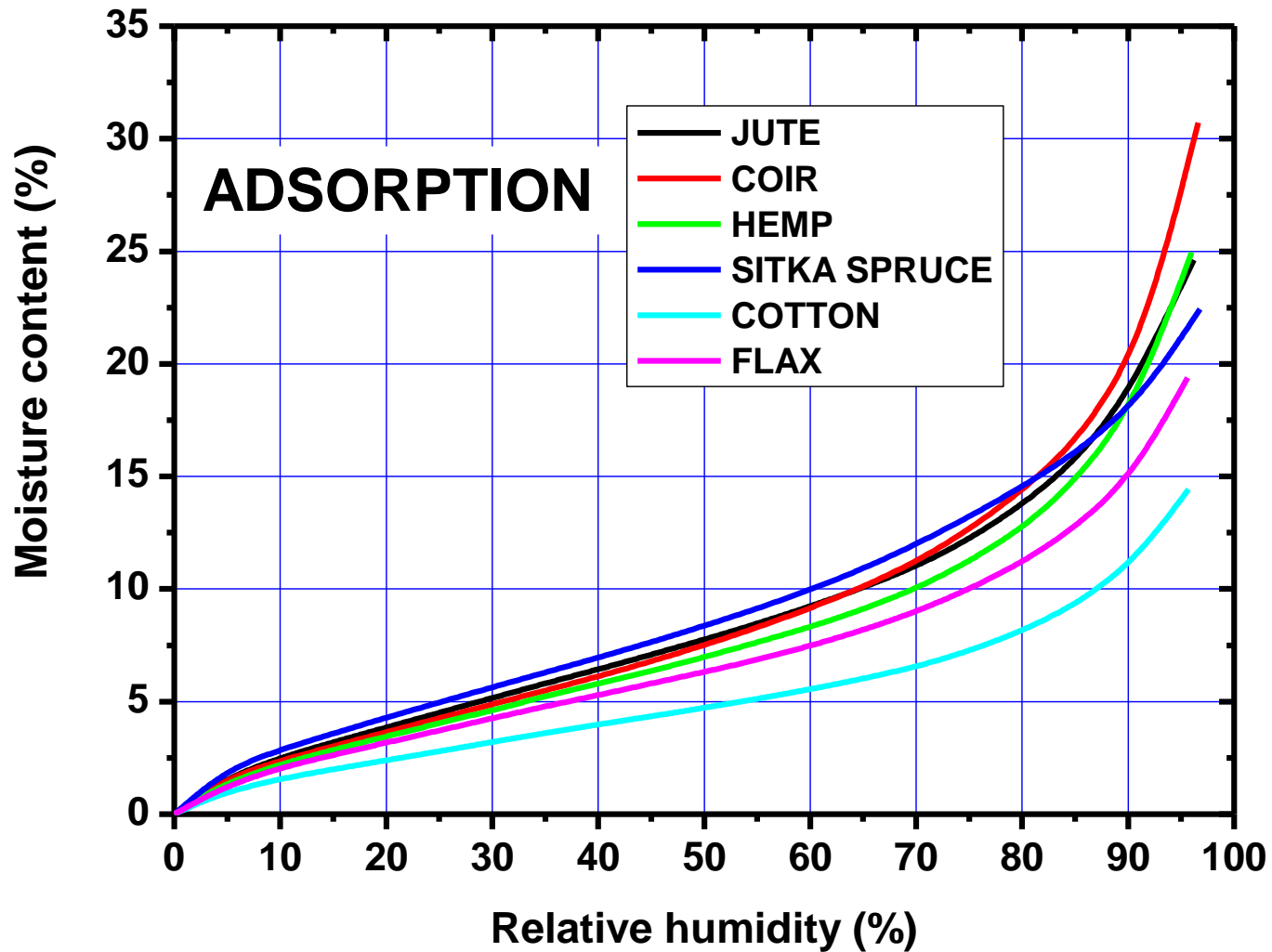
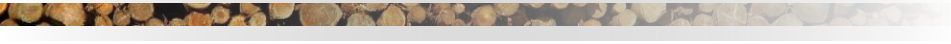


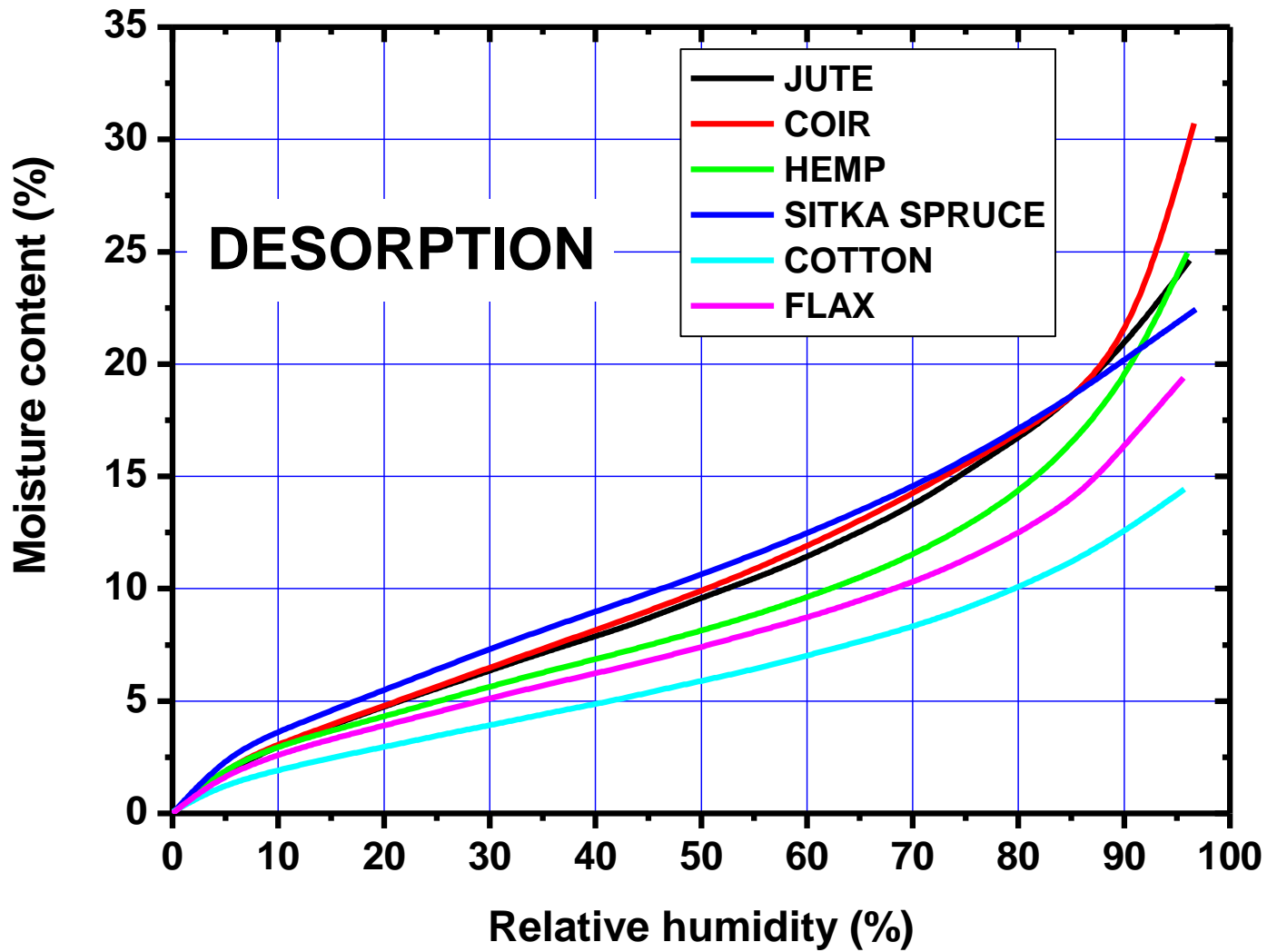
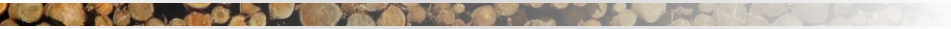
HYSTERESIS

