Review of Polymer Solutions for Near-Field Electrospinning with Spatial Control

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Abstract

Near-field electrospinning (NFES) is identified to be a technique able to fabricate polymer nano and micro fibers with accurate placement. In the past years (2006-2019), several polymer solutions have been successfully electrospun into fibers through several variants of the conventional NFES process. Each NFES variant intents to tailor the process parameters in order to improve the fibers' properties. This paper presents a review on the research and related development of electrospun fibers, emphasizing the used polymers, solvents, and fiber characteristics. Relevant summary of polymer solutions and near-field electrospinning processing conditions is provided in this paper.

1

Keywords: polymer, solvent, near-field electrospinning, NFES, fibers, spatial control

Contents

1	Intr	oduction	1
2	Poly	mer Solution and NFES Param-	
	eter	\mathbf{s}	2
	2.1	Applied Voltage	2
		2.1.1 Electric field	3
	2.2	Substrate	4
	2.3	Polymer Solution	4
3	NFI	ES Variants	5
4	Con	clusion	15
	Refe	erences	15

1. Introduction

Even though electrospinning is an old invention [1], it is currently a trending topic among researchers [2–4]. One of the reasons electrospinning is to be studied is its potential to fabricate

polymer nano-fibers from a variety of polymers. The technique allows the production of thin continuous fibers with ease, with diameters down to 3 nm in some cases, which is something difficult to achieve by other techniques. Furthermore, the basic setup can be modified with ease to fabricate different fibers with diversified functionalities with different materials. The produced fibers can be aligned or unaligned. Besides, the electrospinning equipment is inexpensive and of small size, compared to the equipment of standard spinning techniques. On the other hand, the understanding of the electrospinning process has improved in the last years [5].

The main components of the electrospinning technique are the fluid control unit (e.g. syringe pump) and a voltage power supply. The process also requires a target electrode or combination of electrodes on which the fibers can be collected. Figure 1 describes a typical near-field electrospinning set-up [5]. Two sub-techniques can be derived from electrospinning depending on the distance between the dispensing electrode and the collector. The process in which the electrospun jet can be controlled near the tip is called NFES or near-field electrospinning [6]. Moreover, if the

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distance between the collector and the dispensing needle is greater, the configuration is known as FFES or far-field electrospinning [7]. Near-field electrospinning is considered to be an outstanding technique to fabricate polymer fibers with spatial control and it has suffered several modifications to improve the precision and accuracy of the fiber deposition. This paper intents to collect the NFES variants of electrospunable polymer solutions with spatial control in recent research.

2. Polymer Solution and NFES Parameters

To spin nano fibers at close distances, the initial diameter of the jet is required to be as small as possible since stretching of the thread is limited. Kameoka et al (2003) demonstrated that a small initial spinning radius can be achieved using an atomic force microscope tip with a small polymer solution drop at the tip. Near-field electrospinning, has exhibited to be capable fabricate nano fibers over and nano fiber patterns (Sun et al 2006). Nevertheless, having a small polymer solution drop at the nozzle tip limits the length of the fibers that can be fabricated in a continuous manner. Using a spinneret with a reservoir (e.g. syringe) of solution generally produces fibers with diameter of a few micrometers [Gupta et al 2007; Xue et al 2014] since it creates a limit to which the nozzle inner diameter can be reduced to allow the solution to flow through. Coppola et al (2014) have showned a NFES variant that allows polymer nano fibers to be deposited directly from a polymer drop, averting the issue of nozzle clogging. The fibers are also prone soaking after deposition thus giving the fibers a semi-circular cross-section as depicted in Xue et al's (2014) work. The thinnest nozzles in literature so far are about 100 μm in diameter, for instance Chang et al (2008) used a 100 µm inner diameter needle tip to electrospin poly(ethylene oxide) (PEO) and Camillo et al (2013) used a µm-diameter tip Tungsten spinneret in a 26 gauge needle to electrospin co-polymer, poly[2-methoxy-5-(2ethylhexyloxy)-1,4-phenylenevinylene (MEH-PPV) with poly(ethylene oxide) (PEO).

