Janus nanofiber array pellicle: facile conjugate electrospinning construction, structure and bifunctionality of enhanced green fluorescence and adjustable magnetism

Guoyi Wang, Qianli Ma, Jiao Tian, Libing Fan, Dan Li, Xiangting Dong, Wensheng Yu, Jinxian Wang and Guixia Liu

A \([\text{Fe}_3\text{O}_4/\text{polyvinyl pyrrolidone (PVP)}]//[\text{ Tb(BA)}_3\text{phen/PVP}]\) Janus nanofiber array pellicle (denoted JNAP) was successfully constructed by facile conjugate electrospinning without twisting for the first time. The JNAP offers the dual-functionality of fluorescence and magnetism. This technology entirely solves the dilemma of the magnetic spinning dope and fluorescent spinning dope being easily mixed together during the parallel electrospinning process, as it achieves complete segregation of magnetic nanoparticles and fluorescent molecules. Moreover, conjugate electrospinning without twisting has fewer requirements on the viscosity of the spinning dope compared with parallel electrospinning, in which the two spinning dopes should have the same viscosity. It was satisfactorily found that the JNAP has higher fluorescence intensity than the corresponding non-aligned Janus nanofiber pellicle. The magnetism of the JNAP could be tailored by changing the doping amount of the \(\text{Fe}_3\text{O}_4\) NPs. The JNAP has potential applications in nanotechnology and biomedicine, etc., due to its enhanced green fluorescence and adjustable magnetism. In addition, this design concept and manufacturing process provide a facile way for preparing other one-dimensional Janus nanomaterials with multifunctionality.

Introduction

It is becoming harder to meet the needs of many emerging practical technologies using single-functional materials alone. Therefore, research into the preparation and performance of multifunctional materials has become increasingly important in the realm of materials science and technology.\(^1\) Fluorescent-magnetic bifunctional nanomaterials have attracted much attention of researchers due to their potential uses in cell imaging, cancer studies, sensing and biomolecular detection, etc.\(^2\) For instance, fluorescent-magnetic bifunctional nanomaterials are ideal candidates for drug delivery because the loaded drug can be transported to a specific location using magnetic navigation, and meanwhile, the real-time location of the drug can be monitored by measuring fluorescence signals emitted from the fluorescent-magnetic bifunctional nanomaterial.\(^3\) As another example, fluorescent-magnetic bifunctional nanomaterials possess the property of dual-mode imaging, including fluorescent imaging and magnetic resonance imaging, which is beneficial for biological detection.\(^4\) However, researchers in this field must face the plight that if the magnetic nanoparticles and fluorescent molecules are directly mixed, the fluorescence intensity of the sample will be greatly reduced.\(^14\)\(^,\)\(^15\) Hence, exceptional structures need to be designed and constructed to separate magnetic nanoparticles from fluorescent molecules in an integrated system. Inspired by Janus, a god in Roman mythology, an exception “Janus structure” has been proposed by several research groups.\(^16\)\(^,\)\(^17\) Researchers have proposed that magnetic nanoparticles and fluorescent molecules should be respectively added to both sides of a Janus material in order to isolate two different substances from each other and reduce the detrimental influence of magnetic nanoparticles on the fluorescence intensity.\(^18\)\(^,\)\(^19\) Magnetic-fluorescent bifunctional Janus nanofibers are typical Janus materials that realize separation of magnetic nanoparticles from fluorescent molecules. According to the literature, Janus nanofibers can be gained by parallel electrospinning, where two kinds of spinning dopes are respectively loaded into two syringes to generate magnetic-fluorescent bifunctional Janus nanofibers using a specially-made parallel spinneret under electrospinning.\(^20\)\(^–\)\(^23\) Xi, et al.\(^24\)\(^,\)\(^25\) fabricated flexible magnetic-photoluminescent bifunctional Janus nanofibers by parallel electrospinning. It has been proven that such Janus nanofibers have stronger fluorescence intensity than the counterpart composite nanofibers. However, parallel electrospinning still has a drawback in...
that it is difficult to fulfill complete segregation of the magnetic nanoparticles and the fluorescent molecules, since the two spinning dopes are easily mixed at the outlet of spinneret. Consequently, it is a pressing subject of study to develop a new technique to overcome this drawback.

