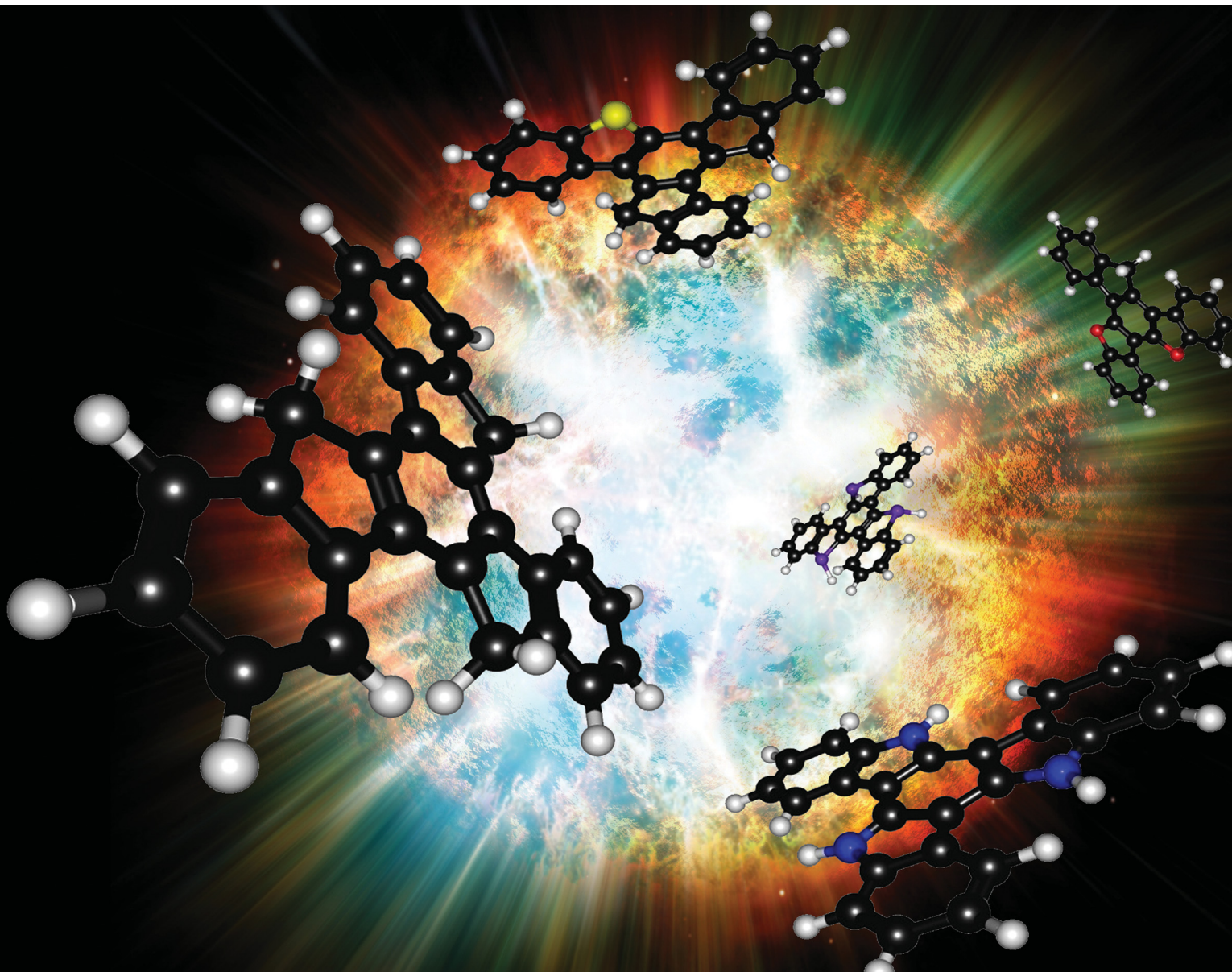


NJC

New Journal of Chemistry
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A journal for new directions in chemistry



ISSN 1144-0546

PERSPECTIVE

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Cite this: *New J. Chem.*, 2022, 46, 8939

Received 17th February 2022,
Accepted 22nd March 2022

DOI: 10.1039/d2nj00816e

rsc.li/njc

From truxenes to heterotruxenes: playing with heteroatoms and the symmetry of molecules

Krzysztof Górski,^a Justyna Mech-Piskorz^b and Marek Pietraszkiewicz^b

The truxene molecule is more versatile and widespread than ever imagined. At the end of the last century, this star-shaped structure was recognized mostly in liquid crystals. However, the development of optoelectronics resulted in a revolution in truxenes. Stable, fast, easily processed and cheap optoelectronic devices are desirable in many areas of industry and science. Consequently, the contribution of truxenes to these fields is manifold, ranging from D–A emitters, semiconductors, and complexes to sensors. They attract attention because of their relatively low mass and expanded π -electron system. Symmetric molecules give rise to dendrimers, bowl-shape molecules and fullerenes. Further advantages are shown by the recently discovered non-symmetric truxenes obtained via one- or two-heteroatom displacement. Thus, the chemistry of new heteroaromatic compounds has been expanded.

1. Introduction

Truxene is a C_{3h} aromatic hydrocarbon possessing seven fused rings, specifically, three five-membered and four six-membered rings, as shown in Fig. 1. Nowadays, this name also refers to its heteroanalogues. Most (hetero)truxenes are cyclotrimerization products of commercially available ketones and oxindoles.

To obtain mono- and diheterotruxenes, 6π -electrocyclization is required. The latter requires the synthesis of specific substrates because they are not commercially available. Truxenes has attracted vast interest, similar to other polycyclic aromatics, in the last few years due to the growing demand for organic emitters and semiconductors. Accordingly, scientific efforts are focused on the photostability of materials, purity of their colours, and increasing their charge carrier mobility. Easy processing is also an essential factor. This trend includes truxenes, as evidenced by numerous review papers.^{1–5} The typical features of truxenes is their significant thermal and

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research interests are focused on the synthesis and physicochemistry of (hetero)aromatic functional dyes.

Dr Krzysztof Górski received his MSc from Warsaw University in 2012 under the supervision of Professor Marek Pietraszkiewicz and Professor Andrzej Kudelski. He continued his research on heterotruxenes at the Institute of Physical Chemistry of the Polish Academy of Sciences, where he obtained his PhD in 2020 under the supervision of Professor Marek Pietraszkiewicz. Currently, he is a Postdoctoral Researcher in Prof. Daniel Gryko's group. His



Justyna Mech-Piskorz

started her own project at the Institute of Physical Chemistry PAS in 2015 concerning the physicochemical aspects of hybrid materials. Currently, she is working on non-linear effects in truxenes and its hybrids.

Dr Justyna Mech-Piskorz was born in Bochnia (Poland). She graduated with her MSc Eng. from Jagiellonian University, specializing in photonics and nanotechnology and supervised by Dr Jerzy Zachorowicz. During her PhD studies, she participated in interdisciplinary studies involving chemistry and solid state physics. She graduated with a PhD from AGH University of Science and Technology, supervised by Prof. Konrad



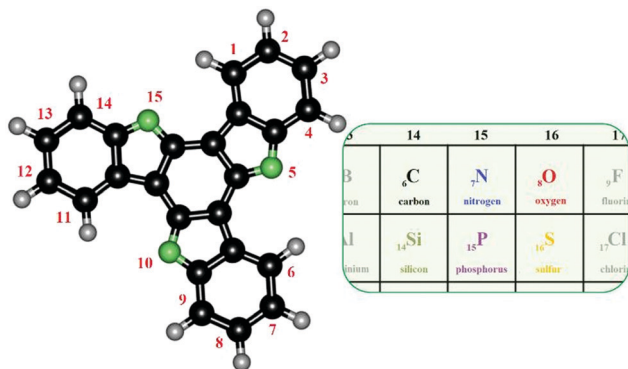


Fig. 1 Possible structures of (hetero)truxenes.

photostability. Truxenes and their heteroanalogues can be utilized in OLEDs,^{6–9} OFETs,^{10–12} DSSCs,^{13–15} NLO,^{16–19} sensors,^{20,21} gases and small molecule storage^{22–24} materials, and many more. The enormous diversity of truxenes is due to the unprecedented possibilities for their functionalization. Their modification can be realized on two levels. The first level is individual molecule derivatization, and the second level is building structured materials from the bottom up. In this manner, materials with designed physicochemical properties and structures can be obtained. Furthermore, heteroatom displacement offers the unique possibility of electronic structure tuning.

One example is 5,10,15-triazatruxene derivatives, which are characterized by a high-lying HOMO of -5.1 eV and exhibit good hole mobility of $8.9 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$.²⁵ Fig. 1 presents the various configurations of heteroatom(s) insertion. Truxenes, due to their specific structure, can be polyradicals.²⁶ They are utilized to synthesize strained aromatic moieties,²⁷ including

fullerenes,²⁸ chiral molecules,^{29,30} and supramolecular units.^{31,32} The main goal of this review is to present the structure-electronic relationship of truxenes. This information can be beneficial for the appropriate design of truxene-like structures. In the present review, truxenes are systematized in terms of their symmetry (presence or absence of a triple-axis of symmetry). Then, in each of these groups, the different heteroatoms are distinguished.

2. C_{3h} -Symmetric truxenes

2.1. Truxene: synthesis and reactivity

Truxene **4**, for the first time, was accidentally obtained in 1889 by Hausmann during the synthesis of indan-1-one **2** from 3-phenylpropanoic acid **1** (Scheme 1). The mechanism of this reaction, with the participation of the α,β -unsaturated intermediate **3**, was elucidated many years later.³³ This cyclotrimerization of aldol is still the most frequently used method for the synthesis of truxenes.

H^+ is necessary to initiate condensation, and hence **4** is formed in strong acidic media, e.g., polyphosphoric acid³⁴ or mixture of conc. AcOH and aqueous HCl.³⁵ The reaction yield of the latter is a record-breaking 98%. Truxenes are also obtained from appropriately functionalized indan-1-ones to synthesize, for example, hexamethoxy derivatives.³⁶ Other truxene derivatives were obtained under harsher conditions but with a lower yield (Scheme 2).³⁷ This can be explained by the negligible solubility of the corresponding α,β -unsaturated ketone.

