## Electrospun Aligned Polyethylene Oxide Nanofibers

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Abstract: An novel and facile electrospinning method, in which two parallel grounded metallic conductors are introduced to change electric field as auxilliary electrodes besides collector, is used to prepare oriented polyethylene oxide (PEO) nanofibers. SEMs of experiment results show that aligned nanofibers which d2 ameters are around 200~ 490 nm can be achieved by this one2step method, and the diameters of nanofibers decrease at first when the distance is smaller than 2 mm, then increase if the distance is larger than 2 mm with the increasing distance between two parallel conductors as the combination of streetching sufficiency and decreasing electric field strength. It also can be verified that the aligned nanofibers mainly caused by changed electrostatic field and not by structure.

Key words: electrospinning; aligned nanofibers; polyethylene oxide; auxilliary electrode EEACC: 0560

# 电纺定向聚氧化乙烯纳米纤维

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**摘 要**: 采用一种新颖的纺丝工艺制备定向聚氧化乙烯纳米纤维. 该工艺就是将两个平行的接地金属导体置于收集板的两侧, 电镜图实验结果表明采用该一步法可以制得 200~490nm 左右的 PEO 定向纳米纤维, 而且随着电极之间距离的增大, 在 拉伸的充分性和电场的共同作用下, 纳米纤维先减小后增大. 实验还证实定向纤维是电场而非结构作用的结果. 关键词: 电纺丝; 定向纤维; 聚氧化乙烯

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Due to their remarkable characteristics of high sur2 face surface to volume ratio, unique optics, chemistry and mechanics<sup>[1]</sup> etc, polymer nanofibers, especially on& dimensional nanofibers, will play important roles in fu2 ture applications, such as reinforced material, high per2 formance filters<sup>[2]</sup>, high2sensitivity chemical sensor<sup>[3]</sup>, tissue scaffolds<sup>[425]</sup>, drug delivery<sup>[6]</sup>, and so on Elec2 trospinning which can be dated back in 1934<sup>[7]</sup> by Form2 hals A., is a straight, cost2efficient and convenient technology to fabricate polymer nanofibers. In a typical procedure, a high direct2current voltage is applied to a metallic needle, which is connected to a syringe filled with a polymer solution with enough viscosity and con2 ductivity. With the high voltage increasing, the electro2 static forces will overcome the surface tension of the poD ymer droplet suspended beneath nozzle of the needle to cause ejection of a fine jet. Undergoing solvent evapora2 tion, stretching and whipping instability, the charged jet turns into many continuous nanometer size fibers when reaching the grounded metallic collector.

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Unfortunately, the nanofibers deposited on the collector are always randomly deposited and act as non2 waven mat. As one2dimensional nanostructures are i2 deal building blocks for hierarchical assembly of func2 tional nanoscale electronic and photonic structure that could overcome fundamental and economic limitations of lithography based fabrication<sup>[8]</sup>, many researchers all over the world have begun to mainly pay their em2 phasises on how to achieve aligned nanofibers for fur2 ther integration by electrospinning. Although the abili2 ty to obtain oriented nanofibers remains a major chal2 lenge in the field, some progress has been made and there are some literatures to cover  $it^{[\mathfrak{D}22]}$ . A classical report example was reported that a thin rotating wheel with sharp edge was used to collect aligned nanofibers in<sup>[9].</sup> The sharp edge is used to change applied electric field, which dominates the electrospinning process. However, the vital disadvantages of the setup are that the sharp edge is difficult to fabricate and the speed of wheel is hard to set because the speed of nanofibers is nonlinear and difficult to determine. Another technique is that 7, 500V AC potential with 60Hz frequency is used instead of DC potential by Royal Kessick et al<sup>[10]</sup>. But the fibers orientation degree is only to some ex2 tent. Consequently, in this paper, a novel technique with electrospinning process is brought out, which is much easier to fabricate massively than previous work.