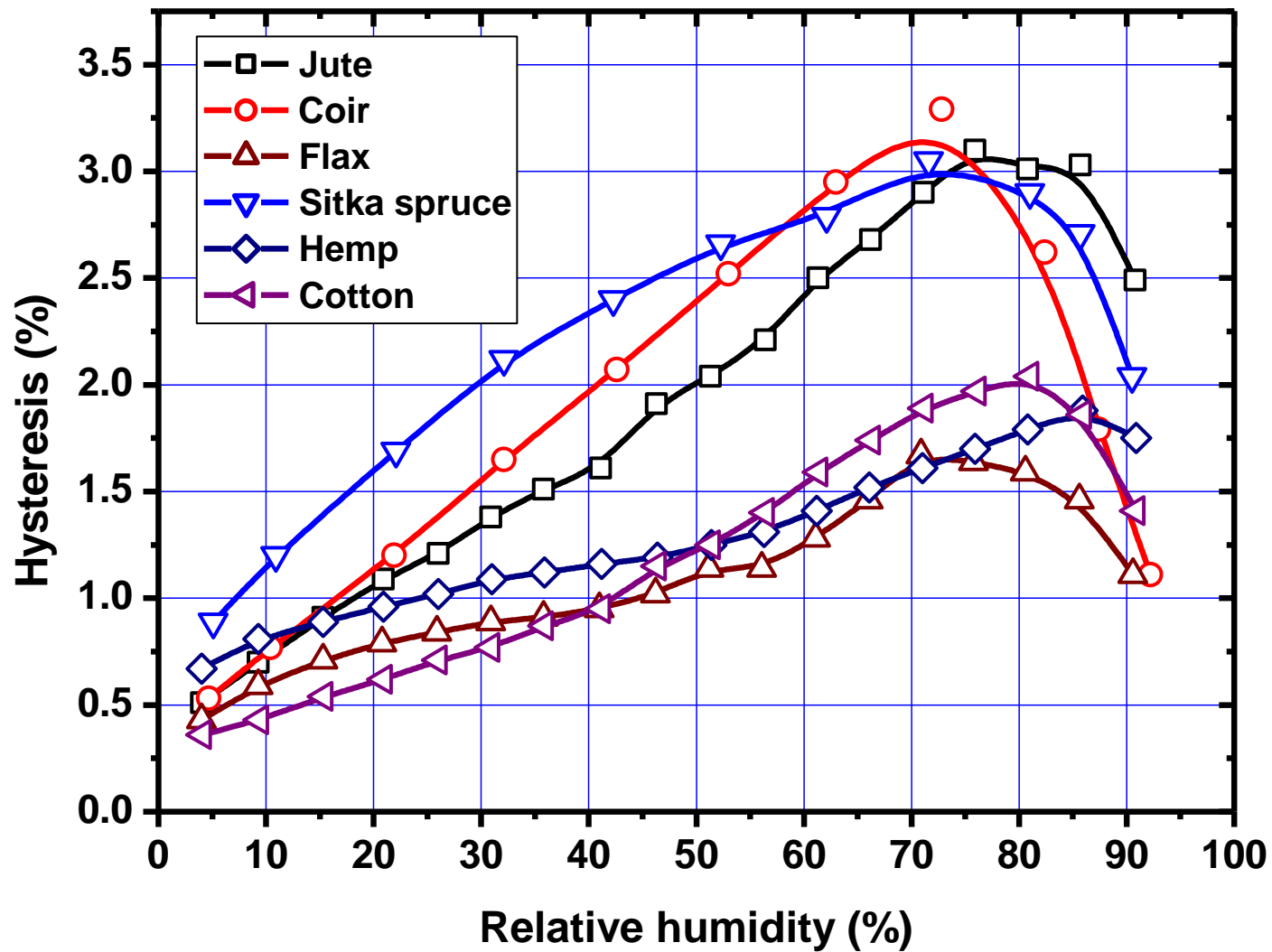


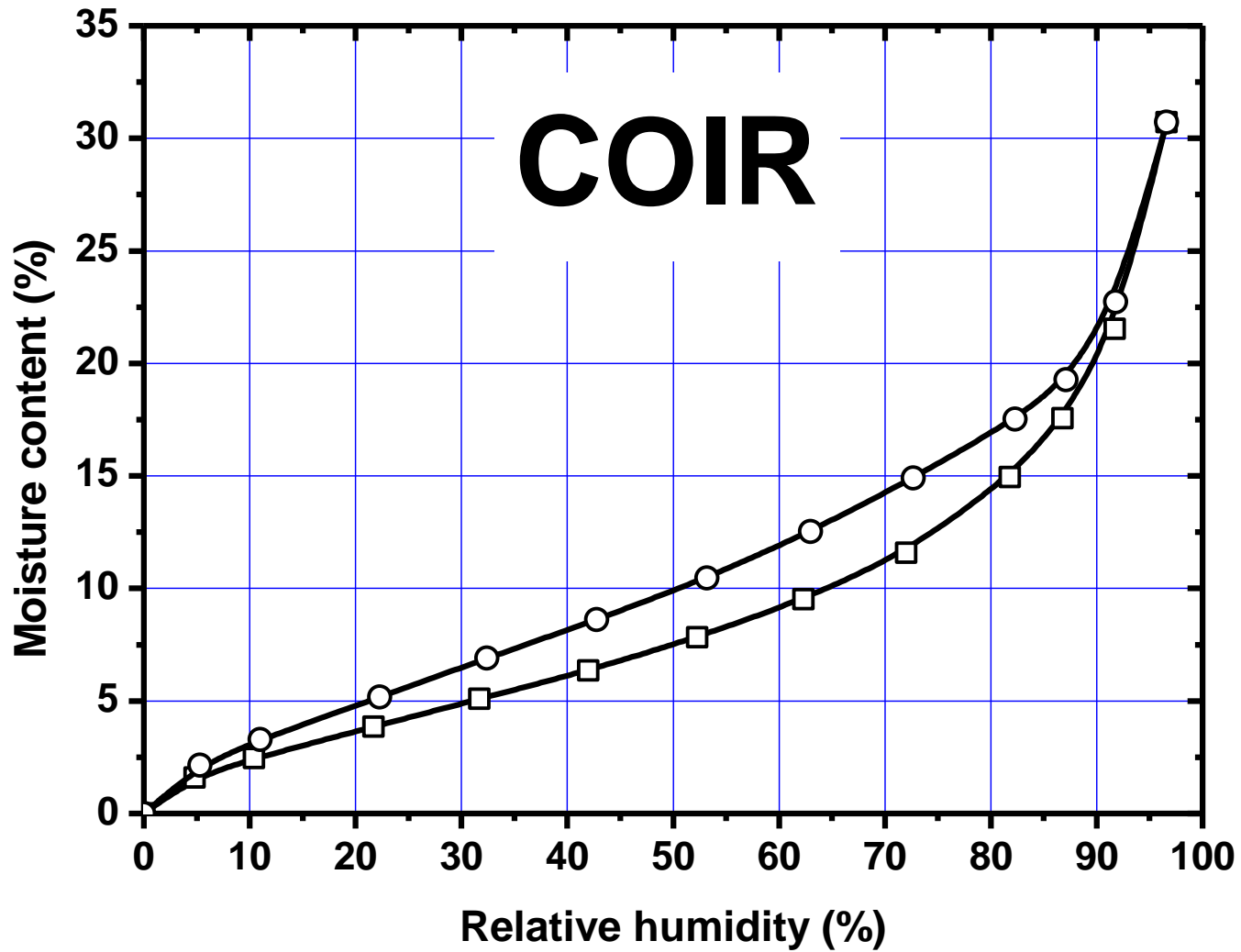
WHY DO WE GET HYSTERESIS?

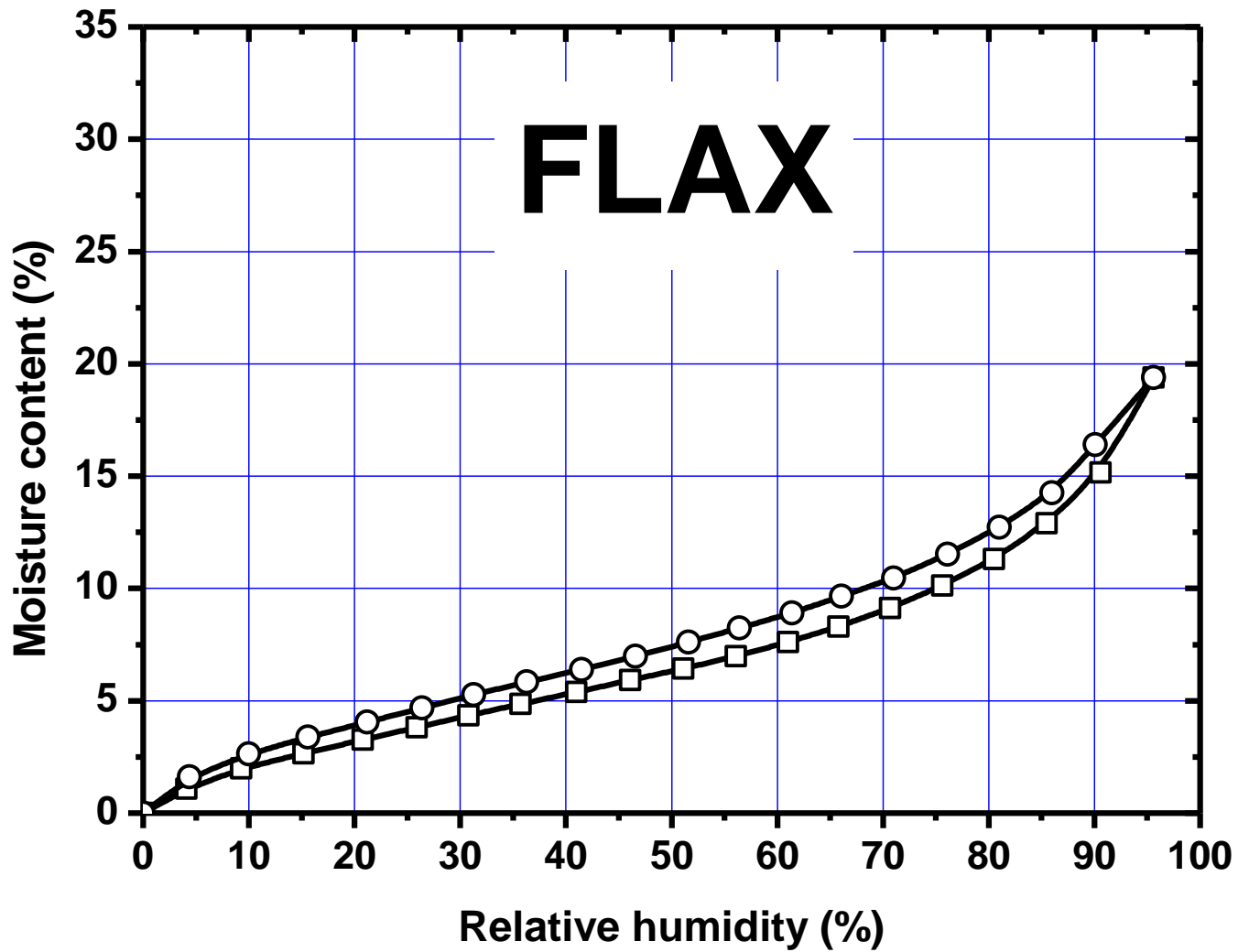
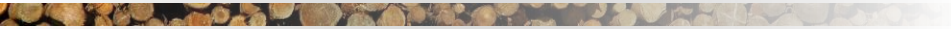
- Questions:
- What is the origin of sorption hysteresis?
- Can the mechanism for hysteresis be linked to the sorption kinetic behaviour?









