

Journal Pre-proof

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PII: S2238-7854(23)00694-4

DOI: <https://doi.org/10.1016/j.jmrt.2023.03.231>

Reference: JMRTEC 6975

To appear in: *Journal of Materials Research and Technology*

Received Date: 5 December 2022

Revised Date: 23 March 2023

Accepted Date: 31 March 2023

Please cite this article as: Daskalakis E, Hassan MH, Omar AM, Cooper G, Weightman A, Bartolo P, Rheological behaviour of different composite materials for additive manufacturing of 3D bone scaffolds, *Journal of Materials Research and Technology*, <https://doi.org/10.1016/j.jmrt.2023.03.231>.

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Rheological behaviour of different composite materials for additive manufacturing of 3D bone scaffolds

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Abstract: The production of scaffolds for bone tissue applications is requiring a combination of physical and biological properties, which are depending on the materials morphology and pro-cessing conditions during the production process. The aim of the paper is the investigation of rheological behaviour of polymer and composite blends regularly used for the production of scaffolds for bone tissue applications with the use of additive manufacturing. Poly-ε-caprolactone (PCL), hydroxyapatite (HA), β-tri-calcium phosphate (TCP) and Bioglass 45S5 blends containing different ceramic concentrations (10wt%, 15wt% and 20wt%) were prepared with the use of melt blending procedure and investigated with the use of oscillation and rotational rheology tests. Results are showing that all blends are presenting viscoelastic behaviour with higher viscous modulus, compared with elastic modulus for low frequencies, with this difference reducing while the frequency is increasing. All blends are presenting shear-thinning behaviour suitable for use with additive manufacturing methods. Viscous and elastic modulus are increasing by adding ceramic particles. Results are presenting that PCL/HA blends of the same material concentration are presenting higher elastic modulus properties compared with the other blends, while PCL/Bioglass blends are presenting lower loss factor, lower relaxation time and lower shear viscosity making them easier to handle during the printing procedure.

Keywords: Additive manufacturing, Polymer-ceramic blends, Printability, Rheology

1. Introduction

Additive manufactured scaffolds represent the most suitable approach for tissue engineering applications [1-6]. Different additive manufacturing techniques such as vat-photopolymerization, binder jetting, powder bed fusion and extrusion-based systems have been used to create scaffolds with well-defined architecture, pore shape, pore size, pore distribution,

pore interconnectivity and porosity [7-14]. Among these techniques extrusion-based processes are the most commonly used additive manufacturing for the fabrication of bone scaffolds [15-19]. Extrusion-based additive manufacturing is based on melting a polymer-based material and printing the molten material in a moveable platform where the material will cool down. During this heating a cooling process there is an inherent morphological development process that will determine the final properties of the printed scaffolds [20-26]. The kinetics of the cooling process also determine the crystalline structure formation (e.g., degree of crystallinity and crystal size) and volumetric changes of the printed filaments [27-31]. The heating process, which depends on the material to be printed, determines the viscosity of the printed material, how easy is the material printed and the shape fidelity of the printed part upon cooling. To guarantee good printability, materials must present a shear thinning behaviour that describes a material in which the viscosity decreases by increasing the shear rate [31-36].

Materials for extrusion-based additive manufacturing can be in the form of pellets, processed through screw-assisted extrusion systems, or in the form of filaments, processed through filament-based extrusion systems also known as Fused Deposition Modelling (FDM) systems [37, 38]. Most studies on bone scaffolds are based on the use of screw-assisted extrusion additive manufacturing [39, 40].

Scaffolds can be made with bioceramic materials, but such scaffolds are usually brittle, or using biodegradable and biocompatible polymers, but usually these scaffolds present limited bioactivity [41-43] 412-414]. These limitations can be addressed through the use of polymer/ceramic composite blends. PCL is one of the most commonly used polymeric material for bone tissue engineering applications [44, 45] 415, 416]. It is biocompatible, biodegradable, easy to process, but presents limited mechanical properties, low bioactivity and long degradation times [46, 47]. To overcome these limitations, PCL is usually combined with bioactive ceramics such as HA, TCP and Bioglass [48-52]. These reinforcements have the potential to increase the overall mechanical properties of the 3D printed composite scaffolds and also cell attachment, proliferation and differentiation. However, the addition of bioceramic particles have also a strong effect on the rheological properties of the blends compromising their printability and eventually clogging the printing head.

This Chapter investigates the effect of different reinforcements (hydroxyapatite, β -tricalcium phosphate and Bioglass 45S5) and level of reinforcement (0wt%, 10wt%, 15wt% and 20wt%) on the rheological and viscoelastic properties of the materials.

2. Materials and Methods

2.1 Materials and Methods

Poly- ϵ -caprolactone (PCL) (melting point = 60 °C, glass transition temperature = -60 °C, molecular weight = 50000Da, CAPA 6500, Perstorp Caprolactones, Cheshire, UK) in pellet shape. Hydroxyapatite (HA) (molecular weight = 502.31 gr/mol, melting point = 1100 °C, Sigma-Aldrich, St. Louis, USA) in nanopowder shape (< 20 nm particle size), β -tricalcium phosphate (TCP) (molecular weight = 310.18 gr/mol, melting point = 1391 °C, Sigma-Aldrich, St. Louis, USA) in powder shape (ranging between 20 μ m to 30 μ m) and Bioglass 45S5 (6wt% P₂O₅, 45wt% SiO₂, 24.5wt% Na₂O and 24.5wt% CaO, CeraDynamics Ltd. James Kent Group, Stoke, UK), in powder shape (< 10 μ m particles size) (Table 1).

Table 1. Composition of the different material blends (values are in wt%).

Material/Configuration	PCL	PCL/HA	PCL/TCP	PCL/Bioglass
0wt%	100	-	-	-
10wt%	-	90/10	90/10	90/10
15wt%	-	85/15	85/15	85/15
20wt%	-	80/20	80/20	80/20

2.2 Morphological analysis

Scanning electron microscopy (SEM) images were taken using FEI Quanta 200 (FEI ESEM Quanta 200, FEI Company, Hillsboro, Orlando, USA) to investigate the morphology of HA, TCP and Bioglass 45S5 particles in powder form. EMITECH K550X sputter coater (Quorum Technologies, Laughton, East Sussex, UK) was used to coat the powder particles (platinum coating).

2.3 Rheology analysis

Rheological analysis including oscillation and rotational studies were investigated with the use of DHR-3 TA Instrument (TA Instruments, New Castle, UK). The materials were tested using a 20 mm (diameter) steel parallel plate with a gap distance of 450 μ m. Three samples of each blend were used, and three different temperatures were considered for the rheology tests: 80 °C, 90 °C and 100 °C, which are within a typical printing temperature range for screw-assisted extrusion-based 3D printing of PCL scaffolds. To simulate the molten state during the printing process, the materials were preheated for 15 min. For the oscillation test a constant strain rate of 1% was considered, while the frequency of logarithmic sweep was between 10 HZ and 0.01

Hz. For the rotational test the shear rate of the logarithmic sweep was between 0.01s^{-1} and 100s^{-1} . The results presented in this research are the average values of the three samples of each blend.

The measurement of the storage modulus (G') and loss modulus (G''), which describes the elastic and viscous behaviour of the material respectively, allows to evaluate the viscoelastic characteristics of the considered blends. Storage modulus and loss modulus are given by the following equations [53]:

$$G' = (\sigma_o/\gamma_o)\cos\delta \quad (1)$$

$$G'' = (\sigma_o/\gamma_o)\sin\delta \quad (2)$$

, where σ_o is the stress amplitude, γ_o is the strain amplitude and δ is the phase angle between 0° to 90° .