2.1. Applied Voltage

Further development in the near field electrospinning spinning process have attempted to improve fiber deposition precision and reducing fiber diameter. Camillo et al (2013) was able to fabricate 100 nm diameter fiber at an applied voltage of 1.5 kV and a tip to collector distance of 500 µm using a modified fine tip spinneret. Separate reports by Chang et al (2008) and Bisht et al (2011) have shown that higher voltage leads to a significant increase in the fiber diameter (in the micrometer range) and loss of jet stability. The remedy is to significantly reduce the voltage used in the electrospinning process to about 200 to 600 V with tip to collector distance at about 0.5 to 1 mm. However, the charges on the solution drop at the tip of the needle were insufficient to break free from the surface tension to initiate electrospinning without assistance. Chang et al (2008) used a tungsten probe tip and Bisht et al (2011) used a glass microprobe tip (1 to 3 µm tip diameter) to mechanically draw the solution at the tip of the needle to initiate electrospinning. In the study by Chang et al (2008), reduction of electrospinning voltage from 1.5 kV (at tip to collector distance of 500 μm) to 600 V reduces the fiber diameter from 3 μm to 50 nm. Using a lower voltage of 200 V with tip to collector distance of 1 mm, Bisht et al (2011) was able to pattern nanofibers (polyethylene oxide) with diameter below 20 nm. Similar to electrospinning with longer tip to collector distance, it is likely that there is an optimum voltage which the fiber diameter obtained will be at its finest. Voltage higher or lower than this value will see an increase in the fiber diameter. Song et al (2015) showed that when the voltage for electrospinning polystyrene was increased from 400 to 500V, at a tip to collector distance of 20 µm, the fiber diameter reduced from close to 160 nm to about 60 nm. Such fiber diameter response to voltage is due to a balance of stretching of the jet and the speed at which it hit the collector. While increasing voltage causes greater stretching which reduces the fiber diameter, this also causes greater jet acceleration where the stretching terminates when the jet hit the collector.

To use a low working voltage in near field elec-

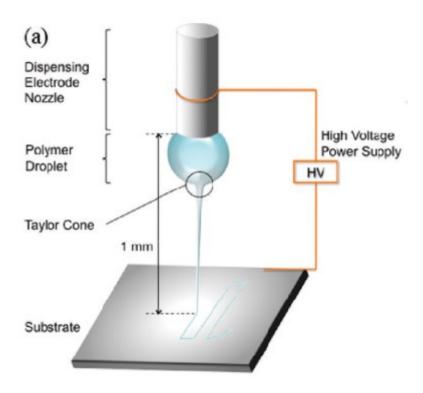


Figure 1: Typical near-field electrospinning set-up [8].

trospinning while eliminating the need to use a physical object to initiate electrospinning, an alternative is to use a higher voltage for initiation of electrospinning and switch to a lower voltage once the jet has erupted from the nozzle. Huang et al (2014) used this concept with a movable stage collector to produce ordered patterns with interfiber pitch of 50 µm. By controlling the height between the nozzle tip and the collector stage and the speed of stage, fibers with different orientation and cross-sectional shape can be obtained. Generally, closer distance between nozzle tip and collector (ranging from 0.5 mm to 2 mm) results in flat fibers due to impaction of the electrospinning jet. A limitation of the setup is that the landing point at electrospinning jet initiation cannot be determined although subsequent adjustment can be made after the jet has landed or the structure can be built up based on the displacement relative to the landing point. To control the landing point of the electrospinning jet, a target point may be set.

Low voltage, near field electrospinning has shown characteristics that differ from conventional near field electrospinning using higher voltage. Fiber diameter has already been shown to be smaller using this technique. With low voltage, near field electrospinning, the fiber diameter was found to be sensitive to the collector stage movement due to mechanical stretching; low velocity giving rise to larger fiber diameter and vice versa [Bisht et al 2011]. Instead of very fine spinneret tip, Bisht et al (2011) showed that it is possible to spin fibers with diameter less than 100 nm using a 27 gauge needle (approx. 200 µm inner diameter).

2.1.1. Electric field

Electrospinning jet can be very sensitive to variation in electric field. Thus a target with electric field profile that attracts the jet may be used to guide the electrospinning jet towards the desired landing point. Bisht et al (2011) demonstrated the precision and accuracy of low voltage, near field electrospinning by suspending fiber across carbon post with diameter of 30 µm and interpostal distance of 100 µm. Min et al (2013) used near field electrospinning to deposit semiconducting poly(3-hexylthiophene) (P3HT):PEO-blend organic nanowire over multiple field-effect transistors on a flexible polyarylate

substrate at a speed of 13.3 cm/s with regular spacing of 50 µm and fiber diameter of 289 nm. They have also demonstrated the ability to spin highly aligned nanowires from other materials such as poly(9-vinyl carbazole) (PVK) and poly{[N,N'-bis(2-octyldodecyl)-naphthalene-1,4,5,8-bis(dicarboximide)-2,6-diyl]-alt-5,5'-(2,2'-bithiophene)}.