#### 1 Experimental

Polyethylene oxide (PEO, average molecular weight M= 300, 000 g/ mol) purchased from Dadi Fine Chemical Co., Ltd. (Changchun, China) was used to prepare solutions that was used as the working fluid. PEO was dissolved in water and ethanol/ water solvent at different concentrations. All solutions were stored at room temperature. All experiments were performed with a setup showed as Fig. 1 at room temperature in air. The solutions were stored in the syringe. The anode of high po2 tential power supply (DW2P40321AC, Tianjin Dongwen) (The voltage carried out is 9,000V) is connected to the needle of syringe which inner di2 ameter is about 0.3 mm and the cathode is connect2 ed to the metallic conductors, which are on the quartz substrate collector. A metallic plate, which is parallel to collector, is connected to needle to make electric field much more even. The distance between the end of needle and two near electrodes is about 9cm. Both cathodes are well grounded. When the high voltage source is on, the solutions can be pulled from the needle by electric field and turned into nanofibers collected on the regions of the quartz substrate and two near electrodes. Elec2 tron Micrographs of the electrospun nanofibers were obtained by Germany LEO1530 scanning elec2 tron microscope and XL30ESEM.



Fig. 1 Schematic diagram of the system setup

#### 2 Results and discussion

As we all know, concentration, which affects viscosity, volume charge density, is one of the most important parameters in electrospinning process <sup>[13]</sup>. Drastic morphological changes were observed when concentration of polymeric solution was changed. From Fig. 2, it can be found that when concentration is smaller than 8%, most of



Fig. 2 SEM of fibers of (a), (b), (c) and (d) collected by conventional electrospinning from concentrations of 8%, 12%, 15% and 18% respectively in pure water.
them are beads and few fibers exist due to its low viscosity of solution. With the concentration in2 creasing, more fibers come into being and beads disappear gradually. The SEM image of nanofibers

in Fig. 2 show that in order to achieve uniform nanofibers, the concentration should be 18% at least when solvent is pure water. But if there are some ethonal, the concentration can be much more smaller. So in the following experiments the solu2 tion of 15% (w/w) in 1/4 ethonal/H<sub>2</sub>O is used as working solution.

Now the orientation morphology of nanofiers is studied. Scanning electron micrographs of nano2 fibers deposited by setup of Fig. 1 are shown in Fig. 3. The images clearly illustrate that the nano2 fibers are well aligned, which are perpendicular to the edge of the trench between two parallel grouned conductors. The results are caused by e2 lectrostatic field, which is different to that of the traditional eletrospinning. As shown FEA diagram in Fig. 4a, the electric field forces on the trench be2 tween two conductors are split into two fractions perpendicular to the edge of the conductors. Fig. 4b illustrates the electrostatic forces acting on a charged fiber. The induced charges on the elec2 trodes mainly caused by high electro field are nega2 tive, so the electric forces F<sub>1</sub> and F<sub>2</sub> are downward and point to electrodes respectively. Then the fiber will goes in the direction that the moment on the fiber is minimized.



Fig. 3 SEMs of oriented PEO nanofibers. The deposition ime is 8 seconds. (a) and (b) are taken from the same sample.



Fig. 4 (a) FEA of the electric field around the trench. In order to make the electric field strength more obvious, the distance between of the auxiliary electrodes and height between anode and collector are set to be 2mm and 5cm respectively. The high potential is set to be 9,000V. (b) Mechanical analysis of nanofiber between two conductors.

Therefore, the as2spun charged fiber therein is pulled along the direction upright to the edge of the  $e^2$ 

lectrodes and aligned with each other when deposited on the quartz substrate. Two pairs of electrodes which is vertical to each other are introduced to prepare crossed nanofibers shown in Fig. 5.



Fig. 5 Crossing PEO nanofiber deposited by two pairs of the electrodes. The distances are both 6mm

From experiments, it can also be found that the diameters of the nanofibers decrease to  $210 \sim 265$  nm at first when the distance of the two elec2 trodes increase to 2 mm and then increase gradually with the distance increasing when the distance is larger than 2 mm, as shown in Fig. 6. we believe the reason is that when the distance is smaller than 2 mm, the stretching sufficiency of the a&spun nanofibers is not complete, then the diameters will decrease in spite of decrease of electric field strength with the distance increasing. But if the distance is larger than 2 mm, the diameters are dominated by electric field strength. Obviously, when the distance is increased, the diameters will increase with the electric field strength decreases.