To investigate the average the ratio of energy dissipated to the maximum energy stored it is important to determine the ratio of the average power loss to the peak load loss by calculating the loss factor ($\tan\delta$) [53]:

$$\tan\delta = G'/G'' \quad (3)$$

Accelerated degradation studies were conducted using sodium hydroxide (NaOH) of 5 mol/l (5 N) in aqueous solution (VWR, Pennsylvania, USA), with a density of 1.185 gr/cm^3 (20°C), solubility of 20°C and 14 pH (H_2O , 20°C). The degradation period took place for 5 days. On each day, 5 samples were used from each considered case and measured using a high precision balance. At each time point, the samples were removed from the NaOH and washed three times with the use of deionised water and left to dry overnight. Once completely dried the samples were measured to determine the weight reduction. The amount of NaOH used for 5 rectangular scaffolds was 15ml and for 5 the anatomically designed bone bricks was 50 ml (due to their size). The pH was monitoring throughout the experimental work and no changes were observed (pH of 14).

$$\tau = \eta \dot{\gamma}^n \quad (4)$$

, where τ is the shear stress (Pa), η is the flow consistency or coefficient (Pa.s), " $\dot{\gamma}$ " is the shear rate (s^{-1}) and n is the power law index that defines the material behaviour [54]:

$n < 1$: shear-thinning material

$n = 1$: Newtonian material

$n > 1$: shear-thickening material.

Moreover, to investigate the homogeneity dispersion of the materials in the blends, the Cole-Cole plot was used considering both the imaginary viscosity and the real viscosity [54]:

$$\eta'' = G'/w \quad (5)$$

$$\eta' = G''/w \quad (6)$$

, where η'' is the imaginary viscosity, η' the real viscosity and w is the oscillation frequency.

3. Results and discussion

3.1 Morphological analysis

Figure 1 presents the ceramic materials used for the production of PCL/HA, PCL/TCP and PCL/Bioglass blends for the rheological tests.

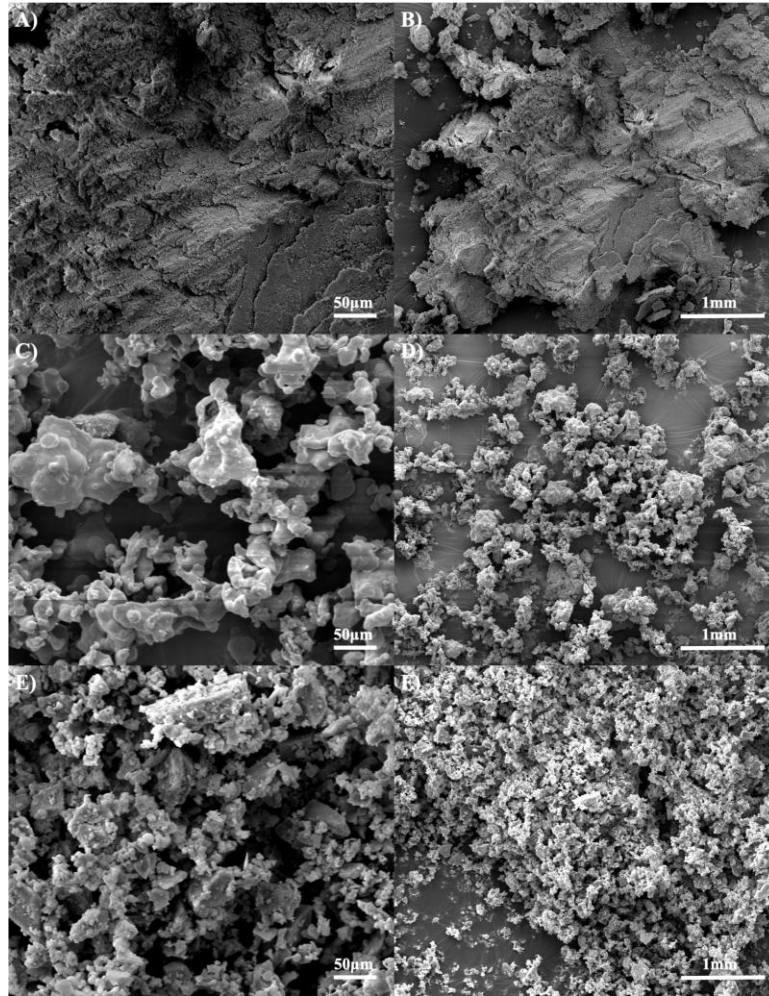


Figure 1. SEM images of A) and B) HA nanoparticles, C) and D) TCP microparticles and E) and F) Bioglass microparticles.

3.2 Oscillation analysis

Figure 2 shows both the storage and the loss modulus for all considered material compositions at different temperatures (80 °C, 90 °C and 100 °C) as a function of frequency. As observed, both storage modulus and loss modulus increase by increasing the frequency from 0.01 Hz to 10 Hz. This can be due to a sufficient relaxation time of polymer chains at low frequency allowing the self-disentanglement [55-64]. Moreover, it can be observed that, for all material compositions, the loss modulus is higher than the storage modulus, indicating that the viscous behaviour is the dominant effect within this temperature range, indicating that 3D printing is a suitable technique to process these blends. Additionally, by increasing the frequency it is possible to observe that the difference between storage modulus and loss modulus decreases, suggesting that composite materials present a more solid-like behaviour at a high deformation rate, which may hinder the flowability of the materials during the printing process. Furthermore, results show that the addition of ceramics into the PCL matrix decreases the storage modulus compared with pure PCL, while loss modulus does not show any significant differences. On the other hand, blends containing PCL/HA (80/20 wt%) and PCL/Bioglass (80/20 wt%) exhibit higher storage modulus in comparison to pure PCL for frequencies between 0.01 Hz to 0.1 Hz.

Figures 3 and 4 show the loss factor ($\tan\delta$) as a function of frequency for all blends at different temperatures. In all cases, results show that the loss factor decreases by increasing frequency indicating enhanced viscous behaviour or weakened elastic behaviour. Moreover, for all considered cases, it can be observed that the loss modulus decreases by increasing the deformation rate, which suggests that the elastic effect of the materials is becoming more important than their viscous effect, meaning that during the printing process the materials are becoming more viscous making them difficult to extrude through the nozzle. Furthermore, results show that for the same level of reinforcement the loss factor increases by increasing the temperature from 80 °C to 100 °C, as this temperature raise promotes a fluid-like behaviour making the materials easier to print. Additionally, for low frequencies and the same material content the loss factor decreases by increasing the material concentration, while by increasing the frequency rate this trend seems to disappear, suggesting that at high frequencies the material concentration effect is dominant compared to the deformation rate. For the same reinforcement concentration and temperature, PCL/TCP blends present higher loss factor value than PCL/HA and PCL/Bioglass blends, while PCL presents the higher loss factor. These results suggest that

for the same level of reinforcement and processing parameters, PCL/TCP blends can be easily printed than PCL/HA and PCL/Bioglass blends.