Conjugate electrospinning is an excellent technique for constructing one-dimensional nanomaterials. This technique can be divided into two kinds: conjugate electrospinning with twisting and that without twisting. Conjugate electrospinning with twisting is used to prepare nanofiber yarns which can be applied to tissue repair, nerve regeneration and electrically conductive material, etc. Conjugate electrospinning with twisting possesses excellent permeability. A [Fe\textsubscript{3}O\textsubscript{4}/PVP]/[Tb(BA)\textsubscript{3}phen/PVP] Janus nanoarray pellicle (abbreviated as JNAP) with magnetic superparamagnetism, good biocompatibility and high permeability was fabricated as follows: Fe\textsubscript{3}O\textsubscript{4} NPs and Tb(BA)\textsubscript{3}phen were respectively used as a template, Fe\textsubscript{3}O\textsubscript{4} NPs (denoted as Fe\textsubscript{3}O\textsubscript{4} NPs for short) and Tb(BA)\textsubscript{3}phen/PVP were prepared according to the literature.46,47

Electrospinning process

Two different types of spinning dopes (named spinning dope A and spinning dope B) were used to prepare the Janus nanofibers. Spinning dope A, with fluorescence properties, was prepared as follows: PVP (1 g) was fully dissolved in ethanol (7 g) under magnetic stirring, and then a certain amount of Fe\textsubscript{3}O\textsubscript{4} NPs (denoted as Fe\textsubscript{3}O\textsubscript{4} NPs for short) and Tb(BA)\textsubscript{3}phen complex was uniformly dispersed in the solution to form spinning dope A. Spinning dope B, with magnetic properties, was fabricated as follows: Fe\textsubscript{3}O\textsubscript{4} NPs were dispersed in ethanol (7 g) under ultrasonication for 20 min, and then PVP (1 g) was added into the above suspension under mechanical stirring. The actual ingredients of spinning dope A and B are respectively shown in Tables 1 and 2.

A device diagram for constructing the [Fe\textsubscript{3}O\textsubscript{4}/PVP]/[Tb(BA)\textsubscript{3}phen/PVP] JNAP by conjugate electrospinning is presented in Table 3. The spinning dopes A and B were separately loaded into two syringes with plastic spinnerets. The angle between the syringe and the horizontal line was ca. 45° in the conjugate electrospinning device. To obtain the array pellicle, a rotating drum was used as a collector. The corresponding spinning dopes and detailed spinning conditions are summarized in Table 3.

For comparison, a [Fe\textsubscript{3}O\textsubscript{4}/PVP]/[Tb(BA)\textsubscript{3}phen/PVP] Janus nanofiber array pellicle (referred to as JNNP) was prepared by conjugate electrospinning with a wire mesh as the collector, and a parallel electrospinning-made [Fe\textsubscript{3}O\textsubscript{4}/PVP]/[Tb(BA)\textsubscript{3}phen/PVP] Janus nanofiber array pellicle (named PJNAP) was constructed by using spinning dopes S\textsubscript{A3} and S\textsubscript{A1}. Another spinning dope was prepared by evenly mixing equal volumes of

