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Temperature and Concentration Dependent Viscosity of Microcrystalline Cellulose in Water

Wan Nor Suhaila Wan Aziz^{1,2}, Shahrul Kadri Ayop^{2,*}, Rosazley Ramly²

¹ Unit Kawalselia Radiasi Perubatan, Pahang State Health Department, Jalan IM/4, Bandar Indera Mahkota, 25582, Kuantan, Pahang, Malaysia
² The Department of Physics, Faculty of Science and Mathematics, Sultan Idris Education University, 35900 Tanjong Malim, Perak, Malaysia

ARTICLE INFO	ABSTRACT
Article history: Received 25 September 2020 Received in revised form 13 January 2021 Accepted 15 January 2021 Available online 15 February 2021	The viscosity of cellulose behaves differently and uniquely in various conditions. In this paper, we aim to report the viscosity measurement and related properties of low concentration of microcrystalline cellulose (MCC) in water using a magnetic bearing rheometer. Dynamic viscosities for MCC diluted in water at varying concentrations were measured using the standard rheometry technique. The viscosity of the MCC solution was found highly dependent on its concentration and the experiment temperature. This varieties behaviour and properties offers benefits to the current growing rapidly technology applications such as in food, pharmaceutical cosmetics and textile.
Keywords:	
Viscosity; magnetic bearing rheometer; cellulose; microcrystalline	

1. Introduction

Microcrystalline cellulose (MCC) is widely used in various industrial fields such as medicine, cosmetics, pharmaceuticals and polymer composite [1-7]. Recently due to its novelty, non-toxicity properties, economic value, biodegradation, mechanical properties, surface area and biocompatibility, the interest on MCC have increased [1-7]. For example, the properties of hydrogels facilitate their usage in bio-related applications, including drug delivery systems, tissue-engineering scaffolds, wound dressing, and biomedical devices [8-12]. The imperfect mechanical properties of hydrogels would give limitations for applications that require high strength properties [8,13]. Therefore, cellulose can be act as a suitable biopolymer for synthesizing a hydrogel to have outstanding mechanical properties [8]. Due to the inherent crystalline structure, cellulose exhibits high strength, high stiffness, and low density [14]. It is potentially possible to synthesize the high-strength microcrystalline cellulose hydrogel by controlling the viscosity of cellulose solutions [8]. The information of viscosity of MCC is essential for a small scale to industrial scale in a diverse field in order to optimize the output quality and can affect directly to the final product [2].

* Corresponding author.

E-mail address: shahrul.kadri@fsmt.upsi.edu.my

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Furthermore, the estimation level of carbohydrate degradation caused by pulping and bleaching steps could use the viscosity information as a control parameter of cellulose pulp quality [15]. The resulting viscosity of a fully bleached cellulose pulp showed that substantially lower than unbleached cellulose pulp, which contributes to the lower paper strength properties [15,16]. The previous study has shown that less degraded bleached cellulose pulps that combined higher viscosities and residual lignin contents would have lower water absorption and vice versa as the determination of water absorption is essential information for packaging and printing industries [15].

The viscosity affects significantly the abundance of air-water with glycerin that clings to the inner walls of the pipe and higher concentration of glycerin in the liquid would increase the wave and stuck as reported in two-phase flow in mini pipe simulation [17]. The consideration of two-phase flow in mini pipe are important in design and development in industrial and medical applications, such as micro heat exchangers, micro cooling electronics and bio-micro-electro mechanical system [17]. In automobile and domestic application, currently the fossil diesel mixed with biodiesel through blending process before any usage. The biodiesel density and viscosity are the influence factors for diesel biodiesel mixture properties [18]. Moreover, it also found that the density and viscosity value are slightly increase due to the biodiesel content and the viscosity increasing proportional in trend to biodiesel composition [18,19]. The initiative is aim to reduce the environment damage from the carbon dioxide produced.

Microcrystalline cellulose (MCC) has a limitation that is needed for some applications compared to other natural polysaccharide polymers such as low wettability, moisture absorption, and limitation in processing temperature [1]. The previous research related to cellulose filled engineering thermoplastics, scientist believed that the use of cellulose as a reinforcement or additive did not give encouraging results as the materials exhibited severe discoloration and cellulose thermal degradation occurs at temperatures needed to process these engineering thermoplastics [20]. However, recently researchers have looked again at cellulose filled engineering thermoplastic suggested that MCC-filled nylon composites relevant in thermally challenging areas due to the higher mechanical properties and lower density such as engine covers, intake manifolds and radiator end tanks [21]. It is also essential to know and observe the temperature dependence due to the limitation of MCC in processing temperature in order to optimize the quality of final product and cost-effectiveness in variety of industrial process [1]. The viscosity value expects to decrease due to the increasing temperature for all liquids. However, modifications of viscosity value are essential as it could influence the operational cost of several stages in the industrial process such as mixing and fluid transport [22].

This study aimed to measure viscosity and observe the related properties of low concentration and temperature of microcrystalline cellulose (MCC) in water. Findings from this study could be fully utilized as a ground comparison in measurement technique between the rheometer and optical trapping technique on rheological properties of low concentration of a polymeric solution. Optical trapping technique offers possible measurement on such a solution at very low sample volume consumption. The previous study had been discussed about the potential of optical tweezer for microrheology measurement of variety polymeric solution [23]. It also highlights the advantages of optical trapping application as compared with the conventional method [23].