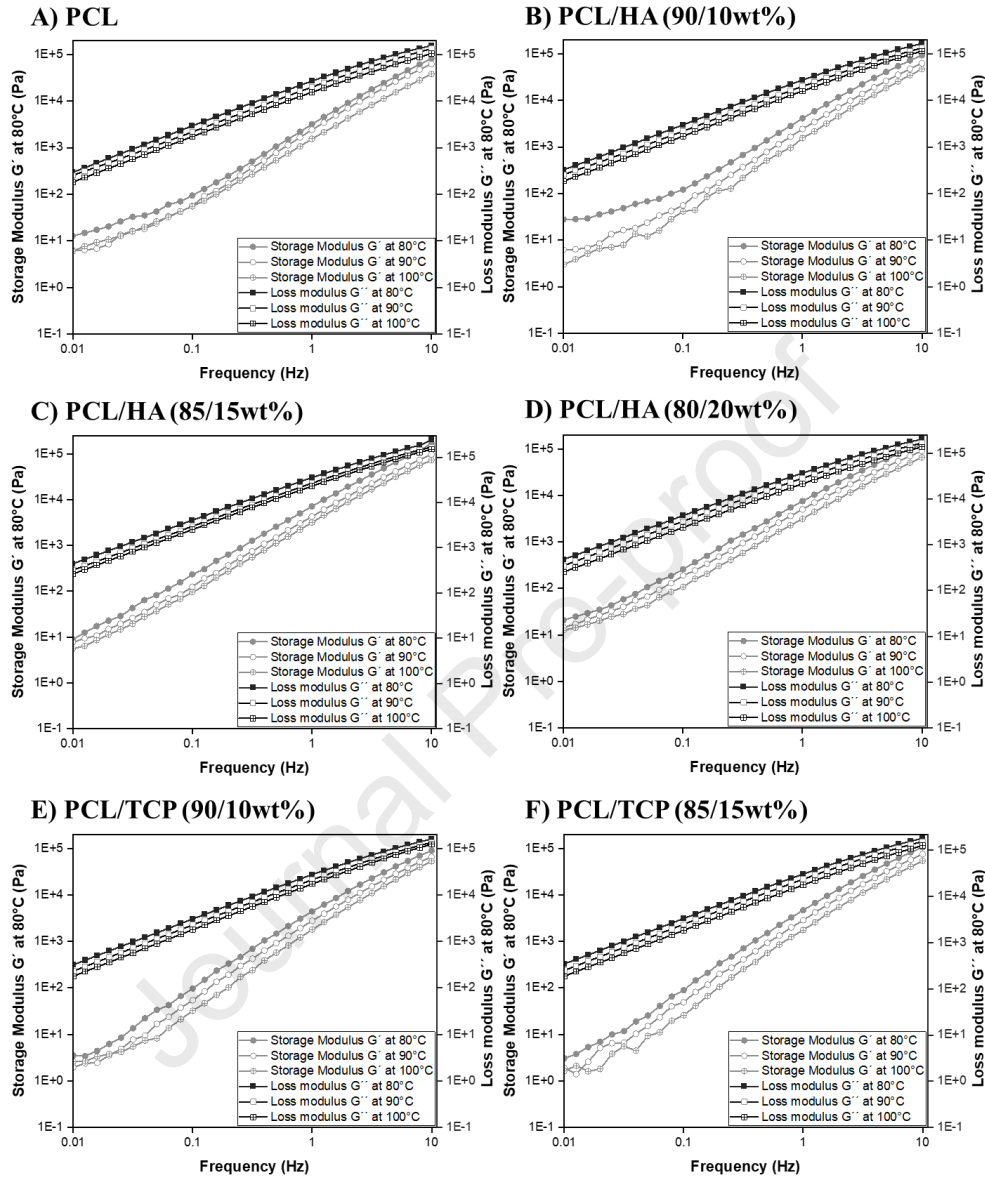


Figure 2. Frequency as a function of storage modulus and loss modulus for the same material concentration at different temperatures of A) PCL, B) PCL/HA (90/10wt%), C) PCL/HA (85/15wt%), D) PCL/HA (80/20wt%), E) PCL/TCP (90/10wt%), F) PCL/TCP (85/15wt%), G) PCL/TCP (80/20wt%), H) PCL/Bioglass (90/10wt%), I) PCL/Bioglass (85/15wt%) and J) PCL/Bioglass (80/20wt%).

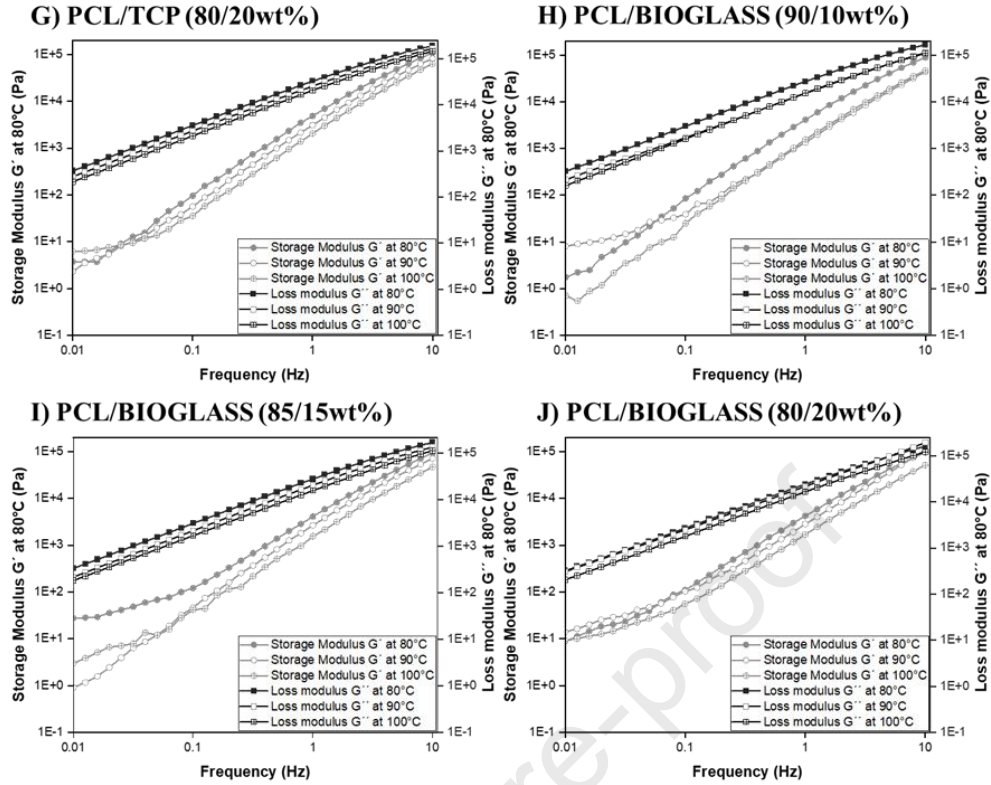


Figure 2 (cont.). Frequency as a function of storage modulus and loss modulus for the same material concentration at different temperatures of A) PCL, B) PCL/HA (90/10wt%), C) PCL/HA (85/15wt%), D) PCL/HA (80/20wt%), E) PCL/TCP (90/10wt%), F) PCL/TCP (85/15wt%), G) PCL/TCP (80/20wt%), H) PCL/Bioglass (90/10wt%), I) PCL/Bioglass (85/15wt%) and J) PCL/Bioglass (80/20wt%).

