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# Preparation and characterization of novel naphthyl epoxy resin containing 4-fluorobenzoyl side chains for low-*k* dielectrics application†

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A new type of epoxy monomer 1,5-bis(4-fluorobenzoyl)-2,6-diglycidyl ether naphthalene (DGENF) was obtained through a three-step procedure involving Friedel–Crafts acylation, demethylation, and followed by nucleophilic reaction. The chemical structures were confirmed by <sup>1</sup>HNMR, <sup>19</sup>FNMR and FT-IR. After curing with methylhexahydrophthalic anhydride (MeHHPA), the properties of DGENF epoxy resin were measured and compared with three other kinds of commercial epoxy resins. As a result, DGENF exhibited excellent thermal stability, hydrophobic and dielectric properties. For example, DGENF had a higher glass transition temperature of 170 °C than the other three commercial epoxy resins. DGENF showed a higher contact angle of 116°, which could satisfy the standard of hydrophobic materials. In addition, DGENF showed significantly lower dielectric constant (2.97 at 1 MHz) and dielectric loss (0.0188 at 1 MHz) than those of the other commercial epoxy resins because of the introduction of fluorine on the side chains, which improved the electronegativity of the epoxy resin and reduced the polarizability of molecules efficiently. Herein, we believe the novel naphthyl epoxy resin (DGENF) has demonstrated potential application in electronic industries.

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## 1. Introduction

With the rapid development of micro-electronics industries and the continuing miniaturization in dimensions of electronic devices, there is a demand for electronic materials to possess more functions to satisfy the requirements for optimal performance and lifetime of electronic components. These functions mainly include electric insulation, mechanical support and overcoming the propagation delay caused by interconnection. Therefore, advanced electronic materials are required to possess better mechanical properties, good thermal stability, excellent chemical and corrosion resistance as well as lower dielectric properties.<sup>1–9</sup>

As one of the most important electrical materials, epoxy resins have been widely used in electrical coatings, electronic packaging, electrical substrates, adhesives of chips and encapsulation.<sup>10–19</sup> However, conventional epoxy resins do not have adequate electrical performance to satisfy the requirements of advanced microelectronic materials, especially dielectric properties.<sup>20</sup> Therefore, the development of novel epoxy resins possessing lower dielectric constants and loss (low-*k*) has attracted more and more attention of researchers around the world.

According to reports, there are various methods to achieve low dielectric constant materials, such as the incorporation of fluorinate groups and stiff non-polar structures, as well as the introduction of controlled porosity in polymer.<sup>21,22</sup> Furthermore, the existence of fluorinate and stiff non-polar structure can also decrease moisture absorption and improve the thermal stability of materials.<sup>23</sup> Sasaki<sup>24</sup> reported that the bisphenol A type epoxy resin cured with fluorinated phthalic anhydride had the lowest dielectric constant (2.77) and dielectric loss factor (0.01) among all cured epoxy resins. Meanwhile, the water absorption of this fluorinated epoxy resins could be reduced by 75%. Tao and coworkers<sup>25</sup> also reported fluorinated epoxy resins with reduced moisture absorption and dielectric constant. They had lower dielectric constant (3.3) and water uptake (0.38%) than diglycidyl ether of bisphenol A (DGEBA) cured with the same curing agent. Xu *et al.*<sup>26</sup> synthesized a series of novel epoxy resins bearing naphthyl and limonene moieties, which showed a remarkably higher *T<sub>g</sub>* (171 °C), lower coefficient (36 ppm °C<sup>−1</sup>) of thermal expansion, better moisture resistance (0.13%) and lower dielectric property (3.7). All of the properties were better than DGEBA cured with the same curing agent. Therefore, we believe that the introduction of fluorinate groups and stiff non-polar structure into polymer is a promising method for developing low dielectric materials with improved thermal stability and excellent moisture resistance.

In this work, we synthesized a novel naphthyl epoxy monomer, 1,5-bis(4-fluorobenzoyl)-2,6-diglycidyl ether naphthalene (DGENF) containing 4-fluorobenzoyl side chains and stiff

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non-polar naphthalene moieties simultaneously. After the addition of methylhexahydrophthalic anhydride (MeHHPA) as the curing agent and 2-methylimidazole (2-MI) as the accelerator with a stoichiometric amount, the cured epoxy resins with low dielectric properties and improved thermal stabilities were obtained. Additionally, we prepared other three kinds of cured commercial epoxy resins in the same method and characterized their properties for comparison, including bisphenol A epoxy resin (BPA-EP), bisphenol F epoxy resin (BPF-EP) and 4,4'-diglycidyl(3,3',5,5'-tetramethylbiphenyl) epoxy resin (TMBP).

