

RESEARCH ARTICLE



Numerical simulation and printability analysis of fused deposition modeling with dual-temperature control

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Abstract

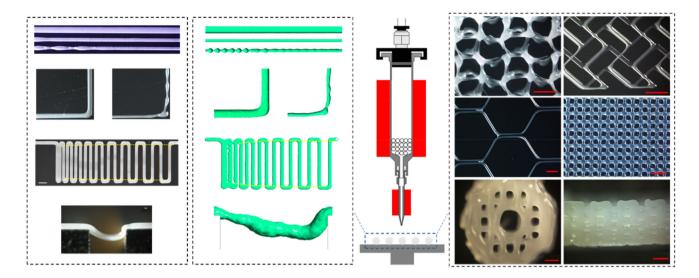
Ideal tissue engineering scaffolds need interconnected pores and high porosity to enable cell survival, migration, proliferation, and differentiation. However, obtaining a high-resolution structure is difficult with traditional one-temperature control fused deposition modeling (FDM). In this study, we propose a dual-temperature control method to improve printability. A numerical model is developed in which the viscosity is a function of temperature and shear rate to study the influence of two different temperature control modes. Quantitative tests are used to assess filament formation and shape fidelity, including one-dimensional filament printing, deposition at corners, fusion, and collapse. By using dual-temperature control, the width of the deposited poly(ϵ -caprolactone) filament is reduced to 50 μ m. The comparative results of both the experimental method and numerical simulation suggest that the dual-temperature control FDM can manufacture spatially arranged constructs and presents a promising application in tissue engineering.

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Graphic abstract



Keywords Fused deposition modeling (FDM) · Dual-temperature control · Filament deposition · Printing resolution · Viscosity

Introduction

Biomedical polymers, namely poly(lactic acid) (PLA) [1], poly(ε-caprolactone) (PCL) [2], and poly(lactic-co-glycolic) acid (PLGA) [3], are popular scaffold materials for tissue engineering. Compared with ceramics and metals [4-6], polymers have better biocompatibility, a more controllable degradation rate, and excellent processability. Conventionally, polymer scaffolds are fabricated by using salt leaching [7], solvent casting [8], gas foaming [9], and phase separation [10]. However, conventional fabrication techniques generally lack the capability to construct regular porous structures, and fully removing toxic organic solvents after fabrication is difficult, resulting in the poor biocompatibility of conventional polymeric scaffolds [11]. Recently, three-dimensional (3D) printing technologies [12], including extrusion-based printing [13], selective laser sintering [14], and melt electrospinning writing [15], have developed rapidly. Compared with conventional manufacturing technologies, 3D printing technologies present several advantages in tissue engineering, including high printing resolution, fast fabrication speed, and the ability to customize objects to meet the demands of specific applications [16–20].

A widely used 3D printing method is fused deposition modeling (FDM), which is easier to implement than other 3D printing methods [21–25]. During FDM, thermoplastic polymeric materials are melted at the nozzle and extruded as filaments, which are deposited and fused layer-by-layer onto the receiving platform to solidify into final parts [26,

27]. FDM can precisely control the diameter and location of deposited filaments, resulting in overall control of scaffold shape and pore morphology [21]. In addition, FDM has no toxic chemical solvents that need removal in tissue scaffold fabrication [28–30]. Based on these aforementioned advantages, FDM has been used to print different polymers, including PLA and its copolymer [31] and PCL [32–36]. Recently, research has increasingly focused on the FDM of PCL as a tissue engineering scaffold. By using two lay-down patterns, 0°/90° and 0°/60°/120°, Zein et al. [34] manufactured PCL scaffolds with different pore shapes and a fully interconnected channel network. Zhang et al. [35] fabricated PCL scaffolds with different pore sizes for meniscus tissue engineering; the PCL scaffold with a mean pore size of 215 µm demonstrated optimized cell behaviors, extracellular matrix (ECM) production and deposition, and the resultant mechanical properties. Manjunath et al. [37] used FDM to fabricate PLA/PCL composites to improve the Young's modulus of bone tissue engineering scaffolds.

The ideal tissue engineering scaffolds must have interconnected pores and high porosity to ensure cell migration, nutrition diffusion, and metabolic waste removal [38–41]. Larger pores in scaffolds are conducive to cell diffusion and migration, whereas smaller pores provide a higher scaffold surface area for cell adhesion [34, 35]. Therefore, the accurate control of the porosity and pore size of tissue engineering scaffolds is of great importance. Previous literature rarely reports achieving FDM-printed PCL filaments with a diameter smaller than 200 µm, which is much larger than the



dimension of cells (less than $50~\mu m$) [42, 43]. Thus, printing PCL tissue engineering scaffolds with a smaller size and high porosity using FDM is difficult. Efforts have been made to increase the printing resolution of FDM. Kumar et al. [44] reduced the width of FDM-printed filaments by 42.42% when the printing speed was increased from 700 to 1300~mm/min, but the filament diameter was still larger than 1 mm. Hıra et al. [45] used the optimized nozzle channel geometries to improve the dimensional accuracy (deviation of width is within 6%–8%) of the FDM-printed acrylonitrile butadiene styrene (ABS), PLA, and chlorinated polyethylene (CPE) parts, but the optimization mechanism of the nozzle geometry was difficult to extend to other printers.

