

DERIVING A PROCESS VISCOSITY FOR COMPLEX PARTICULATE NANOFIBRILLAR CELLULOSE GEL-CONTAINING SUSPENSIONS

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ABSTRACT:

Phase-separable particulate-containing gel structures constitute complex fluids. In many cases they may incorporate component concentration inhomogeneities within the ensemble matrix. When formulated into high consistency suspensions, these can lead to unpredictable time-dependent variations in rheological response, particularly under shear in simple parallel plate and cylindrical rotational geometries. Smoothing function algorithms are primarily designed to cope with random noise. In the case studied here, namely nanocellulose-based high consistency aqueous suspensions, the system is not randomised but based on a series of parallel and serial spatial and time related mechanisms. These include: phase separation, wall slip, stress relaxation, breakdown of elastic structure and inhomogeneous time-dependent and induced structure rebuild. When vacuum dewatering is applied to such a suspension while under shear, all these effects are accompanied by the development of an uneven solid content gradient within the sample, which further adds to transitional phenomena in the recorded rheological data due to spatial and temporal differences in yield stress distribution. Although these phenomena are strictly speaking not noise, it is nevertheless necessary to apply relevant data smoothing in order to extract apparent/process viscosity parameters in respect to averaging across the structural ensemble. The control parameters in the measurement of the rheological properties, to which smoothing is applied, are focused on parallel plate gap, surface geometry, shear rate, oscillation frequency and strain variation, and relaxation time between successive applications of strain. The smoothing algorithm follows the Tikhonov regularisation procedure.

KEY WORDS:

data smoothing, rheology of gel suspension, dewatering, immobilisation, nanocellulose, phase separable process

1 INTRODUCTION

Gel structures may exhibit very different rheological behaviour depending on the mechanism of gelation. For example, bridging structures between particles may undergo viscoelastic response to strain in respect to the flocculating mechanism [1, 2]. In some gels, for example those based on mineral nanoclay particles, the gelation may relate to a direct swelling of the particles due to intercalate exchange and exfoliation [3, 4]. Polymer gels, on the other hand develop a structural integrity due to liquid uptake potential and diffusion between the constituent polymer molecules in the matrix [5]. An example material which is undergoing intensive research currently is nanocellulose and nanofibrillar cellulose, derived from the refining and/or oxidation of tree fibres [6–9]. This highly charged hydrophilic material behaves similarly to a superabsorbing polymer, in that the nano-

particulate matter itself does not swell, but the attraction to water results in a high osmotic pressure leading to bound and interstitial water molecules clustered within a swollen gel [10, 11]. This behaviour, when introduced in mixes with other suspension particles, including, for example, micro and macroscopic cellulose fibres and inert pigment fillers, results in challenging rheological properties when considered for applications in a wide range of manufacturing processes, ranging from paper and board manufacture, as well as composites, to food, pharmaceuticals and cosmetics [12–16]. Such materials form phase-separable gel structures, which may incorporate inhomogeneities within the ensemble matrix when incorporated in high consistency suspensions, and so constitute complex fluids, whose rheological response can lead to unpredictable time-dependent variations under shear in simple parallel plate and cylindrical rotational geometries [17–19].

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