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<thead>
<tr>
<th>Table 1</th>
<th>Ingredients of spinning dope A</th>
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<tbody>
<tr>
<td>Spinning dope A</td>
<td>Tb(BA)\textsubscript{3}phen : PVP (wt%)</td>
</tr>
<tr>
<td>S\textsubscript{A1}</td>
<td>5%</td>
</tr>
<tr>
<td>S\textsubscript{A2}</td>
<td>10%</td>
</tr>
<tr>
<td>S\textsubscript{A3}</td>
<td>15%</td>
</tr>
<tr>
<td>S\textsubscript{A4}</td>
<td>20%</td>
</tr>
<tr>
<td>S\textsubscript{A5}</td>
<td>25%</td>
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<table>
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<tr>
<th>Table 2</th>
<th>Ingredients of spinning dope B</th>
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</thead>
<tbody>
<tr>
<td>Spinning dope B</td>
<td>Fe\textsubscript{3}O\textsubscript{4} NPs/PVP (mass ratio)</td>
</tr>
<tr>
<td>S\textsubscript{B1}</td>
<td>0.5 : 1</td>
</tr>
<tr>
<td>S\textsubscript{B2}</td>
<td>1 : 1</td>
</tr>
<tr>
<td>S\textsubscript{B3}</td>
<td>1.5 : 1</td>
</tr>
<tr>
<td>S\textsubscript{B4}</td>
<td>2 : 1</td>
</tr>
</tbody>
</table>

**Experimental**

**Chemicals**

Tb\textsubscript{3}O\textsubscript{7} (99.99%), concentrated nitric acid (HNO\textsubscript{3}), 1,10-phenanthroline (phen), benzoic acid (BA), FeSO\textsubscript{4}-7H\textsubscript{2}O, FeCl\textsubscript{3}-6H\textsubscript{2}O, polyethylene glycol (PEG, M\textsubscript{w} ≈ 20,000), NH\textsubscript{4}NO\textsubscript{3}, ammonia (NH\textsubscript{3}·H\textsubscript{2}O), oleic acid (OA), polyvinyl pyrrolidone (PVP) and ethanol (CH\textsubscript{3}CH\textsubscript{2}OH) were used to prepare the samples. All chemicals were of analytical grade.

**Syntheses of OA-modified Fe\textsubscript{3}O\textsubscript{4} NPs and terbium complexes**

OA-modified Fe\textsubscript{3}O\textsubscript{4} NPs (denoted as Fe\textsubscript{3}O\textsubscript{4} NPs for short) and Tb(BA)\textsubscript{3}phen were prepared according to the literature.46,47
Table 3 Electrospinning device diagrams, corresponding spinning dopes and electrospinning conditions of samples

<table>
<thead>
<tr>
<th>Samples</th>
<th>Electrospinning device diagrams</th>
<th>Spinning dopes</th>
<th>Electrospinning conditions</th>
</tr>
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<tbody>
<tr>
<td>JNAP</td>
<td><img src="image" alt="JNAP Diagram" /></td>
<td>Spinning dope A: $S_{A1}, S_{A2}, S_{A3}, S_{A4}$ and $S_{A5}$, Spinning dope B: $S_{B1}$</td>
<td>Collector: rotating drum (10 cm in diameter), rotation speed: 300 rpm, distance between the two spinnerets: 15 cm, curing distance: 25 cm, applied voltages: $\pm 7$ kV, temperature: $15 \pm 5^\circ$C, relative humidity: 25 $\pm$ 5%</td>
</tr>
<tr>
<td>JNNP</td>
<td><img src="image" alt="JNNP Diagram" /></td>
<td>Spinning dope A: $S_{A3}$, Spinning dope B: $S_{B1}$</td>
<td>Collector: wire mesh, distance between the two spinnerets: 15 cm, curing distance: 25 cm, applied voltages: $\pm 7$ kV, temperature: $15 \pm 5^\circ$C, relative humidity: 25 $\pm$ 5%</td>
</tr>
<tr>
<td>P-JNAP</td>
<td><img src="image" alt="P-JNAP Diagram" /></td>
<td>Spinning dope A: $S_{A3}$, Spinning dope B: $S_{B1}$</td>
<td>Collector: rotating drum (10 cm in diameter), rotation speed: 300 rpm, distance between the two spinnerets: 15 cm, curing distance: 25 cm, applied voltages: $\pm 7$ kV, temperature: $15 \pm 5^\circ$C, relative humidity: 25 $\pm$ 5%</td>
</tr>
<tr>
<td>HNAP</td>
<td><img src="image" alt="HNAP Diagram" /></td>
<td>Spinning dope: mixing isopyknic $S_{A1}$ and $S_{B1}$</td>
<td>Collector: rotating drum (10 cm in diameter), rotation speed: 300 rpm, distance between the two spinnerets: 15 cm, curing distance: 25 cm, applied voltages: $\pm 7$ kV, temperature: $15 \pm 5^\circ$C, relative humidity: 25 $\pm$ 5%</td>
</tr>
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</table>
spinning doped NPs, and then a [Fe3O4/Tb(BA)3phen/PVP] homogeneous nanofiber array pellicle (called HNAP) was also fabricated by conjugate electrospinning. The device diagrams and specific conditions for the comparative samples are also displayed in Table 3.