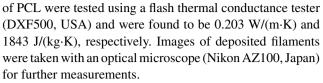
Generally, in most FDM printers, the one-temperature control system is implemented, and the temperature control is set on the metal printing syringe (Fig. 1a). Thus, the temperature gradient clearly decreases from the syringe to the nozzle, leading to a much lower temperature at the nozzle outlet than the setting temperature. Therefore, controlling the needle temperature by regulating the syringe temperature is highly difficult. Moreover, the viscosity of the melting polymer fluid is sensitive to printing temperature, and a higher viscosity results in poor or even failed extrusion, while a lower viscosity leads to the melt spreading of the polymeric filament. Therefore, accurate control of the printing temperature is of great importance for FDM of thermoplastic polymers. Based on this contradiction, this study proposes dual-temperature control of FDM printing. A higher heating temperature is applied at the syringe to improve the fluidity of the upper part of the melt PCL, while a lower temperature is applied at the needle to achieve better formability of the extruded PCL.

In this study, we present a viable, cost-effective, and simple process modification to decrease the filament width to 50 μ m and below by changing the traditional one-temperature control to dual-temperature control. Using experimental methods and numerical simulations, this study explores the printability with different temperature control modes and examines the mechanism by which dual-temperature control has better printability than one-temperature control. By using dual-temperature control, the width of the deposited PCL filament is reduced, and the printability of the FDM is improved, thereby providing a new idea for further application of PCL scaffolds in tissue engineering.

Materials and methods

Materials and characterization

Melted PCL (Mw=150,000) with a density of 1146 kg/m³ was purchased from Daigang Biomaterial (Jinan, China). The thermal conductivity coefficient and specific heat capacity



The rheological properties of the melted PCL were determined using a fast-rotating rheometer (MCR102, Anton Paar, Austria). The experiments were performed by using a 25-mm diameter parallel-plate configuration (diameter 25 mm, gap 1 mm). Experiments on the temperature and viscosity of PCL samples were carried out at 60 °C and 140 °C in a step of 2 °C as the shear rates ranged from 0.1 to 4 s⁻¹. Subsequently, experiments on the shear rate and viscosity were carried out on PCL samples with temperatures ranging from 65–140 °C and shear rates of 0.1–30 s⁻¹.

FDM with dual-temperature control

To achieve an appropriate temperature at the printing nozzle outlet, we propose dual-temperature control (Fig. 1a). A higher heating temperature was applied at the syringe (high-temperature heating zone) for PCL melting, while a lower temperature was applied at the needle (low-temperature heating zone). When the melted PCL flowed from the high-temperature heating zone to the low-temperature heating zone, the printing temperature gradually decreased, and the rheology of melted PCL became appropriate for extrusion.

The FDM printing platform consists of a 3D motion platform, a system of air pressure mediation and a temperature controller. Figure 1a shows the schematic diagram of the printing head. In dual-temperature control, a long needle (17 mm) was used and wrapped with a syringe with an aluminum heating block. By comparison, the one-temperature control used a short needle (7 mm), and only the syringe was wrapped with an aluminum heating block. The temperature sensor was used to provide an accurate actual temperature for proportional-integral-derivative (PID) control to precisely control the heating temperature. Needles with different inner diameters (100 and 200 µm) were purchased from Deli Precision (Shenzhen, China). Under the control of computer software, the nozzle can move vertically (x-y plane), and the substrate can move horizontally (z-direction), as shown in Fig. 1b. All printing processes were performed at room temperature.

Numerical model of dual-temperature control FDM

In this study, the computational fluid dynamics approach was conducted to evaluate the printability of FDM printing with two temperature control modes using the software ANSYS (Ansys, Canonsburg, PA, USA). To guarantee the precision of the computational simulation, the geometry was meshed



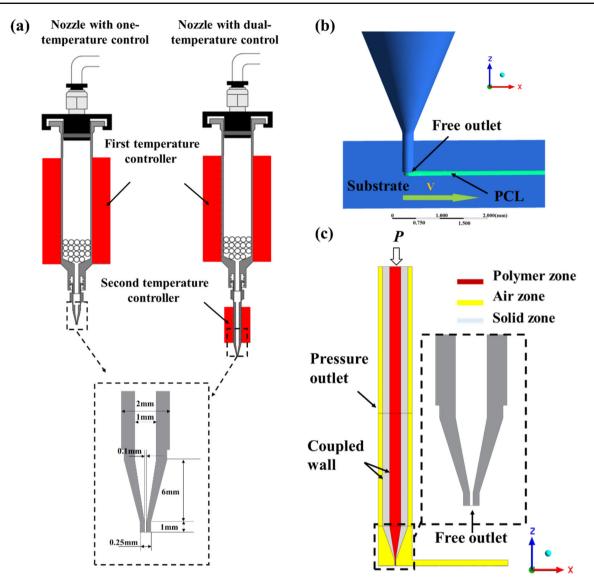


Fig. 1 Fused deposition modeling (FDM) system: **a** schematic diagram with one- and dual-temperature control and the nozzle geometry; **b** front view of a simulation model of filament deposition; **c** geometry and boundary conditions of the numerical model (PCL: poly(ϵ -caprolactone))