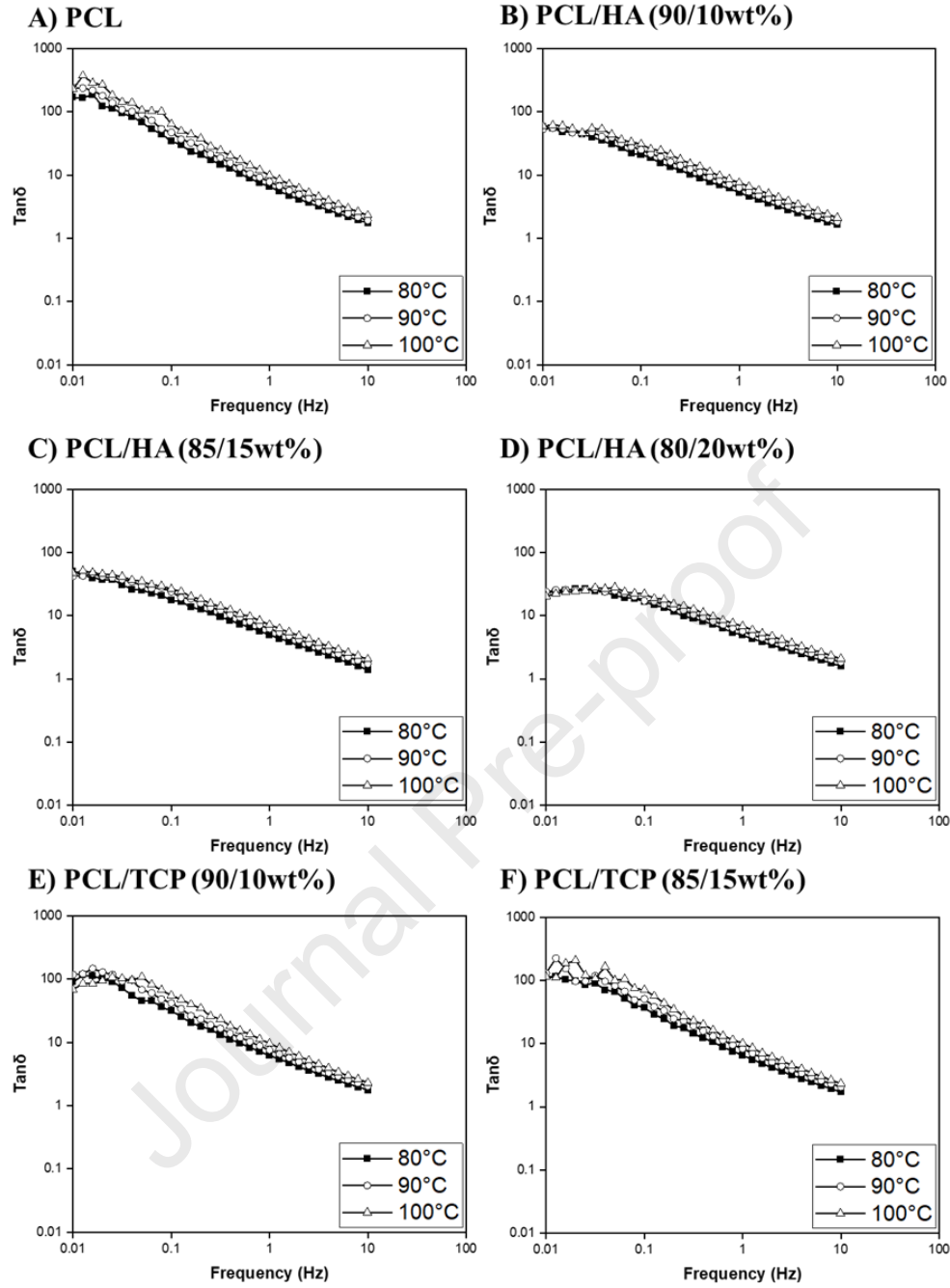


Figure 3. Frequency as a function of Loss factor for the same material concentration at different temperatures of A) PCL, B) PCL/HA (90/10wt%), C) PCL/HA (85/15wt%), D) PCL/HA (80/20wt%), E) PCL/TCP (90/10wt%), F) PCL/TCP (85/15wt%), G) PCL/TCP (80/20wt%), H) PCL/Bioglass (90/10wt%), I) PCL/Bioglass (85/15wt%) and J) PCL/Bioglass (80/20wt%).

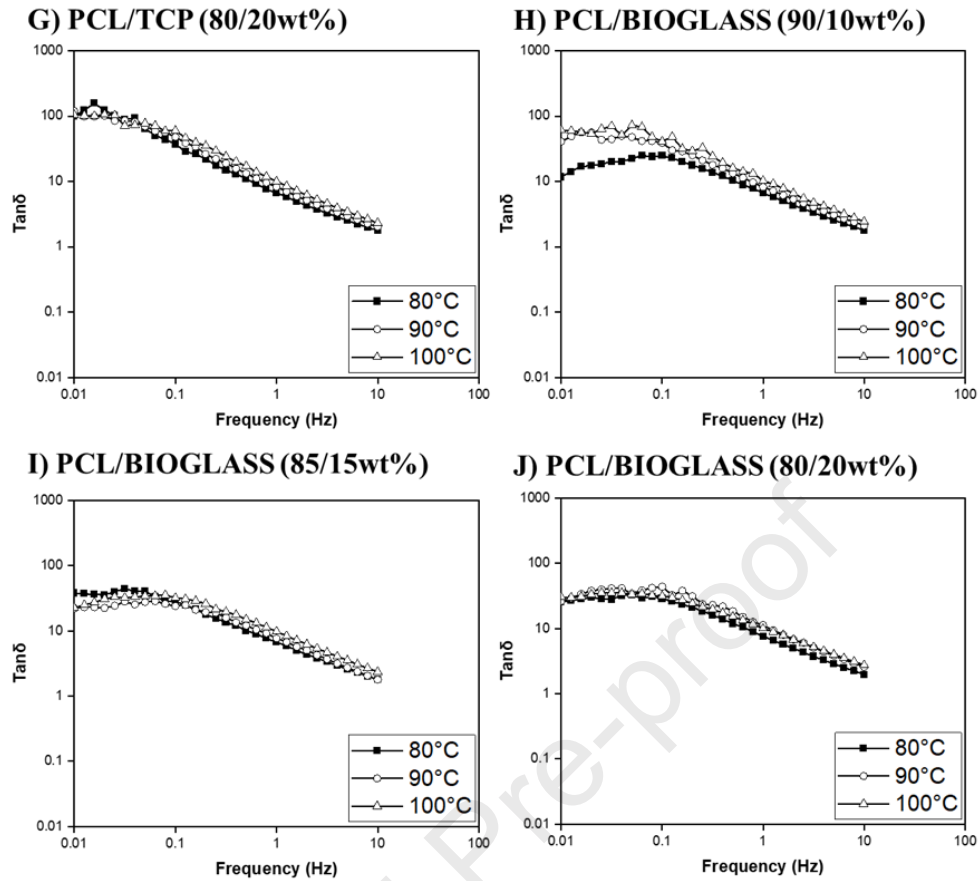


Figure 3 (cont.). Frequency as a function of Loss factor for the same material concentration at different temperatures of A) PCL, B) PCL/HA (90/10wt%), C) PCL/HA (85/15wt%), D) PCL/HA (80/20wt%), E) PCL/TCP (90/10wt%), F) PCL/TCP (85/15wt%), G) PCL/TCP (80/20wt%), H) PCL/Bioglass (90/10wt%), I) PCL/Bioglass (85/15wt%) and J) PCL/Bioglass (80/20wt%).