**Characterization methods**

The phase compositions of Fe3O4 NPs, JNAP, JNNP, HNAP and P-JNAP were analyzed using X-ray power diffraction (XRD) with Cu Kα radiation. The operation voltage and current were respectively kept at 40 kV and 20 mA. The morphology of the JNAP was observed using a field-emission scanning electron microscope (FESEM), equipped with energy-dispersive X-ray spectroscopy (EDS). A Hitachi fluorescence spectrophotometer F-7000 was used to investigate the fluorescence of different samples when the excitation and emission slits were 2.5 nm and 720 nm. The magnetic properties of the samples were measured using a vibrating sample magnetometer (VSM).

**Results and discussion**

**Phase compositions**

Fig. 1 exhibits the XRD patterns of the Fe3O4 NPs, JNAP, JNNP, HNAP and P-JNAP. The XRD patterns of the Fe3O4 NPs conform to the cubic structure of Fe3O4 (PDF#75-0499), and no characteristic peaks of other impurities are found. The corresponding diffraction peaks of Fe3O4 are also found in the four kinds of nanofiber pellets, which indicates that the nanofiber pellets contain Fe3O4 NPs. Meanwhile, the diffraction peak of the amorphous PVP (2θ ≈ 22°) can also be observed in the four kinds of nanofiber pellets.

**Morphological and structural analyses**

SEM images, EDS spectra and diameter distribution histograms of the Janus nanofibers in JNAP are given in Fig. 2. As seen in Fig. 2a, the nanofibers are in good directional alignment and form an array pellicle. It can be seen in Fig. 2b that exceptive-structured Janus nanofibers are obtained, with every single Janus nanofiber being composed of two nanofibers bound together side-by-side. As shown in Fig. 2b and c, the diameter of each single nanofiber in the Janus nanofibers is almost equal (the average diameter of a single nanofiber is ca. 720 nm). The EDS spectrum of the JNAP (Fig. 2d) indicates that the Janus nanofibers are composed of the elements C, N, O, Fe, Tb and Pt. The Pt peak is attributed to the conductive film sprayed on the surface of the sample for SEM observation. Fig. 2c presents the EDS line-scan analysis results for the Janus nanofibers, where Fe and Tb elements indicate Fe3O4 and Tb(BA)3phen, respectively. The Fe and Tb elements are respectively found in two single nanofibers, further demonstrating that the Janus nanofibers were successfully constructed by conjugate electrospinning and the goal of segregating magnetic nanoparticles and fluorescent molecules has been realized.

**Fluorescence performance**

A series of JNAP samples (the mass ratio of Fe3O4 NPs to PVP was 0.5 : 1, and the mass percentages of Tb(BA)3phen to PVP were 5%, 10%, 15%, 20% and 25%, respectively) were prepared to find the optimal mass percentage of Tb(BA)3phen to PVP. Fig. 3a and b respectively exhibit the excitation and emission spectra of the above-described JNAP samples. A broad excitation band from 200 nm to 400 nm of the JNAP was observed when the monitoring wavelength was 545 nm. The strongest excitation peak at 275 nm can be put down to the π → π* electron transition of the ligands. The strong characteristic emission peaks of Tb3+ were found at 490, 545, 586, 622 nm under 275 nm excitation. These peaks are attributed to the 5D4 → 7F6 (490 nm), 5D4 → 5D5 (545 nm), 5D4 → 5D4 (586 nm), 5D4 → 7F4 (622 nm) energy level transitions of Tb3+, respectively. The 5D4 → 7F5

