Crystal structure of bis(1,8-dibenzoyl-7-methoxynaphthalen-2-yl) terephthalate: Terephthalate phenylene moiety acts as bidentate hydrogen acceptor of bidirectional C-H···π non-classical hydrogen bonds

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ABSTRACT

The title compound lies about a crystallographic inversion centre located at the terephthalate moiety. The two peri-benzenoarylphenylene units having atrope chirality are also situated centrosymmetrically. In the two peri-benzenoarylphenylene moieties, two benzoyl groups are substituted at 1 and 8 carbons of the naphthalene ring in anti-orientation. Then two absolute configurations of peri-benzenoarylphenylene moieties are consequently assigned as complementary to each other, i.e. one unit has R,R-configuration and the other S,S-one, respectively. The two benzoyl groups in peri-benzenoarylphenylene moiety and the terephthalate phenylene ring are non-coplanarly located against the naphthalene ring. The dihedral angles of each benzene ring of two benzoyl groups and terephthalate unit with the naphthalene ring are 73.73° and 75.96°, and 71.79°. In molecular packing, several kinds of weak interactions are responsible to induce three-dimensional molecular network. Especially, the synergetic effect realized through the bidentate hydrogen acceptor function in bidirectional C-H···π non-classical hydrogen bonds by the terephthalate phenylene ring moiety plausibly plays the determining role.

1. Introduction

Weak interactions in the solid of organic molecular substances have acquired interests and the reevaluation in the scientific area over the last two decades [1-3]. Indeed, weak interactions have been established as essential features in influencing the chemical, physical, and biological activity properties of materials [4-21], by determining their three-dimensional single molecular structure and spatial organization of molecular assembly. A number of papers and reviews have been dedicated to the crystallographic and theoretical study and consequently the analysis of halogen interactions and nonclassical hydrogen bonds such as C-H···X (X = Polar atom) and C-H···π hydrogen bonds have drastically increased over recent years, emphasizing the applicability and usefulness of weak interactions to scientific disciplines [22-30].

Congested molecules of accumulated aromatic rings have unique spatial structural restriction that the aromatic rings are compellingly arranged in non-coplanar manners. This structural constraint suggests that the contribution of parallel overlap of aromatic rings, i.e., π···π stacking, is extremely small. In this consequence, other weak interactions play relatively leading roles in the stabilization of the molecular packing. From this point of view, crystals of compounds of non-coplanarly accumulated aromatic rings can be expected to emphasize the contributions of rather weak non-covalent bonding interactions other than π···π stacking interactions or classical hydrogen bonds as structure determining factors. Based on the observation results of crystal structural analysis, the method for evaluating the actual stabilizing interactions of aromatic-ring-bearing molecules in solid states should be developed. The authors have investigated synthesis, reaction behavior, and analysis of spatial organization of single molecular and accumulation states of peri-aroylnaphthalene homologous compounds in which two aryl groups are substituted at the adjacent inner carbons of the naphthalene ring [31-34], accompanying with various related compounds [35,36].
In particular, the authors have reported the crystal structures of these compounds as nearly a hundred articles. peri-Aroylnaphthalene compounds have non-coplanarly accumulated structure of aromatic rings and the dihedral angles between aromatic rings are influenced by substituents on the aromatic rings [31]. As a part of the authors’ ongoing studies, herein, the X-ray crystal structure of the title compound, an aromatic-ring-accumulated organic substance composed of two 1,8-dibenzoylnaphthalene units and a terephthalate moiety, is reported and consequently the correlation among spatial organization of single molecule, molecular aggregation structure, and non-covalent bonding interactions via comparison with 1,8-dibenzoyl-2,7-dimethoxynaphthalene [37] is discussed (Figure 1).