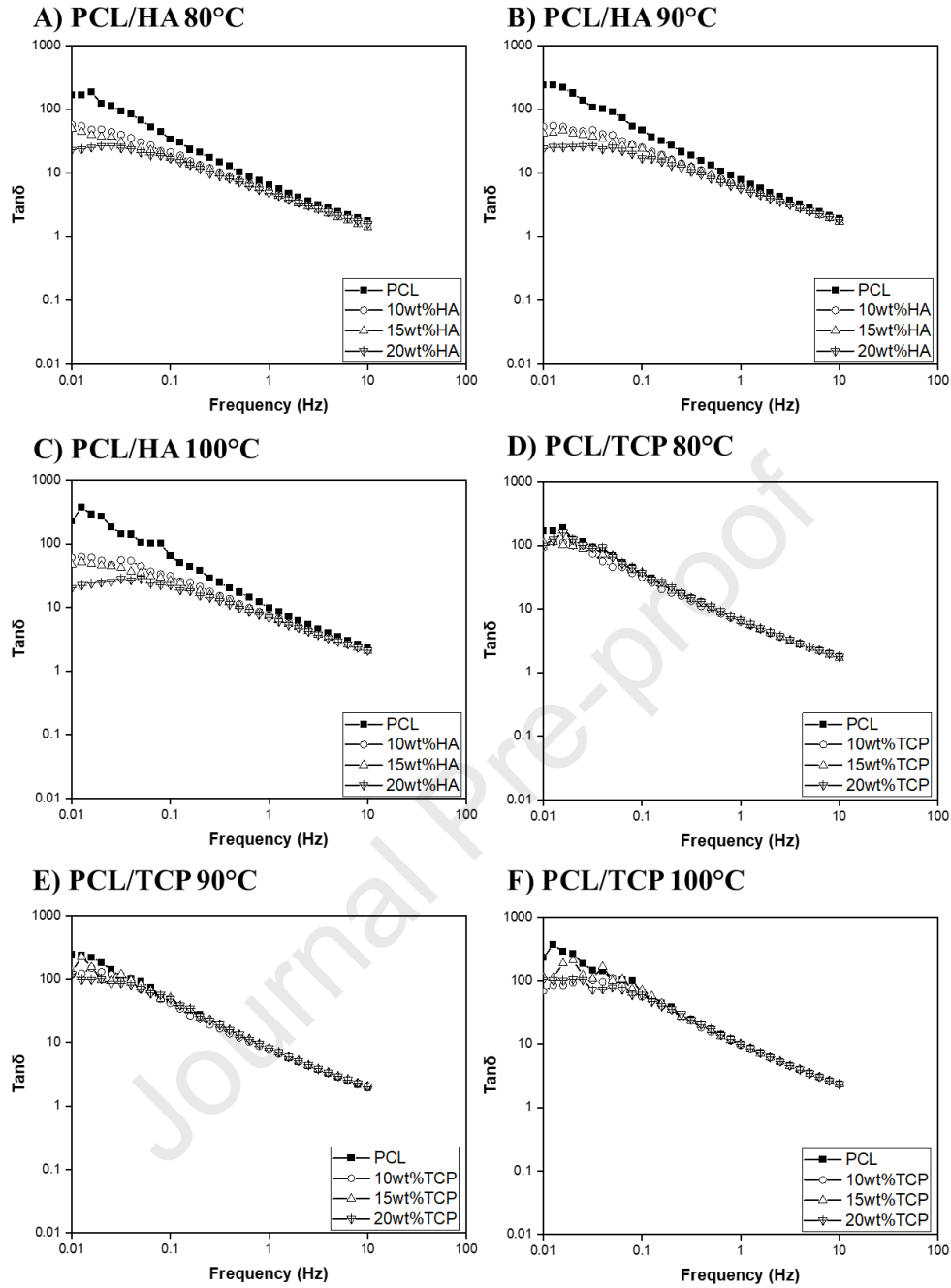


Figure 4. Frequency as a function of Loss factor for the same material content of A) PCL/HA (90/10wt%), B) PCL/HA (85/15wt%), C) PCL/HA (80/20wt%), D) PCL/TCP (90/10wt%), E) PCL/TCP (85/15wt%), F) PCL/TCP (80/20wt%), G) PCL/Bioglass (90/10wt%), H) PCL/Bioglass (85/15wt%) and I) PCL/Bioglass (80/20wt%).

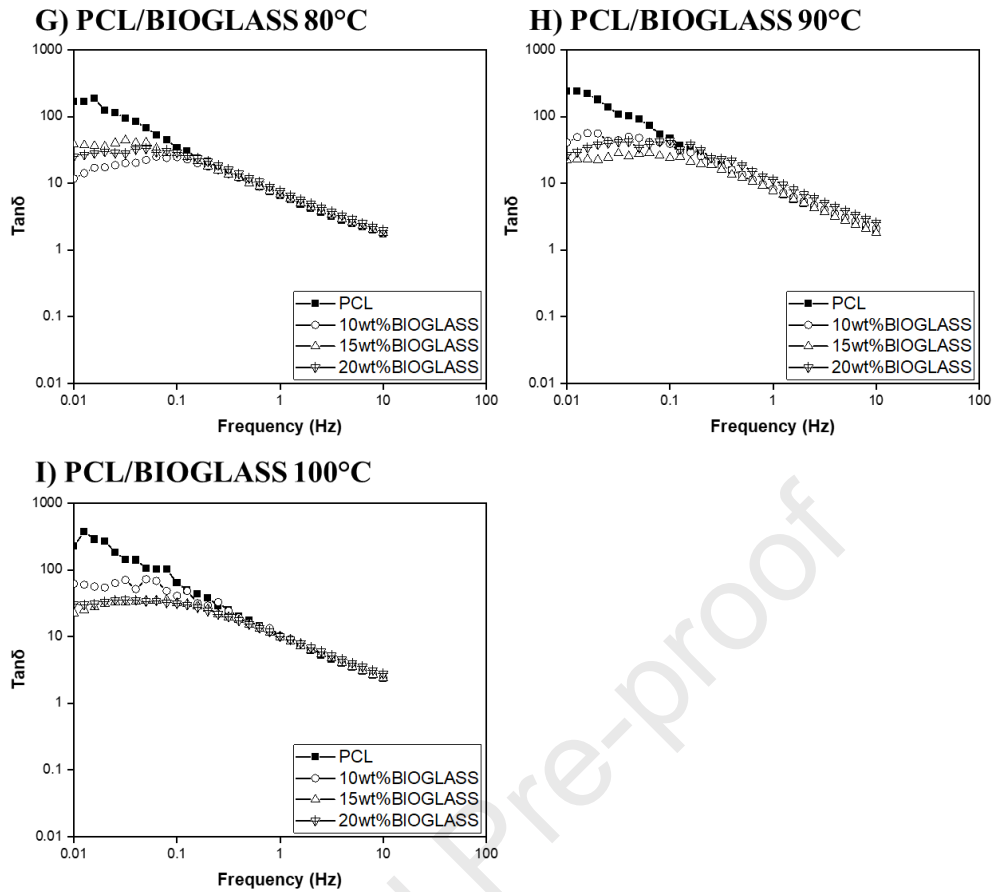


Figure 4 (cont.). Frequency as a function of Loss factor for the same material content of A) PCL/HA (90/10wt%), B) PCL/HA (85/15wt%), C) PCL/HA (80/20wt%), D) PCL/TCP (90/10wt%), E) PCL/TCP (85/15wt%), F) PCL/TCP (80/20wt%), G) PCL/Bioglass (90/10wt%), H) PCL/Bioglass (85/15wt%) and I) PCL/Bioglass (80/20wt%).

3.3 Cole-Cole plot analysis

The Cole-Cole plot can be used to define the dispersion of a non-homogeneous material and the dielectric relaxation of polymer blends [65, 66]. Moreover, through a Cole-Cole plot it is able to detect the relaxation time (τ_c) and zero shear viscosity (η_0) and the compatibility of the different materials in the blends [65-68]. Figure 5 presents the semi quarter-circle plots for considered blends. Results show that the arc created from PCL/HA, PCL/TCP and PCL/Bioglass blends is larger compared to PCL, indicating that the relaxation time and zero shear viscosity of composite blends are higher, leading to longer re-orientation times and higher stresses during the printing process. Additionally, for the same bioceramic material by increasing the fillers concentration the elastic behaviour of the blends increases leading to the reduction of material flow during the printing process. In addition, results show that for the same material content, an increase in temperature reduces the arc, which becomes flatter,

reducing the elasticity of the materials, indicating that the printing process is strongly determined by the viscous characteristics of the blends. Tables 2 and 3 present the zero-shear viscosity and relaxation time for the different blends at different temperatures. As observed the zero-shear viscosity increases by increasing the bioceramic content and decreases by increasing the temperature. The relaxation time depicts a similar trend. In addition, results show that for the same ceramic content and temperature, PCL/Bioglass blends present lower zero-shear viscosity, while PCL/TCP blends present lower relaxation time. In both cases PCL exhibits lower zero-shear viscosity and relaxation times than the polymer-ceramic blends.

Table 2. Zero shear viscosity (Pa.s) results of polymer blends at different temperatures.

Temperature/Material Composition	80 °C	90 °C	100 °C
PCL	2,922.4	2,335.4	1,766
PCL/HA (90/10wt%)	3,671.4	2,699.1	2,140.6
PCL/HA (85/15wt%)	4,174	2,895.9	2,462.7
PCL/HA (80/20wt%)	4,250.4	3,038.4	2,438.6
PCL/TCP (90/10wt%)	3,113.1	2,473.3	2,034.8
PCL/TCP (85/15wt%)	3,610.8	2,617.1	2,183.1
PCL/TCP (80/20wt%)	3,908.6	3,165.3	2,538.7
PCL/Bioglass (90/10wt%)	3,267.3	2,438.1	1,970.8
PCL/Bioglass (85/15wt%)	3362.9	2,480.2	1,843.9
PCL/Bioglass (80/20wt%)	3,445.5	2,950	1,975

Table 3. Relaxation time (s) results of polymer blends at different temperatures.