## 2. Experimental

### 2.1 Materials

2,6-Dimethoxynaphthalene (99.0%), 4-fluorobenzoyl chloride (98.0%), boron tribromide ( $\text{BBr}_3$ , 99.99%) and ferric chloride ( $\text{FeCl}_3$ , 97.5%) were purchased from Sigma-Aldrich Chemical Co., Ltd. Epichlorohydrin (ECH, 98.0%), MeHHPA (98.0%), 2-MI (99.0%), NaOH, tetrabutylammonium bromide (purity 98.0%) were obtained from Sinopharm Chemical Reagent Co., Ltd. BPA-EP (KDS-8128 EEW = 172 g per eq.) was purchased by Kukdo Chemical (Kunshan) Co., Ltd and BPF-EP (NPEF-170 EEW = 170 g per eq.) were obtained by Nanya plastics (Taiwan) Co., Ltd. TMBP was prepared according to our method reported previously.<sup>27</sup> All other solvents were obtained commercially and used without further purification.

### 2.2 Synthesis of 1,5-bis(4-fluorobenzoyl)-2,6-dimethoxynaphthalene (DMNF)

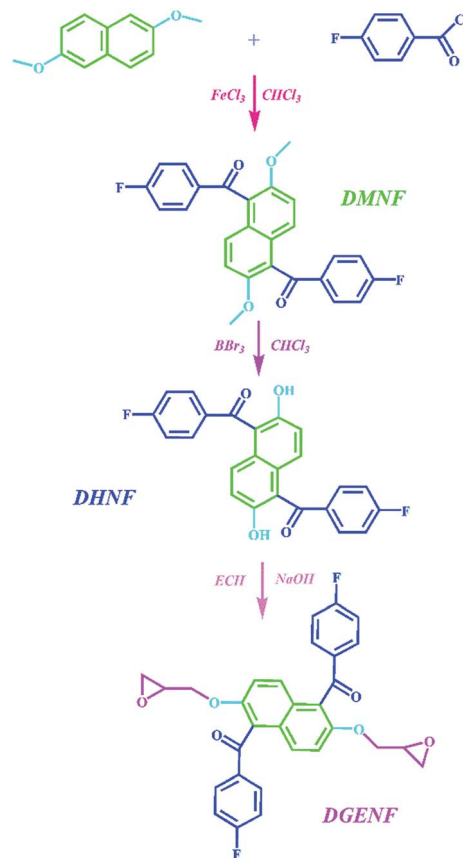
The synthetic procedure of DMNF is described in Scheme 1. 2,6-Dimethoxynaphthalene (9.4 g, 0.05 mol) and 4-fluorobenzoyl chloride (17.4 g, 0.11 mol) were dissolved in chloroform ( $\text{CHCl}_3$ ). Then anhydrous ferric chloride (1.65 g, 0.01 mol) was added to the mixture under 5 °C and the reaction was then kept at ambient temperature for 24 h. The resulting mixture was poured into hydrochloric acid. The liquid phase was removed by decantation and the brown solid was precipitated in methanol. The crude product was then recrystallized from dimethylformamide (yield: 78%).

### 2.3 Synthesis of 1,5-bis(4-fluorobenzoyl)-2,6-dihydroxynaphthalene (DHNF)

DMNF (12.975 g, 0.03 mol) obtained from above procedure was added into 300 mL  $\text{CHCl}_3$  and mixed under 5 °C (ice bath). Then 1 mol  $\text{L}^{-1}$   $\text{CHCl}_3$  solution of  $\text{BBr}_3$  (300 mL) was added into the mixture dropwise during 2 h. The mixture was stirred at room temperature for 48 h before pouring into ice-water to absorb excess  $\text{BBr}_3$ , and then washed with deionized water. The resulting yellow powder (DHNF) was dried under vacuum at 60 °C for 24 h (Scheme 1) (yield: 80%).

### 2.4 Synthesis of epoxy monomer 1,5-bis(4-fluorobenzoyl)-2,6-diglycidyl ether naphthalene (DGENF)

The synthesis route of the epoxy monomer DGENF is illustrated in Scheme 1. A mixture of DHNF (4.05 g, 0.01 mol), ECH (16 mL,



Scheme 1 The synthesis routes of DMNF, DHNF and DGENF monomers.