using hexahedrons, and the maximum size of the control volumes was set as 0.02 mm. Figure 1b shows a representative image of the printed whole filament. In this numerical model, the melted PCL fluid at the nozzle is assumed to be laminar and fully developed.

Governing equations

In the numerical simulation, the mass conservation of the incompressible flow is given as

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \, \overrightarrow{u}) = 0 \tag{1}$$

and the Navier–Stokes equation is used to describe the melted PCL fluid flow

$$\frac{\partial}{\partial t}(\rho \overrightarrow{u}) + \rho(\overrightarrow{u} \cdot \nabla) \overrightarrow{u} = \nabla \cdot [-pI + \mu(\nabla \overrightarrow{u} + (\nabla \overrightarrow{u})^{\mathrm{T}})] + \rho \overrightarrow{g} + \overrightarrow{F}_{\mathrm{st}},$$
(2)

where ρ is the density, u is the flow velocity, p is the pressure applied to the fluid, I is the identity matrix, μ is the dynamic viscosity of the fluid, g is the gravity field, $F_{\rm st}$ represents the surface tension, and t is time. The temperature distribution of PCL and air can be obtained by solving the energy equation:

$$\rho C_{\mathbf{p}} \vec{u} \cdot \nabla T + \nabla \cdot (-k \nabla T) = \dot{Q}, \tag{3}$$



where C_p is the specific heat capacity at constant pressure, k is the thermal conductivity, \dot{Q} is the heat flow rate, and T is temperature. The volume of fluid method is used for tracking the interface between the liquid and gas [46–50].

Geometric model and boundary conditions

In the simulation model, plane symmetry is used on the middle plane (x-z plane) of the nozzle in Fig. 1c. The nozzle is a cylindrical tube with inner diameter D and a fixed position, while the substrate moves with a controlled (printing) speed. Below the printing head, the moving substrate is modeled by a perpendicular stiff plane, with a gap distance H to the nozzle outlet. The long needle that is used in dual-temperature control has a heating section L_1 . The syringe and the needle are filled with the melted PCL at the beginning of the simulation. With a constant inlet pressure, the melted PCL fluid is extruded and deposited onto a moving substrate. The deposited PCL gradually cooled and began to solidify when the polymer temperature was below the melting point of PCL. Notably, fixed temperature boundary conditions are used at the inlet. The temperatures of the syringe and needle of the dual-temperature control FDM were 140 and 120 °C, respectively, while those of the one-temperature control FDM were set to 120 °C. There is convective heat transfer between the wall and the air and thermal conduction between the wall and the fluid; thus, the nozzle wall is set to be coupled under thermal boundary conditions (coupled wall). Table 1 lists the necessary material properties for computational simulations.

Constitutive equations of PCL

According to the Cross-Williams-Landel-Ferry (Cross-WLF) viscosity model [51, 52], an empirical model is proposed to describe the rheological behavior of melted PCL under different temperatures. The Cross-WLF model can be described as

$$\eta(T, \gamma) = \frac{\eta_0(T, \gamma)}{1 + \left(\frac{\eta_0 \dot{\gamma}}{\tau^*}\right)^{1-n}} \tag{4}$$

and

$$\eta_0(T, \gamma) = D_1 \exp \left[-\frac{A_1(T - T_g)}{A_2 + (T - T_g)} \right],$$
(5)

where viscosity (η) is a function of temperature (T) and shear rate $(\dot{\gamma})$, τ^* is the critical shear stress for the fluid to change into a pseudoplastic fluid, n is a non-Newtonian index, η_0 is the zero-shear viscosity and is defined by Eq. (5), and T_g is the glass transition temperature. When T is higher than the glass transition temperature (T_g) , the zero-shear viscosity is considered a function of temperature (Eq. (5)). T_g is