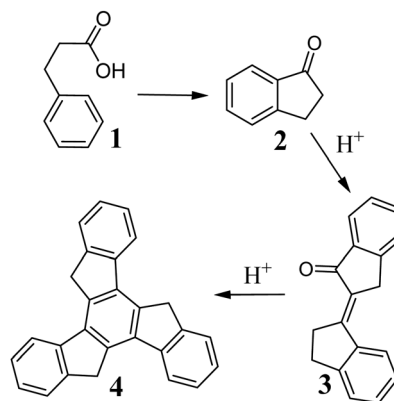
The described methods require a large excess of acid relative to ketone, but this is not necessary for efficient cyclotrimerization. The use of 3.5 equiv. of p-TsOH and propionic acid in 1,2-dichlorobenzene at 105 °C allows the efficient condensation (59–89%) of indan-1-ones to the corresponding truxenes.³⁸ Further, using 1 equiv. of MeSO_3H with $\text{Si}(\text{OEt})_4$ in DCM at room temperature allows the cyclotrimerization of **2** with 80% yield.³⁹ Indan-1-ones also undergo condensation in the presence of a catalytic amount of $\text{Rh}(\text{III})$ complex at 150–200 °C without solvent.⁴⁰ The cyclotrimerization of more complexed indanones in the presence of POCl_3 is worth mentioning.⁴¹ Isomeric compounds **2** and **8** undergo cyclotrimerization to **4** in the presence of SiCl_4 in EtOH (Scheme 3).⁴²



Marek Pietraszkiewicz

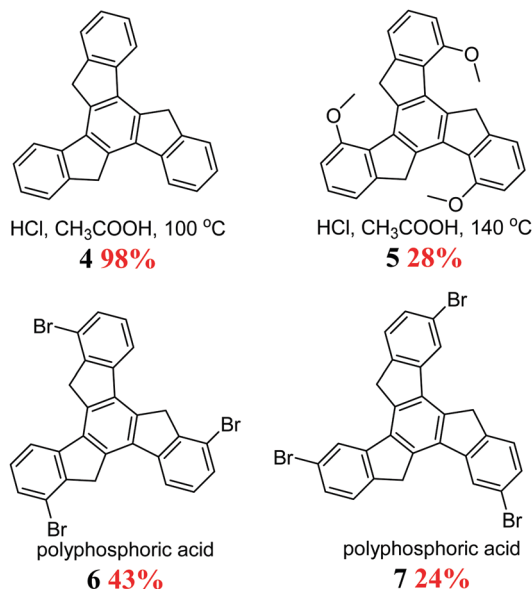
Prof. Marek Pietraszkiewicz was born in Warsaw (Poland). He obtained his PhD at the Institute of Organic Chemistry, Polish Academy of Sciences, in 1978, with Prof. Osman Achmatowicz as his supervisor. He was a Post-Doc Researcher at Orlean University, supervised by Prof. Pierre Sinay, working on the total synthesis of cerulenin antibiotics. He held a second Post-Doc position at University Louis Pasteur, Strasbourg, France, supervised by

Prof. Jean-Marie Lehn, working on photoluminescent lanthanide complexes. He obtained his DSci (habilitation) at the Institute of Physical Chemistry, Polish Academy of Sciences. His scientific research interests include organic synthesis, supramolecular chemistry, lanthanide photoluminescent complexes, and materials for organic electronics. He is the founder and chairman of the Polish and Global Supramolecular Chemistry Networks.



Scheme 1 First truxene synthesis.

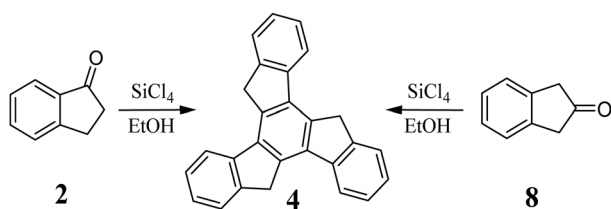




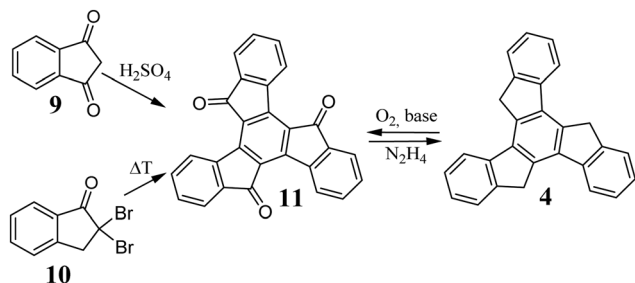
Scheme 2 Indan-1-one cyclotrimerization products, conditions and reaction yields.

However, the broader use of indan-2-ones for the synthesis of truxenes is limited, mainly due to their availability.

The described cyclotrimerization processes allow truxenes that are unfunctionalized in positions 5, 10, and 15 to be obtained. These positions are important for modifying the electronic structure and improving the solubility of truxenes. Indan-1,3-dione **9** in the presence of concentrated H₂SO₄ forms truxene-5,10,15-trione **11** (Scheme 4).³⁵ Microwave-assisted cyclotrimerization significantly reduces the reaction time while maintaining a high yield of 77%.⁴³ Likewise, 2,2-dibromoindan-1-one **10**, under pyrolysis condition, undergoes cyclotrimerization to form **11**.⁴⁴ The main advantage of using **9** and **10** is the direct synthesis of



Scheme 3 Cyclotrimerization in the presence of $\text{SiCl}_4/\text{EtOH}$.



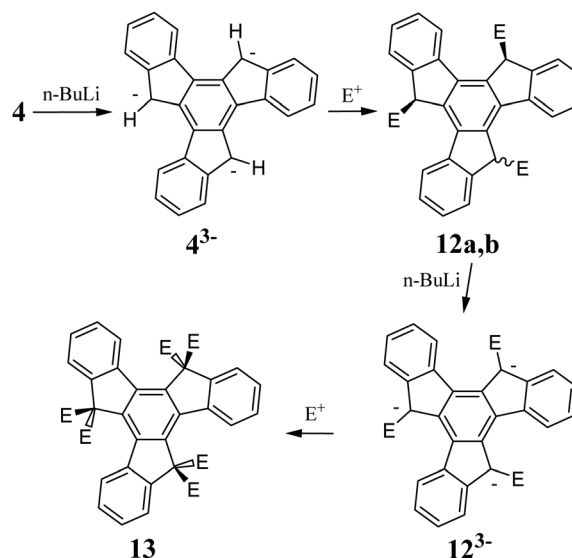
Scheme 4 Synthesis of truxene-5,10,15-trione.

functionalized truxene in positions 5, 10, and 15. The carbonyl groups in **11** undergo ketone-specific reactions, and hence N_2H_4 can convert it to **4**.⁴⁵ Reverse conversion, *i.e.*, oxidation of the methylene moieties of **4** in positions 5, 10, and 15, is also possible. The use of a composite of KOH and graphene⁴⁶ or a suspension of KOH in THF⁴⁷ in the presence of O_2 efficiently oxidizes **4** to **11**. The base used in this process must be strong enough to ionize the 5, 10, and 15 positions; therefore, KOH can be replaced by Me_4NOH or K_2CO_3 .⁴⁸ Recent studies indicate that in the presence of light and O_2 , **4** can be oxidized to a monocarbonyl derivative, namely truxene-5-on.⁴⁹

Unsubstituted truxenes display low solubility, which is a direct consequence of their strong π -stacking interactions. Thus, to reduce this intermolecular effect, alkyl substituents are introduced in the 5, 10, and 15 positions. Strong bases can ionize the acidic hydrogen atoms in a five-membered ring. The produced aromatic anions 4^{3-} and 12^{3-} can serve as valuable substrates for further derivatization of the truxene core, for example, *C*-alkylation (Scheme 5).

The alkali metal hydrides NaH^{50} or KH^{51} and organolithium compounds are the most commonly used bases to generate 4^{3-} . In the reaction between **4** and *n*-BuLi, the orange fluorescent 4^{3-} is formed. Subsequent reaction of 4^{3-} with, *e.g.*, alkyl halide, results in a mixture of two diastereomers, containing mainly the anti-product **12a**. The syn-product **12b** can be obtained from **12a** after reacting with *t*-BuOK in *t*-BuOH.⁵² In the case that the solubility of the reactants appears to be insufficient, DCM is added to increase their solubility and the isomerization yield. A mixture of **12a** and **b** can be ionized and converted to the corresponding hexa-substituted truxenes **13**. Alkyl or benzyl halides are the most commonly used electrophilic agents. However, by using appropriate dibromoalkanes, spiro systems **15** can be obtained (Scheme 6).⁵³

The isolation of products can be troublesome due to their similar polarity after alkylation is performed. Thus, for



Scheme 5 Triple lithiation of **4**.

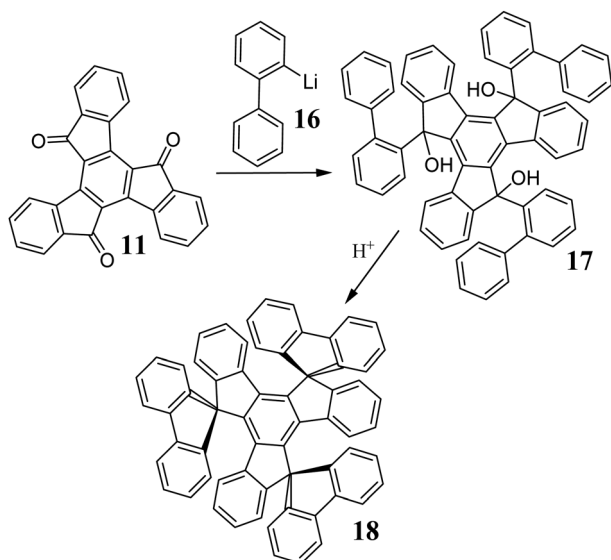
Chemical reaction scheme showing the synthesis of compound **21** from compound **19**.

Compound **19** (a triphenylmethane derivative with three ethyl ester groups) reacts with ArLi to form intermediate **20** (a triphenylmethane derivative with three hydroxyl groups and three Ar groups).