The combination of near-field electrospinning and a guiding electrode has the potential to obtain precise and accurate fiber deposition. Xu et al (2014) used a guiding electrode behind the collector to create a direct line from the nozzle tip to it. This significantly dampens the deviation of the electrospinning jet from its original path as a result of electrostatic repulsion from the preceding deposited nano fiber. Without the guiding electrode, the near-field electrospun fibers have a spread of 74 µm. With the guiding electrode, the spread was reduced to just 7 µm. This raises the possibility of building up 3D structures using electrospinning. Kim et al (2018) used inkjet printing with conductive Ag nanoparticles loaded ink to form patterns on a paper as a target for near field electrospun fibers. The conductive printed pattern served as a guiding electrode for the electrospinning jet. Poly(vinylidene fluoride) (PVDF) solution was electrospun from a height of 750 μm and a 150 µm offset from the edge of the pat-The sensitivity of the electrospinning jet towards the electric field can be seen as the fibers are stacked on the edge of the conductive pattern where the relative electric field was much higher at the edge than at its center. When the pattern lines formed acute angle, right-angle or obtuse angle, the accuracy of the deposited fibers are influenced by slight changes in the relative electric field. From acute angle to right-angle, the electric field singularity increases from the edge to the intersection between the lines. In this case, the fibers were stacked directly on the edge of the line and to the middle of the intersection. For lines forming obtuse angle, the deposited fibers followed the edge of the lines by veered off the line at the intersection.

2.2. Substrate

In near field electrospinning, one of the risks is electrical shorting due to the proximity of the charged needle tip to the grounded collector. Any electrical shorting will disrupt the electrospinning process and result in discontinuous fiber. While using a lower voltage may reduce this risk, an alternative is to use a less conductive collector. Liu et al (2014) used a rotating glass tube with a copper foil lining at the inner surface of the tube for the collection of electrospun oriented polyvinylidene fluoride (PVDF) fiber. While initial fiber alignment was excellent, the alignment starts to deteriorate after prolonged fiber deposition which can be attributed to the presence of residual charges.

The influence of residual charges on the precision of fiber deposited is more pronounced when an insulating surface and a conductive surface were used as collectors. Choi et al (2017) used a hydrophobic and insulating acrylic substrate as collector. To increase conductivity of selected region of the collector, plasma treatment was carried out to render those region hydrophilic. The collector was placed in a high humidity environment such that the hydrophilic region will be slightly conductive due to the presence of water molecule attached to it. Near field electrospinning of polyurethane showed that on the insulating hydrophobic surface, the fibers were twisted and curved due to weak electric field profile between the emitter and the collector surface and the inability of charges to escape. In contrast, on the hydrophilic region, the fiber were placed in accordance to the movement of the emitter relative to the collector. This shows that electrical charges on the electrospinning jet needs to escape for precise deposition and even an insulating substrate with slight conductivity is crucial for ordered fiber arrangement.

2.3. Polymer Solution

In electrospinning, it is generally agreed that with higher concentration, the diameter of the fibers increased due to greater viscosity which resist stretching. In near field electrospinning, similar observations have been reported where concentration increases, fiber diameter increased [Chang et al 2008; Zheng et al 2012]. However, in separate studies by Pan et al (2014) using poly(γ -benzyl α , l-glutamate) and Pan et al (2015) using polyvinylidene fluoride (PVDF) reported reduction in fiber diameter with increasing concentration. Pan et al (2015) attributed this to a higher charge accumulation in higher concentration PVDF solution. However, more studies need to be carried out to verify this.

3. NFES Variants

Low-Voltage NFES (LV NFES) [8]

Scanning Tip Electrospinning [9]

3D Electrospinning [10]

Electrohydro-dynamic 3D Print-patterning or Electrohydro-dynamic Jetting [11]

Multinozzle NFES [12–14]

Electrohydro-dynamic Writing or Mechanoelectrospinning (MES) [15]

Electrohydro-dynamic Direct-Write (EDW) [16]

Mechano-Electrospinning [17]

Suspension NFES [18]

Helix Electrohydro-dynamic Printing (HE-printing) [19]

Electrohydro-dynamic (EHD) jet printing [20]

Airflow-assisted Electrohydro-dynamic Directwriting (EDW) [21]

Tethered Pyro-Electrohydro-dynamic Spinning (TPES) [22]