Fig. 6 Diameter of nanofiber as function of distance of two parallel conductors

In this paper, the experiments of other mate2 rial of conductors are carried out too. The elec2 trodes, which shape is as same as of the Fig. 1, made of quartz or silicon are tested. The nanof2 bers achieved by quartz electrodes and silicon elec2 trodes are chaos and oriented respectively, as seen from Fig. 7. So it can be concluded that the orien2 ted nanofibers are caused by the changed electro2 static field and not by structure effect of elec2 trodes.



Fig. 7 SEM of nanofibers on quartz substrate between two parallel structure of (a) glass (b) silicon

#### 3 ConclusionsP

In summary, a facile technique to fabricate o2 riented nanofiber is reported in this paper, which is that the separate and parallel conductors are atta2 ched between the collector. Expreriment results show that <sup>1</sup> The diameters decrease at first within the distance of 2 mm due to the stretching suffien2 cy when the distance of the two electrodes is smal2 ler than 2 mm and then increase due to the decrea2 sing electric field strength when it is larger than 2mm. ° The aligned nanofibers is mainly caused by changed electric field, but not by structrue of e2 lectrodes. This technique offers a cost2effective technique to fabricate on 2 dimensional nanoscale " building blocks" that have potential in bottom2up assembly applications in such field as nanoelectron2 ics and photonics.

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#### 参考文献:

- Yuris Dzenis. Spinning Continous Fiber for Nanotechnology
   [J]. Science, 2004, 304: 191721919.
- Phillip Gibson, Heidi Schreude2Gibson, Donald Rivin. Transport Properties of Porous Membranes Based on Electro2 spun Nanofibers[J]. Colloids and Surfaces A: Physicochem. Eng. Aspects 1872188, 2001: 4692481.
- [3] Haiqing Liu, Jun Kameoka, David A. Czaplewski, and H. G. Craighead. Polymeric Nanowire Chemical Sensor [J]. Nano Letter s. 2004, 4: 6712675.
- [4] C. Y. Xu, R. Inai, M. Kotaki, S. Ramakrishna. Aligned Biode2 gradable Nanofibrous Structure: a Potential Scaffold for Blood Vessel Engineering[J]. Biomaterials, 2004, 25: 8772886.
- [5] Jamil A. Matthews, Gary E. Wnek, David G. Simpson, and Gary L. Bowlin. Electrospinning of Collagen Nanofibers [J]. Biomacromolecules, 2002, 3: 2322238.
- [6] Jing Zeng, Xiaoyi Xu, Xuesi Chen, Qizhi Liang, Xin chao B2 an, Lixin Yang, Xiabin Jing. Biodegradable Electrospun F2 bers for Drug Delivery [J]. Journal of Controlled Release, 2003, 92: 227231.
- [7] Formhals A. US Patent[P] 1, 975, 504, 1934.
- [8] Yu Huang, Xiangfeng Duan, Qingqiao Wei, Charles M. Lieber. Directed Assembly of On& Dimensional Nanostructures into Fun& tional Networks[J]. Science, 2006, 291: 6302633.
- [9] Theron A, Zussman E and Yarin AL, Electrostatic Fiel&AS sisted Alignment of Electrospun Nanofibres[J]. NanotechnoD ogy, 2001: 3842390.
- [10] Royal Kessick, John Fenn, Gary Tepper. The use of AC P& tentials in Electrospraying and Electrospinning Processes[J]. Polymer, 2004, 45: 298122984.
- [11] Paul D. Dalton, Doris Klee, Martin Moller. Electrospinning with Dual Collection Rings[J]. Polymer, 2005, 46: 612614.
- [12] Daoheng Sun, Chieh Chang, Sha Li and Liwei Lin. Nea Field Electrospining[J]. Nano Letters. 2006, 16: 8392842.
- [13] Xinhua Zong, Kwangsok Kim, Dufei Fang, Shaofeng Ran, Benjamin S. Hsiao, Benjamin Chu. Structure and Process Relationship of Electrospun Bioabsorbable Nanofiber Mem2 branes [J]. Polymer, 2002, 43: 44024412.