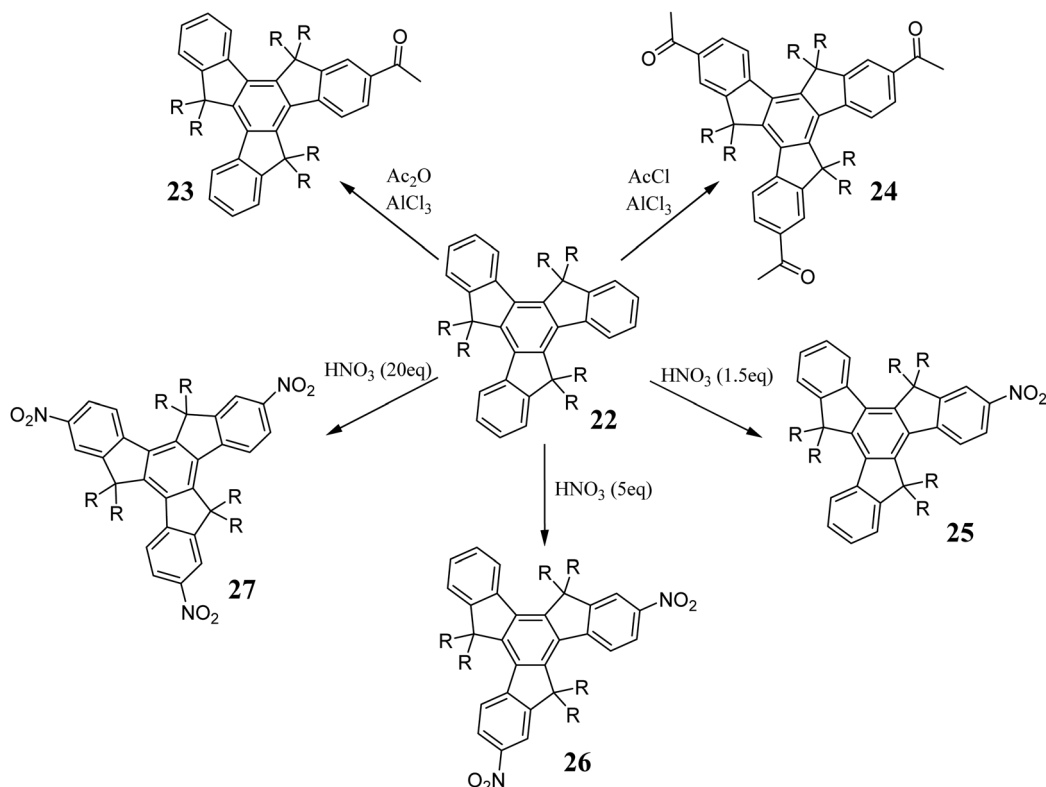
Intermediate **20** then undergoes acid-catalyzed cyclization (H^+) to form the final product **21** (a triphenylmethane derivative with three Ar groups).

The main limitation of the described methods is the low solubility of **11**. Therefore, some modifications may prove challenging to implement. An alternative approach to introduce aryl

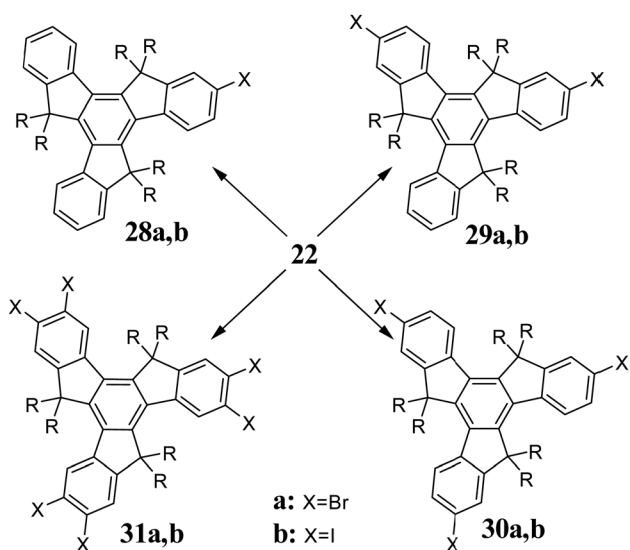
Halogenated truxenes undergo lithium-halogen exchange, which enables the introduction of a carboxyl⁷⁶ or formyl^{77,78} group or their conversion to appropriate boronic acids.⁷⁹



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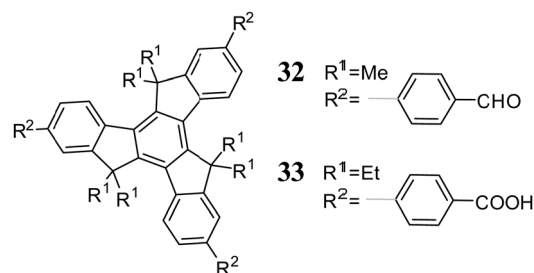
Scheme 9 Electrophilic substitution of truxene.



Scheme 10 Halogenation of truxenes.

Halogen atom replacement with a boronic ester was also carried out catalytically using $\text{Pd}(\text{dppf})_2\text{Cl}_2$.⁸⁰ Truxenes also undergo direct triple borylation in the presence of $[\text{Ir}(\text{OCH}_3)(\text{COD})]_2$.⁸¹ Under Ulman⁸² or Buchwald-Hartwig⁸³ reaction, halogenotruxenes are converted into the corresponding 3° amines. However, the most crucial halogenotruxene transformation is Pd-catalysed coupling. The peripheral functionalization *via* Suzuki reaction allows easy

π -electron expansion of the truxene core.⁸⁴ Halogenotruxenes react with simple boronic acids^{85–88} and π -expanded derivatives.^{89–96} The connection of benzoic acid and benzaldehyde is also possible *via* this process.⁹⁷ The mentioned Suzuki transformation allows the synthesis of yellow⁹⁸ and white⁹⁹ polymeric emitters, as well as the creation of ordered porous materials, namely, two-dimensional COFs (covalent organic frameworks) **32**¹⁰⁰ and three-dimensional MOFs (metal-organic frameworks) **33**¹⁰¹ (Scheme 11). The microwave-assisted coupling of halogenotruxenes with boronic acids allows the introduction of up to six oligofluorene moieties with high reaction yield.¹⁰² The Suzuki reaction allows the direct connection of truxene subunits, leading to truxene-only dendrimers¹⁰³ and polymers.¹⁰⁴ An alternative coupling often used to expand the π -electron truxene system is the Sonogashira reaction.^{105–110} Other Pd-catalyzed couplings, such as the Heck⁶⁷ and Negishi¹¹¹

Scheme 11 Structural subunits **32** – COF and **33** – MOF.

34a: R = 
34b: R = 
34c: R = 

Scheme 12 NLO liquid crystalline truxenes.

by Baunsgaard *et al.*¹²³ The $T_1 \rightarrow S_0$ transition is forbidden due to multiplicity and symmetry, resulting in a very long phosphorescence lifetime of about 16 s.¹²⁴ The truxene moiety exhibits a reversible oxidation process and irreversible reduction.¹²⁵ In contrast, in its derivatives, both redox processes are reversible,^{47,55,126} which is beneficial for optoelectronics.

Truxenes having long alkyl chains were the first derivatives thoroughly investigated. They exhibit liquid crystalline properties and rich polymorphism.^{127–132} Due to their strong π -stacking interactions, truxenes exhibit a tendency to undergo columnar ordering in the solid state. The aggregation combined with the reversible reduction process of truxene-5,10,15-trione makes it promising for the synthesis of new n-type semiconductors.¹³³ Recent studies also suggest the possibility of using truxenes decorated with long alkoxy chains in NLO (Scheme 12).¹³⁴

Truxene **35** has been the subject of many application studies (Scheme 13).^{135–139} Regardless of the substituent type, **35** has very high fluorescence quantum yields up to 99%.¹⁴⁰ The π -electron system expansion upon oligoaryl chain elongation is visible by the redshift in its absorption and emission spectra.^{141,142} **35** is thermally and electrochemically stable,¹⁴³ even exhibiting chemiluminescence.¹⁴⁴ These properties make **35** and its derivatives valuable compounds for LEDs,¹⁴⁵ OPVs,¹⁴⁶ and OFETs.¹¹ They also act as good emitters¹⁴⁷ and light-harvesting dyes.^{148,149}

35b has also been tested for amplified spontaneous emission (ASE).^{150,151} Similar truxene structures substituted by oligofluorenes¹⁵² and oligofluorenes terminated with pyrenes¹⁵³ or diphenylamines^{154,155} were tested in a blue laser. The non-linear properties of truxenes **35b** and **e** reveal significant two-photon absorption (TPA).^{156,157} Theoretical studies suggest that **35c** should also exhibit similar properties.¹⁵⁸ Besides optoelectronic applications, truxenes,¹⁵⁹ truxenes modified by thienyl chains,^{160,161} and dendrimeric structures made of them,¹⁶² have been used as stationary phases in gas chromatography.

The presence of aromatic subunits enables tuning of the physicochemical properties of the π -electron system.¹⁶³ The introduction of donating subunits increases the electron density significantly in the truxene core, increasing its hole conductivity (Scheme 14).^{164–171} Its conducting properties can also be tuned differently. The presence of peripheral long alkoxy chains increases the intermolecular interactions, changing the conductivity of substituted truxenes, resulting in hole¹⁷² or ambipolar¹⁷³ conductivity. Donor subunits result in a red shift in the absorption and fluorescence spectra of truxene derivatives.¹⁷⁴ Consequently,