Parameter	Definition	Value	Unit	
$C_{\mathrm{p,a}}$	Specific heat capacity of the air	1000	J/(kg·K)	
$C_{\mathrm{p,n}}$	Specific heat capacity of the nozzle	500	J/(kg·K)	
$C_{\mathrm{p,p}}$	Specific heat capacity of the PCL	1843	J/(kg·K)	
$C_{\mathrm{p,s}}$	Specific heat capacity of the substrate	800	J/(kg·K)	
D	Inner diameter of the nozzle	100/200	μm	
Н	Gap distance between nozzle and substrate	80	μm	
$k_{\rm a}$	Thermal conductivity coefficient of the air	0.034	$W/(m \cdot K)$	
$k_{ m n}$	Thermal conductivity coefficient of the nozzle	17	W/(m·K)	
$k_{ m p}$	Thermal conductivity coefficient of the PCL	0.203	W/(m·K)	
k_{s}	Thermal conductivity coefficient of the substrate	2.5	W/(m·K)	
L_1	Heating length	10	mm	
L_2	Taper length	6	mm	
L_3	Capillary length	1	mm	
P	Air pressure	0.4	MPa	
T	Syringe/needle temperature	140/120	°C	
$T_{\rm s}$	Atmosphere temperature	25	°C	
v	Printing speed	0.2-2.1	mm/s	
$\eta_{ m a}$	Viscosity of the air	2.2×10^{-5}	Pa·s	
$\eta_{ m p}$	Viscosity of the PCL	$\eta \left(\mathrm{T,}\gamma \right) ^{\ast }$	Pa·s	
$ ho_{ m a}$	Density of the air	0.9	kg/m ³	
$ ho_{ m n}$	Density of the nozzle	7800	kg/m ³	
$ ho_{ m p}$	Density of the PCL	1160	kg/m ³	
$ ho_{ m s}$	Density of the substrate	2200	kg/m ³	
σ	Surface tension coefficient of the PCL in air	1.00	mN/m	

^{*} This parameter is described in Section "Constitutive equations of PCL". PCL: poly(ε-caprolactone)

the reference temperature, mainly related to the glass transition temperature of the melted PCL. D_1 , A_1 , and A_2 are the material constants of melted PCL to be fitted.



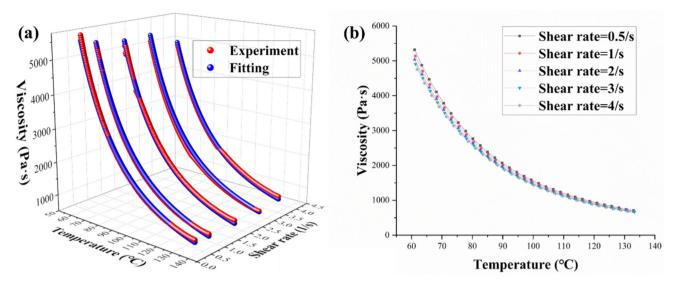


Fig. 2 Constitutive model of melted poly(ε-caprolactone) (PCL): a experimental and fitted data; b viscosity–temperature curves of PCL at different shear rates

Table 2 Parameters of the cross-Williams–Landel–Ferry (cross-WLF) model

Parameter	n	τ* (Pa)	D_1 (Pa·s)	$T_{g}(K)$	A_1	A ₂ (K)
Value	0.9495	9.97×10^{3}	7.429×10^9	213.15	21.51	63.49

Results and discussion

Determining melted PCL constitutive equations

Figure 2a shows the experimental results of rheological characterization of melted PCL, and the fitting data based on the constitutive model (Eqs. (4) and (5)) are found to be in good agreement with the experimental data (Fig. 2a). Table 2 lists the fitting parameters of the PCL constitutive model.

The glass transition temperature ($T_{\rm g}$) of melted PCL is approximately -60 °C. Based on the constitutive model (Eq. (4)), the melted PCL fluid demonstrates shear-thinning behavior under different temperatures. The viscosity–temperature curve under a series of shear rates is shown in Fig. 2b.

Printing of one-dimensional (1D) filament

To compare the printability of 1D filaments with two different temperature control modes, we deposited linear fibers with different printing speeds to explore the mechanisms of formability, fiber width, and cross-sectional variation [53]. Figure 3a shows that when the printing speed was small, the melted PCL was deposited on the substrate to form continuous filaments. However, as the printing speed increased to a critical value, the filament became unstable, and continuous filaments could not be formed. Notably, the critical printing speed of the dual-temperature control FDM (2 mm/s) is greater than that of the one-temperature control FDM

(1.7 mm/s). In addition, at the same printing speed, the width of the filament printed by the dual-temperature control FDM is smaller than that printed by the one-temperature control FDM, as shown in Fig. 3b. Figure 3c shows that with increasing printing speed, the cross section of the deposited filaments gradually changes from a rectangular to a circular shape. At the same printing speed, the deposited filaments printed by the dual-temperature control FDM had a higher shape fidelity, and the cross section of deposited filaments printed by one-temperature control FDM was easier to collapse.