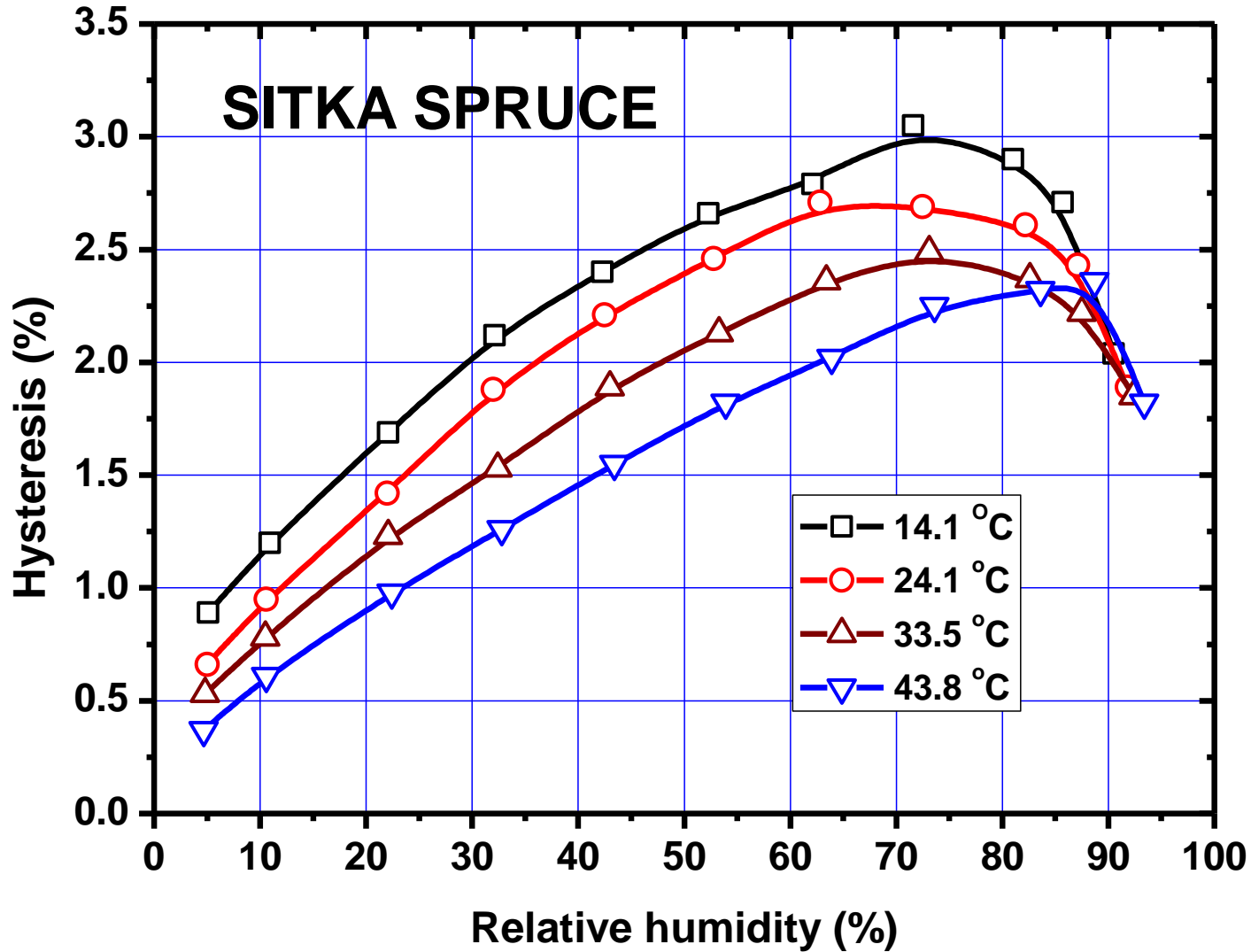


HYSTERESIS

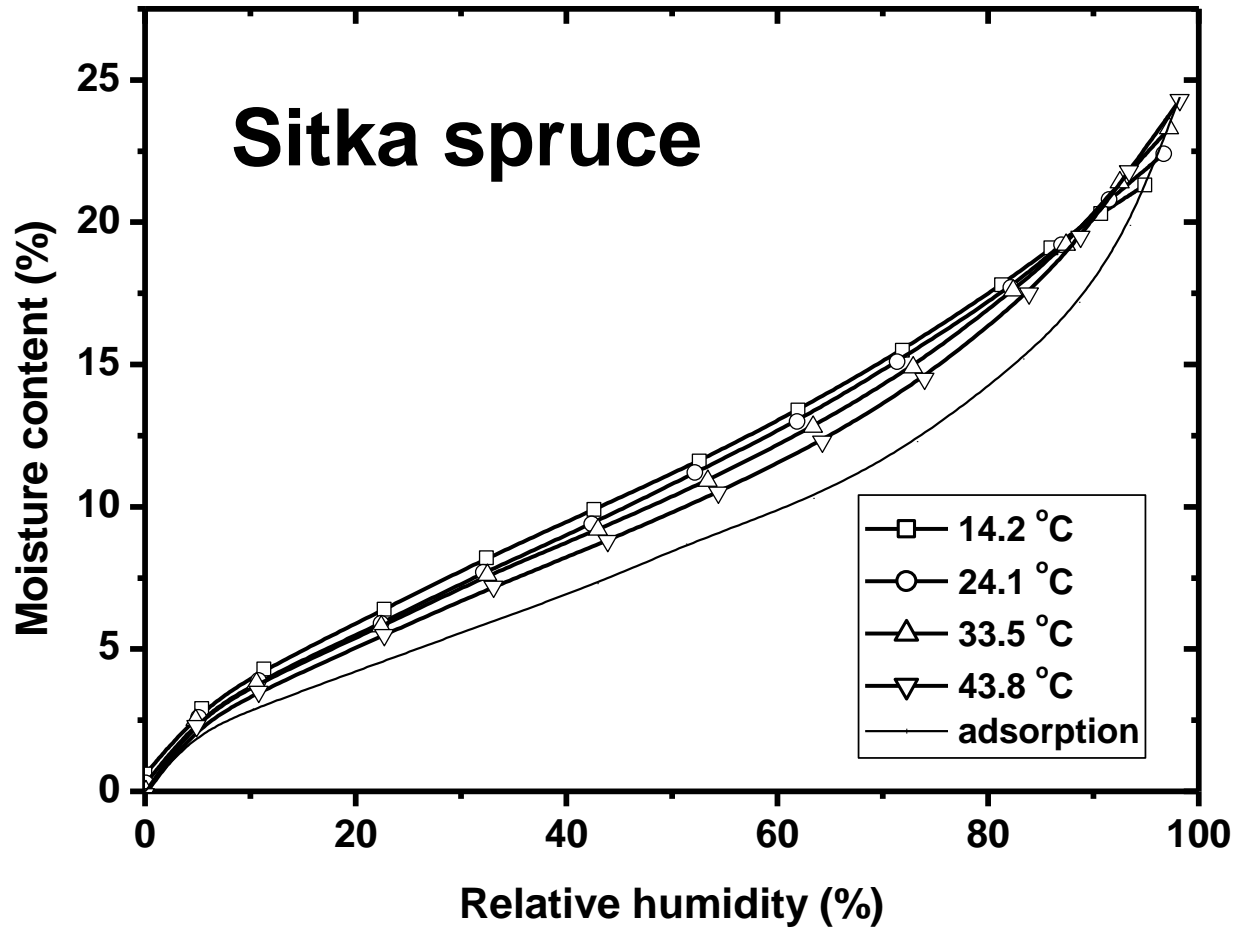


- Model by Vrentas and Vrentas
- Sorption onto a glassy solid below the glass transition temperature
- Hysteresis arises due to adsorption and desorption into/out of a material in different states