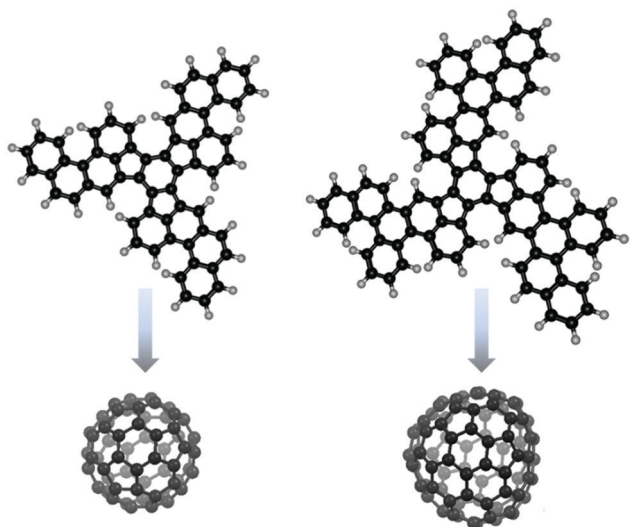
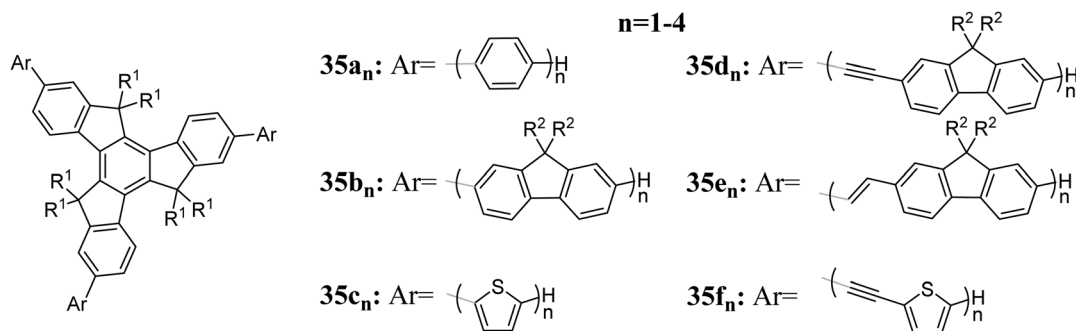
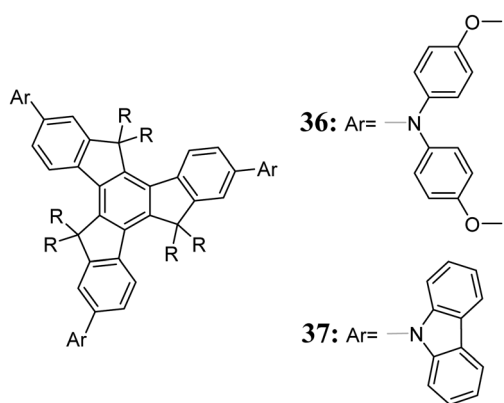


Fig. 3 Truxene-based fullerenes precursors: left – C₆₀ and right – C₈₄.



Scheme 13 Oligoaryl truxenes.

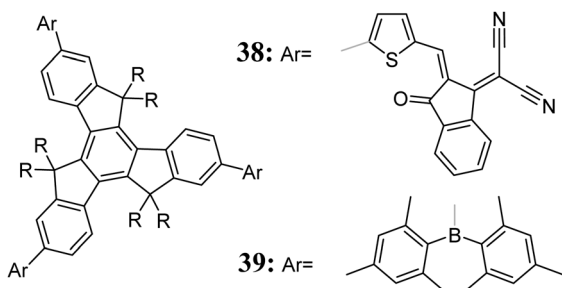


Scheme 14 Examples of electron-rich truxenes.

they can be used as blue emitters in OLEDs⁶ and lasers.¹⁷⁵ Recent research indicates that the presence of the truxene core significantly impacts the light-harvesting properties of materials.¹⁵

Alternatively, an acceptor unit, such as fullerene,¹⁷⁶ attached to truxene, increases its electron conductivity. Current photo-voltaic electron-deficient systems do not contain fullerene molecules.^{126,133,177–182} However, better electron conductivity is not the only advantage of electron-withdrawing substituents. π -Electron systems equipped with diarylboron substituents **39** (Scheme 15) are characterized by more than 4-fold higher fluorescence quantum yield compared to that with **4**.¹⁸³

The donor properties of dithiole-modified truxene and its characteristic concave shape can be used in noncovalent interaction with fullerene C₆₀, which is known for its accepting



Scheme 15 Examples of electron-deficient truxenes.

behavior. This noncovalent donor–acceptor pair is presented in Fig. 4.

Despite the already discussed donor or acceptor substituents for truxene, it is worth paying more attention to covalent D–A systems.^{184–188} The common feature of these systems is solvatochromism, resulting from excited-state charge separation.¹⁸⁹ The presence of D–A subunits in OPVs inhibits charge recombination.¹⁹⁰ This feature makes D–A truxenes applicable in dye-sensitized solar cells (DSSC).^{191,192}

Due to the high affinity of F[−] to the boron atom, some truxenes can act as fluorescent switches.¹⁹³ As the concentration of F[−] increases, the CT state fluorescence intensity decreases in favour of local emission, changing the colour of the emitted light. Besides, some truxenes can detect cations,^{194,195} nitro compounds,^{196,197} and even proteins containing metalloporphyrin subunits.¹⁹⁸ Recent studies showed that their sensing property is directly related to the presence of the truxene core and the substitution position.¹⁹⁹

Truxene-based MOFs and COFs can be used for the storage of gases or small molecules,²² as well as for HCl sensing.²⁰ Surprisingly, an adequately designed structure may lead to electron conductive materials²⁰⁰ or water desalination membranes.²⁰¹ Some truxene-based polymers show interesting sorption properties, *e.g.*, for CH₄.²⁰² Another truxene-based polymeric material exhibited photocatalytic properties²⁰³ and could be applied as solar fuels^{204,205} or for the homocoupling of amines.²⁰⁶

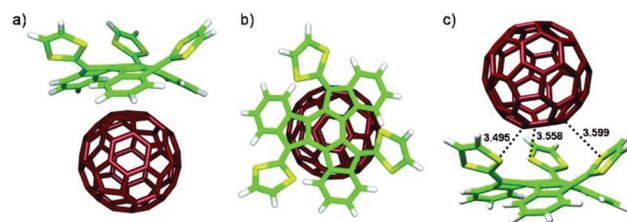


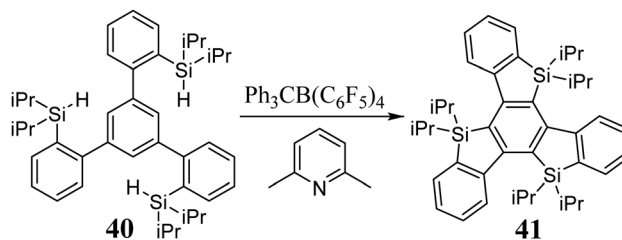
Fig. 4 Structures of the 3 a-C₆₀ complex calculated at the MPWB1K/6-31G** level. (a) Side view of the preferred configuration. (b) Top view of the same configuration, showing the stack between the central benzene ring of the truxene core and one of the hexagonal rings of C₆₀. (c) Side view of the other possible configuration, with C₆₀ approaching 3 a on the dithiole side. S–C short contacts [Å] are also shown. The carbon atoms of the fullerene are depicted in red for clarity. Reproduced from E. M. Pérez, M. Sierra, L. Sánchez, R. Torres, R. Viruela, P. M. Viruela, E. Orti, N. Martín, *Angew. Chem., Int. Ed.*, 2007, **46**, 1847–1851, with permission from John Wiley and Sons.

Every truxene monomer can be applied to create 2D or 3D materials in multiple ways, where DFT calculations can enable the preselection of suitable monomers. It is worth mentioning that truxene derivatives can be used for the amplification of chirality in discrete molecular assemblies, where one chiral molecule can control the chirality of the entire assembly.⁷⁸

The preparation of heavier analogues of truxenes has been unattainable for years. However, the silole ring formation development, including the sila-Friedel-Crafts reaction²⁰⁸ and catalytic silylation,²⁰⁹ enabled real attempts for the synthesis of 5,10,15-trisilatruxene. In 2017, Ogaki *et al.* synthesized an alkyl derivative of 5,10,15-trisilatruxene **41** (Scheme 16).¹²⁴

The three silicon atoms present in the truxene core do not change the symmetry of the system, and hence the first two-electron transitions, as in truxene, are forbidden. The energy of the HOMO is slightly lower, while that of the LUMO is significantly lower compared to truxene. This is due to the $\sigma^*-\pi^*$ interaction in the silole ring. Consequently, a redshift in the absorption and emission spectra is observed.

Generally, the presence of silicon atoms does not affect the fluorescence quantum yield but visibly changes the phosphorescent properties. Two intense peaks are present in the spectrum and the phosphorescence lifetime is four-times shorter, *i.e.*, $\tau_{\text{phos}} \approx 4$ s, than in truxene. Due to the relatively high energy of its T₁ state, **41** can be used as a sensitizer for blue phosphorescent OLEDs. In comparison to their carbon analogues, siloles have



Scheme 16 Synthesis of 5,10,15-trisilatruxene.

better electron conductivity,²¹⁰ and hence 5,10,15-trisilatruxenes can be considered semiconductors.

The 5,10,15-triazatruxene **47** was presented for the first time by Bergman *et al.* concerning the condensation of indole **42** and 2-oxindole **43** (Scheme 17).²¹¹ **47** is a product of a multistep synthesis. It begins from bisindole **44**, which is obtained during thermolysis of 2-iodoindole **45** or *via* the Vilsmeier reaction between **42** and **43**. Another Vilsmeier reaction between **43** and **44** gives trimer **46**, followed by conversion to **47** by cationic radical cyclization, with a total yield of 30.6%. Using activated Cu bronze at 205 °C allowed the conversion of **45** to **47** with 81% yield.²¹²

The electron-rich indoles **48a-e**, when treated with $(\text{NH}_4)_2\text{S}_2\text{O}_8$ ²¹³ or $\text{Na}_2\text{S}_2\text{O}_8$,²¹⁴ undergo oxidative cyclotrimerization, leading to isomeric aromatic compounds, *i.e.*, triazatruxene (**49a-e**) and triaisotruexene (**50a-e**) (Scheme 18). Trimerization in the presence of $\text{S}_2\text{O}_8^{2-}$ is strongly dependent on the solution pH; hence, the best reaction yields ($\sim 30\%$) are obtained at pH = 1.4. It should also be mentioned that **48f** and **g** under the same reaction conditions lead to triaisotruexene **50f** and **g** as the