![Fig. 1 XRD patterns of the Fe3O4 NPs, JNAP, JNNP, HNAP and P-JNAP.](image)

![Fig. 2 SEM images at low magnification (a) and high magnification (b) of the JNAP. Diameter distribution histogram of the nanofibers in Janus nanofibers in the JNAP (c). EDS spectrum of the JNAP (d). EDS line-scan analysis of a single Janus nanofiber in the JNAP (e).](image)
transition at 545 nm (green light) is the predominant emission peak. The fluorescence intensity of the JNAP first enhances and then weakens with the increase of Tb(BA)₃phen content, as revealed in Fig. 3. When the mass percentage of Tb(BA)₃phen to PVP is 15%, the JNAP achieves the highest fluorescence intensity. When the mass percentage of Tb(BA)₃phen to PVP exceeds 15%, the distribution of the Tb(BA)₃phen becomes denser in the polymer matrix, resulting in stronger non-radiative transitions amongst the Tb³⁺ ions, which causes obvious reductions in the fluorescence intensity. Therefore, the optimum mass percentage of Tb(BA)₃phen to PVP is 15%.

Physical photographs of the JNAP are displayed in Fig. 4. From Fig. 4a–c, it can be seen that the JNAP can be easily bent by hand and also possesses the ability of self-recovery, which proves that the JNAP is flexible. Fig. 4d shows a camera photograph of the JNAP under 275 nm UV illumination in darkness, indicating that the JNAP can emit green fluorescence.

The fluorescence decay curves of Tb³⁺ ions in the JNAP samples doped with different amounts of Tb(BA)₃phen are shown in Fig. 5. The excitation wavelength is set to be 275 nm and the monitoring wavelength is 545 nm. It is generally known that the fluorescence decay curves follow the single-exponential decay:

\[ I_t = I_0 \exp(-t/\tau) \]

where \( I_t \) signifies the intensity at time \( t \), \( I_0 \) represents the intensity at \( t = 0 \) and \( \tau \) symbolizes the decay lifetime. The average lifetime values (\( \tau \)/ms) of the JNAP are revealed in Fig. 5. As the content of Tb(BA)₃phen increases, the fluorescence lifetime of the \( ^{5}D_{4} \rightarrow ^{7}F_{5} \) (545 nm) transition is gradually decreased. The introduction of more Tb(BA)₃phen leads to a reduced distance between Tb³⁺ ions in Tb(BA)₃phen molecular clusters in the JNAP. Thus, the energy transfer among Tb³⁺ ions is increased, and the fluorescence lifetime of Tb³⁺ is shortened.

A series of JNAP samples (the mass percentage of Tb(BA)₃phen to PVP was 15% and the mass ratios of Fe₃O₄ NPs to PVP were 0.5 : 1, 1 : 1, 1.5 : 1, and 2 : 1, respectively) were prepared to explore the effect of different amounts of Fe₃O₄ NPs on the fluorescence intensity. The intensities of the excitation and emission peaks gradually decrease along with increasing content of Fe₃O₄ NPs added into the JNAP, as illustrated in Fig. 6. As previously reported, dark-colored Fe₃O₄ NPs can absorb visible light (400 nm < \( \lambda \) < 700 nm) and ultraviolet light (\( \lambda \) < 400 nm). Therefore, Fe₃O₄ NPs can absorb the excitation...
light and emission light of the JNAP, and the degree of absorption becomes stronger as the content of Fe₃O₄ NPs is increased. Moreover, the more Fe₃O₄ NPs added into the JNAP, the darker the color of the products and the more intense the absorption of excitation emission light by the Fe₃O₄ NPs, as revealed in Fig. 7, giving rise to lower fluorescence intensity.