2. Experimental

2.1. Instrumentation and materials

$^1$H NMR spectra were recorded on a JEOL JNM-AL300 spectrometer (300 MHz) and a JEOL JNM-ECA500 spectrometer (500 MHz). $^{13}$C NMR spectra were recorded on a JEOL JNM-AL300 spectrometer (75 MHz) and a JEOL JNM-ECA500 spectrometer (125 MHz). Chemical shifts were reported as the delta scale in ppm relative to CHCl$_3$ ($\delta = 7.29$ ppm) for $^1$H NMR and CDCl$_3$ ($\delta = 77.7$ ppm) for $^{13}$C NMR. Chemical shifts were reported as the delta scale in ppm relative to DMSO ($\delta = 2.50$ ppm) for $^1$H NMR and DMSO-d$_6$ ($\delta = 39.7$ ppm) for $^{13}$C NMR. Elemental analyses were carried out by using a SUMIGRAPH NCH-22 elemental analyzer. IR spectra were recorded on a JASCO FT/IR-4100 spectrometer. High-resolution FAB mass spectra were recorded on a JEOL MSStation (MS700) ion trap mass spectrometer in positive ion mode. All reagents were of commercial quality and were used as received. Solvents were dried and purified using standard techniques [38].

2.2. Synthesis

The title compound was prepared the following synthetic procedures presented in Scheme 1. Synthetic procedures of 2,7-dimethoxynaphthalene [39], 1-benzoyl-2,7-dimethoxynaphthalene [40] and 1-benzoyl-2-hydroxy-7-methoxynaphthalene [41] have been reported in the literatures.

2.2.1. Synthesis of 1,8-dibenzoyl-2-hydroxy-7-methoxynaphthalene

Benzoyl chloride (0.44 mmol, 0.05 mL), TiCl$_4$ (1.32 mmol, 0.14 mL) and dichloromethane (0.5 mL) were placed into a 10 mL flask, followed by stirring at room temperature. To the reaction mixture thus obtained, 1-benzoyl-2-hydroxy-7-methoxynaphthalene (0.2 mmol, 56 mg) was added. After the reaction mixture was stirred under reflux conditions for 3 h, it was poured into iced water (30 mL) and the mixture was extracted with chloroform (10 mL × 3). The combined extracts were washed with brine. The organic layers thus obtained were dried over anhydrous MgSO$_4$. The solvent was removed under reduced pressure to give a cake (yield 57%). The crude product was purified by column chromatography (ethyl acetate:hexane, v:v, 1:2, isolated yield 36%).

$^1$H NMR (CDCl$_3$, $\delta$): 7.90 (d, 2H, J = 8 Hz), 7.74 (d, 2H, J = 8 Hz), 7.62 (m, 4H), 7.34 (t, 1H, J = 8 Hz), 7.24 (d, 2H, J = 8 Hz), 7.15 (d, 2H, J = 8 Hz), 7.00 (m, 4H), 6.87 (d, 2H, J = 8 Hz), 6.75 (d, 2H, J = 8 Hz), 6.54 (d, 2H, J = 8 Hz), 6.37 (d, 2H, J = 8 Hz), 6.27 (d, 2H, J = 8 Hz), 6.00 (s, 2H), 3.74 (s, 3H), 3.71 (s, 3H).
The title compound was prepared by treatment of a mixture of 1,8-dibenzoyl-2-hydroxy-7-methoxynaphthalene (0.2 mmol, 0.11 equivalent) with triethylamine (0.22 mmol, 0.03 mL) in dichloromethane (0.5 mL). After the reaction mixture was stirred at rt for 3 h, it was poured into water (30 mL) and the mixture was extracted with chloroform (10 mL x 3). The combined extracts were washed with brine. The organic layers thus obtained were dried over anhydrous MgSO4. The solvent was removed under reduced pressure to give cake (yield 83%). The crude product was purified by crystallization from acetonitrile and colorless platelet crystals suitable for X-ray diffraction were obtained (isolated yield 30%).

**Table 2. Hydrogen bond geometry (Å, °) .**

| D···H-H | H···A-H | D···A | Z D···H-H | A
|----------|---------|-------|----------|------|
| 0.95     | 2.36    | 3.3052 | 175       |
| 0.95     | 2.59    | 3.4513 | 151       |
| 0.95     | 2.78    | 3.6247 | 149       |

* Symmetry code: (i) -1+x, y, z; (ii) 1-x, 1/2+y, 1/2-z; (iii) x, 1/2+y, 1/2-z; Cg is C27–C29-C29-C27-C29-C27 ring.