0.20 mol) and tetrabutylammonium bromide (0.096 g, 0.3 mmol) were stirred in a three-neck flask at 90 °C for 6 h. After cooling to room temperature, the excess ECH was removed at a reduced pressure. Subsequently, 30 mL toluene and 2.88 g NaOH (50 wt% aqueous solution) were added and the mixture was kept at 90 °C for another 3 h. Then the mixture was washed and distilled under reduced pressure. After drying in a vacuum oven at 60 °C for 12 h, the white crystalline product DGENF was finally obtained (yield: 88%).

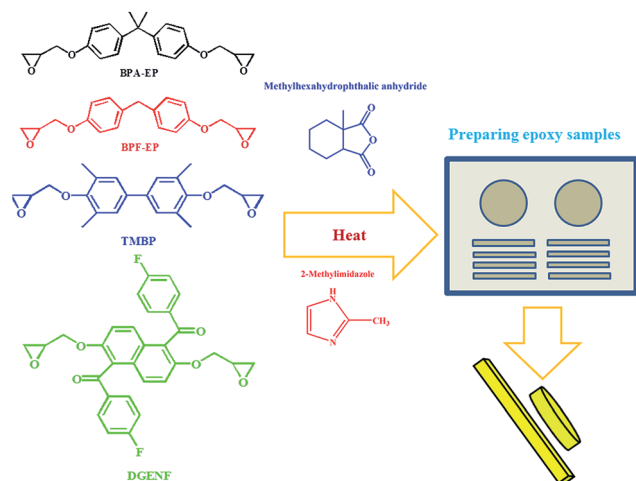
### 2.5 Preparation of cured epoxy resin samples

As shown in Scheme 2, a stoichiometric amount of MeHHPA was added into DGENF, BPA-EP, BPF-EP and TMBP, respectively. The mixtures were stirred continuously for 1 h, and 2-MI was then added into the mixtures as the curing accelerator. After stirred for 30 min, the mixtures were put into the vacuum oven in order to remove the entrapped air. Finally, the mixtures were poured into a polish mould and cured at 120 °C for 2 h, 160 °C for 2 h and 200 °C for 2 h to obtain the epoxy resins.

### 2.6 Characterization

$^1\text{H-NMR}$  spectra were measured on a 500 MHz Bruker Avance 510 spectrometer at 298 K with  $\text{DMSO}-d_6$  as the solvent and tetramethylsilane as the standard.





Scheme 2 Procedure of preparing epoxy resins.

FT-IR spectra were obtained with a Nicolet Impact 410 Fourier transform spectrometer using KBr pellets.

Dynamic mechanical analysis (DMA) was performed on a DMA Q800 analyzer (TA Instruments, New Castle, DE, USA) using a single cantilever clamp over a temperature range of 40–200 °C at a heating rate of 3 °C min<sup>-1</sup> with a constant frequency of 1 Hz.

Thermogravimetric analysis (TGA) was conducted on a Pyris 1 TGA analyzer (Perkin-Elmer, USA). The samples were heated from 100 °C to 700 °C at a rate of 10 °C min<sup>-1</sup> under nitrogen atmosphere. Before the heating scan, all the samples were pre-dried under N<sub>2</sub> at 100 °C for 10 min to remove the residual water and solvent thoroughly.

Dielectric constants of the samples were measured using a Hewlett-Packard 4294A apparatus at room temperature over a frequency range of 0–10<sup>6</sup> Hz.

Contact angle measurements were carried out on a JC2000C2 contact angle goniometer (Shanghai Zhongchen Powereach Company, China) by the sessile drop method with a micro syringe at room temperature.

## 3. Results and discussion

### 3.1 Synthesis and characterization of monomers

As depicted in Scheme 1, the epoxy monomer DGENF was synthesized *via* a three-step synthetic procedure. Firstly, the monomer DMNF was synthesized by a Friedel-Crafts acylation reaction between 2,6-dimethoxynaphthalene and 4-fluorobenzoyl chloride using anhydrous ferric chloride as the catalyst. Then, DMNF containing two methoxyl groups was demethylated to DHNF bearing hydroxyl groups by using BBr<sub>3</sub> solution. Finally, DGENF containing epoxy groups was obtained by a nucleophilic reaction between DHNF and ECH under the catalyst of NaOH. The structure of DMNF and DHNF intermediates as well as the epoxy monomer DGENF was confirmed by their <sup>1</sup>H NMR spectra in DMSO-*d*<sub>6</sub>. As displayed in Fig. 1, the proton peak at 3.7 ppm was assigned to the protons on –OCH<sub>3</sub> groups of DMNF (Fig. 1a). After the demethylation reaction, the