Polymer(s)	Solvent(s)	NFES Variant	Process Parameters and Fiber Characterization	Ref.
Poly(ethylene ox-	Deionized wa-	Low-Voltage NFES	Solution Concentration: 1, 2, and 3 $wt\%$ PEO	[8]
ide) (PEO; $MW =$	ter	(LV NFES)	Nozzle: 27 gauge type 304; stainless steel needle	
4,000,000)			Solution deposition rate: lower than $1\mu L/h$	
•			Nozzle-to-substrate distance: 1mm	
			Substrate composition: Pyrolyzed SU-8 carbon and Si	
			Applied voltage: polymer jet initiated at 400-600 V and	
			dispensed at 200-400 V	
			x-y stage velocity: $10-40mm/s$	
			Fiber Diameter: 50-425nm	
			Distance between adjacent fibers: Not determined	
Poly[2-	acetonitrile	Typical NFES pro-	Solution Concentration:	[23]
methoxy-5-(2-	toluene mixture	cess	10mg of MEH-PPV in $2mL$ of toluene; $500mL$ of MEH-PPV	
ethylhexyloxy)-1,4-	(65/35); acetic		solution with $250mg$ of PEO in $3.5mL$ of acetonitrile; $500mL$	
phenylenevinylene]	acid toluene		of MEH-PPV solution with $250mg$ of PEO in $3mL$ of acetic	
(MEH-PPV; MW	(17/83); pure		acid / toluene (17 / 83). The resulting MEH-PPV/PEO	
= 380,000) with	toluene		concentration is 1:100	
Poly(ethylene ox-			Nozzle: mm-diameter tip Tungsten spinneret in a 26 gauge	
ide) (PEO; $MW =$			needle	
300,000)			Solution deposition rate: $50\mu L/h$	
			Nozzle-to-substrate distance: $500\mu m$	
			Substrate composition: SiO2/Si (oxide thickness = 800	
			nm)	
			Applied voltage: around $1.3kV$	
			x-y stage velocity: $50cm/s$	
			Fiber Diameter: 100nm	
			Distance between adjacent fibers: around $100\mu m$	

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6

Poly(ethylene	Water	Scanning Tip Elec-	Solution Concentration: 7wt% PEO	[9]
oxide) (PEO)		trospinning and	Nozzle: Needle outer diameter of $200\mu m$ and inner diame-	
, , ,		NFES	ter of $100\mu m$	
			Solution deposition rate: $0.1\mu L/h$	
			Nozzle-to-substrate distance: $500 \mu m$	
			Substrate composition: Not determined	
			Applied voltage: polymer jet initiated at $1.5 \ kV$ and dis-	
			pensed at $600V$	
			x-y stage velocity: $120mm/s$	
			Fiber Diameter: $709\pm131nm$; $49-74nm$ when applied volt-	
			age is $800V$	
			Distance between adjacent fibers: Not determined	
			Notes: 108m yield in 15min with a fiber diameter of	
			$709 \pm 131 nm$	
Poly(vinylidine flu-	N,N Dimethyl-	Helix Electrohydro-	Solution Concentration: 1.8 <i>g</i> PVDF in 4.1 <i>g</i> of DMF and	[19
orid) (PVDF)	formamide	dynamic Printing	4.1g of acetone. The resulting concentration is 18% PVDF.	
	(DMF)	(HE-printing)	Nozzle: Needle outer diameter of $510\mu m$ and inner diameter	
			of $260\mu m$	
			Solution deposition rate: $400nL/min$	
			Nozzle-to-substrate distance: 10-50mm	
			Substrate composition: Poly(dimethylsiloxane) (PDMS)	
			on Ecoflex	
			Applied voltage: $1.5-3kV$	
			x-y stage velocity: 0-400mm/min	
			Fiber Diameter: about $1.5-3\mu m$	
			Distance between adjacent fibers: Not determined	

Continued on next page

Polyhedral	Dimethyl	Electrohydro-	Solution Concentration: POSS-PCU and POSS-PCL-	[11]
Oligomeric	acetamide	dynamic 3D Print-	PCU used in $20\%w/w$ concentration in DMAC	[++]
Silsesquioxane-	(DMAC) and	patterning or	Nozzle: needle of 750 μm in diameter	
Poly(Carbonate-	1-Butanol	Electrohydro-	Solution deposition rate: less than $1\mu L/min$	
Urea)Urethane	1 D 0.001101	dynamic Jetting	Nozzle-to-substrate distance: about between $500\mu m$ to	
(POSS-PCU)			2mm	
and Polyhe-			Substrate composition: Not determined	
dral Oligomeric			Applied voltage: $8.0-10.0kV$	
Silsesquioxane			x-y stage velocity: $10mm/s$	
Poly(Caprolactone-			Fiber Diameter: $5-50\mu m$	
Poly(Carbonate-			Distance between adjacent fibers: $250\mu m$	
Urea)Urethane)			· ·	
(POSS-PCL-PCU)				
Poly(ethylene	Distilled water	Electrohydro-	Solution Concentration: $6wt\%$ PEO	[15]
oxide) (PEO)		dynamic Writing	Nozzle: Not determined	
		or Mechanoelectro-	Solution deposition rate: $1200nL/min$	
		spinning (MES)	Nozzle-to-substrate distance: 7.5mm	
			Substrate composition: Not determined	
			Applied voltage: polymer jet initiated at $2 kV$ and dis-	
			pensed at $0.8-1kV$	
			x-y stage velocity: around $400mm/s$	
			Fiber Diameter: 200-350nm	
			Distance between adjacent fibers: $5\mu m$	
Poly(ethylene	Deionized wa-	Airflow-assisted	Solution Concentration: $8wt\%$ PEO	[21]
oxide) (PEO)	ter and the	Electrohydro-	Nozzle: Outer airflow passage diameter: 1mm Airflow	
	ethanol with a	dynamic Direct-	gas pump pressure: $25kPa$ Inner liquid passage diameter:	
	volume ratio of	writing (EDW)	0.21mm	
	3:1		Solution deposition rate: $30\mu L/h$	
			Nozzle-to-substrate distance: 2mm	
			Substrate composition: Silicon	
			Applied voltage: about $2kV$	
			x-y stage velocity: 1-20mm/s	
			Fiber Diameter: $3.73 \pm 1.37 \mu m$	
			Distance between adjacent fibers: $5.13 \pm 6.67 \mu m$	