Furthermore, the fluorescence intensities of samples with the same ingredients and content prepared by different methods were compared to highlight the superiority of the JNAP. The JNAP has the highest fluorescence intensity compared with three comparative samples. It has stronger fluorescence intensity than the P-JNAP, and HNAP possesses the lowest fluorescence intensity, as shown in Fig. 8. This outcome can be explained by three aspects: the special Janus structure, the arrangement of Janus nanofibers, and the electrospinning method. Fig. 9 presents schematic graphs of the excitation and emission light for the JNAP, JNNP, P-JNAP and HNAP. As seen from Fig. 9a, the JNAP is composed of aligned Janus nanofibers, and the Janus nanofibers are closely arranged. The Janus nanofibers in the upper layer can directly absorb excitation light and the produced emission light can be directly emitted without refraction or reflection, which is the most effective mode for fluorescence. Furthermore, the Fe₃O₄ NPs and Tb(BA)₃phen molecules are absolutely segregated, leading to a reduced effect of the Fe₃O₄ NPs on the fluorescence performance. As depicted in Fig. 9b, the Fe₃O₄ NPs and Tb(BA)₃phen molecules are also completely segregated in the JNNP. The surface of the JNNP is loose due to the unordered arrangement of Janus nanofibers. In this case, some of the excitation light can pass through the gaps
in the upper layer and thereby excite the Janus nanofibers in lower layer. In this process, the Janus nanofibers in the upper layer can absorb some of the excitation light, and thus, the excitation light arriving to the Janus nanofibers in the lower layer is weakened. For the same reason, the emission light emitted from the lower layer is also absorbed by the upper layer in the JNNP. Hence, the fluorescence intensity of the JNNP is lower than that of the JNAP. P-JNAP was prepared by parallel electrospinning. Fig. 10 describes the actual situation of two spinning dopes in parallel spinneret during the parallel electrospinning process. Inside the parallel spinneret, two spinning dopes mutually diffuse at the contact interfaces, so that diffusion regions exist at the contact interfaces between nanofibers in the Janus nanofibers. This results in an incomplete separation of Fe$_3$O$_4$ NPs from Tb(BA)$_3$phen molecules in the Janus nanofibers. Thus, the fluorescence intensity of P-JNAP is lower than that of JNAP. As seen in Fig. 9d, the HNAP is totally made up of homogeneous nanofibers where Fe$_3$O$_4$ NPs and Tb(BA)$_3$phen molecules are directly mixed, so that the fluorescence intensity of the HNAP is the lowest. The above new findings thoroughly prove that conjugate electrospinning has more advantages than parallel electrospinning in the fabrication of Janus nanofibers.

**Magnetic properties**

Typical hysteresis loops of the Fe$_3$O$_4$ NPs, the JNAP with different mass ratios of the Fe$_3$O$_4$ NPs and comparative samples are exhibited in Fig. 11, and corresponding saturation magnetization results are summarized in Table 4. It is known that the saturation magnetization of a magnetic compound material depends on the doping mass ratio of the magnetic material. As can be seen in Table 4, the saturation magnetization of the Fe$_3$O$_4$ NPs reaches 37.99 emu g$^{-1}$. As the content of the Fe$_3$O$_4$ NPs increases in the JNAP, the saturation magnetization of the JNAP also increases from 8.93 to 18.99 emu g$^{-1}$. Thus, the JNAP possesses adjustable magnetism via changing the doping amount of the Fe$_3$O$_4$ NPs. The magnetism of the comparative samples is similar to that of the JNAP (prepared by S$_{Sa}$/S$_{B1}$) owing to the same content of Fe$_3$O$_4$ NPs in the samples.

![Digital photograph of parallel spinneret during parallel electrospinning process.](image)

**Fig. 10** Digital photograph of parallel spinneret during parallel electrospinning process.

![Hysteresis loops of the Fe$_3$O$_4$ NPs (a), the JNAP doped with different amounts of Fe$_3$O$_4$ NPs (b–e), JNNP (f), P-JNAP (g) and HNAP (h).](image)

**Fig. 11** Hysteresis loops of the Fe$_3$O$_4$ NPs (a), the JNAP doped with different amounts of Fe$_3$O$_4$ NPs (b–e), JNNP (f), P-JNAP (g) and HNAP (h).