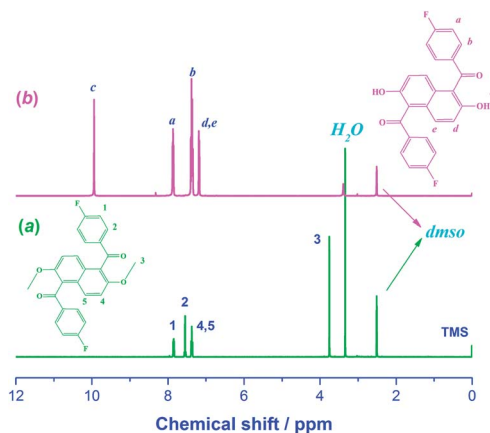


Fig. 1 <sup>1</sup>H-NMR spectra of DMNF and DHNF monomers.

signal of –OCH<sub>3</sub> at 3.7 ppm disappeared completely in the spectrum of DHNF (Fig. 1b). Instead, the characteristic peak for the proton of –OH can be observed at 10.0 ppm. This variation indicated that methoxyl groups in DMNF were completely converted into hydroxyl groups in DHNF.

Fig. 2 shows the <sup>1</sup>H-NMR spectrum of the epoxy monomer DGENF. Compared to DHNF, the proton peak of –OH at 10.0 ppm disappeared, while the proton peaks (*H<sub>e</sub>* and *H<sub>g</sub>*) for the methylene group (–CH<sub>2</sub>–) appeared at 4.30 ppm, 3.98 ppm, 2.62 ppm and 2.48 ppm due to the incorporation of epoxy groups. The results indicated that the hydroxyl groups were converted to epoxy groups thoroughly by reacting with ECH.

Fig. 3 shows the <sup>19</sup>F NMR of DGENF. The only one peak at –105 ppm corresponds to the terminal fluorine atom on 4-fluorobenzoyl side chains. Therefore, we could confirm that 4-fluorobenzoyl side chains exist in the structure of our target DGENF product.

### 3.2 FT-IR spectra of cured epoxy resins

DGENF epoxy resin was prepared using MeHHPA as the curing agent and 2-MI as the accelerator. Another three kinds of cured commercial epoxy resins including BPA-EP, BPF-EP and TMBP were also prepared at the same condition for comparison. Fig. 4

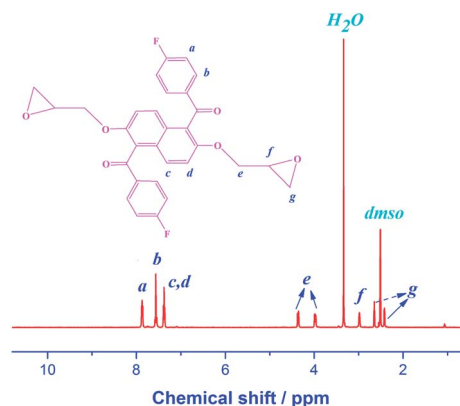


Fig. 2 <sup>1</sup>H-NMR spectra of epoxy monomer DGENF.



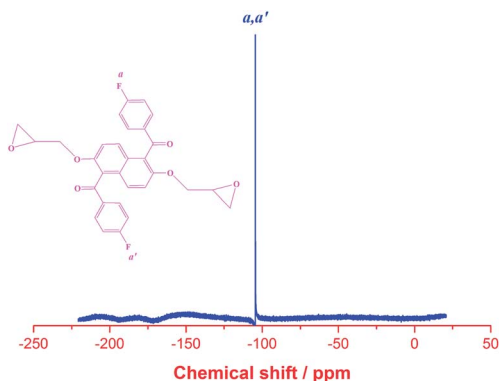


Fig. 3  $^{19}\text{F}$  NMR of the epoxy monomer DGENF.