Hysteresis



HYSTERESIS



HYSTERESIS



The Water Vapor Sorption Behavior of a Galactomannan Cellulose Nanocomposite Film Analyzed Using Parallel Exponential Kinetics and the Kelvin-Voigt Viscoelastic Model

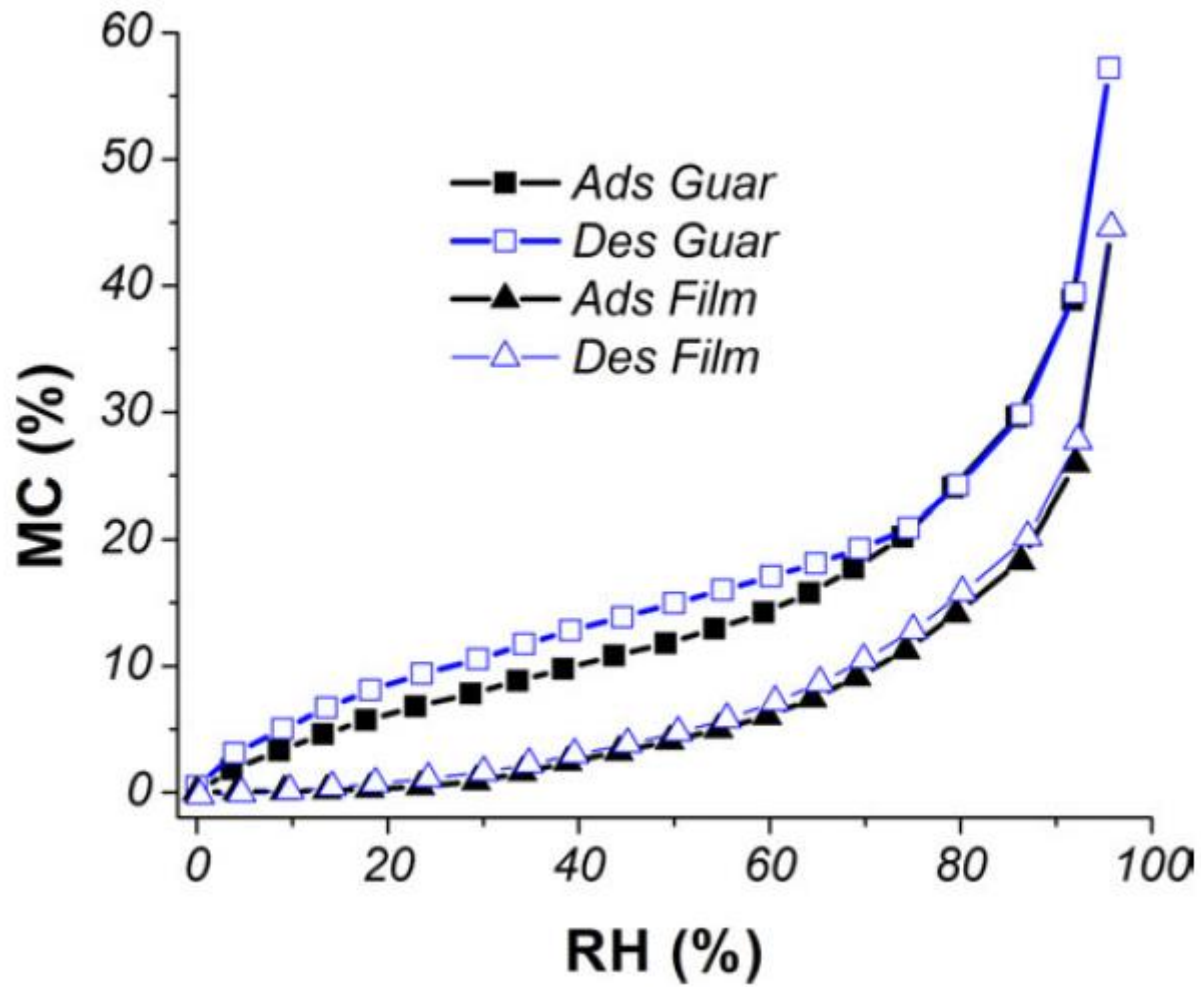
Barbara A. Keating,¹ Callum A. S. Hill,² Dongyang Sun,² Rob English,² Phil Davies,³ Charles McCue³

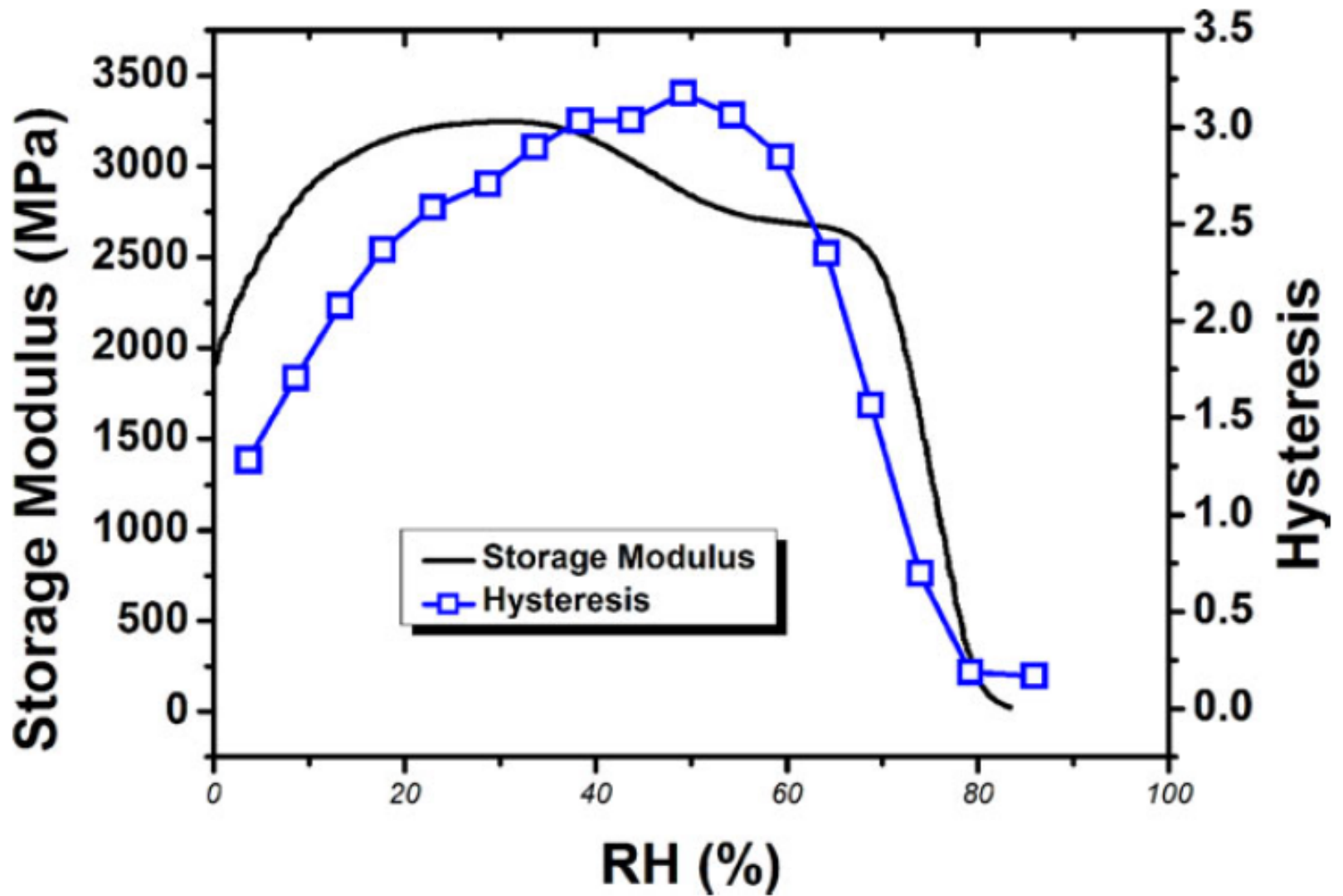
¹Department of Mechanical and Aerospace Engineering, The University of Strathclyde, Glasgow G1 1XJ, United Kingdom

²Forest Products Research Institute, Joint Research Institute for Civil and Environmental Engineering, Edinburgh Napier University, Edinburgh EH10 5DT, United Kingdom