2.2.2. Synthesis of bis(1,8-dibenzoyl-7-methoxynaphthalen-2-yl)terephthalate

The title compound was prepared by treatment of a mixture of 1,8-dibenzoyl-2-hydroxy-7-methoxynaphthalene (0.2 mmol, 76 mg), terephthaloyl dichloride (0.11 mmol, 22 mg), and triethylamine (0.22 mmol, 0.03 mL) in dichloromethane (0.5 mL). After the reaction mixture was stirred at rt for 3 h, it was poured into water (30 mL) and the mixture was extracted with chloroform (10 mL x 3). The combined extracts were washed with brine. The organic layers thus obtained were dried over anhydrous MgSO4. The solvent was removed under reduced pressure to give cake (yield 83%). The crude product was purified by crystallization from acetonitrile and colorless platelet crystals suitable for X-ray diffraction were obtained (isolated yield 30%).

**Bis(1,8-dibenzoyl-7-methoxynaphthalen-2-yl)terephthalate:**
Color: Colorless. Yield: 85%. M.p.: 316.0-316.5 °C. FT-IR (KBr, ν, cm−1): 1739 (C=O) (ester), 1658 (C=O, Ar) (ketone, naphthalene), 1612 (Ar) (phenyl), 1598 (Ar) (phenyl), 1582 (Ar) (phenyl), 1510 (Ar), 1452 (CH=O) (ether), 1260 (Ar) (ether), 1175 (CH=O) (ether), 1015 (CH=O) (ether). 1H NMR (300 MHz, CDCl3), δ ppm: 7.33-7.40 (m, 14H, Ar-H), 7.49 (t, J = 7.2 Hz, 2H, Ar-H), 7.65 (d, J = 7.2 Hz, 4H, Ar-H), 7.71 (d, J = 7.2 Hz, 4H, Ar-H), 8.02 (d, J = 9.0 Hz, 2H, Ar-H). 8.07 (d, J = 9.3 Hz, 2H, Ar-H). 13C NMR (125 MHz, CDCl3), δ ppm: 155.0 [1C, Ar-C(OH)], 156.4 [1C, Ar-C(OCH3)], 163.0 [1C, Ar-C(O-C=O)], 76.84; H, 4.28. Found: C, 77.82; H, 4.36%.

2.3. X-ray diffraction analysis

For the crystal structure determination, the single crystal of the title compound was used for data collection on a four circle Rigaku RAXIS RAPID diffractometer (equipped with a two-dimensional area IP detector). The graphite-monochromated MoKα radiation (λ = 0.7070 Å) was used for data collection. The lattice parameters were determined by the least-squares method on the basis of all reflections with F2 > 2σ(F). Crystal data, data collection and structure refinement details are summarized in Table 1. The data collection and cell refinement were performed using PROCESS-AUTO software [42]. The data reduction was performed using CrystalStructure [43]. The structures were solved by direct methods using SHELXL04 [44] and refined by a full-matrix least-squares procedure using the program SHELX97 [45]. All H atoms were found in a difference map and were subsequently refined as riding atoms, with the aromatic C-H = 0.95 Å and methyl C-H = 0.98 Å, and with Uiso(H) = 1.2Ueq (C). The hydrogen bond geometries of the title compound are listed in Table 2. Crystallographic data of title compound has been deposited with the Cambridge Crystallographic Data Centre as Supplementary publication no. CCDC-2063371.

3. Results and discussion

Single molecular structure of the title compound is exhibited in Figure 2.
The dinaphthyl terephthalate molecule lies about a crystallographic inversion centre located at the terephthalate phenylene ring so that the asymmetric unit contains one-half of the molecules. Thus, the two peri-benzoylnaphthalene units are situated centrosymmetrically with respect to the centre of terephthalate phenylene ring. The spatial configurations of the two peri-benzoylnaphthalene moieties are assigned to $R,R$-form and $S,S$-form, respectively. The two benzoyl groups in peri-benzoylnaphthalene moiety and the terephthalate phenylene ring are non-coplanarly located to the naphthalene ring. The dihedral angles of each benzene ring with the naphthalene ring are $73.73^\circ$ for the outer benzoyl group, $75.96^\circ$ for the inner benzoyl group, and $71.79^\circ$ for the central terephthalate phenylene moiety. The five benzene rings in the molecule align in stacking manner toward alternative sides against naphthalene planes.