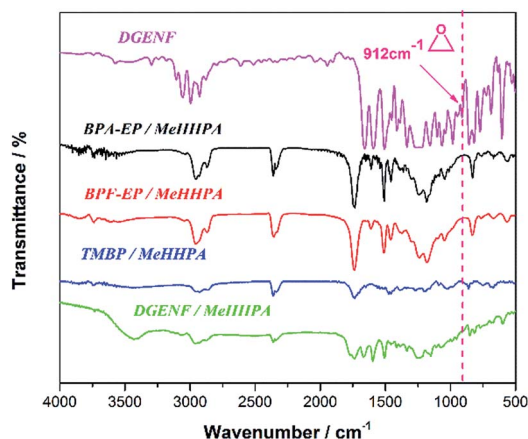


Fig. 4 FT-IR spectra of epoxy monomer DGENF and cured epoxy resins.

shows the FT-IR spectra of the epoxy monomer DGENF and four kinds of cured epoxy resins. For the DGENF monomer, the absorption bands observed from  $1500\text{ cm}^{-1}$  to  $1650\text{ cm}^{-1}$  can be attributed to the skeleton vibration of naphthalene rings. A sharp and strong characteristic band at  $1740\text{ cm}^{-1}$  was assigned to the  $\text{C}=\text{O}$  absorption on the side chains. The characteristic absorption peak of oxirane ring vibration was observed at  $912\text{ cm}^{-1}$ , which demonstrated that the epoxy group had been successfully introduced into DGENF monomer. In order to confirm that the curing reaction was completed, we also used FT-IR to characterize the cured epoxy samples. As shown in Fig. 4, the characteristic peak of epoxy group ( $912\text{ cm}^{-1}$ ) disappeared after curing with MeHHPA for all the samples. It is believed that the anhydride group of MeHHPA reacted with epoxy ring thoroughly to form ester groups in the presence of cured reagent and accelerator at high temperature.

### 3.3 Mechanical and thermal properties

The mechanical property is essential for epoxy materials for applications in electronic field. In this study, we used DMA to investigate the mechanical properties and glass transition temperature ( $T_g$ ) of the cured epoxy resins. As shown in Fig. 5,

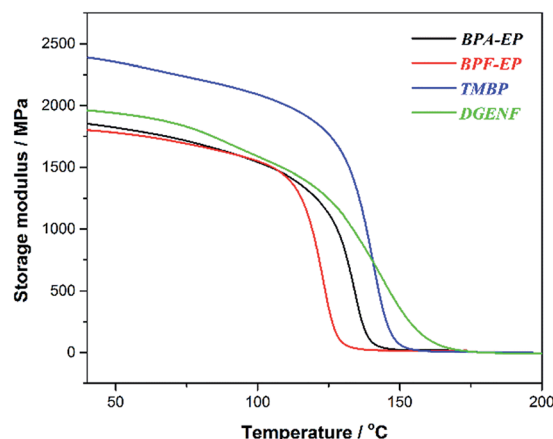


Fig. 5 Storage modulus curves of cured epoxy resin samples.

TMBP had a highest storage modulus (2375 MPa) among all epoxy resin samples within the experimental temperature range due to the biphenyl structure in TMBP, which improved the stiffness of the molecule chains and enhanced the mechanical strength. Compared to BPA-EP and BPF-EP, DGENF possessed a higher storage modulus (1976 MPa) at  $40\text{ }^{\circ}\text{C}$  and showed a slower decreasing rate with the temperature increasing. This result was ascribed to the naphthyl group with sufficient stiffness in DGENF, which also provided good thermal stability for the cured epoxy resin. Although DGENF possessed rigid naphthyl group, the existence of 4-fluorobenzoyl side chains decreased the cross-linking density of the cured epoxy resin, thus resulting in a relatively lower storage modulus than TMBP.

As shown in Fig. 6, the  $T_g$  of DGENF resin samples derived from  $\tan \delta_{\text{max}}$  is  $170\text{ }^{\circ}\text{C}$ , which is much higher than those of other three kinds of epoxy resin materials (Table 1). The result can be explained by the existence of stiff naphthyl group in DGENF, thus improving the rigid nature of polymer chains. Meanwhile, DGENF resin shows a relatively broad  $\tan \delta$  peak, which is indicative of the chain branching within the epoxy resin due to the introduction of 4-fluorobenzoyl side chains. DMA results indicated that this novel naphthyl epoxy

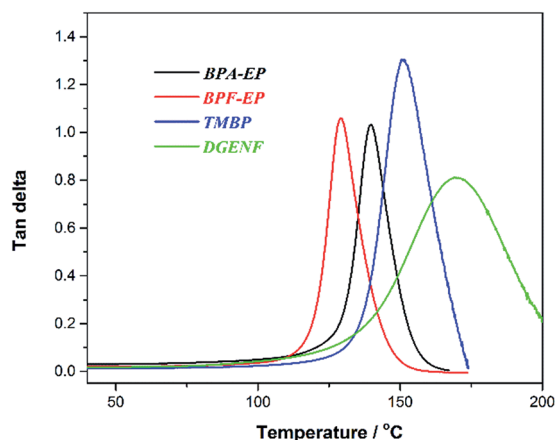


Fig. 6 Tan delta curves of cured epoxy resin samples.