Figures 4a1 and a2 show the radial temperature distribution at the nozzle outlet. Compared with the dual-temperature control FDM, the one-temperature control FDM has a less uniform radial temperature distribution, which leads to a less uniform viscosity distribution of the melted PCL at the nozzle outlet (Fig. 4a3). Therefore, the radial velocity distribution at the outlet is not uniform (Fig. 4a4), and the shear rate at the outlet of the one-temperature control FDM is approximately 10 times that of the dual-temperature control FDM (Fig. 4a5). Figure 4b2 shows that the viscosity of the melted PCL fluid at different positions is significantly different and smaller than that of the dual-temperature control FDM. Given the lower viscosity at the outlet of one-temperature control FDM, the outlet flow rate is larger at the same printing speed, which causes the accumulation of the melted PCL at the outlet (Fig. 3c). Therefore, the one-temperature control FDM has wider filaments during printing, meaning a lower printing resolution. The viscosity distribution at the nozzle outlet is



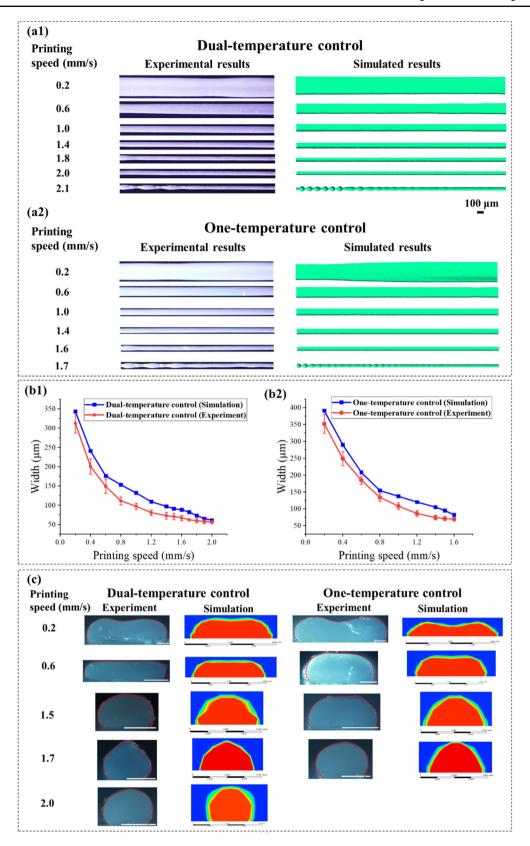


Fig. 3 Filament printing (nozzle with an inner diameter of $100~\mu m$): a experimental results and simulated results with dual- and one-temperature control; b filament width as a function of printing speed with dual- and one-temperature control (the blue line represents the

simulated result and the red line represents the experimental result); \bm{c} exploration of the shape of the cross section of the filament, scale bar=100 μm



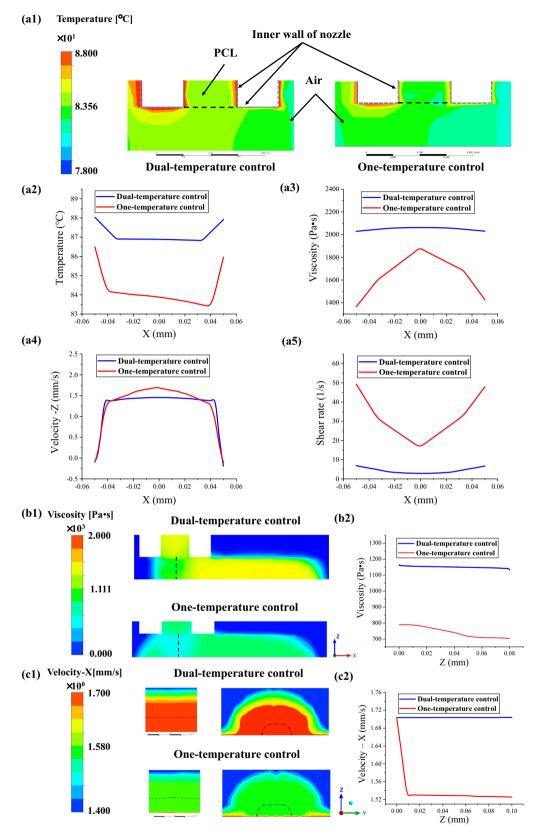


Fig. 4 Mechanistic analysis of filament printing: **a1** contour of temperature (the black dotted line represents the diagram of the exported data line), **a2** temperature, **a3** viscosity, **a4** velocity at z-direction, and **a5** shear rate at the nozzle outlet; **b1** contour of viscosity at t=0.5 s and **b2**

viscosity distribution at *z*-direction; **c1** contour of velocity at *x*-direction and **c2** velocity distribution of *x*-direction of *z*-direction melted poly(ε -caprolactone) (PCL) when printing speed v=1.7 mm/s



less uniform with one-temperature control (Fig. 4b2), resulting in a velocity gradient in the *x*-direction of the *z*-direction deposited PCL filament (Fig. 4c2). The velocity gradient leads to the shear rate between layers, which leads to unstable filament formation (Fig. 3a).