In the molecular packing, peri-benzoylnaphthalene units are stacked forming $R,R-R,R-R,R$- and $S,S-S,S-S,S$-configured columns (blue and red molecules) along the $a$-axis direction (Figure 3a). The naphthalene ring interdigitates with the oppositely configurated one in the adjacently aligned column, and vice versa, to make both configurated naphthalene moieties overlap alternatively. In addition, each column has been found to possess C-H(sp$^2$)···O(=C) non-classical hydrogen bonds between identically configurated dibenzoylnaphthalene units situated in adjacent columns to the bundle pillars. These C-H(sp$^2$)···O(Me) non-classical hydrogen bonds are formed between aromatic hydrogen atoms of the inner benzoyl group and the oxygen atom of the methoxy group ($C15-H15$···O5; 2.25 Å). On the other hand, C-H(sp$^2$)···π (terephthalate) non-classical hydrogen bonds link the oppositely configurated isomeric diaroylnaphthalene units between columns (Figure 4). Moreover, the C-H(sp$^2$)···π (terephthalate) nonclassical hydrogen bonds are found between aromatic hydrogen atoms of outside benzoyl groups on the dibenzoylnaphthalene moiety and the terephthalate phenylene ring ($C22-H22$···π; 2.78 Å).

The authors’ group has already reported the crystal structure of a homologous compound, i.e., 1,8-dibenzoyl-2,7-dimethoxynaphthalene, one of the simplest homologues in the authors’ series of peri-aroylnaphthalene compounds (Figure 5) [37]. In the crystal of the homologous compound, the molecules show chirality caused by atrope isomerization as well as the title compound. Furthermore, molecules are located on a twofold rotation axis. Therefore, the two benzoyl groups are situated in an anti-orientation and the dihedral angle between the mean planes of the benzene ring and the naphthalene ring system is $80.25(6)^\circ$. The phenyl and carbonyl groups are arranged almost coplanarly. Figure 6a shows the molecular packing structure in which $R,R$- and $S,S$-isomeric molecules are expressed as blue and red molecules, respectively. The crystal packing is stabilized by (methoxy)C-H···C=O and C-H(sp$^2$)···π (terephthalate) nonclassical hydrogen bonds (C14-H14B···O1 = 2.39 Å; C12-H12···O1 = 2.60 Å) and π···π stacking interaction between benzoyl benzene rings [centroid-centroid and interplanar distances are 3.6383 (10) and 3.294 Å, respectively].
The homologous compound has a sole value for the dihedral angle between each benzene ring and naphthalene and then lies on a two-fold rotation axis. The dihedral is closer to 90° than those of title compound. The high symmetricity of the spatial organization of the homologous compound indicates that two aryl groups take suitably effective positions for forming π···π stacking interactions. On the other hand, in title compound, the π···π stacking interaction formation between aryl groups is disturbed by steric hindrance brought about through the combination of bulkiness of largely congested two peri-benzoylnaphthalene units with terephthalate moiety. The benzoyl groups of peri-benzoylnaphthalene moiety in the title compound form effective C-H(sp²)···O=C and C-H(sp²)···O(-Me) non-classical hydrogen bonds instead of π···π stacking interactions (Figures 3b and 4). Furthermore, the terephthalate moiety on the inversion centre behaves as bidentate hydrogen acceptor of bidirectional C-H···π non-classical hydrogen bonds (Figure 4). This bidentate acceptant function of the terephthalate benzene ring is very unique behavior and substantially determines the whole crystal structure of the title compound. On the comparison with the homologous compound, the bidentate interaction is considered as transformation of the

Figure 4. C-H(sp²)···O(-Me) and C-H(sp²)···π(terephthalate) non-classical hydrogen bonds between the columns along c-axis. R,R- and S,S-isomers are expressed as blue and red molecules, respectively.

Figure 5. Single molecular structure of 1,8-dibenzoyl-2,7-dimethoxynaphthalene.

Figure 6. Molecular packing structures of (a) 1,8-dibenzoyl-2,7-dimethoxynaphthalene and (b) title compound with symmetry elements. R,R- and S,S-isomers are expressed as blue and red molecules, respectively.
π···π stacking interaction observed in the homologous compound through slipping of the two parallel benzene rings into the coplanar position accompanying with insertion of the benzene ring of terephthalate to result in achievement of synergetic enhancement of this type of noncovalent bonding interaction (Figures 6a and 6b). Consequently, different types of rather weak nonclassical hydrogen bonds cooperatively induce three-dimensional molecular network in the absence of determining support of effective non-covalent bonding interactions.