Temperature/Material Composition	80 °C	90 °C	100 °C
PCL	0.0011	0.0005	0.0003
PCL/HA (90/10wt%)	0.0022	0.0021	0.001
PCL/HA (85/15wt%)	0.0026	0.0023	0.0014
PCL/HA (80/20wt%)	0.0045	0.0036	0.0016
PCL/TCP (90/10wt%)	0.0011	0.0004	0.0002
PCL/TCP (85/15wt%)	0.0011	0.0006	0.0003
PCL/TCP (80/20wt%)	0.0012	0.0009	0.0003
PCL/Bioglass (90/10wt%)	0.0008	0.0006	0.0004
PCL/Bioglass (85/15wt%)	0.0012	0.0006	0.0004
PCL/Bioglass (80/20wt%)	0.0013	0.0007	0.0007

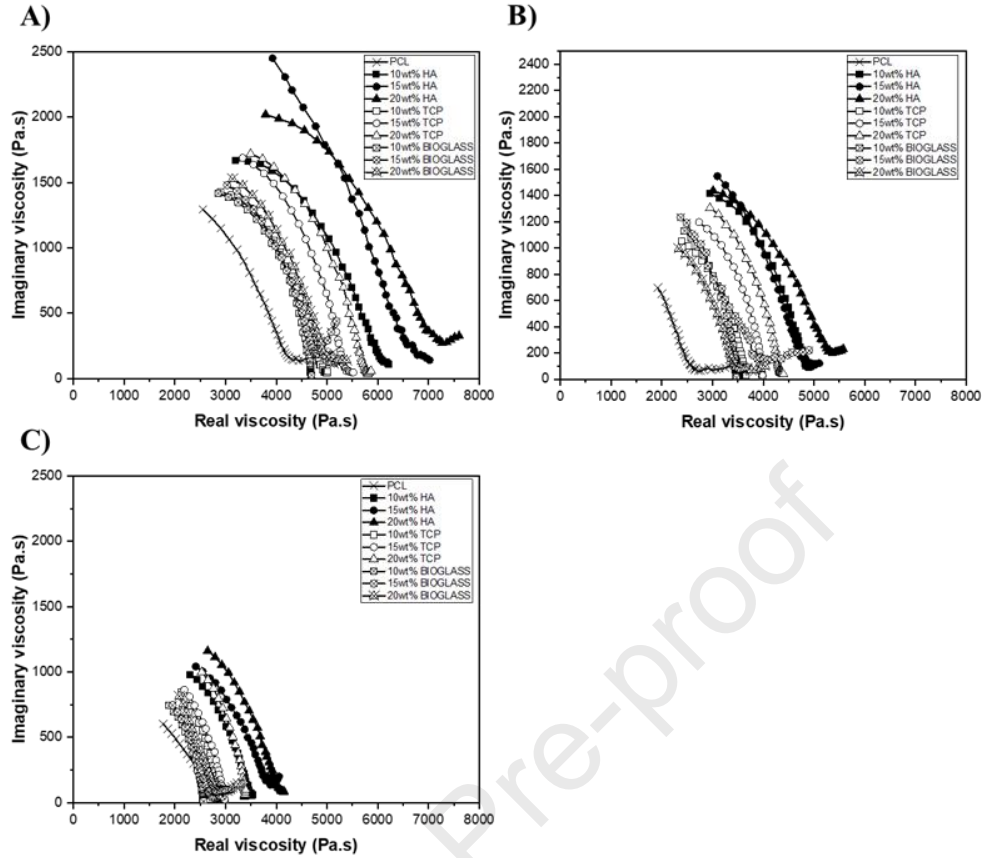


Figure 5. Cole-Cole plot of real viscosity as a function of imaginary viscosity for all material contents of A) 80°C, B) 90°C and C) 100°C.

3.4 Rotational analysis

Figures 6 and 7 present the variation of the dynamic viscosity as a function of the shear rate for the different blends and different temperatures as a function of the shear rate. Based on these results it was possible to calculate the power law index (n) (Equation 4) as shown in Table 4, which indicates that all blends exhibit a shear thinning behaviour being suitable for extrusion printing. Results show that at low shear rate values, PCL/HA, PCL/TCP and PCL/Bioglass blends present higher viscosity values than PCL. However, by increasing the shear rate up to a certain threshold value, the addition of ceramics and their concentration seems to have no major impact and a clear shear-thinning behaviour is observed for all considered samples [60-63, 65, 66, 69, 70]. Moreover, it is possible to observe that, for the same bioceramic content, the viscosity of the composite material increases with the concentration of the inorganic reinforcements. In the Newtonian region (flat plateau) the viscosity of the different blends decreases by increasing temperature, which can be attributed to an increase on the polymer chains movement [65, 66, 69, 70]. Additionally, for the same ceramic material and the same

temperature by increasing its concentration the dynamic viscosity is increasing. For the same material concentration and temperature, PCL/HA blends exhibit higher dynamic viscosity than PCL/TCP and PCL/Bioglass blends, while the lowest dynamic viscosity, between blends with the same ceramic concentration, was observed for PCL/Bioglass.

Table 4. Power law index (n) and Adjusted (R²) results of polymer blends.

Temperature/Material Composition	80 °C	90 °C	100 °C
PCL	0.452	0.534	0.561
PCL/HA (90/10wt%)	0.376	0.31	0.425
PCL/HA (85/15wt%)	0.408	0.348	0.39
PCL/HA (80/20wt%)	0.289	0.276	0.37
PCL/TCP (90/10wt%)	0.455	0.452	0.617
PCL/TCP (85/15wt%)	0.494	0.528	0.626
PCL/TCP (80/20wt%)	0.508	0.629	0.681
PCL/Bioglass (90/10wt%)	0.539	0.42	0.57
PCL/Bioglass (85/15wt%)	0.413	0.258	0.478
PCL/Bioglass (80/20wt%)	0.523	0.478	0.546