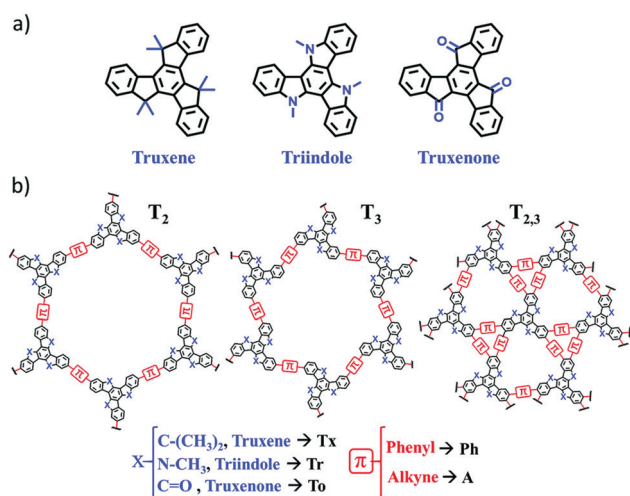
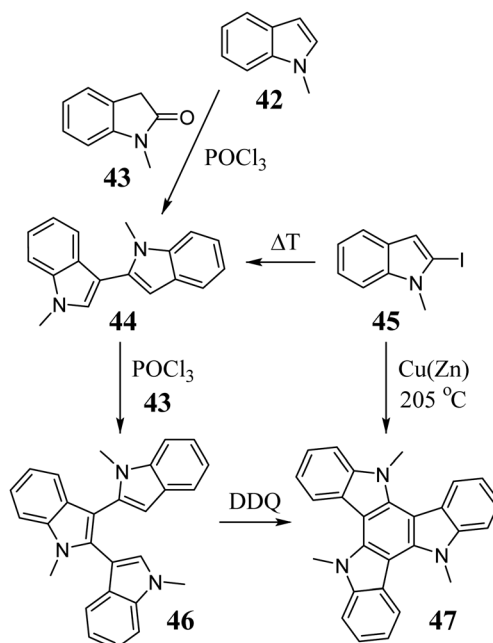
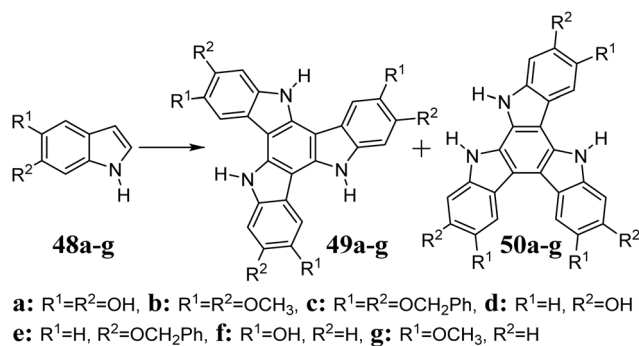


Fig. 5 General chemical structures of the conjugated cores (a) and 2D polymers (b) examined in this study with the notation used throughout the text. Note that the T_2 and T_3 polymers refer to tri-substituted materials with different linkage positions (2, 7, and 12 in T_2 and 3, 8, and 13 in T_3), while $T_{2,3}$ corresponds to hexa-substituted materials in the 2, 3, 7, 8, 12 and 13 positions. Reproduced from S. Gámez-Valenzuela, M. Echeverri, B. Gómez-Lor, J. I. Martínez, M. C. Ruiz Delgado, *J. Mater. Chem. C*, 2020, **8**, 15416–15425, with permission from The Royal Society of Chemistry.



Scheme 17 Synthetic approaches to **47**.

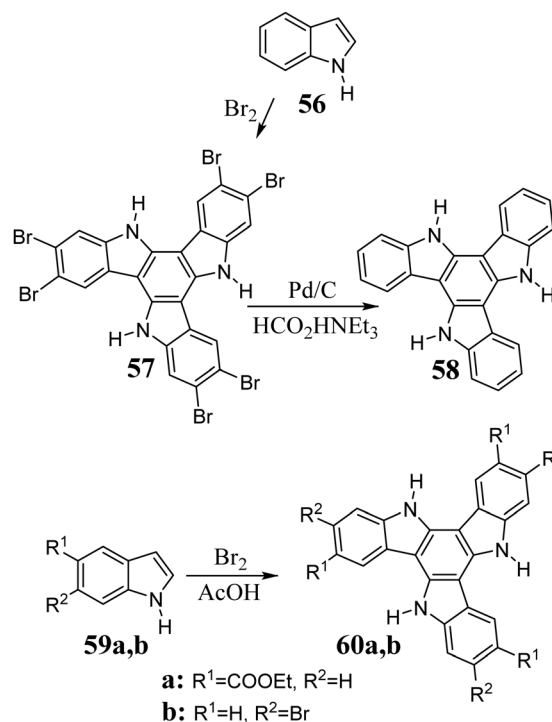
Scheme 18 Oxidative cyclotrimerization of **48**.

only cyclotrimer, respectively. However, due to the large share of byproduct **50** (yield ~20%), oxidative cyclotrimerization may not be an attractive method for the synthesis of 5,10,15-triazatruxenes.

In 2009, Ginnari-Satriani *et al.* reported the one-step synthesis of 5,10,15-triazatruxene **52** from 2-oxindole **51** (Scheme 19).²¹⁵ The result of this reaction is highly dependent on both the temperature and the concentration of $POCl_3$. As the concentration of $POCl_3$ increases, the yield of cyclotrimer **52** decreases in favour of cyclotetramer formation **53**.²¹⁶ The discovery of this method initiated further development, which allows the condensation of substituted 2-oxindoles **54**.^{16,217,218}

Indole **56** in the presence of Br_2 undergoes cyclotrimerization to 5,10,15-triazatruxene. The triazatruxene core is highly reactive towards electrophilic substitution and with excess Br_2 , leads to the hexabromo derivative **57**.²¹⁹ After the debromination of **57** in the presence of Pd/C and HCO_2HNEt_3 , **58** can be obtained (Scheme 20).²²⁰ Alternatively, some deactivated indoles can be synthesized *via* cyclotrimerization in $AcOH$ in the presence of Br_2 .²²¹

As a result of the reaction between indole and NBS, polybrominated 5,10,15-triazatruxenes are obtained. After debromination,

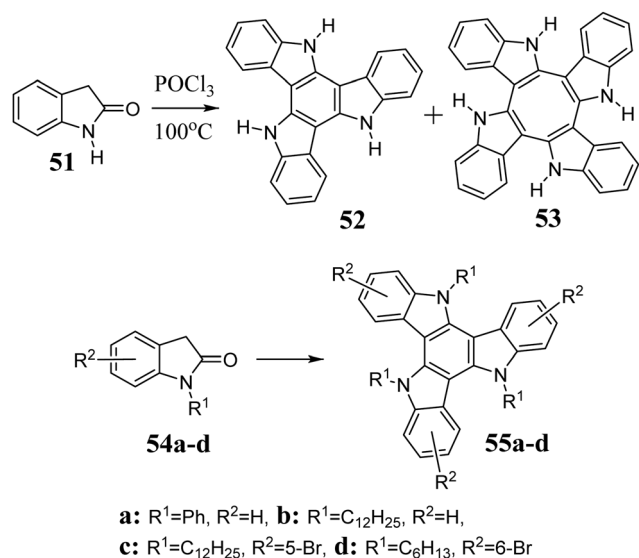


Scheme 20 Bromine-promoted indole cyclotrimerization.

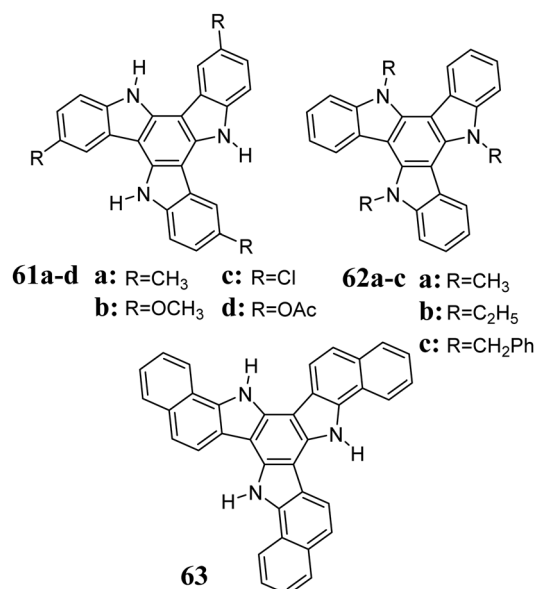
they lead to functionalized derivatives **61**, **62** and **63** (Scheme 21).²²²

5,10,15-Triazatruxene is very reactive toward electrophilic substitution. The most reactive positions are located opposite the nitrogen atom (3, 8, and 13 – green dots, Fig. 6). Alternatively, in the presence of excess electrophile, the substitution may proceed further at positions 2, 7 and 12 (blue dots).

5,10,15-Triazatruxene, in the presence of $TfOH$, reacts with 3° aromatic alcohols *via* triple electrophilic substitution.²²³



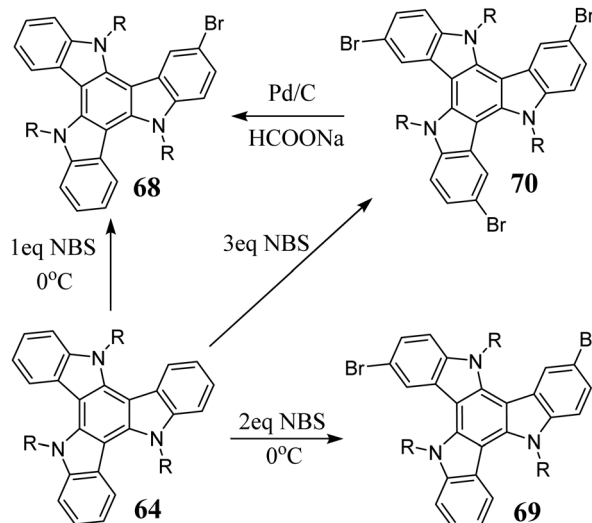
Scheme 19 2-Oxindole condensation.



Scheme 21 Indole condensation products.