PCL deposition at corners

The deposition of ink at the corners significantly affects the geometric quality of parts fabricated by extrusion printing. To quantify the corner filling performance, we propose a dimensionless number to evaluate the deviation between the design and the actual path as (Fig. 5b1),

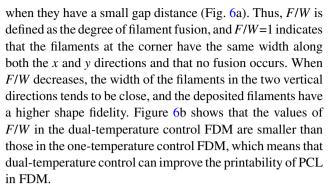
$$A_{\rm d} = \frac{A_{\rm deviation}}{A_{\rm design}},\tag{6}$$

where $A_{\rm deviation}$ = $A_{\rm overfill}$ + $A_{\rm underfill}$, and $A_{\rm design}$ is the designed area. The deviation area ($A_{\rm deviation}$) consists of two parts: one part is the amount of overfill ($A_{\rm overfill}$), and the other is that of underfill ($A_{\rm underfill}$). As $A_{\rm d}$ decreases to 0, the deposition of PCL at the corner is closer to the designed path, resulting in an excellent geometric quality of the corner. Figure 5a shows that the filling quality of the corner worsens with increasing printing speed. At the same printing speed, the temperature control of the corner filling performance is better with the dual-temperature control FDM than with the one-temperature control FDM (Fig. 5b2). In addition, the numerical results agree well with the experimental results (Fig. 5a).

To understand the mechanism of the smaller deviation of dual-temperature control FDM, we use a numerical model with a printing speed of 1.6 mm/s for analysis. Notably, regardless of moving in the *x*-direction (Fig. 5c1) or the *y*-direction (Fig. 5c2), the velocity gradient is smaller in the dual-temperature control FDM. Based on the viscosity distribution at the outlet, the viscosity of the melted PCL in dual-temperature control FDM is larger than that of the one-temperature control FDM; when the viscosity of the melted PCL is larger, the speed of filament deposition is closer to the printing speed. When printing from the *x*- to *y*-direction, the velocity gradient in the latter bends the filament, and the bending degree increases with the velocity gradient. Therefore, the design and the actual path show a slight difference in the dual-temperature control FDM.

Filament fusion test

Instead of maintaining the 3D constructs, the two adjacent filaments are fused, which significantly influences the printing resolution. The resolution of the printed filaments in the x-y plane was assessed by measuring their width (W), gap (G), and fusion length (F) (Fig. 6c). Notably, filaments fuse



The filament fusion test is performed by printing the 90° corner, and its quality is assessed by using the corner length (L_c) (Fig. 6c1), which refers to the deviation between the actual and designed paths. Notably, when G < W, the filaments completely fused (Fig. 6c2); when $W < G < L_c$, the filaments only fused at the corner (Fig. 6c3); when $G>L_c$, the filament did not fuse, and $F/W \approx 1$ (Fig. 6c4). When the printing speed is 1.0 mm/s, W_1 is 138 μ m and L_{c1} is 630 μ m with the onetemperature control FDM, while W_2 is 132 μ m and L_{c2} is 310 µm with the dual-temperature control FDM (Fig. 6d1). Changes in G lead to four results (Fig. 6d2): (1) when $L_{c1} < G$ $< L_{c2}$, the filament does not fuse; (2) when $W_1 < G < L_{c1}$, and $G > L_{c2}$, filament fusion occurs with one-temperature control but not with dual-temperature control; (3) when $W_1 < G < L_{c1}$, and $W_2 < G < L_{c2}$, filament fusion occurs in both temperature modes; and (4) when $G < W_1$, and $G < W_2$, complete filament fusion is found in two modes.

Filament collapse test

To quantify the effects of different printing speeds and spanning distances on filament collapse, a filament collapse test was performed (Fig. 7a). Here, this study introduces the collapse distance, which is the distance between the lowest point of collapse and the substrate. The collapse distance gradually increases with the spanning distance at the same printing speed (Fig. 7c1) but gradually decreases with increasing printing speed at the same spanning distance (Fig. 7c2). In addition, the collapse distance in the one-temperature control FDM is always larger than that in the dual-temperature control FDM with the same printing parameters (Fig. 7c). Therefore, the dual-temperature control FDM of PCL provided a better shape fidelity of PCL scaffolds than the one-temperature control FDM.

The deformation of the suspended filaments is caused by the weight of the material. Given that the high viscosity of melted PCL leads to a smaller deformation caused by gravity and that the viscosity of melted PCL in one-temperature control FDM is lower than that in dual-temperature control FDM (Fig. 7d), the collapse distance is larger in one-temperature control FDM.