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Table 1 continued				
Poly(Vinylidene	Acetone and	3D Electrospinning	Solution Concentration: $17wt\%$ PVDF; $1.7g$ of PVDF,	[10]
Fluoride) (PVDF)	Dimethyl Sul-		5g of acetone, $0.5g$ of Capstone FS-66, $5g$ of DMSO	
	foxide (DMSO)		Nozzle: Needle inner diameter of $100\mu m$	
			Solution deposition rate: $14 nL/min$	
			Nozzle-to-substrate distance: $750\mu m$	
			Substrate composition: A4 size commercial printing pa-	
			per (Double A)	
			Applied voltage: $1.9kV$	
			x-y stage velocity: $10mm/s$	
			Fiber Diameter: Not determined	
			Distance between adjacent fibers: Not determined	
Poly(9-Vinyl Car-	Styrene	Typical NFES pro-	Solution Concentration: 3.96wt% PVK in styrene	[24]
bazole) (PVK)		cess	Nozzle: Needle inner diameter of $100\mu m$	
			Solution deposition rate: $500nL/min$	
			Nozzle-to-substrate distance: around 2.5mm	
			Substrate composition: Si/SiO2	
			Applied voltage: $3-4kV$	
			x-y stage velocity: $13.3cm/s$	
			Fiber Diameter: $289.26 \pm 35.37nm$	
			Distance between adjacent fibers: $50\mu m$	
			Notes: 15m yield in 2min	
Polystyrene (PS)	1,2,4-Trichloro	Electrohydro-	Solution Concentration: 1 to $5wt\%$ PS	[20]
	benzene	dynamic (EHD) jet	Nozzle: Glass nozzle inner diameter of $2\mu m$ and outer di-	
		printing	ameter of $2.66\mu m$	
			Solution deposition rate: Si	
			Nozzle-to-substrate distance: 20, 30, $40\mu m$	
			Substrate composition:	
			Applied voltage: $500 \text{ to } 400V \text{ in } 25V \text{ increments}$	
			x-y stage velocity: $0.01-10mm/s$	
			Fiber Diameter: about $60-170\mu m$	
			Distance between adjacent fibers: Not determined	
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9

June 24, 2019

Table 1 continued				
Poly(ethylene	Not determined	Typical NFES pro-	Solution Concentration: $3wt\%$ PEO	[25]
oxide) (PEO)		cess	Nozzle: Not determined	
			Solution deposition rate: Not determined	
			Nozzle-to-substrate distance: $500\mu m$	
			Substrate composition: Si	
			Applied voltage: $1000V$	
			x-y stage velocity: $20cm/s$	
			Fiber Diameter: 300nm	
			Distance between adjacent fibers: $25\mu m$	
Poly(ethylene	Distilled water	Multinozzle NFES	Solution Concentration: $5wt\%$	[12]
oxide) (PEO)			Nozzle: four-nozzle and six-nozzle array with needle spacing	
			changes from $1.5mm$ to $3.5mm$	
			Solution deposition rate: $1-3\mu L/min$	
			Nozzle-to-substrate distance: 2mm	
			Substrate composition: Not determined	
			Applied voltage: $1.7-2.7kV$	
			x-y stage velocity: Not determined	
			Fiber Diameter: $5.47\mu m$	
			Distance between adjacent fibers: 3-5 mm	
Poly(ethylene	Distilled water	Multinozzle NFES	Solution Concentration: $5wt\%$	[13]
oxide) (PEO)			Nozzle: Dual-28G-needle array with needle inner diameter	
			of $0.18mm$ and outer diameter of $0.36mm$; with needle spac-	
			ing changes from $2.0mm$ to $3.0mm$	
			Solution deposition rate: $0.2\mu L/min$	
			Nozzle-to-substrate distance: 3.0-4.0mm	
			Substrate composition: Not determined	
			Applied voltage: $2.0-3.0kV$	
			x-y stage velocity: $20mm/s$	
			Fiber Diameter: Not determined	
			Distance between adjacent fibers: $218-326\mu m$	