2. Methodology

The microcrystalline cellulose MCC in powder form (Sigma-Aldrich[®], Product No.310697) was dissolved in deionised water (DI water) with 4 different concentration; 0% w/w (DI water), 1% w/w,



3% w/w, 5% w/w and 10% w/w as Figure 1. The MCC solution was sonicated using a sonicator (Q sonica) and bathed using bath sonicator (Branson 2800) for process duration of 60 minutes. Sonication process is important in order to achieve a uniform condition of MCC solution [24].

Viscosity measurement in this study was conducted using the rheometer type AR-G2 series (TA Instrument) and 60 mm 1° steel cone type of geometry (code: 992176). According to the manufacturer recommendation, the gap between geometry and Peltier plate, also known as truncation height, should be more or equal to tenth times particle sizes, as illustrated in Figure 2. In this study, the microcrystalline cellulose is 20 μ m in length, and the gap has been set up consistently for 200 μ m.



Fig. 1. Microcrystalline cellulose (MCC) diluted in deionised water with concentration of 0% w/w, 1% w/w, 3% w/w, 5% w/w and 10% w/w

The rheometer is used to monitor the displacement in time and converts it to shear rate. This rheometer is connected to the water circular in order to control the sample temperature by setting the required temperature from the AR Instrument software. In order to have a right bearing of the moving geometry and to prevent friction, the installed geometry should go through the mapping and calibration process before taking any measurement [24].



Fig. 2. The truncation height between cone type geometry and Peltier plate. Appropriate sample filling is advisable, avoid under or overfilling in order to have the correct measurement

The temperature of the sample condition was varied from 20 °C to 30 °C, whose parameters of interest were being measured. A minimum of 1 ml sample was required for each measurement and the measurement time was 120 seconds for each sample then repeated three times using a new sample. The deionized water was chosen as a benchmark to ensure the magnetic bearing rheometer worked appropriately during the measurement [25]. The viscosity measurement was



validated using water as water has a well-known viscosity value then proceeded to the MCC sample. This validation aims to minimize fluctuation and reading errors.

3. Results

3.1 Concentration Dependent

The viscosities of the MCC solutions were measured at different concentrations and temperatures. It is observed that the rheological behaviour is highly influenced by the concentration of microcrystalline cellulose [26,27]. This viscosity value increased in trend to the concentration, as shown in Figure 3.



Fig. 3. The viscosity of microcrystalline cellulose solution at 0, 1, 3, 5 and 10% w/w as measured using a rheometer

Deionized water itself has a very low viscosity. However, an addition of a small amount of microcrystalline cellulose, which was well dispersed in the deionized water showed a logical trend even though cellulose hardly dissolved in water or oil. This indicated that the sonication process of MCC solution achieves uniform condition. A slight change in the viscosity value of MCC solution can be observed at a concentration of 1 to 3% w/w. However, it drastically increased by 3% w/w and upward. According to the literature on the preparation, structure, and properties of microcrystalline cellulose, the aqueous suspension of a particular concentration of MCC could form a gel under the effect of shear force [28]. Besides, it also stated that the viscosity of MCC gel increased with increasing content of MCC in water and sharply increase of viscosity occurred in the 3-6% w/w concentration range [28]. This phenomenon showed that the internal resistance of the solution has increased as the solution more viscous [29].

3.2 Temperature Dependent

Findings for each concentration amount of MCC diluted in deionized water; the viscosity decreases as the temperature increased is illustrated in Figure 4. Even though the range of temperature varying is not too wide, but the effect can be seen as when the temperature increased



the viscosity of MCC solution becomes decreased. The phenomenon is due to the MCC having higher thermal energy, and the binding forces are more easily able to weaken the attractive forces between them. According to the previous literature on the investigation of temperature-dependent structural changes in hydrogen bonds (H-bonds) in microcrystalline cellulose (MCC) by infrared (IR) and near-infrared (NIR) spectroscopy suggest that structural changes in the H-bonds in the O_3 – H_3 ^{...} O_5 intrachain H-bonds in MCC gradually occur in the temperature region of 25° C to 130° C, and it becomes continuously greater above 130° C [30].



Fig. 4. Variations of viscosity with MCC concentration at different temperature

The viscosity values of MCC solution varying with concentration and temperature is in Table 1. The viscosity value summarised in the table is an average from the three times repeated data in cP unit while the temperature is represented in Celsius and MCC concentration in percentage.

Table 1							
The viscosity values of MCC solution at different temperatures and concentrations of MCC							
Temperature, T (°C)	Viscosity (cP) Value for concentration, % (w/w)						
	0	1	3	5	10		
20.7	0.8953	1.1093	1.1327	1.2397	1.4550		
21.6	0.8767	0.9765	0.9994	1.1107	1.3070		
22.6	0.8686	0.9305	0.9565	1.0673	1.2503		
23.6	0.8636	0.9106	0.9370	1.0470	1.2190		
24.5	0.8598	0.8987	0.9254	1.0327	1.1973		
25.4	0.8563	0.8899	0.9164	1.0213	1.1790		
26.3	0.8501	0.8820	0.9084	1.0107	1.1630		
27.1	0.8499	0.8745	0.9008	1.0008	1.1480		
28.1	0.8468	0.8677	0.8936	0.9914	1.1337		
29.0	0.8439	0.8609	0.8867	0.9822	1.1203		
30.0	0.8410	0.8545	0.8801	0.9734	1.1077		



4. Conclusions

The viscosity behaviour of microcrystalline cellulose (MCC) solution is highly dependent on the concentration and temperature. The viscosity was found to be drastically increased when the concentration was increased. However, the viscosity value decreased when the sample temperature was increased and trends differently as concentration before. In conclusion, the viscosity value is highly dependent on sample concentration and temperature within the described experimental conditions.

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