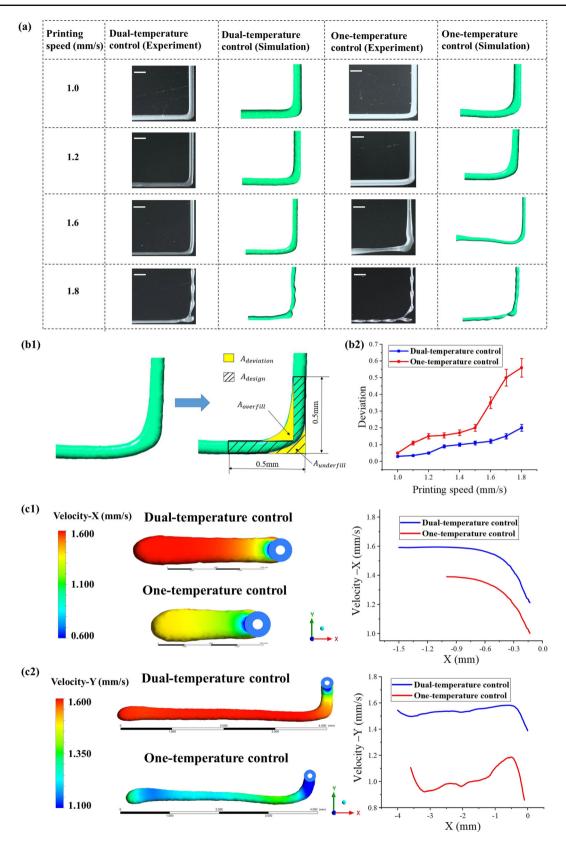


Fig. 5 Filament deposition at corners: a experimental and simulated results; b1 schematic diagram of the calculation of the deviation amount, scale bar=200 μ m, and b2 plot of deviation as a function of printing

speed. When printing speed v=1.6 mm/s: **c1** velocity in the *x*-direction when t=1 s and **c2** velocity in the *y*-direction when t=2.8 s



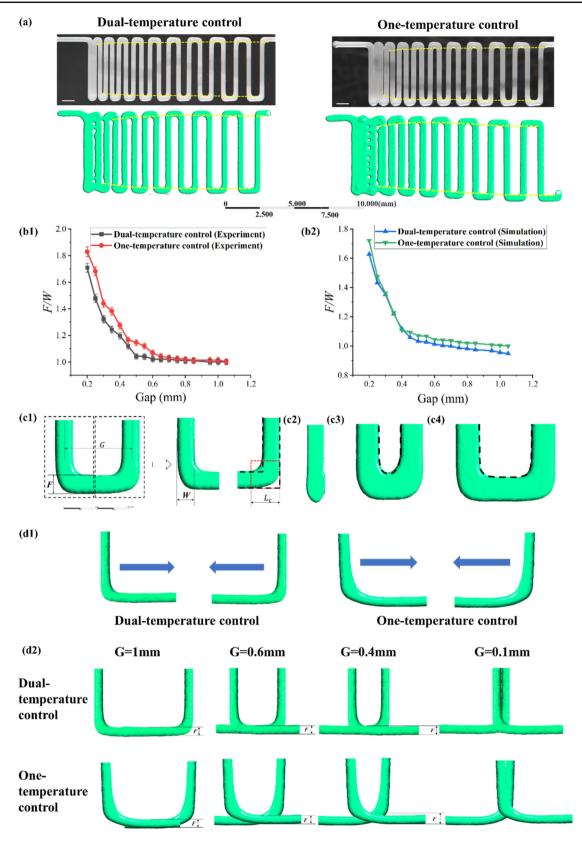


Fig. 6 Exploration of the filament fusion test: **a** experimental and simulated results when printing speed v=1.0 mm/s, scale bar=300 μ m; F/W as a function of gap with dual- and one-temperature control: **b1** experimental results and **b2** simulated results; **c1** exploded view, **c2** complete

fusion between filaments, ${\bf c3}$ fusion between filaments at the corners, ${\bf c4}$ no fusion between filaments; ${\bf d1}$ schematic diagram of filament fusion test explained by deposition at corners and ${\bf d2}$ filament fusion test with gap changes



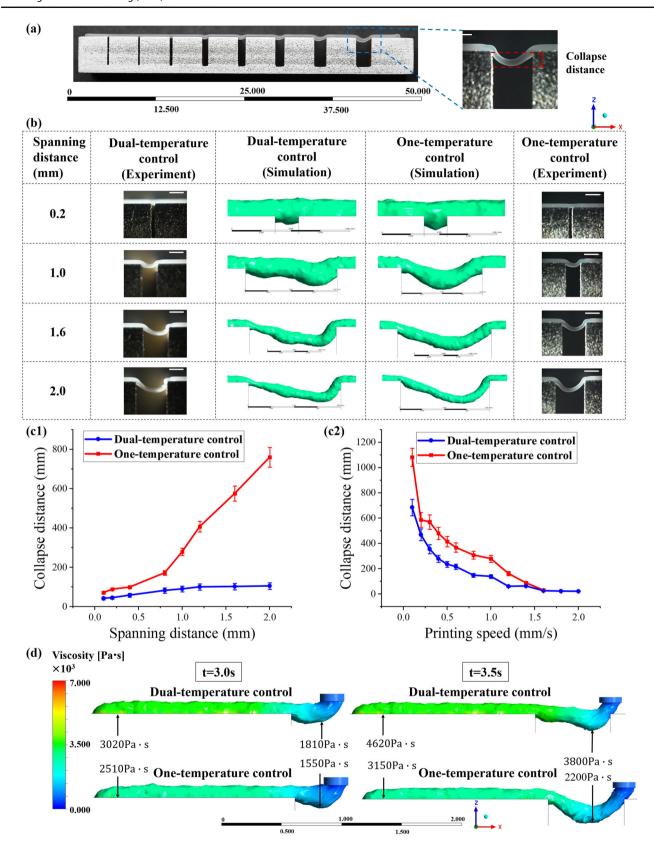


Fig. 7 Exploring the mechanism of filament collapse: a substrate model, scale bar=600 $\mu m;\,b$ results based on dual- and one-temperature control, scale bar=1 mm; c1 when printing speed=1.0 mm/s, the collapse distance changes with the spanning distance; c2 when spanning distance=1.0 mm, the collapse distance changes with the speed of the