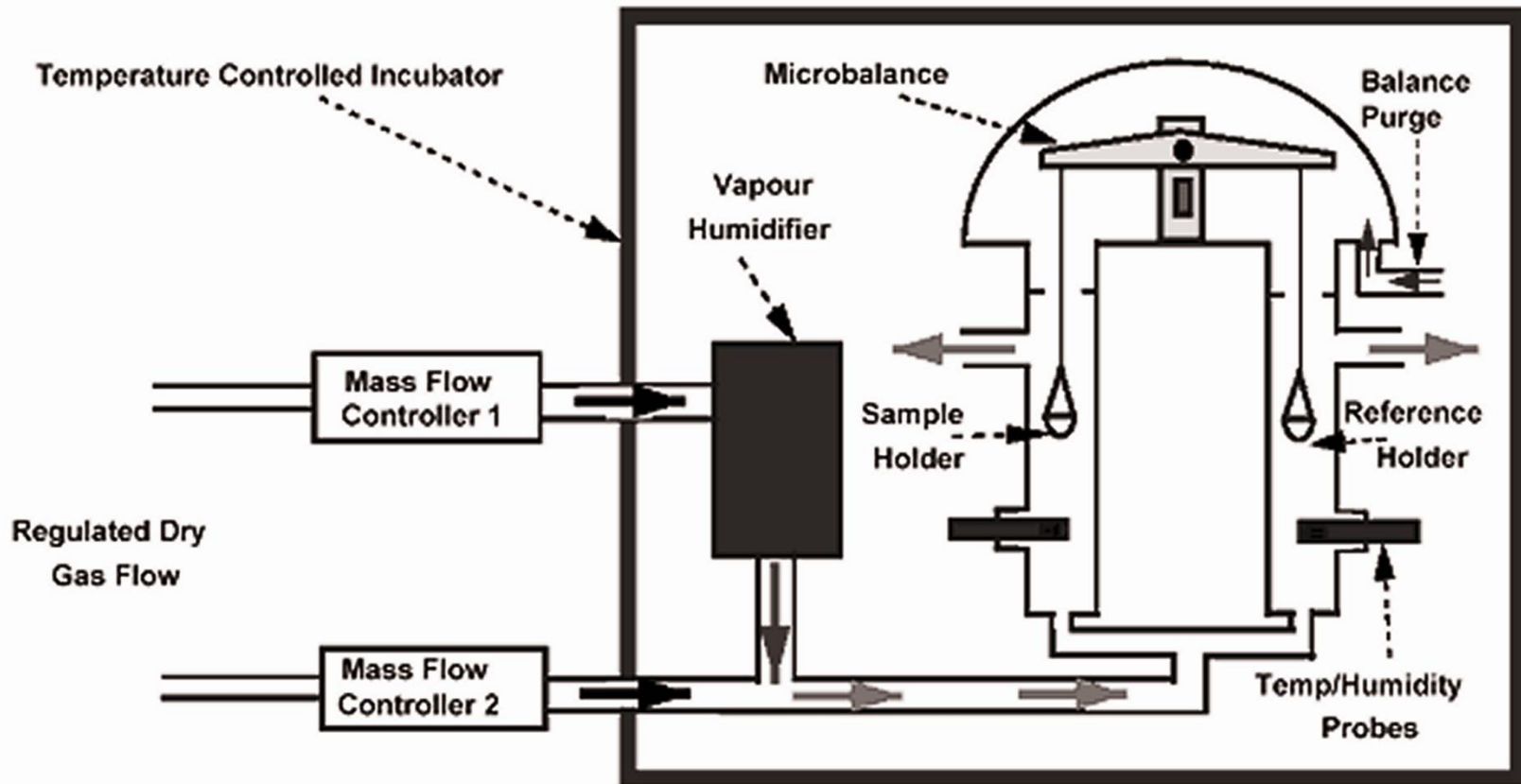
³Waters, Atlas Park, Simonsway, Manchester M22 5PP, United Kingdom

Correspondence to: B. Keating (E-mail: barbara.keating@strath.ac.uk)