Table 1 continued				
Poly(ethylene	Distilled water	Multinozzle NFES	Solution Concentration: $5 wt\%$	[14]
oxide) (PEO)			Nozzle: Dual-28G-needle array with needle inner diameter	
			of $180\mu m$ and outer diameter of $360\mu m$; with needle spacing	
			changes of $2.0mm$	
			Solution deposition rate: $0.2\mu L/min$	
			Nozzle-to-substrate distance: 4.0mm	
			Substrate composition: chromium-plated glass	
			Applied voltage: $2.5kV$	
			x-y stage velocity: $20mm/s$	
			Fiber Diameter: Not determined	
			Distance between adjacent fibers: 2.3002-2.7224mm	
Poly(ethylene	Not determined	Typical NFES pro-	Solution Concentration: $2wt\%$	[26]
oxide) (PEO)		cess	Nozzle: G30 needle with inner diameter of $0.15mm$	
			Solution deposition rate: Not determined	
			Nozzle-to-substrate distance: 1-3mm	
			Substrate composition: Silicon	
			Applied voltage: $1250V$	
			x-y stage velocity: Not determined	
			Fiber Diameter: Not determined	
			Distance between adjacent fibers: $20\mu m$	
Gelatin	Acetic Acid and	Typical NFES pro-	Solution Concentration: $11wt\%$ gelatin, $30wt\%$ water,	[27]
(porcine skin)	Ethyl Acetate	cess	35.4wt% acetic acid, $23.6wt%$ ethyl acetate	
			Nozzle: 19G needle tip with outer diameter of 1.08mm	
			Solution deposition rate: Not determined	
			Nozzle-to-substrate distance: 1.25mm	
			Substrate composition: Poly(Dimethylsiloxane) (PDMS)	
			films	
			Applied voltage: $1000V$	
			x-y stage velocity: Not determined	
			Fiber Diameter: around $2-3\mu m$	
			Distance between adjacent fibers: $40\mu m$	

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June 24, 2019

Table 1 continued				
Poly(ethylene	Water/Ethanol	Typical NFES pro-	Solution Concentration: PEO concentrations of 16% adn	[28]
oxide) (PEO)	$(\mathrm{v}/\mathrm{v}=60/40)$	cess	18%	
			Nozzle: $40\mu m$	
			Solution deposition rate:	
			Nozzle-to-substrate distance: 1mm	
			Substrate composition: Planar silicon	
			Applied voltage: $1.7kV$	
			x-y stage velocity: $0.36m/s$	
			Fiber Diameter: $5.15\mu m$	
			Distance between adjacent fibers: Not determined	
Poly(ethylene	Water/Ethanol	Electrohydro-	Solution Concentration: 14wt% PEO	[16]
oxide) (PEO)	$(\mathrm{v}/\mathrm{v}=3/1)$	dynamic Direct-	Nozzle: Stainless needle with inner diameter of $210\mu m$ and	
		Write (EDW)	outer diameter of $400\mu m$	
			Solution deposition rate: $50\mu L/h$	
			Nozzle-to-substrate distance: 2mm	
			Substrate composition: Poly(ethylene terephthalate)	
			(PET)	
			Applied voltage: $3kV$	
			x-y stage velocity: $700mm/s$	
			Fiber Diameter: $15-35\mu m$	
			Distance between adjacent fibers: $70\mu m$	
Poly(ethylene	Deionized wa-	Mechano-	Solution Concentration: $3wt\%$ PEO	[17]
oxide) (PEO)	ter	Electrospinning	Nozzle: Stainless steel nozzle with inner diameter of $160\mu m$	
			and outer diameter of $310\mu m$	
			Solution deposition rate: $50nL/min$	
			Nozzle-to-substrate distance: 2-5mm	
			Substrate composition: Silicone	
			Applied voltage: polymer jet initiated at $2kV$ and dis-	
			pensed at $1kV$	
			x-y stage velocity: $200-400mm/s$	
			Fiber Diameter: from 344 ± 32 to $214\pm27nm$	
			Distance between adjacent fibers: Not determined	