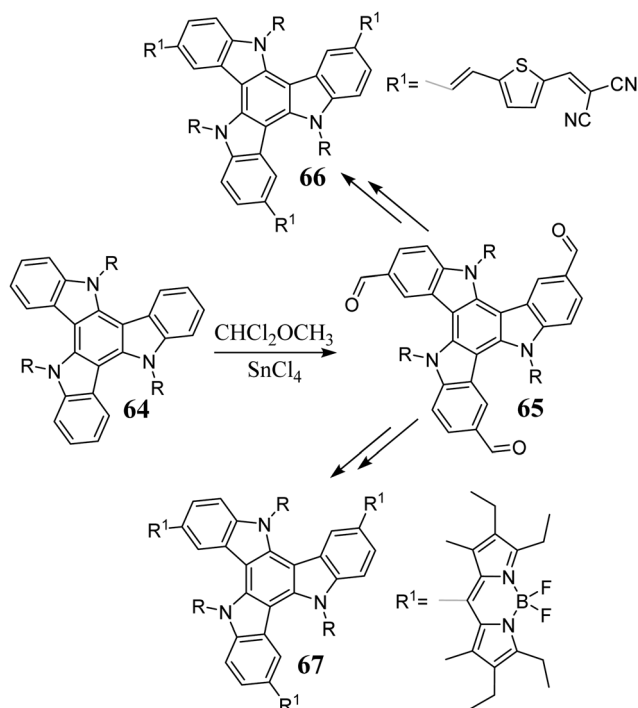


Suzuki coupling allows the synthesis of robust 5,10,15-triazatruxenes.²³³ However, desirable products for optoelectronics

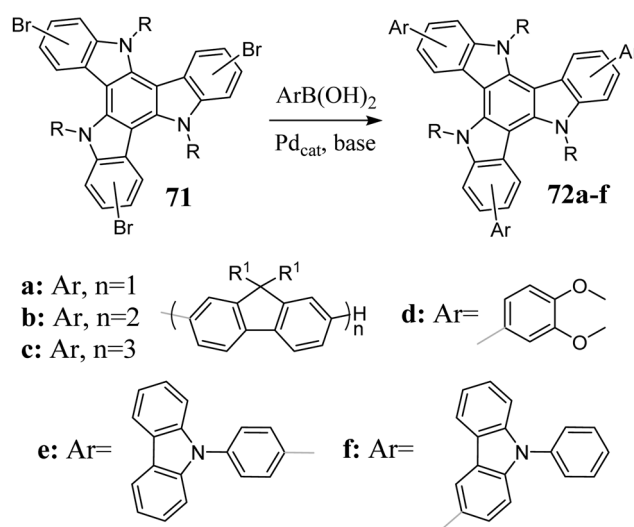


Scheme 23 Bromination of 5,10,15-triazatruxene

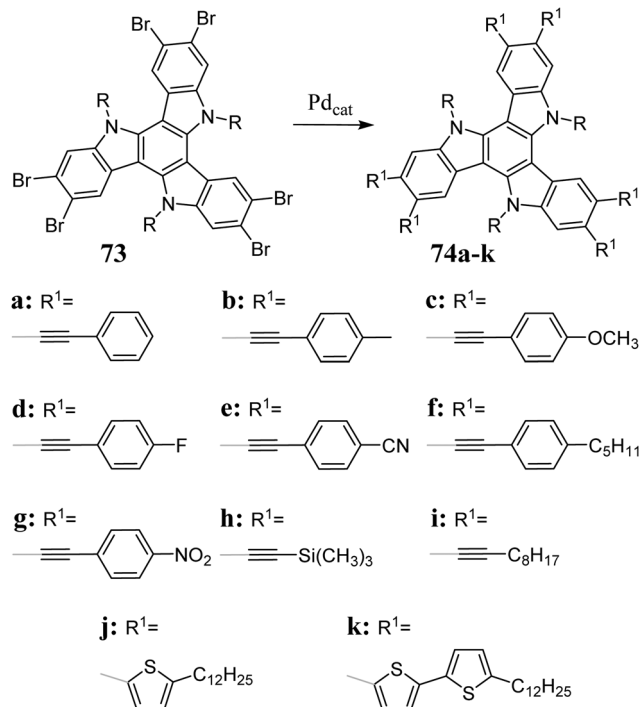
Various types of chlorides, bromides, and iodides can serve as alkylating reagents.²⁴⁰ Benzyl moieties are attractive for protecting nitrogen atoms because they can be easily removed afterward.²³⁴ It is also worth mentioning that the use of an



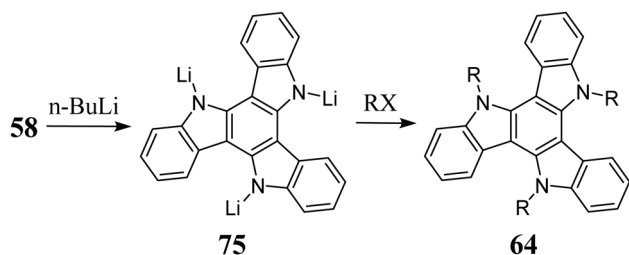
Scheme 22 Synthesis of 5,10,15-triazatruxene **65** and its reactivity.



Scheme 24 5,10,15-Triazatruxenes obtained by the Suzuki reaction.



Scheme 25 Sonogashira and Stille coupling products.



Scheme 26 N-Alkylation of 5,10,15-triazatruxene.

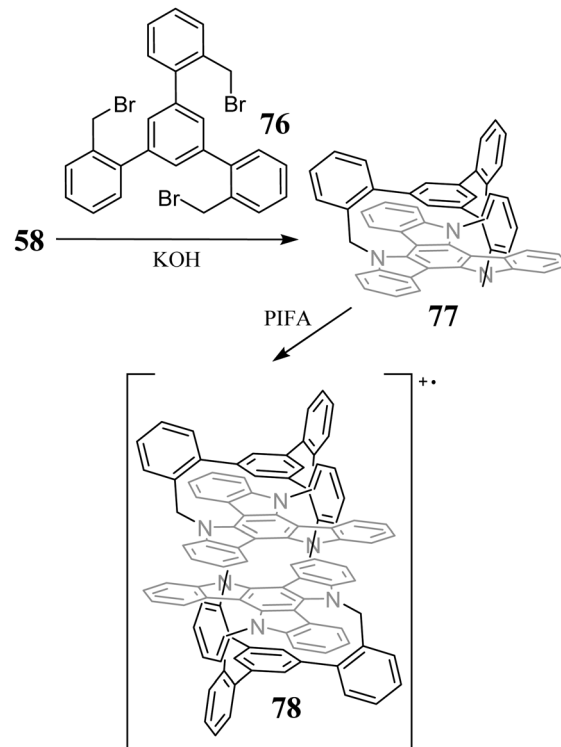
appropriate alkylating agent increases the product solubility in organic solvents and water.²⁴¹

An exciting example of **58** N-alkylation is its reaction with benzyl bromide **76**, resulting in cyclophane **77**. Further reaction with PIFA leads to the stable cation radical **78**, located at two π -stacking triazatruxene subunits (Scheme 27).

N-Alkylation was used to expand the aromatic system. Compounds **79**, due to Pd-catalyzed intramolecular arylation, are transformed into π -expanded systems **80** (Scheme 28).²²⁰ It is worth mentioning that **80** adsorbed on the surface of Pt at 750 °C undergoes cyclodehydrogenation, leading to the formation of the spherical molecule $C_{57}N_3$, triazafullerene.²⁴²

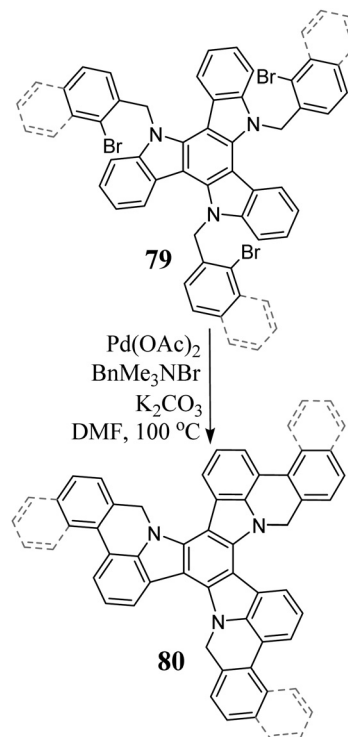
2.5. 5,10,15-Triazatruxene: physicochemical properties and applications

In 5,10,15-triazatruxene, the same as in truxene, the first two-electron transitions are symmetry forbidden, reflecting a low fluorescence quantum yield (QY = 3%).²⁴³ A reduction in its HOMO–LUMO gap causes a clear redshift in its absorption and



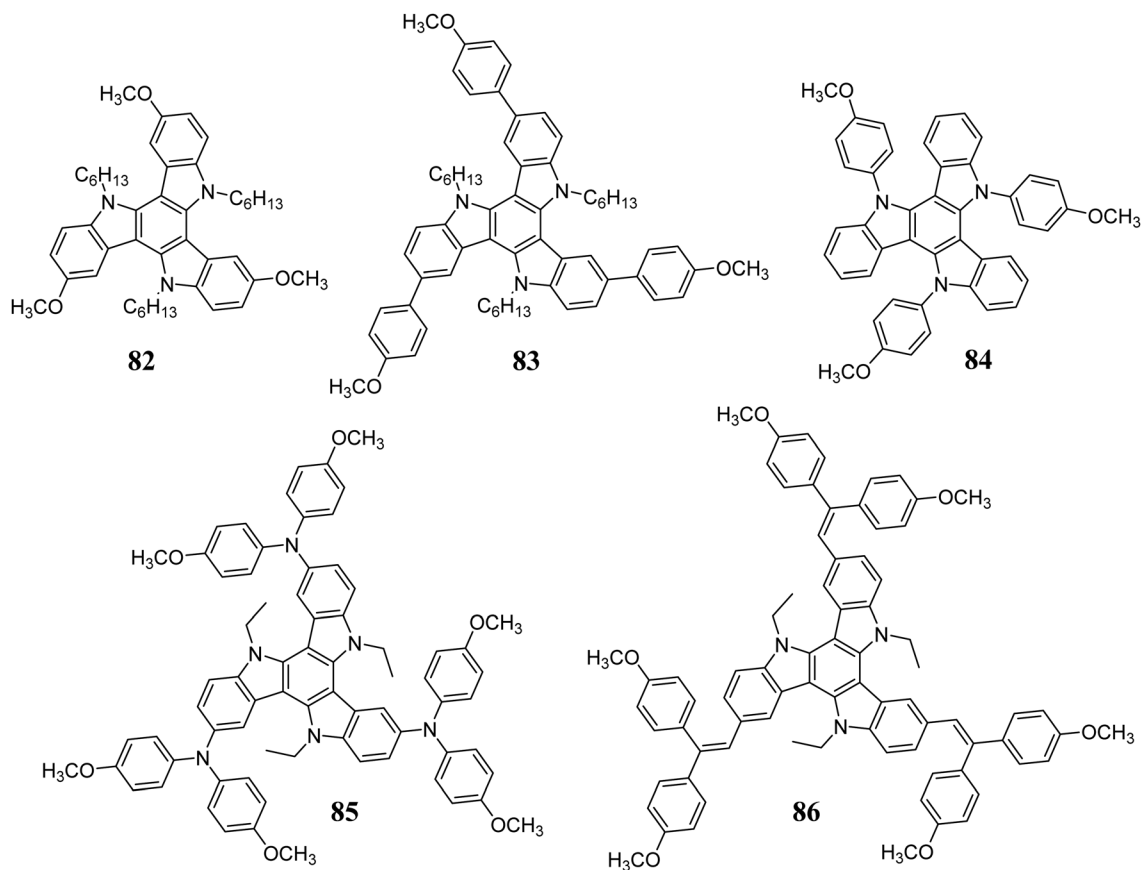
Scheme 27 Synthesis of cyclophane and stable cation radical.

emission spectra compared to truxene. The relatively high energy of its HOMO level is responsible its reversible oxidation, which combined a with its significant tendency to aggregate,

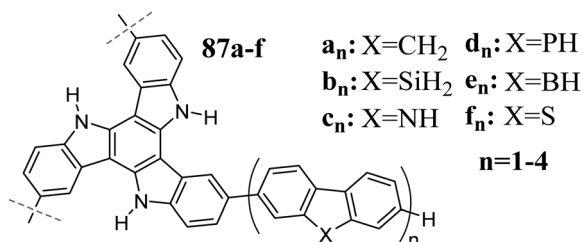
Scheme 28 Synthesis of π -expanded 5,10,15-triazatruxenes.