Table 1 TGA, DMA and dielectric properties of cured epoxy resins

Samples	TGA (°C)			Char yield [%] At 700 °C	DMA		Dielectric properties at 1 MHz	
	$T_{5\%}$	$T_{10\%}$	$T_{50\%}$		Storage modulus (MPa)	$T_g$ (°C)	$D_k$	$D_f$
BPA-EP	334	384	453	14.5	1852	129	3.81	0.0219
BPF-EP	334	381	446	26.8	1800	139	3.95	0.0205
TMBP	313	340	450	7.7	2375	150	3.77	0.0244
DGENF	288	313	532	34.5	1976	170	2.97	0.0188

material could satisfy the demand of thermal and mechanical stabilities to be used as the electronic materials.

Fig. 7 shows the TGA curves of the cured epoxy resin samples measured under  $N_2$  atmospheres. The relevant thermal decomposition data, including the decomposition temperature of 5% weight loss ( $T_{5\%}$ ), 10% weight loss ( $T_{10\%}$ ) and 50% weight loss ( $T_{50\%}$ ), and the char residues at 700 °C are listed in Table 1. It was obviously found that DGENF exhibited a lower 5% weight loss temperatures ( $T_{5\%}$ ) of 288 °C than those of other kinds of epoxy resin materials (BPA-EP 334 °C, BPF-EP 334 °C and TMBP 313 °C). This result was mainly caused by two reasons. Firstly, the existence of 4-fluorobenzoyl side chains decreased the cross-linking density of the cured resins. Secondly, the side chains group could be degraded prior to the rigid main chain. This is also demonstrated by its tan delta curve with a relatively broad peak due to the introduction of 4-fluorobenzoyl side chains. However, as the temperature increased, the degradation rate of DGENF became much slower than other three kinds of epoxy resin samples. For example, as the temperature reached up to 400 °C, DGENF displayed a slow decreasing tendency in the thermal degradation, while the other epoxy resins maintained the continuous process of thermal degradation. Consequently,  $T_{50\%}$  of DGENF (532 °C) became much higher than BPA-EP (453 °C), BPF-EP (446 °C) and TMBP (450 °C). Meanwhile, the char yield of the epoxy resin samples at 700 °C listed in Table 1 indicated DGENF showed more residual (34.5%) than other three kinds of epoxy resins. It can therefore be deduced that the introduced naphthyl group improves the thermal stability of

cured epoxy resins at high temperatures efficiently and delays the degradation process of the material.

### 3.4 Contact angle and dielectric properties

Water absorption as one of the important property of materials not only greatly impair the mechanical and thermal properties of epoxy resins, but also increase relative dielectric constant even at small quantities, which widely limit the usage of epoxy resins as low- $k$  materials. Therefore we measured contact angles of all epoxy samples to determine the relative wettability of the surface. Fig. 8 exhibits the photographs and the result of contact angles. Photos represented the contact angles of BPA-EP ( $\vartheta = 78.2^\circ$ ), BPF-EP ( $\vartheta = 80.2^\circ$ ), TMBP ( $\vartheta = 83.2^\circ$ ) and DGENF ( $\vartheta = 116^\circ$ ), respectively. All the contact angles were less than  $90^\circ$  except DGENF, which was consistent with the standard of hydrophobic materials. This result was attributed to the stiff naphthalene moieties as the skeleton of the molecular chains and the hydrophobic fluorine in the side chains, thus decreasing the surface wettability of the material.