4. Conclusion

Conclusively, crystal structure of bis[(L8-dibenzoyl-7-methoxynaphthalen-2-yl) terephthalate and relationship among single molecular structure, accumulation structure, and effective influential non-covalent bonding interactions have been revealed. Two peri-benzoylnaphthalene units are situated centroaxially due to the inversion centre on the terephthalate moiety. The molecular packing was stabilized by several weak non-covalent bonding interactions, i.e., C=H(sp2)···O=C, C-H(sp3)C-H···OMe, and C-H(sp2)···π non-classical hydrogen bonds. The terephthalate moiety forms bidirectional C-H(sp2)···π non-classical hydrogen bonds as bidentate hydrogen acceptor. Comparison of spatial alignment in crystal with the homologous compound indicates that benzoyl groups in peri-benzoylnaphthalene unit could take a suitable conformation for forming several effective non-classical hydrogen bonds substitutable for impracticable π···π stacking interactions. The correlation among molecular structure, molecular packing structure, and weak interactions revealed in this study might be useful for the design of organic and polymer materials and understanding their properties.

Acknowledgements

The authors express their gratitude to Professor Keichi Noguchi, Instrumentation Analysis Center, Tokyo University of Agriculture and Technology, for technical advice. This work was partially supported by Tokyo Ohka Foundation for The Promotion of Science and Technology and JSPS KAKENHI Grant No. JP20K05473.

Supporting information

CCDC-2063371 contains the supplementary crystallographic data for this paper. These data can be obtained free of charge via https://www.ccdc.cam.ac.uk/structures/ or by e-mailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44(0)1223-336033.

Disclosure statement

Conflict of interests: The authors declare that they have no conflict of interest. Author contributions: All authors contributed equally to this work. Ethical approval: All ethical guidelines have been adhered.

Funding

Tokyo Ohka Foundation for The Promotion of Science and Technology https://doi.org/10.10139/501100010795