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Table 1 continued Poly(co-Glycolic)	Dimethyl Car-	Tethered Pyro-	Solution Concentration: Not determined	[22]
acid (PLGA)	bonate (DMC)	Electrohydro-	Nozzle: nozzle-free	
acid (LGA)	bollate (DMC)	dynamic Spinning	Solution deposition rate: The drop reservoir is placed	
		(TPES)	directly on a flat substrate	
		(IFES)	Nozzle-to-substrate distance: Taylor's cone is focused	
			and put in direct contact with the collector	
			-	
			Substrate composition: Poly(tetrafluoroethylene) (PTFE) coated glass slide	
			, , ,	
			Applied voltage: pyro-electric field of between 2.7 $x10^7 V/m$ and $5.5x10^7 V/m$	
			x-y stage velocity: Not determined	
			Fiber Diameter: 304.7nm	
			Distance between adjacent fibers: Not determined	
Poly(ethylene ox-	N,N Dimethyl-	Typical NFES pro-	Solution Concentration: SU-8/PEO/TBF blend with	[6]
ide) (PEO) with	formamide	cess	0.75wt% PEO, $1wt%$ TBF; the blend is diluted with $30vol%$	[O]
Tetrabutylammo-	(DMF)	CCSS	DMF	
nium tetrafluorob-	(DIVII)		$\mu m \mu m$	
orate (TBF) and			Solution deposition rate: Not determined	
SU-8 2002			Nozzle-to-substrate distance: Not determined	
			Substrate composition: Brass disk with a diameter of	
			38mm	
			Applied voltage: 980V	
			x-y stage velocity: Not determined	
			Fiber Diameter: Not determined	
			Distance between adjacent fibers: Not determined	

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4. Conclusion

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References

- [1] F. Anton, Process and apparatus for preparing artificial threads (1930). doi:https://patents.google.com/?q=D01D5%2f0076.
- [2] Z.-M. Huang, Y. Z. Zhang, M. Kotaki, S. Ramakrishna, A review on polymer nanofibers by electrospinning and their applications in nanocomposites, Composites Science and Technology 63 (15) (2003) 2223–2253. doi:10.1016/S0266-3538(03)00178-7.
- [3] D. H. Reneker, A. L. Yarin, Electrospinning jets and polymer nanofibers, Polymer 49 (10) (2008) 2387–2425. doi:10.1016/J.POLYMER.2008.02.002.
- [4] J. D. Schiffman, C. L. Schauer, A Review: Electrospinning of Biopolymer Nanofibers and their Applications, Polymer Reviews 48 (2) (2008) 317–352. doi:10.1080/15583720802022182.
- [5] Q. Li, Chapter 7: Liquid Crystal-Functionalized Nano- and Microfibers Produced by Electrospinning - Liquid Crystals Beyond Displays: Chemistry, Physics, and Applications, John Wiley & Sons, 2012. doi:9781118078617.
- [6] A. Cisquella-Serra, M. Magnani, Álvaro Gual-Mosegui, S. Holmberg, M. Madou, M. Gamero-Castaño, Study of the electrostatic jet initiation in near-field electrospinning, Journal of Colloid and Interface Science 543 (2019) 106–113. doi:10.1016/J.JCIS.2019.02.041.
- [7] S. K. Nataraj, K. S. Yang, T. M. Aminab-Polyacrylonitrile-based nanofibers—A havi, state-of-the-art review, Progress in Poly-37 (2012)Science (3)487 - 513.mer doi:10.1016/J.PROGPOLYMSCI.2011.07.001.
- [8] G. S. Bisht, G. Canton, A. Mirsepassi, L. Kulinsky, S. Oh, D. Dunn-Rankin, M. J. Madou, Controlled Continuous Patterning of Polymeric Nanofibers on Three-Dimensional Substrates Using Low-Voltage Near-Field Electrospinning, Nano Letters 11 (4) (2011) 1831–1837. doi:10.1021/nl2006164.
- [9] C. Chang, K. Limkrailassiri, L. Lin, Continuous near-field electrospinning for large area deposition of