The dibenzo[*b,d*]thiophene sulfone-based D-A system **90** exhibits a very short delayed fluorescence decay time.⁸ This phenomenon is attributed to about 30 different couplings between the S₁ and T₁ states. Very fast T₁ → S₁ conversion results in slight energy loss, and thus the photoluminescence quantum efficiency is close to 100%. Penfold *et al.* explained the efficient TADF process in D-A systems based on **90**.²⁷⁷ D-A systems can be utilized as TADF emitters or NLO materials.^{18,19} These compounds show increased TPA, resulting from the spatial separation between the excited and ground states. It was found that the cross-section value for TPA also depends on the position of the acceptor subunit, where **92** exhibits stronger TPA than **93** (Scheme 32).²⁷⁸ It is worth mentioning that the NLO properties of 5,10,15-triazatruxenes have been successfully used in cell imaging.²⁷⁹



Scheme 29 5,10,15-Triazatruxene-based HTMs.



Scheme 30 Tri(oligoheterofluorene) 5,10,15-triazatruxenes.

5,10,15-Triazatruxene was also used to create macromolecular systems such as COFs,²⁸⁰ MOFs,^{281,282} and polymers.^{283,284} 5,10,15-Triazatruxene-based polymers can be used as capacitor materials.²⁸⁴ Some COFs²⁸⁰ and polymers²⁸⁵ exhibit higher selectivity towards CO₂ adsorption compared to N₂ or can store H₂ (Fig. 8).²³ The storage capacity of triazatruxene polymers increases in the presence of Li⁺ ions. Thus, it is crucial to redistribute Li⁺ ions homogeneously in the material, which is achieved by cation- π interactions.

Some of these substances exhibit fluorescence quenching upon interaction with electron-deficient systems.

This property inspired the synthesis of low molecular weight and polymeric materials for the detection of explosives.^{286–288} Adequate peripheral functionalization allowed the detection of

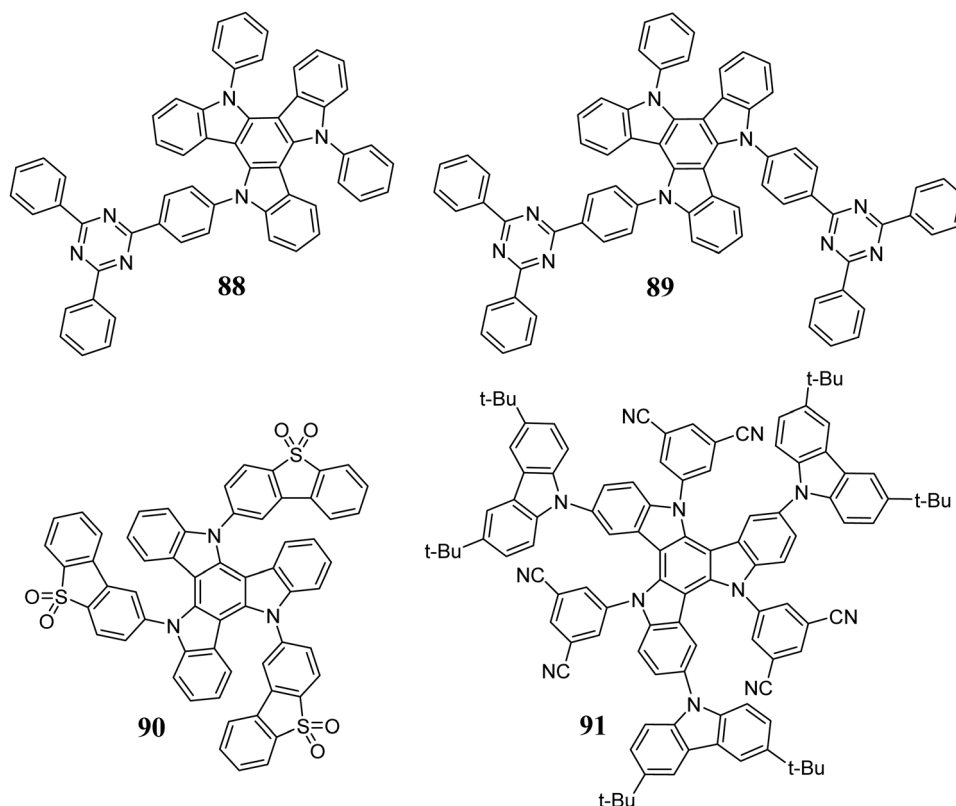
HCl²⁸⁹ and N₂H₄ vapors.²⁹⁰ 5,10,15-Triazatruxene, due to its semiconductive properties, has been used in the synthesis of new, thermally stable polymeric HTMs.^{291–293} A 5,10,15-triazatruxene-based microporous polymer could capture heavy metals.²⁹⁴ A similar interaction with Li⁺ cations resulted in a significant increase in the electrical capacity of the polymeric anode.²⁴ Interactions between 5,10,15-triazatruxene and Fe²⁺/²⁹⁵ or Zn²⁺/²⁹⁶ connected with iminopyridyl led to interesting chiral, tetrahedral molecules. Alternatively, the rational design of the molecule may lead to Al³⁺, Cr³⁺ and Fe³⁺ sensors.²⁹⁷ 5,10,15-Triazatruxene peripherally decorated with carboxyl groups can be used to modify the surface of TiO₂, enhancing the electron injection in DSSCs.²⁹⁸

5,10,15-Triazatruxene modified by mannose²¹ or lactose²⁹⁹ was used for sensing concanavalin A. 5,10,15-Triazatruxene, upon *N*-alkylation with alkylpiperidine chains, becomes soluble in water and shows amphiphilic properties.^{215,300} These substances also have enormous affinity and selectivity toward DNA G-quadruplexes, creating new anti-cancer drugs.^{301,302}

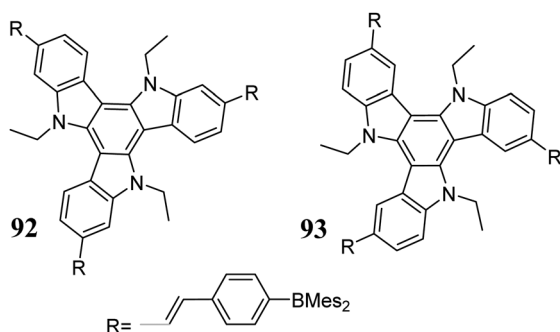
2.6. 5,10,15-Triphosphatruxene

5,10,15-Triphosphatruxene has been synthetically unavailable for years. One of the problems arising during the formation of the phosphole ring is that the known methods are not stereo-selective. The phosphorus atom in the phosphole ring, unlike the nitrogen atom in pyrrole, has sp³ hybridization, becoming a





Scheme 31 Triazatruxene-based D–A systems.



Scheme 32 Isomeric D–A systems exhibiting NLO properties.

potential stereogenic centre. The 5,10,15-triphosphatruxene molecule has three stereogenic phosphorus atoms, but due to its symmetry, the number of possible stereoisomers is reduced to four (Scheme 33).

In 2014, Kojima *et al.*³⁰³ synthesized this complex system *via* aromatic nucleophilic substitution to obtain a mixture of syn **94** and anti **95** isomers in a ratio of about 1 : 15. However, due to the similar properties of these diastereomers, their separation was complicated. Therefore, their crude mixture was subjected to subsequent oxidation. This procedure enabled crystallization of the pure oxidized isomer anti **98** (Scheme 34). Alternatively, upon sublimation of a crude mixture of **94** and **95**, it underwent complete conversion to the thermodynamically more stable isomer **94**. 5,10,15-

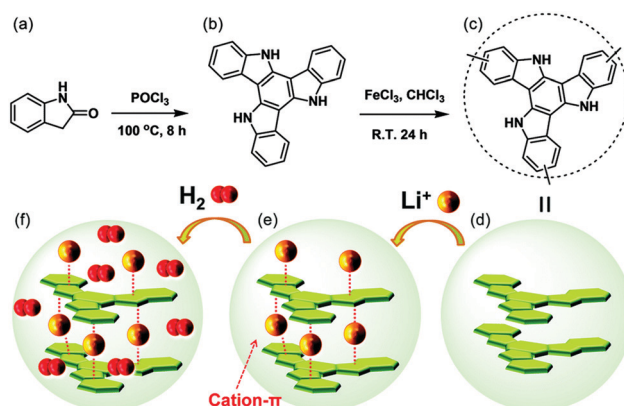


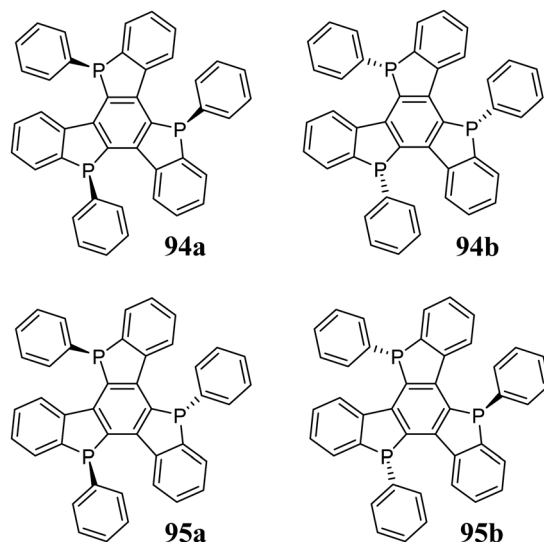
Fig. 8 (a)–(c) Synthetic route for the conjugated microporous polymer PTAT. Schematic representation showing the installation of lithium-doped CMP Li⁺–PTAT through point-to-face cation– π interactions (d and e) and its hydrogen adsorption (f). Reproduced from L. Yang, Y. Ma, Y. Xu, G. Chang, *Chem. Commun.*, 2019, **55**, 11227–11230 with permission from The Royal Society of Chemistry.