Japan Society for the Promotion of Science https://doi.org/10.10139/50110001691

References

[1]. Desiraju, G. R. Acc. Chem. Res. 1991, 24 (10), 290–296.
[2]. Desiraju, G. R. Acc. Chem. Res. 1996, 29 (9), 441–449.
[3]. Steiner, T. Chem. Commun. (Camb.) 1997, No. 8, 727–734.
[4]. Wang, J.-L.; Peng, H.-J.; Lin, Y.-J.; Jin, G.-X. J. Am. Chem. Soc. 2020, 142 (44), 18946–18954.
[5]. Gao, Y.; Liu, Q.; Wang, Q.; Li, Z.; Cai, J.; Zhao, T.; Lei, H.; Wang, S.; Zhang, Y.; Shen, B. Adv. Mater. 2020, 32 (40), e2005229.
[6]. Desiraju, G. R. J. Mol. Struct. 2001, 586 (1-3), 5–15.
[7]. Desiraju, G. R. Cryst. Growth Des. 2011, 11 (4), 896–898.
[8]. Aakeröy, C. B.; Srdižan, K. R. Chem. Rev. 1993, 22 (6), 397–407.
[9]. Desiraju, G. R. C. E. The Design of Organic Solids: Elsevier: Amsterdam, 1989.
[10]. Desiraju, G. R. Angew. Chem. Int. Ed. Engl. 1995, 34 (21), 2311–2327.
[11]. Iida, K.; Sakagawa, S.; Ichikawa, N.; Imamura, Y.; Kato, N.; Tashiro, M.; Tsuchida, H.; Ogushi, T.; Sato, H.; Tohnai, N.; Miyata, M. J. Am. Chem. Soc. 2016, 138 (20), 6617–6620.
[12]. Iida, K.; Iba, H.; Utsui, T.; Tohnai, N.; Coquereul, G.; Sato, H.; Miyata, M. J. Crystal Growth Des. 2016, 16 (3), 1626–1635.
[13]. Budiman, Y. P.; Jayaraman, A.; Friedrich, A.; Kerner, F.; Radius, U.; Marder, T. B. J. Am. Chem. Soc. 2020, 142 (13), 6036–6050.
[14]. Bunde, C.; Koper, M. T. M. J. Am. Chem. Soc. 2019, 141 (30), 12071–12078.
[15]. Elsberg, J. G. D.; Anderson, S. N.; Tierney, D. L.; Reinheimer, E. W.; Berreau, L. Dalton Trans. 2021, 50 (5), 1712–1720.
[16]. Wozniak, D. L.; Hick, A. J.; Sabbers, W. A.; Dobereiner, G. E. Dalton Trans. 2019, 48 (37), 14138–14155.
[17]. Kang, C.; Zhang, Z.; Wee, V.; Usadi, A. K.; Calabro, D. C.; Baugh, L. S.; Wang, S.; Wang, Y.; Zhao, D. J Am. Chem. Soc. 2020, 142 (30), 12995–13002.
[18]. Dionne, E. R.; Dip, C.; Troader, V.; Badia, A. J. Am. Chem. Soc. 2018, 140 (32), 10063–10066.
[19]. Tian, X.; Xin, X.; Gao, Y.; Han, Z. Crystallogr. Commun. 2020, 20 (11), 1580–1596.
[20]. Gargallo, R.; Aviñó, A.; Eritja, R.; Jarosova, P.; Mazzini, S.; Scaglioni, L.; Taborsky, P.; Ertl, U.; Zahn, M.; Tsuchida, H.; Ogushi, T.; Sato, H.; Tohnai, N.; Miyata, M. J. Am. Chem. Soc. 2016, 138 (20), 6617–6620.
[21]. Budiman, Y. P.; Jayaraman, A.; Friedrich, A.; Kerner, F.; Radius, U.; Marder, T. B. J. Am. Chem. Soc. 2020, 142 (13), 6036–6050.
[22]. Bunde, C.; Koper, M. T. M. J. Am. Chem. Soc. 2019, 141 (30), 12071–12078.
[23]. Elsberg, J. G. D.; Anderson, S. N.; Tierney, D. L.; Reinheimer, E. W.; Berreau, L. Dalton Trans. 2021, 50 (5), 1712–1720.
[24]. Wozniak, D. L.; Hick, A. J.; Sabbers, W. A.; Dobereiner, G. E. Dalton Trans. 2019, 48 (37), 14138–14155.
[25]. Kang, C.; Zhang, Z.; Wee, V.; Usadi, A. K.; Calabro, D. C.; Baugh, L. S.; Wang, S.; Wang, Y.; Zhao, D. J Am. Chem. Soc. 2020, 142 (30), 12995–13002.
[26]. Dionne, E. R.; Dip, C.; Troader, V.; Badia, A. J. Am. Chem. Soc. 2018, 140 (32), 10063–10066.
[27]. Tian, X.; Xin, X.; Gao, Y.; Han, Z. Crystallogr. Commun. 2020, 20 (11), 1580–1596.
[28]. Gargallo, R.; Aviñó, A.; Eritja, R.; Jarosova, P.; Mazzini, S.; Scaglioni, L.; Taborsky, P.; Ertl, U.; Zahn, M.; Tsuchida, H.; Ogushi, T.; Sato, H.; Tohnai, N.; Miyata, M. J. Am. Chem. Soc. 2016, 138 (20), 6617–6620.
[29]. Budiman, Y. P.; Jayaraman, A.; Friedrich, A.; Kerner, F.; Radius, U.; Marder, T. B. J. Am. Chem. Soc. 2020, 142 (13), 6036–6050.
[30]. Bunde, C.; Koper, M. T. M. J. Am. Chem. Soc. 2019, 141 (30), 12071–12078.
[31]. Elsberg, J. G. D.; Anderson, S. N.; Tierney, D. L.; Reinheimer, E. W.; Berreau, L. Dalton Trans. 2021, 50 (5), 1712–1720.
[32]. Wozniak, D. L.; Hick, A. J.; Sabbers, W. A.; Dobereiner, G. E. Dalton Trans. 2019, 48 (37), 14138–14155.
[33]. Kang, C.; Zhang, Z.; Wee, V.; Usadi, A. K.; Calabro, D. C.; Baugh, L. S.; Wang, S.; Wang, Y.; Zhao, D. J Am. Chem. Soc. 2020, 142 (30), 12995–13002.