**Table 4** Saturation magnetization results of the Fe$_3$O$_4$ NPs, the JNAP doped with different amounts of Fe$_3$O$_4$ NPs and comparative samples

<table>
<thead>
<tr>
<th>Samples</th>
<th>Saturation magnetization (emu g$^{-1}$)</th>
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<tbody>
<tr>
<td>Fe$_3$O$_4$ NPs</td>
<td>37.99</td>
</tr>
<tr>
<td>JNAP</td>
<td></td>
</tr>
<tr>
<td>Fe$_3$O$_4$ : PVP = 2 : 1</td>
<td>18.99</td>
</tr>
<tr>
<td>Fe$_3$O$_4$ : PVP = 1.5 : 1</td>
<td>15.86</td>
</tr>
<tr>
<td>Fe$_3$O$_4$ : PVP = 1 : 1</td>
<td>13.50</td>
</tr>
<tr>
<td>Fe$_3$O$_4$ : PVP = 0.5 : 1</td>
<td>8.93</td>
</tr>
<tr>
<td>Comparative samples</td>
<td></td>
</tr>
<tr>
<td>JNNP</td>
<td>8.77</td>
</tr>
<tr>
<td>P-JNAP</td>
<td>8.07</td>
</tr>
<tr>
<td>HNAP</td>
<td>7.91</td>
</tr>
</tbody>
</table>
Formation mechanism for the JNAP

Schematic graphs of the formation mechanism of Janus nanofibers and the JNAP are presented in Fig. 12. Under the effect of an electrostatic field, the spinning dopes in the two spinnerets form two bundles of continuous nanofibers with positive and negative charges after positive and negative power are applied to the two spinnerets, respectively, as seen in Fig. 12a. Here, an electric field is formed between the two spinnerets when positive and negative charges accumulate at the tip of the two spinnerets, and the directions of the electric field lines are indicated by the dotted line (from the positive pole to the negative pole), as shown in Fig. 12b. Magnetic nanofibers and fluorescent nanofibers attract each other along the path of the electric field. After the two kinds of nanofiber bundles meet in the middle of the two spinnerets, the electrical charges in the two kinds of nanofiber bundles are neutralized, and the formed Janus nanofibers move downward by gravity. Due to electrostatic repulsion, the same nanofibers (e.g. magnetic nanofiber and fluorescent nanofiber) cannot get together to form a Janus nanofiber in this process. Therefore, [Fe₃O₄/PVP]/[Tb(BA)₃-phen/PVP] Janus nanofibers are obtained. Fig. 12c describes the process of collecting an electrically neutral Janus nanofiber bundle with the rotating drum. At this point, the Janus nanofibers are affected by the vortex generated by the rotating drum in the experimental environment, so that the Janus nanofibers shake slightly and are drawn to and wrap around the rotating drum to form the JNAP.

Conclusions

In summary, a [Fe₃O₄/PVP]/[Tb(BA)₃-phen/PVP] magnetic-fluorescent bifunctional Janus nanofiber array pellicle was successfully fabricated via facile conjugate electrospinning with a rotating drum as a collection device. The JNAP is made up of aligned Janus nanofibers. Each Janus nanofiber is composed of a magnetic [Fe₃O₄/PVP] nanofiber and a fluorescent [Tb(BA)₃-phen/PVP] nanofiber. The diameter of each single nanofiber in the Janus nanofibers is ca. 720 nm. It is noteworthy that the magnetism of the JNAP can be adjusted by adding diverse amounts of Fe₃O₄ NPs and the JNAP can emit green light under the excitation of ultraviolet light. Compared with three comparative samples, the JNAP has the highest fluorescence intensity owing to its peculiar Janus structure, which achieves effective segregation of Fe₃O₄ NPs and Tb(BA)₃-phen molecules, as well as directional alignment of the Janus nanofibers. More importantly, conjugate electrospinning can be extended to manufacture other Janus nanofibers with multifunctionality.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

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