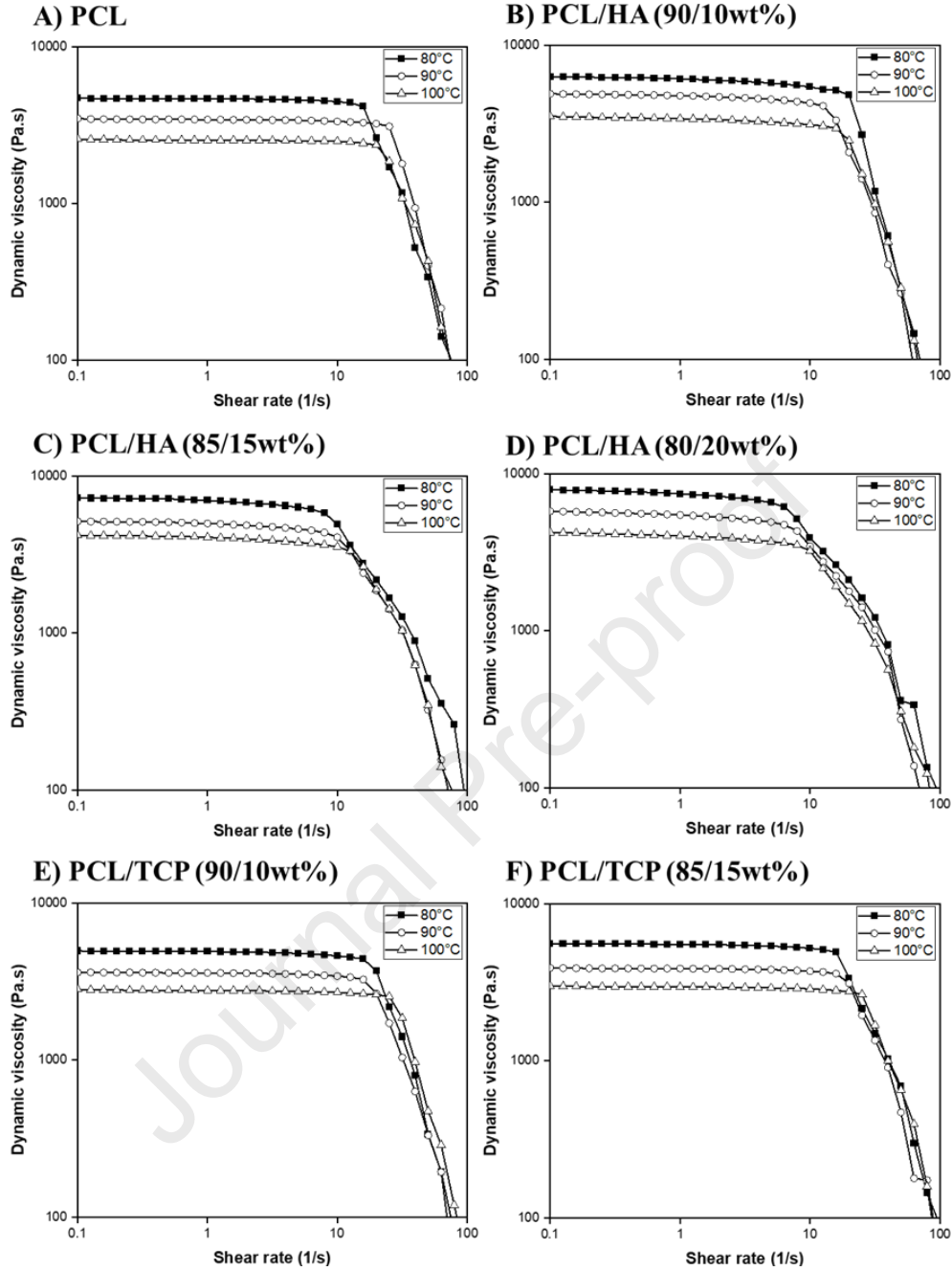


Figure 6. Shear rate as a function of dynamic viscosity for the same material concentration at different temperatures of A) PCL, B) PCL/HA (90/10wt%), C) PCL/HA (85/15wt%), D) PCL/HA (80/20wt%), E) PCL/TCP (90/10wt%), F) PCL/TCP (85/15wt%), G) PCL/TCP (80/20wt%), H) PCL/Bioglass (90/10wt%), I) PCL/Bioglass (85/15wt%) and J) PCL/Bioglass (80/20wt%).

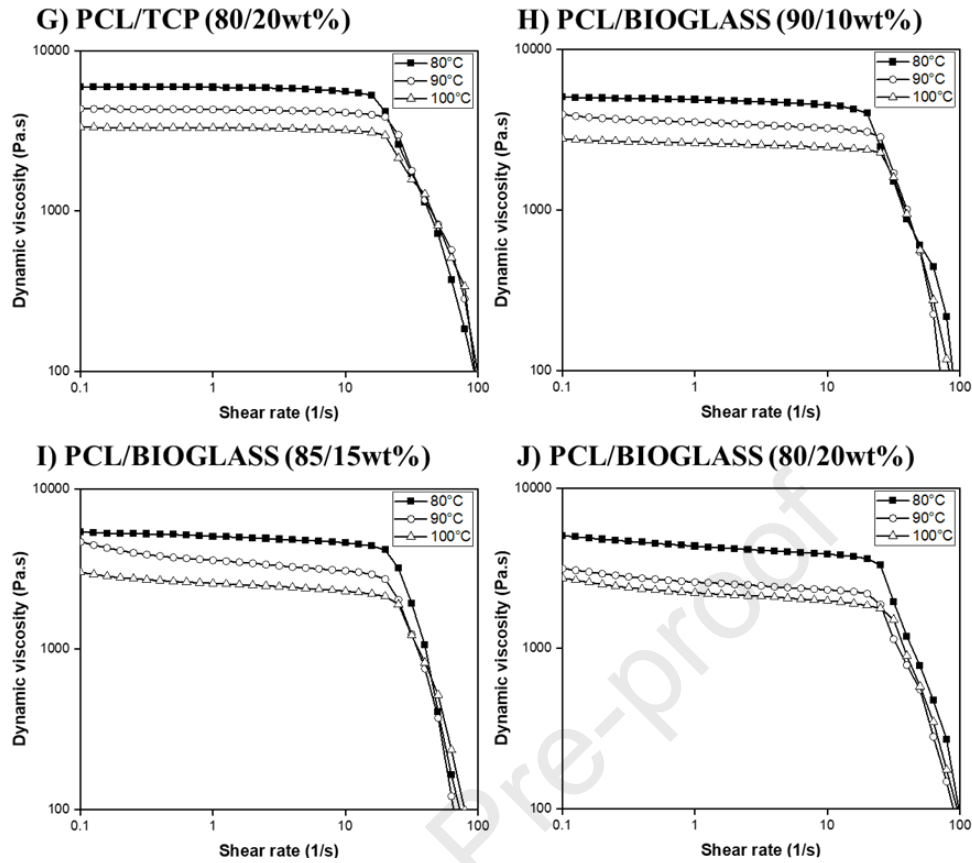


Figure 6 (cont.). Shear rate as a function of dynamic viscosity for the same material concentration at different temperatures of A) PCL, B) PCL/HA (90/10wt%), C) PCL/HA (85/15wt%), D) PCL/HA (80/20wt%), E) PCL/TCP (90/10wt%), F) PCL/TCP (85/15wt%), G) PCL/TCP (80/20wt%), H) PCL/Bioglass (90/10wt%), I) PCL/Bioglass (85/15wt%) and J) PCL/Bioglass (80/20wt%).