substrate (the blue line represents the result with dual-temperature control, and the red line represents the result with one-temperature control); **d** viscosity contours at different times



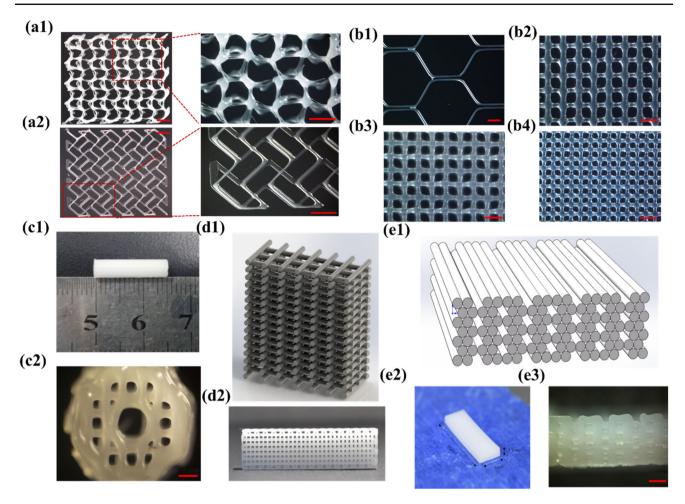


Fig. 8 Structures printed with dual-temperature control: a1 hexagonal architecture, scale bar=600 μm and a2 zigzag architecture, scale bar=600 μm ; b1 hexagonal scaffold with one layer when the printing speed is 1.0 mm/s, scale bar=200 μm . Rectangular scaffold when printing speed is b2 0.2 mm/s, b3 0.3 mm/s, and b4 0.5 mm/s, scale bar=200 μm ; c1 length of the tube scaffold and c2 micrograph of the

cross section of the tube scaffold, scale bar=500 μ m; **d1** schematic diagram of the square scaffold and **d2** image of the PCL square scaffold; **e1** schematic of the PCL tendon scaffold, **e2** image of the PCL tendon scaffold, and **e3** micrograph of the PCL tendon scaffold, scale bar=600 μ m. PCL: poly(ϵ -caprolactone)

Demonstration of dual-temperature control FDM

All structures in Fig. 8 are printed by the dual-temperature control FDM. Figure 8a1 shows the structure of the hexagonal network, and Fig. 8a2 shows the architecture formed by zigzag mesh (inner diameter of the nozzle D=100 μ m). Figure 8b also shows the PCL scaffolds printed by the dual-temperature control FDM using the nozzle with an inner diameter of 60 μ m. The hexagonal scaffold was printed when the printing speed was 1.0 mm/s (Fig. 8b1). Figure 8c shows a tube scaffold with a diameter of 3.8 mm and a height of 15 mm (inner diameter of the nozzle D=200 μ m). Figure 8d is a PCL square scaffold (10 mm \times 5 mm \times 5 mm, D=10 μ m). Figure 8e shows the schematic diagram and image of the tendon scaffold (15 mm \times 5 mm \times 1.5 mm, D=200 μ m). The

dual-temperature control FDM shows precise manufacturing and spatially arranged constructs and is a highly efficient tool for generating bioengineered structures. Therefore, dual-temperature control FDM significantly extends 3D printing in future scaffold applications.

Conclusions

Scaffolds with high porosity and specific surface area used in tissue engineering significantly influence the physiological activities of cell survival, migration, proliferation, and differentiation. In this study, a dual-temperature control method is proposed to improve the resolution of FDM printing. The deposited PCL filament with a width of 50 μ m is achieved by using a nozzle with an inner diameter of 100 μ m. More



importantly, a numerical model is developed by implementing the constitutive model, in which the viscosity is a function of temperature and shear rate, and both numerical and experimental investigations are carried out to study the influence of two different temperature control modes on viscosity. By comparing the results of 1D, 2D, and 3D PCL printing, the dual-temperature control mode provides a higher viscosity and lower velocity gradient of deposited PCL during FDM printing, leading to better printing with satisfactory accuracy. Therefore, the dual-temperature control FDM presents a promising application of PCL FDM in tissue engineering.

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Author contributions XDH and BZ performed 3D printing experiments; XDH, JY, and QLH developed numerical model; YH and JY wrote the original draft and revised the manuscript.

Declarations

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval This study does not contain any studies with human or animal subjects performed by any of the authors.

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