- orderly nanofiber patterns, Appl Phys Lett (2008) 3doi:10.1063/1.2975834.
- [10] J. Kim, B. Maeng, J. Park, Characterization of 3D electrospinning on inkjet printed conductive pattern on paper, Micro and Nano Systems Letters 6 (1) (2018) 12. doi:10.1186/s40486-018-0074-1.
- [11] A. Gupta, A. M. Seifalian, Z. Ahmad, M. J. Edirisinghe, M. C. Winslet, Novel Electrohydrodynamic Printing of Nanocomposite Biopolymer Scaffolds, Journal of BIOACTIVE AND COMPATIBLE POLYMERS 22 (2007). doi:10.1177/0883911507078268.
- [12] H. Wang, S. Huang, F. Liang, P. Wu, M. Li, S. Lin, X. Chen, Research on Multinozzle Near-Field Electrospinning Patterned Deposition, Journal of Nanomaterials 2015 (2015) 1–8. doi:10.1155/2015/529138.
- [13] Z. Wang, X. Chen, J. Zeng, F. Liang, P. Wu, H. Wang, Controllable deposition distance of aligned pattern via dual-nozzle near-field electrospinning, AIP Advances 7 (3) (2017) 035310. doi:10.1063/1.4974936.
- [14] Z. Wang, X. Chen, J. Zhang, Y.-J. Lin, K. Li, J. Zeng, P. Wu, Y. He, Y. Li, H. Wang, Fabrication and evaluation of controllable deposition distance for aligned pattern by multi-nozzle near-field electrospinning, AIP Advances 8 (7) (2018) 075111. doi:10.1063/1.5032082.
- [15] Y. Huang, Y. Duan, Y. Ding, N. Bu, Y. Pan, N. Lu, Z. Yin, Versatile, kinetically controlled, high precision electrohydrodynamic writing of micro/nanofibers, Scientific Reports 4 (1) (2015) 5949. doi:10.1038/srep05949.
- [16] J.-Y. Zheng, H.-Y. Liu, X. Wang, Y. Zhao, W.-W. Huang, G.-F. Zheng, D.-H. Sun, Electrohydrodynamic Direct-Write Orderly Micro/Nanofibrous Structure on Flexible Insulating Substrate, Journal of Nanomaterials 2014 (2014) 1–7. doi:10.1155/2014/708186.
- [17] N. Bu, Y. Huang, X. Wang, Z. Yin, Materials and Manufacturing Processes Continuously Tunable and Oriented Nanofiber Direct-Written by Mechano-Electrospinning Continuously Tunable and Oriented Nanofiber Direct-Written bv Mechano-Electrospinning (2012).doi:10.1080/10426914.2012.700145.
- [18] A. R. Nagle, C. D. Fay, Z. Xie, G. G. Wallace, X. Wang, M. J. Higgins, A direct 3D suspension nearfield electrospinning technique for the fabrication of polymer nanoarrays, Nanotechnology 30 (19) (2019) 195301. doi:10.1088/1361-6528/ab011b.
- [19] Y. Duan, Y. Ding, Z. Xu, Y. Huang, Z. Yin, Helix Electrohydrodynamic Printing of Highly Aligned Serpentine Micro/Nanofibers., Polymers 9 (9) (sep 2017). doi:10.3390/polym9090434.
- [20] C. Song, J. A. Rogers, J.-M. Kim, H. Ahn, Patterned polydiacetylene-embedded polystyrene

- nanofibers based on electrohydrodynamic jet printing, Macromolecular Research 23 (1) (2015) 118–123. doi:10.1007/s13233-015-3024-2.
- [21] J. Jiang, X. Wang, W. Li, J. Liu, Y. Liu, G. Zheng, J. Jiang, X. Wang, W. Li, J. Liu, Y. Liu, G. Zheng, Electrohydrodynamic Direct-Writing Micropatterns with Assisted Airflow, Micromachines 9 (9) (2018) 456. doi:10.3390/mi9090456.
- [22] S. Coppola, V. Vespini, G. Nasti, O. Gennari, S. Grilli, M. Ventre, M. Iannone, P. A. Netti, P. Ferraro, Tethered Pyro-Electrohydrodynamic Spinning for Patterning Well-Ordered Structures at Microand Nanoscale, Chem. Mater 26 (2014) 3360. doi:10.1021/cm501265j.
- [23] D. D. Camillo, V. Fasano, F. Ruggieri, S. Santucci, L. Lozzi, A. Camposeo, D. Pisignano, Near-field electrospinning of conjugated polymer light-emitting nanofibers, Nanoscale 5 (2013) 11637–11642. doi:10.1039/C3NR03094F.
- [24] S.-Y. Min, T.-S. Kim, B. J. Kim, H. Cho, Y.-Y. Noh, H. Yang, J. H. Cho, T.-W. Lee, Large-scale organic nanowire lithography and electronics, Nature Communications 4 (1) (2013) 1773. doi:10.1038/ncomms2785.
- [25] D. Sun, C. Chang, S. Li, L. Lin, Near-Field Electrospinning (2006). doi:10.1021/nl0602701.
- [26] J. Xu, M. Abecassis, Z. Zhang, P. Guo, J. Huang, K. Ehmann, J. Cao, Accuracy Improvement of Nano-fiber Deposition by Near-Field Electrospinning, International Workshop on Microfactories IWMF2014 (9th) (2014).
- [27] N. Xue, X. Li, C. Bertulli, Z. Li, A. Patharagulpong, A. Sadok, Y. Y. S. Huang, Rapid Patterning of 1-D Collagenous Topography as an ECM Protein Fibril Platform for Image Cytometry, PLoS ONE 9 (4) (2014) e93590. doi:10.1371/journal.pone.0093590.
- [28] G. Zheng, W. Li, X. Wang, D. Wu, D. Sun, L. Lin, Precision deposition of a nanofibre by nearfield electrospinning, Journal of Physics D: Applied Physics 43 (41) (2010) 415501. doi:10.1088/0022-3727/43/41/415501.
- [29] D. Shin, J. Kim, J. Chang, Experimental study on jet impact speed in near-field electrospinning for precise patterning of nanofiber, Journal of Manufacturing Processes 36 (2018) 231–237. doi:10.1016/J.JMAPRO.2018.10.011.