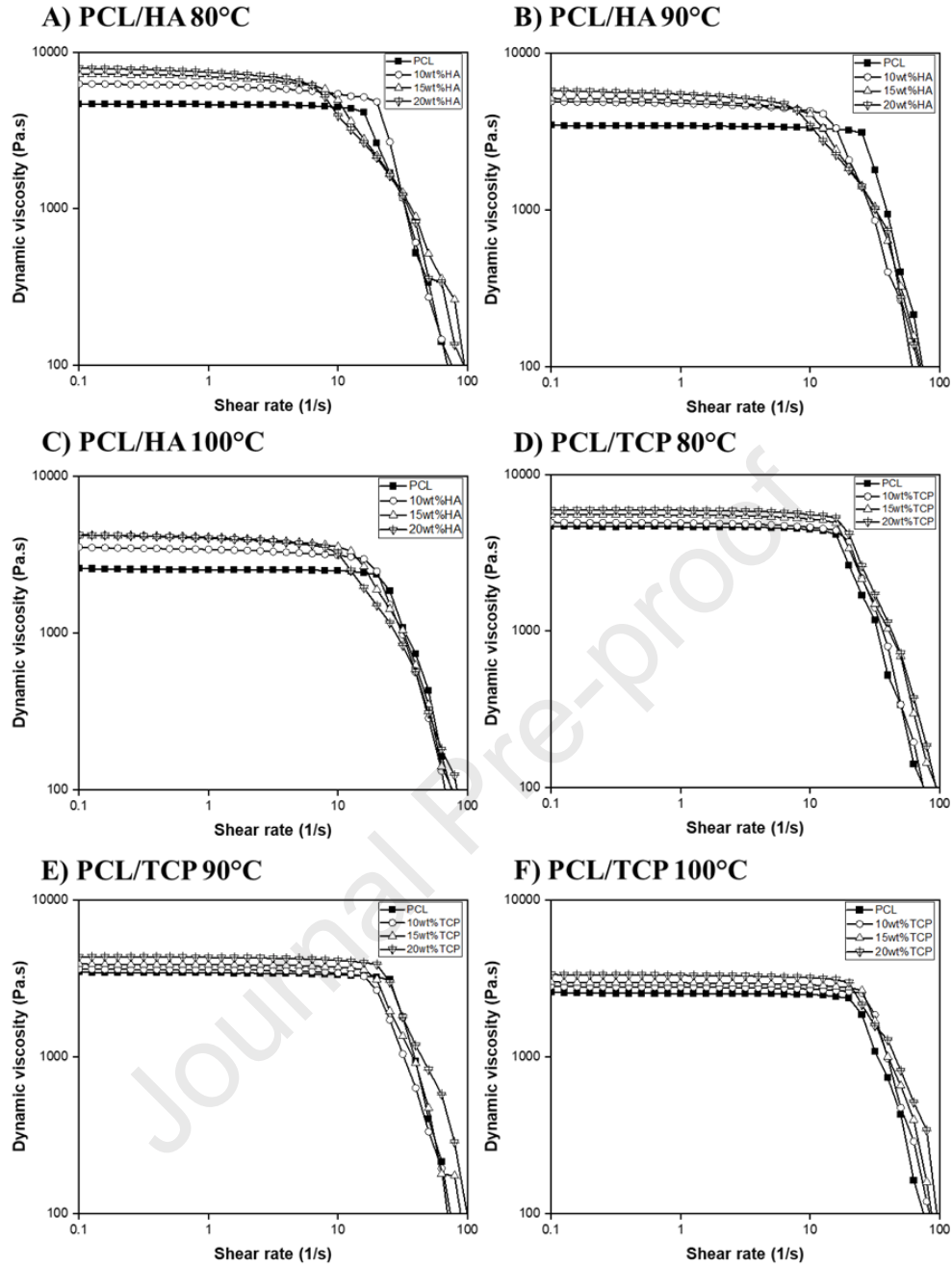


Figure 7. Shear rate as a function of dynamic viscosity for the same material content of A) PCL/HA (90/10wt%), B) PCL/HA (85/15wt%), C) PCL/HA (80/20wt%), D) PCL/TCP (90/10wt%), E) PCL/TCP (85/15wt%), F) PCL/TCP (80/20wt%), G) PCL/Bioglass (90/10wt%), H) PCL/Bioglass (85/15wt%) and I) PCL/Bioglass (80/20wt%).

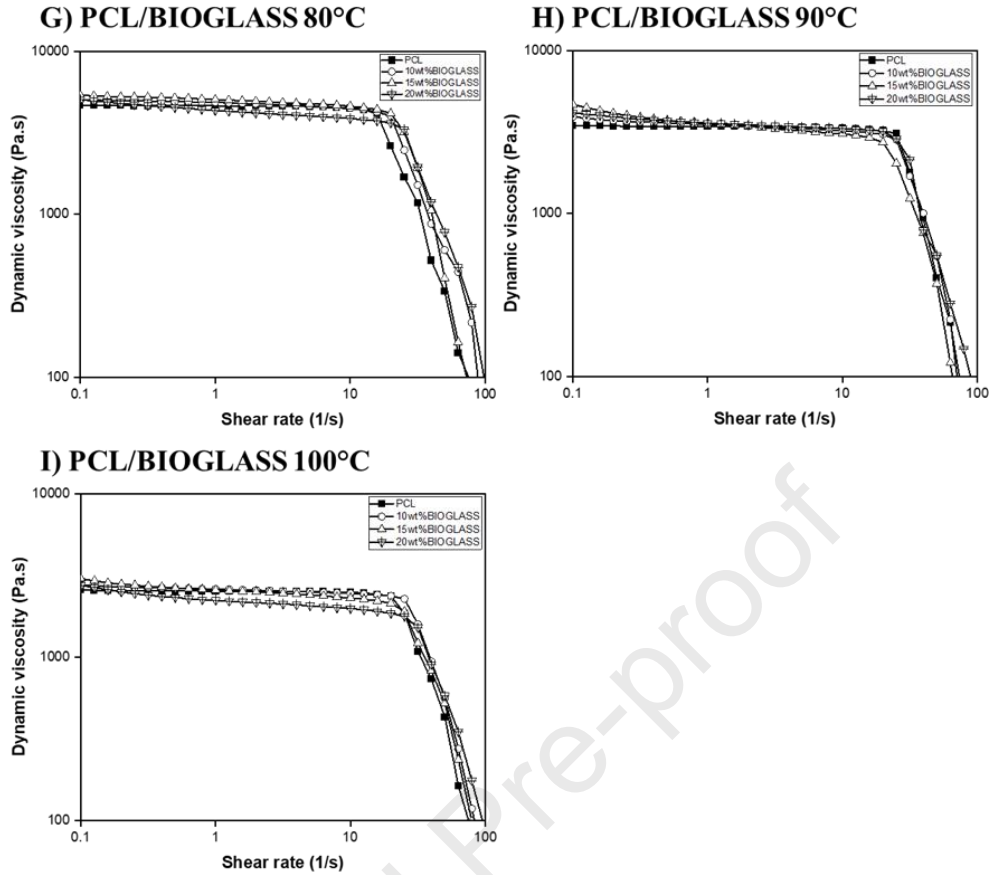


Figure 7 (cont.). Shear rate as a function of dynamic viscosity for the same material content of A) PCL/HA (90/10wt%), B) PCL/HA (85/15wt%), C) PCL/HA (80/20wt%), D) PCL/TCP (90/10wt%), E) PCL/TCP (85/15wt%), F) PCL/TCP (80/20wt%), G) PCL/Bioglass (90/10wt%), H) PCL/Bioglass (85/15wt%) and I) PCL/Bioglass (80/20wt%).

4. Conclusions

The viscoelastic properties of PCL and PCL-based composite blends (PCL/HA, PCL/TCP and PCL/Bioglass) containing different levels of reinforcements (10 wt%, 15 wt% and 20 wt%) was investigated to determine the suitability of these blends for screw-assisted extrusion-based additive manufacturing. At low frequencies all blends present higher viscous modulus than elastic modulus, but these differences tend to reduce by increasing the frequency values. Moreover, the addition of ceramic materials increases both viscous modulus and elastic modulus. Moreover, PCL/HA present higher elastic modulus than PCL/TCP and PCL/Bioglass ones. Composite blends also exhibit longer relaxation times compared with PCL. Overall, for the same material concentration and temperature, blends containing Bioglass particles present slightly lower elastic modulus than PCL/HA blends, and lower loss factor, relaxation times and

shear viscosities than PCL, PCL/HA and PCL/TCP, indicating that PCL/Bioglass blends are more suitable and easier to handle during the printing process.

Author Contributions: Conceptualization, E.D. and P.B.; methodology, E.D., M.H.H. and A.M.O.; software, E.D.; validation, E.D., M.H.H, A.M.O. and P.B.; formal analysis, E.D. and P.B.; investigation, E.D., M.H.H., A.M.O. and P.B.; resources, All Authors.; data curation, All Authors; writing—original draft preparation, E.D.; writing—review and editing, All Authors; visualization, E.D. and P.B.; supervision, G.C, A.W. and P.B.; project administration, P.B.; funding acquisition, P.B. All authors have read and agreed to the published version of the manuscript.

Funding: This project has been supported by the University of Manchester and the Engineering and Physical Sciences Research Council (EPSRC) of the UK, the Global Challenges Research Fund (GCRF), grant number EP/R01513/1.

Acknowledgments: The authors would like to thank the support received from both the University of Manchester and the Engineering and Physical Sciences Research Council (EPSRC) of the UK. This manuscript is the result of the work completed by Evangelos Daskalakis toward a Ph.D. in Mechanical Engineering, granted by the University of Manchester in September of 2022.

Conflicts of Interest: The authors declare no conflict of interest.

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Declaration of interests

☐ The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

☒ The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

Evangelos Daskalakis reports financial support was provided by Engineering and Physical Sciences Research Council.