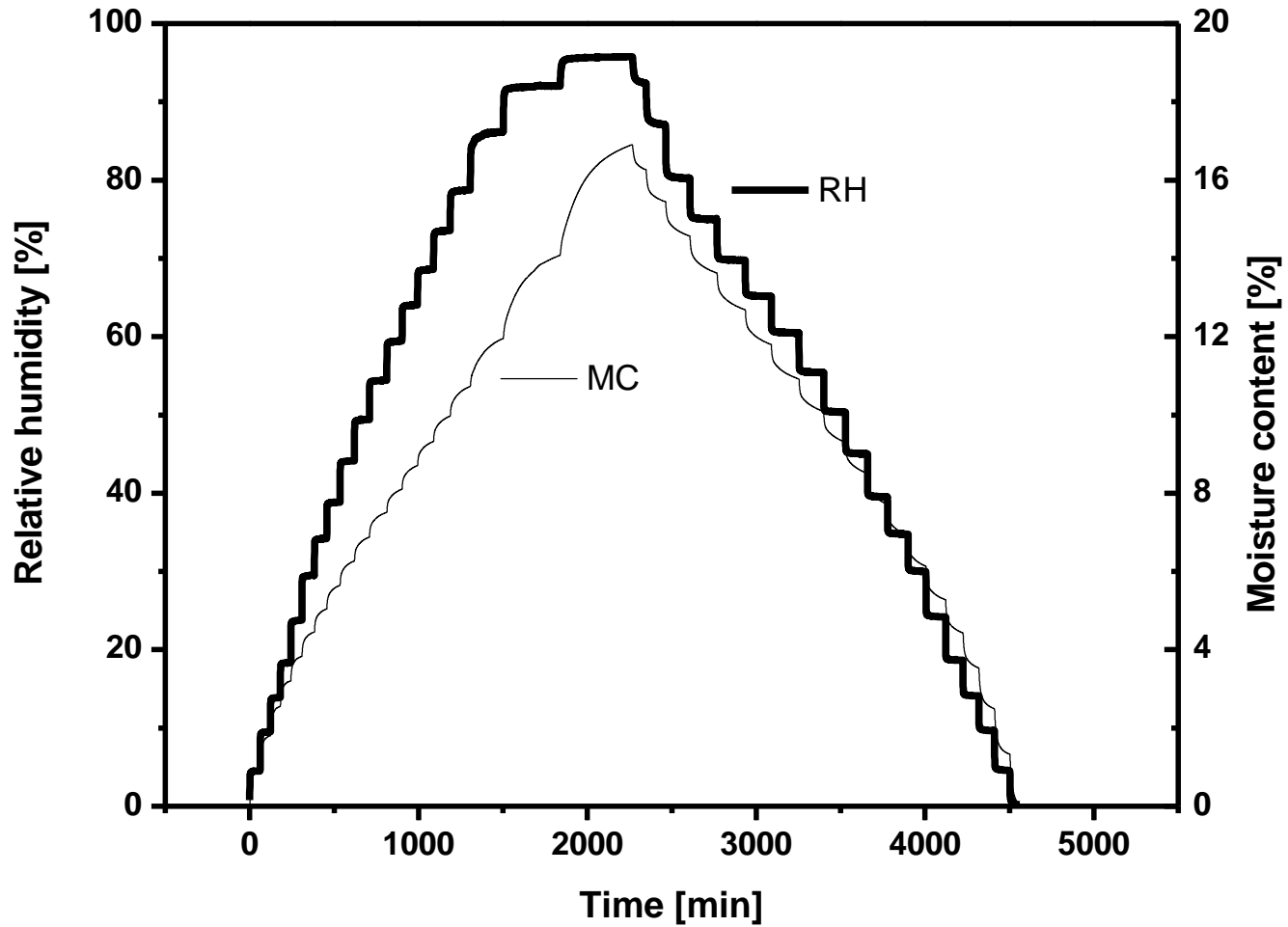




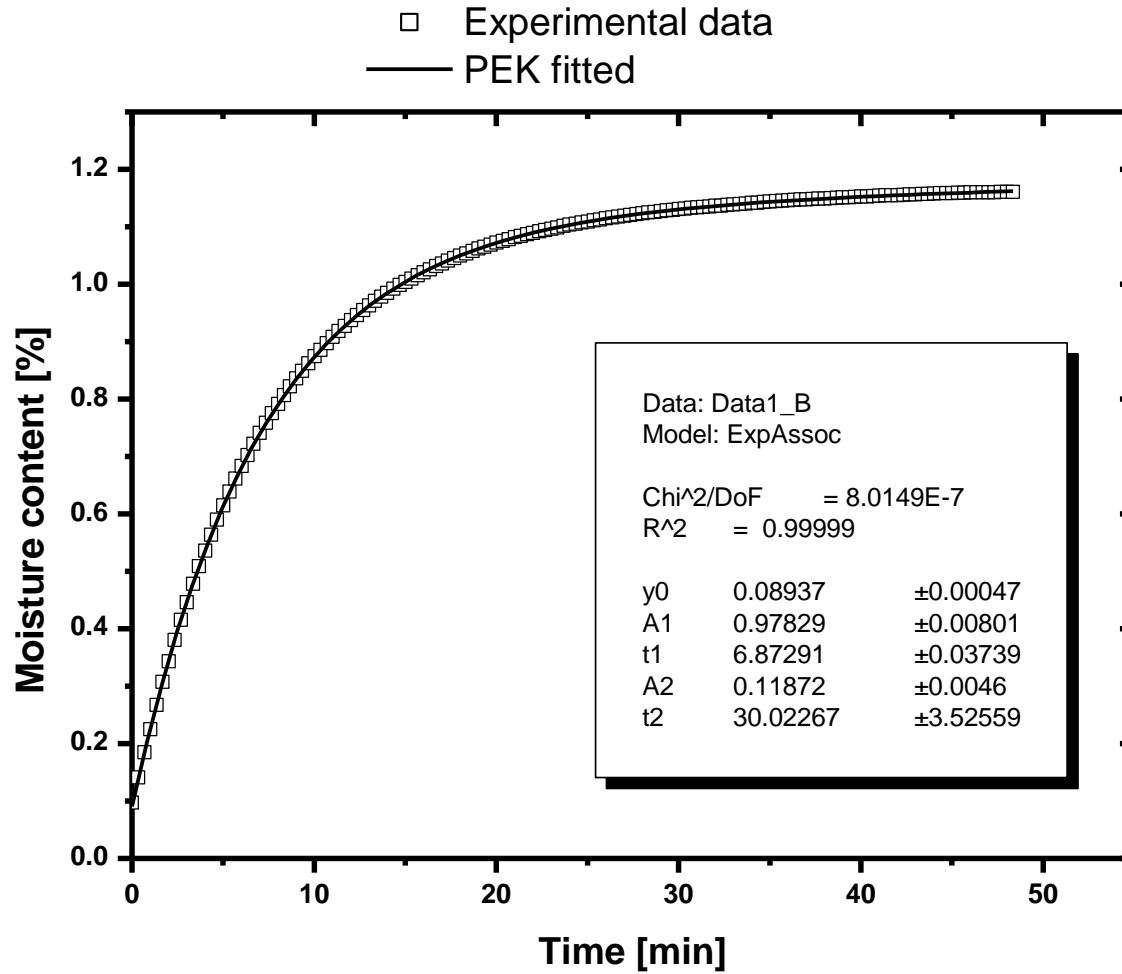
DVS APPARATUS



DVS RUN



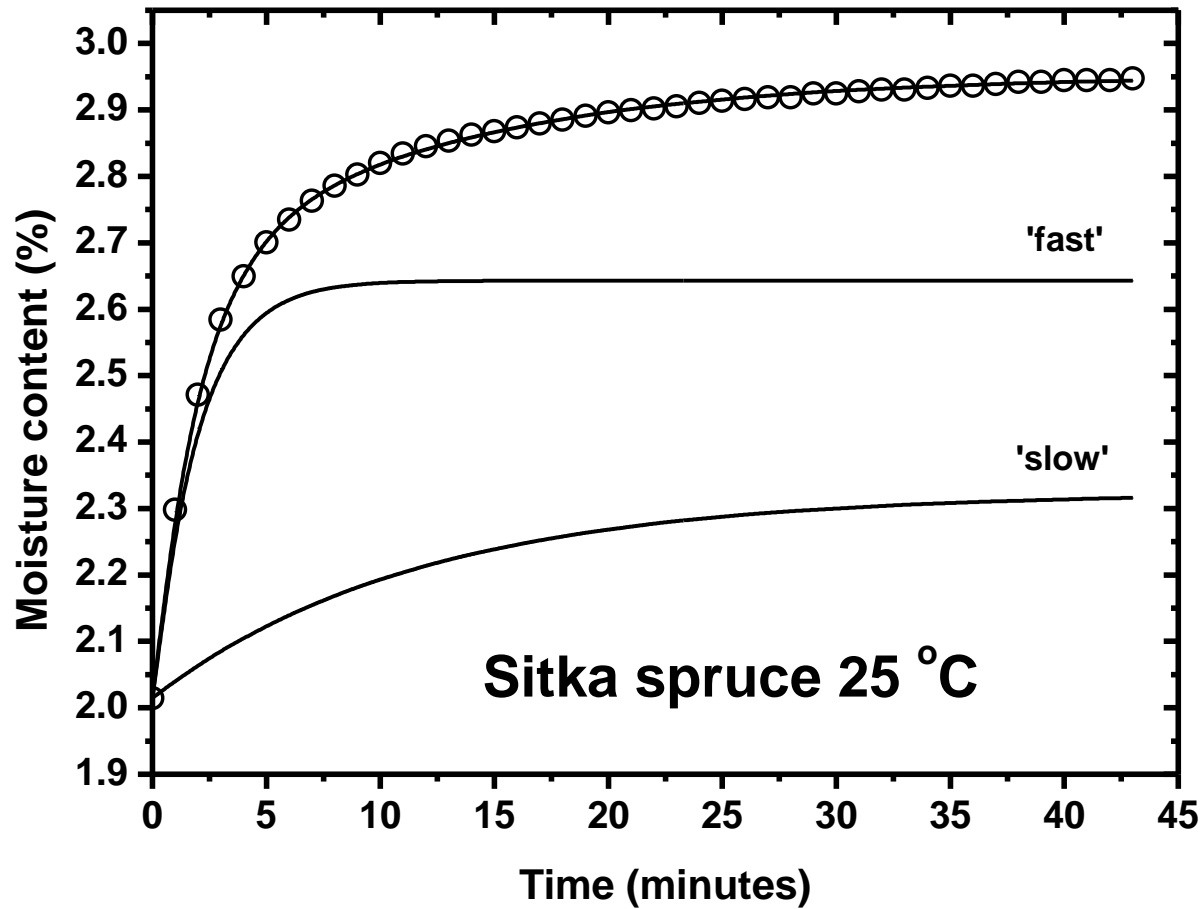
SORPTION KINETICS



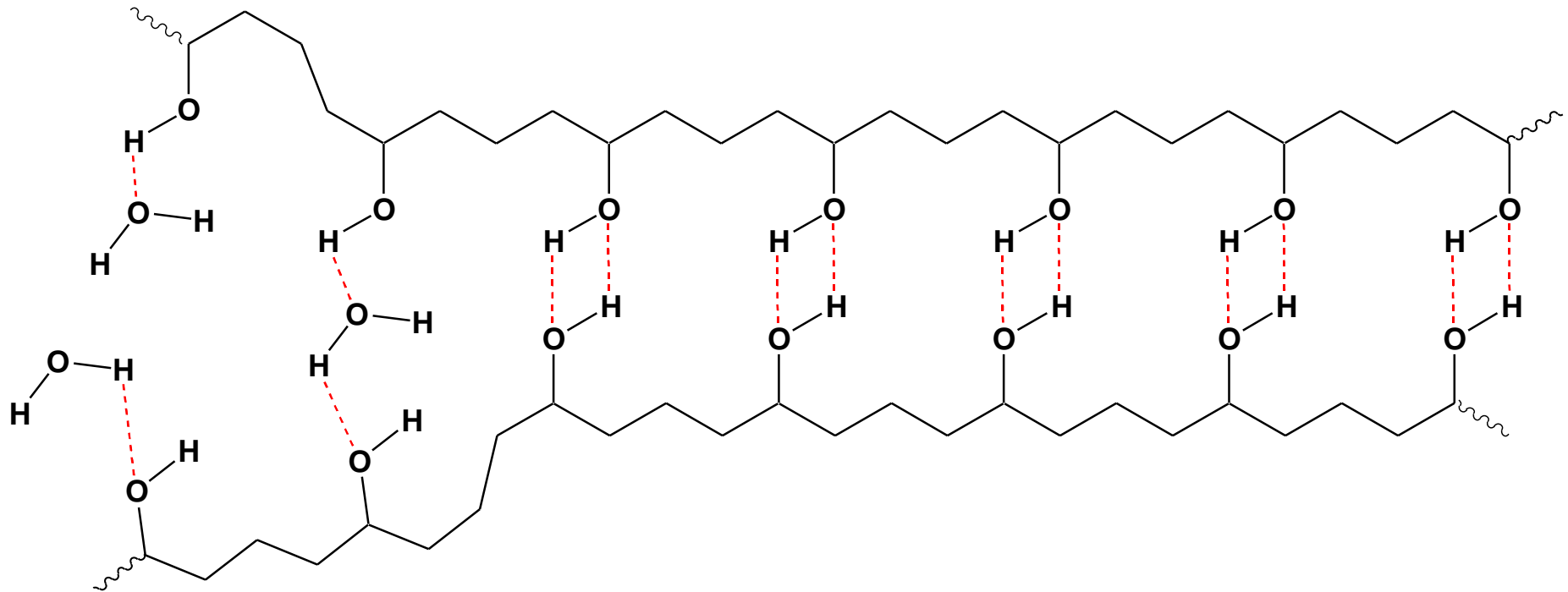
SORPTION KINETICS - PEK MODEL

- $MC = MC_0 + MC_1(1 - e^{-t/t_1}) + MC_2(1 - e^{-t/t_2})$

PEK MODEL



KINETICS

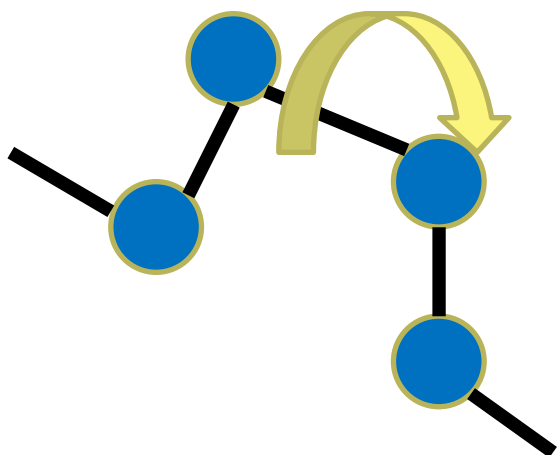


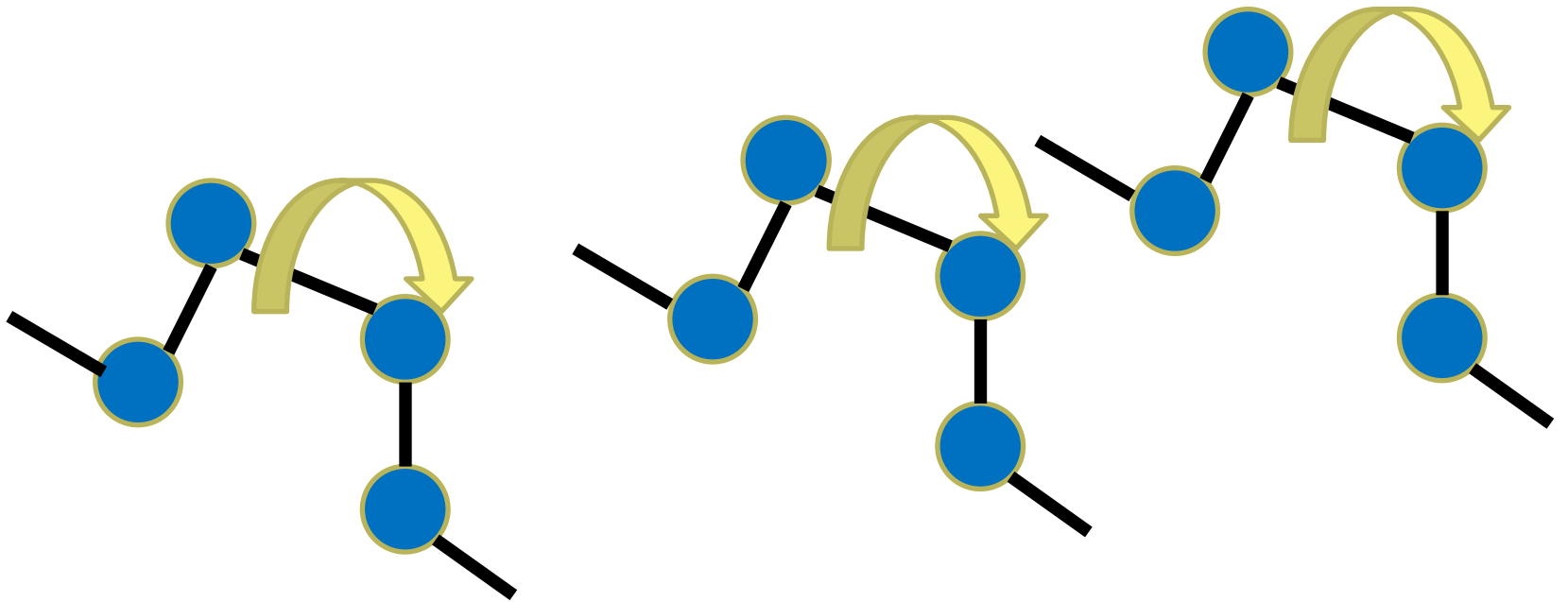
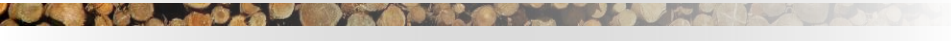


Cooperative Relaxation Processes in Polymers