To the best of our knowledge, the signal propagation loss is related to the dielectric constant ( $D_k$ ) and the dielectric loss factor ( $D_f$ ), which are the most important parameters for epoxy resins used for the electronic materials. Therefore, selecting lower  $D_k$  and  $D_f$  material is beneficial for improving propagating speed and reducing the propagation loss. Fig. 9 and 10 showed the dielectric constant and dielectric loss of all these epoxy samples. It was clearly observed that DGENF had a remarkably lower dielectric constant ( $D_k = 2.97$  at 1 MHz) than that of other

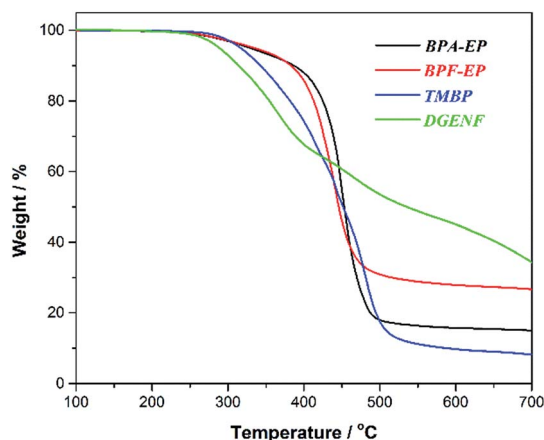


Fig. 7 TGA curves of cured epoxy samples.

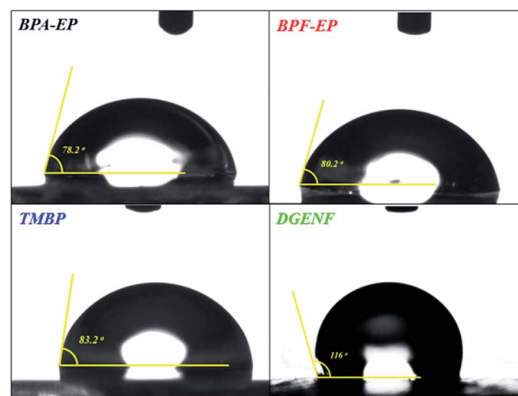


Fig. 8 Contact angles of cured epoxy samples.



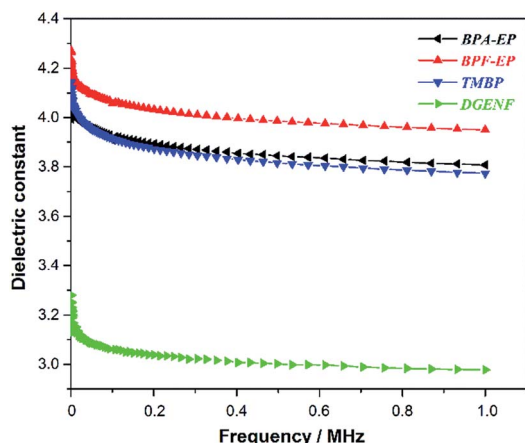


Fig. 9 Dielectric constants of cured epoxy samples.

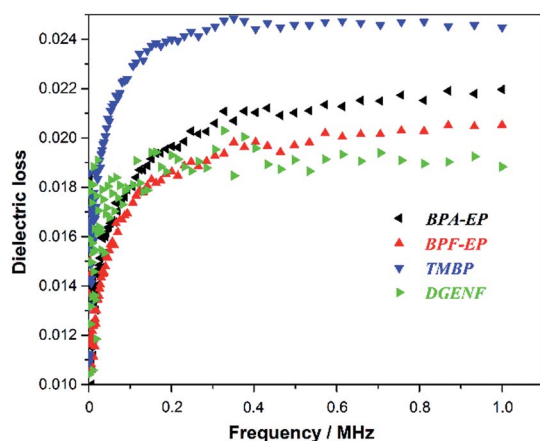


Fig. 10 Dielectric loss of cured epoxy samples.

kinds of conventional epoxy resins (BPA-EP, BPF-EP and TMBP). The result was owing to the existence of non-polar naphthalene moieties in the polymer main chains and 4-fluorobenzoyl groups in the side chain, which provided the small dipole of C-F bonds and thus making the molecule difficult to be polarized. As shown in Fig. 10, we could obviously observe that the dielectric loss of all of these epoxy materials showed an increasing tendency with the frequency increasing. Moreover, the  $D_f$  of DGENF (0.0188 at 1 MHz) was slightly lower than other three kinds of epoxy resin materials (BPA-EP: 0.0219, BPF: 0.0205 and TMBP: 0.0244). It can therefore be deduced that the introduction of non-polar naphthalene group and C-F bond into the polymer could modify the dielectric properties significantly. Thus we believed that this novel naphthyl epoxy resin could satisfy the demand to be used as potential electronic materials.