J. Appl. Polym. Sci., 64, 77, (1997)

SHIRO MATSUOKA,¹ ARTURO HALE²



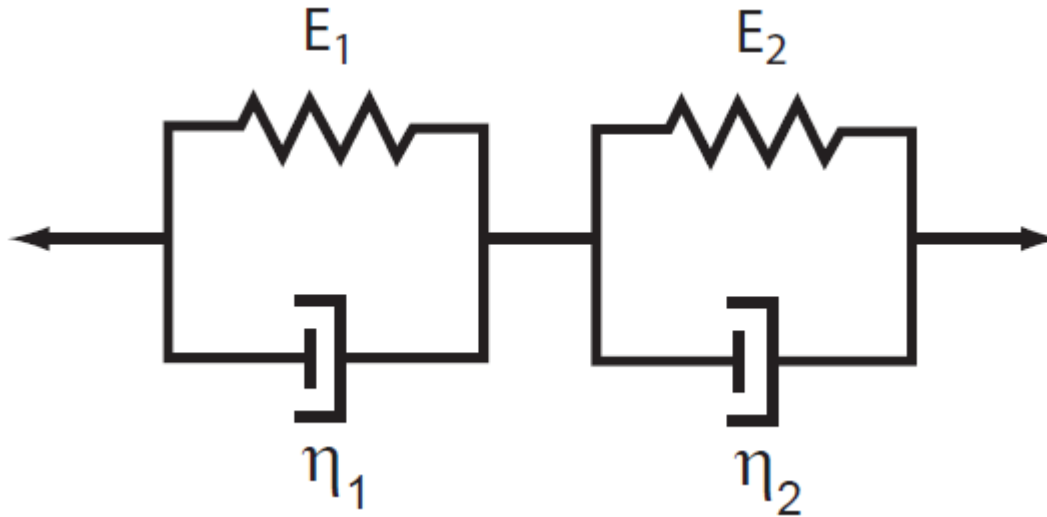


KELVIN-VOIGT



- Mechanical interpretation using Kelvin-Voigt viscoelastic model
- Rate limiting step of sorption process is polymer relaxation

KELVIN-VOIGT MODEL



$$\varepsilon = (\sigma_0 / E) [1 - \exp(-t/\varphi)]$$



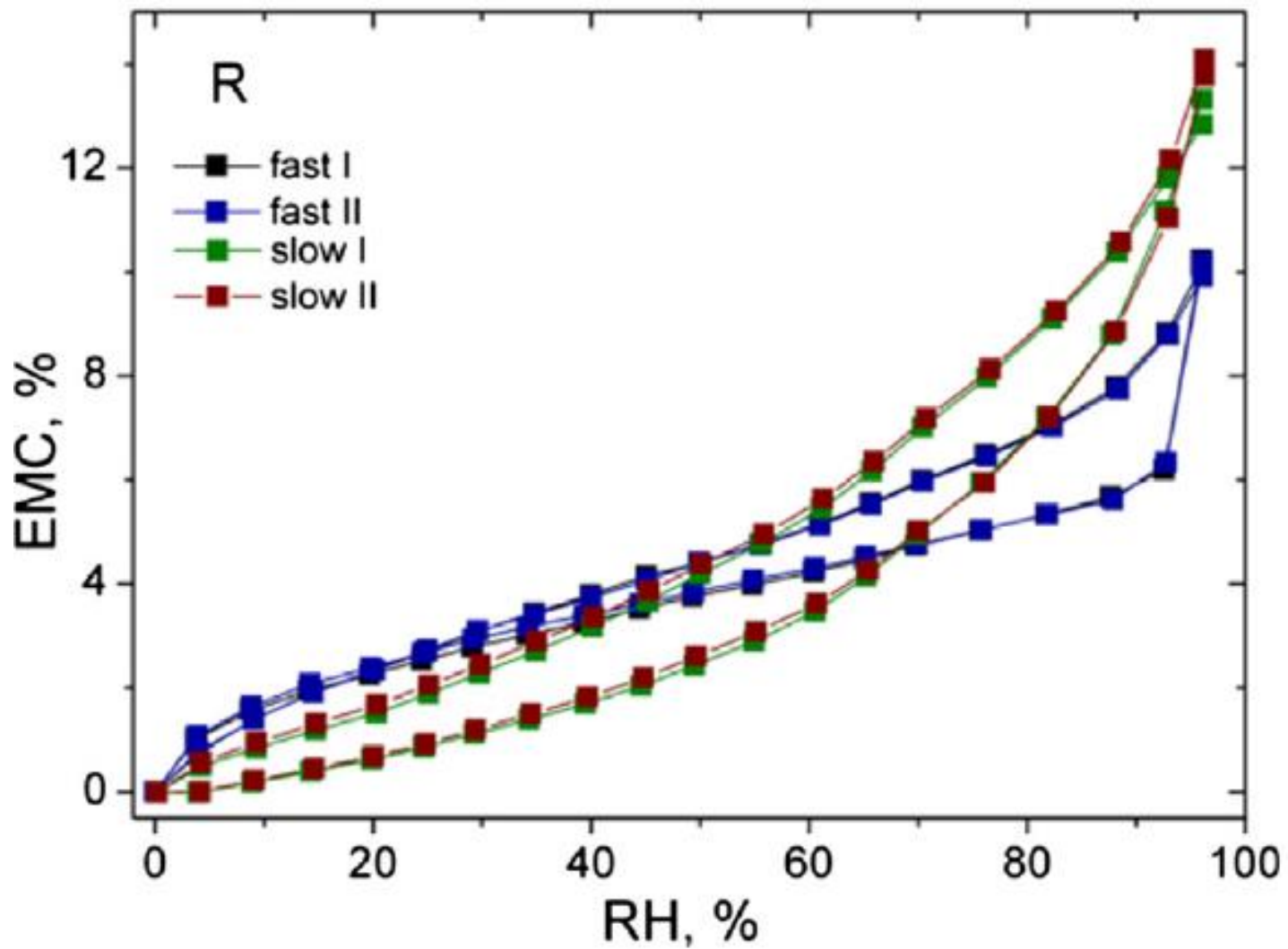
The water vapour adsorption–desorption behaviour of naturally aged *Tilia cordata* Mill. wood

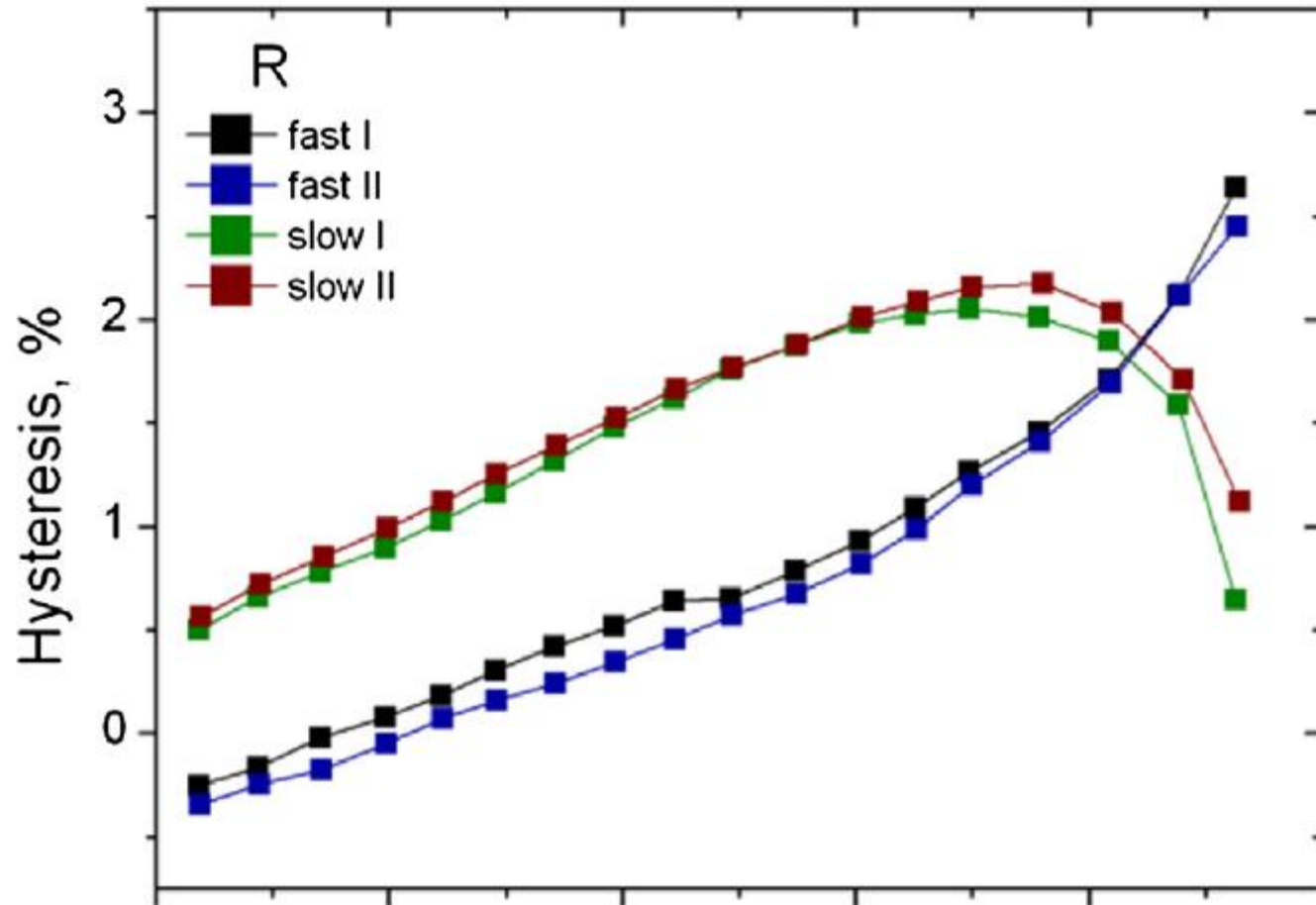
Carmen-Mihaela Popescu^{a,*}, Callum A.S. Hill^b

^a*“Petru Poni” Institute of Macromolecular Chemistry of Romanian Academy, 700487 Iasi, Romania*

^b*Forest Products Research Institute, Edinburgh Research Partnership Joint Research Institute in Civil and Environmental Engineering, Edinburgh Napier University, Edinburgh EH10 5DT, UK*

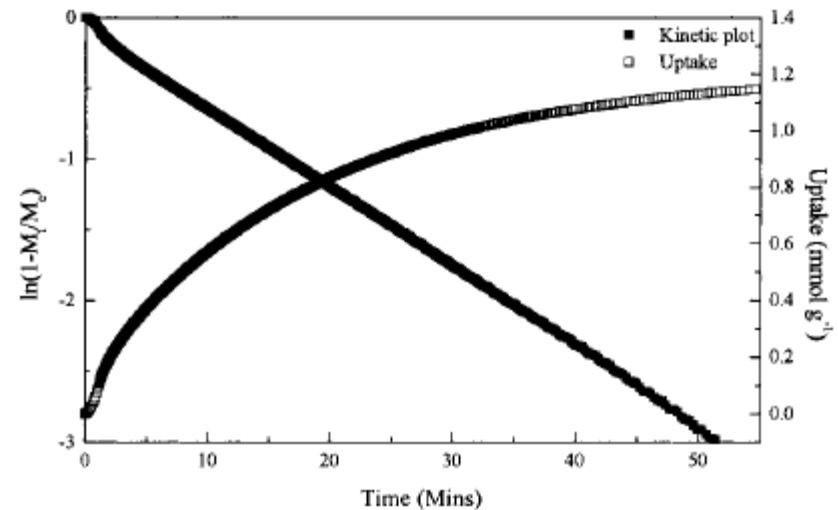
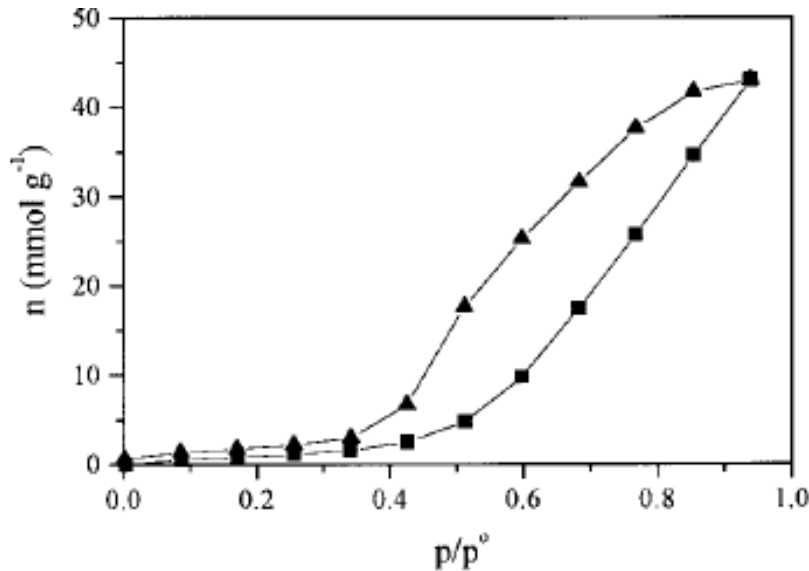
Polymer Degradation and Stability 98 (2013) 1804–1813





Diffusion Barriers in the Kinetics of Water Vapor Adsorption/Desorption on Activated Carbons

A. W. Harding, N. J. Foley, P. R. Norman,[†] D. C. Francis,[†] and K. M. Thomas*



3.2. Adsorption Kinetics. The kinetics of water vapor adsorption on carbon C1 have been shown to follow a linear driving force mass transfer (LDF) model.⁹

$$M_t/M_e = 1 - e^{-kt}$$

CONCLUSIONS



- A 'mechanical' interpretation gives results that are reasonable
- Fast process is probably diffusion-limited and slow process relaxation-limited
- Still to be investigated is the link between sorption hysteresis and kinetics
- Sorption is controlled by energy criteria
- So what is the role of OH groups?