## Conclusions

A novel type of 1,5-bis(4-fluorobenzoyl)-2,6-diglycidyl ether naphthalene (DGENF) was synthesized *via* a three-step procedure involving Friedel-Crafts acylation, demethylation,

followed by nucleophilic reaction with ECH. Then a novel naphthyl epoxy resin containing 4-fluorobenzoyl side chains were prepared by curing with MeHHPA at high temperature. By comparing the properties of DGENF with other three kinds of conventional epoxy resins (BPA-EP, BPF-EP and TMBP), this novel naphthyl epoxy resin showed more hydrophobic and much lower dielectric properties. Therefore, DGENF is considered to be one of the most promising candidates to substitute future low- $k$  dielectrics in microelectronic. These results could motivate the research on the design of lower dielectric constant materials and the investigation of the relationship between the structure and properties of epoxy resins to be used as thermoset materials.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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## References

- 1 J. Hao, Y. Wei and J. Mu, *RSC Adv.*, 2016, **6**, 90.
- 2 J. Mo, W. Ma, W. Zhang, *et al.*, *Mater. Des.*, 2017, **128**, 56.
- 3 G. Maier, *Prog. Polym. Sci.*, 2001, **26**(1), 3.
- 4 Y. Huang, S. Zhang, H. Hu, *et al.*, *J. Polym. Sci., Part A: Polym. Chem.*, 2017, **55**(11), 1920.
- 5 W. Volksen, R. D. Miller and G. Dubois, *Chem. Rev.*, 2010, **110**(1), 56.
- 6 X. Zeng, L. Ye, K. Guo, *et al.*, *Adv. Bioelectron. Mater.*, 2016, **2**(5), 1500485.
- 7 T. Fukumaru, T. Fujigaya and N. Nakashima, *Polym. Chem.*, 2012, **3**(2), 369.
- 8 C. Yuan, J. Wang, K. Jin, *et al.*, *Macromolecules*, 2014, **47**(18), 6311.
- 9 Y. Wang, J. Yu, J. Zhu, *et al.*, *J. Polym. Sci., Part A: Polym. Chem.*, 2016, **54**(11), 1623.
- 10 M. Dogan and S. Unlu, *Polym. Degrad. Stab.*, 2014, **99**, 12.
- 11 J. Wang, L. Qian, B. Xu, W. Xi and X. Liu, *Polym. Degrad. Stab.*, 2015, **122**, 8.
- 12 B. Kandola, B. Biswas, D. Price and A. Richard, *Polym. Degrad. Stab.*, 2010, **95**, 144.
- 13 S. Tang, J. Qian, X. Liu and Y. Dong, *Polym. Degrad. Stab.*, 2016, **133**, 350.
- 14 Z. Chen, J. Yang, Q. Ni, S. Fu and Y. Huang, *Polymer*, 2009, **50**, 4753.
- 15 X. Wang, Y. Hu, L. Song, W. Xing, H. Lu, P. Lv and G. Jie, *Polymer*, 2010, **51**, 2435.
- 16 A. Konuray, X. Fernández-Francos and X. Ramis, *Polymer*, 2017, **116**, 191.
- 17 B. Yang, W. Wang and J. Huang, *Polymer*, 2015, **77**, 129.
- 18 D. Carolan, A. Ivankovic, A. Kinloch, S. Sprenger and A. Taylor, *Polymer*, 2016, **97**, 179.



- 19 S. Yang, Q. Zhang and Y. Hu, *Polym. Degrad. Stab.*, 2016, **133**, 358.
- 20 L. Luo, Y. Meng, T. Qiu, *et al.*, *Polym. Compos.*, 2013, **34**(7), 1051.
- 21 L. Kong, Y. Cheng, Y. Jin, *et al.*, *J. Appl. Polym. Sci.*, 2016, **133**(21), 43456.
- 22 C. Yuan, K. Jin, K. Li, *et al.*, *Adv. Mater.*, 2013, **25**(35), 4875.
- 23 Z. Liu, G. Zhang, Z. Liu, *et al.*, *Polym. Degrad. Stab.*, 2012, **97**(5), 691.
- 24 S. Sasaki, *J. Polym. Sci., Polym. Lett. Ed.*, 1986, **24**(6), 249.
- 25 Z. Tao, S. Yang, Z. Ge, *et al.*, *Eur. Polym. J.*, 2007, **43**(2), 550.
- 26 K. Xu, M. Chen, K. Zhang, *et al.*, *Polymer*, 2004, **45**(4), 1133.
- 27 Z. Liu, G. Zhang, K. Tu, *et al.*, *Polym. Int.*, 2011, **60**(10), 1556.