Triphosphatruxene isomers differ in the symmetry of their molecule. The syn-isomer has C_3 symmetry, whereas the anti-isomer C_1 . Therefore, different spectroscopic properties are expected.

2.7. 5,10,15-Trioxatruxene

The first report on the synthesis of 5,10,15-trioxatruxenes is dated 1984, when Destwde *et al.* obtained several derivatives of





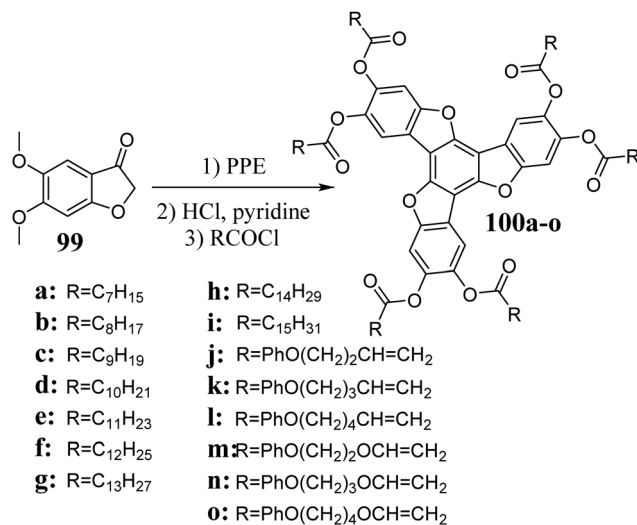
Scheme 33 5,10,15-Triphosphatruxene stereoisomers.

this heterocyclic system.³⁰⁴ Compounds **100** are products of benzo[*b*]furan-3-one **99** cyclotrimerization in the presence of PPE (Scheme 35). After deprotection of the hydroxyl groups, 5,10,15-trioxatruxene can be further modified to increase its solubility.

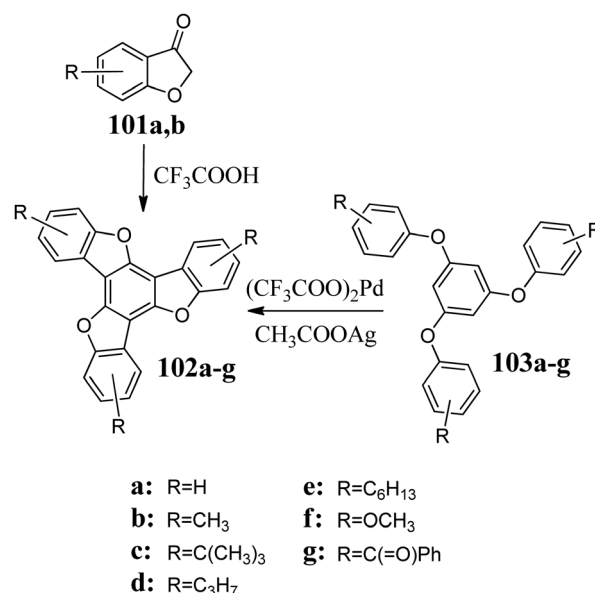
The 5,10,15-trioxatruxenes **102a** and **b** were synthesized using a similar procedure. To carry out the cyclotrimerization of **101a** and **b**, CF₃COOH³⁰⁵ was used instead of PPE, as mentioned above (Scheme 36).

The described methods were used to synthesize several 5,10,15-trioxatruxene monomers, **100j–o**, containing peripheral polymerizable moieties, making it possible to produce a cross-linked polymer (Scheme 35).³⁰⁶

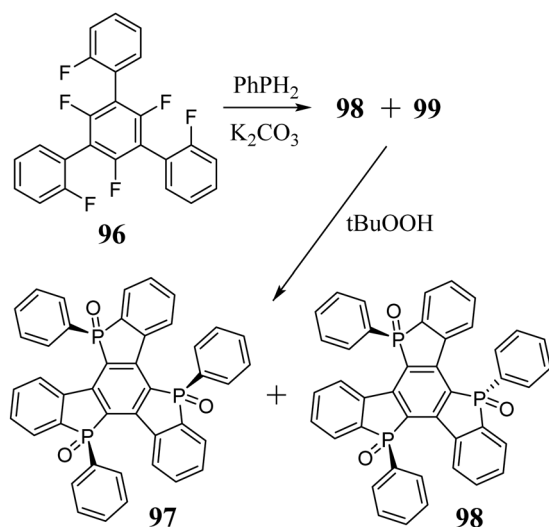
Sato *et al.* refined the synthesis of hexahydroxy-5,10,15-trioxatruxene, enabling the isolation of the condensation products and avoiding chromatography. Thus, this method is



Scheme 35 Synthesis of 5,10,15-trioxatruxene ester derivatives.



Scheme 36 Synthesis of 5,10,15-trioxatruxenes.



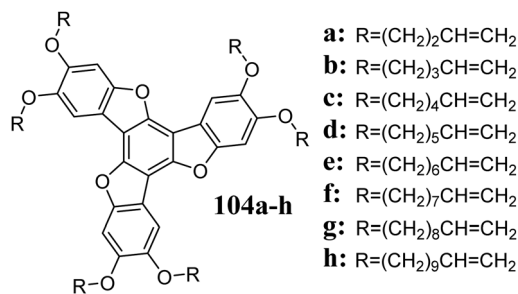
Scheme 34 Synthesis of 5,10,15-triphosphatruxenes.

attractive for industrial applications.³⁰⁷ Sato observed that during deprotection, partial oxidation to quinone occurs. 1,3,5-Triaryloxybenzenes **103a–g**, in the presence of (CF₃COO)₂Pd/CH₃COOAg, are converted to **102** in high yield (Scheme 36).³⁰⁸ However, for derivatives containing donor or acceptor groups, the efficiency of oxidative coupling is relatively low. An exciting example of the efficient synthesis of **102a** is the use of aromatic nucleophilic substitution.¹²⁴

Similar to the previously described systems, the first two electronic transitions in 5,10,15-trioxatruxene are symmetry forbidden.¹²⁴

5,10,15-Trioxatruxene is characterized by a relatively low fluorescence quantum yield of ~17%. Alternatively, the shape of its phosphorescence spectrum suggests the presence of





Scheme 37 5,10,15-Trioxatruxene-based photopolymerization monomers.

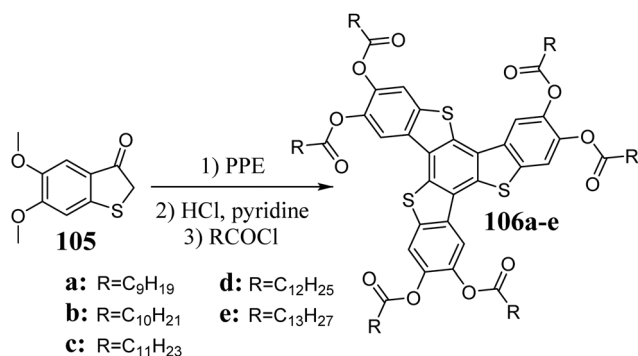
strong coupling between the T_1 and S_0 oscillation states. Moreover, the phosphorescence lifetime of about 5.4 s indicates the forbidden nature of the $T_1 \rightarrow S_0$ transition due to the spin multiplicity and the symmetry of the orbitals.¹²⁴ **102c** was examined towards electron mobility.³⁰⁹ The presence of *t*-butyl groups limits the intermolecular interactions, increasing the solubility of the system and making the oxidation process reversible.³⁰⁸

Alkylated 5,10,15-trioxatruxenes exhibit liquid-crystal properties and rich polymorphism.³⁰⁴ The self-assembly of **102a** and its liquid crystalline derivatives toward columnar ordering, combined with reversible oxidation, allows their use as p-type semiconductors. Substituted 5,10,15-trioxatruxenes, containing double bonds on their peripheral alkyl chains, have been used to synthesize cross-linked polymers,³⁰⁶ while **104** were used as photopolymerization agents, creating so-called “self-healing” LCDs (Scheme 37).³¹⁰

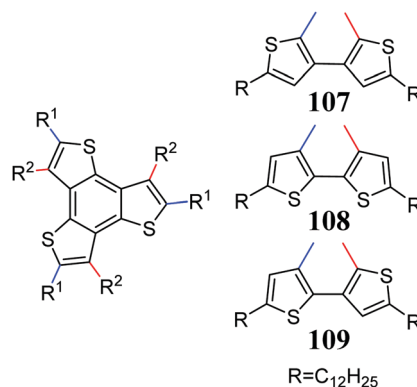
2.8. 5,10,15-Trithiatruxene

Peripherally substituted 5,10,15-trithiatruxenes **106** were described for the first time in 1985 by Nguyen *et al.*³¹¹ Their synthesis was based on the acidic condensation of substituted benzo[*b*]thiophen-3-one, **105**, in the presence of PPE (Scheme 38).

Egestad *et al.* showed that the treatment of benzo[*b*]thiophen-3-one with HCl in AcOH leads to a complex mixture of dimeric, trimeric, and tetrameric systems.³¹² The same research group performed the cyclotrimerization of 3-acetoxybenzo[*b*]thiophene in CF_3COOH . They obtained 5,10,15-trithiatruxene in a yield of 13%.³⁰⁵ 5,10,15-Trithiatruxenes are



Scheme 38 Synthesis of 5,10,15-trithiatruxene ester derivatives.



Scheme 39 5,10,15-Trithiatruxene-based ambipolar semiconductors.

cyclization products of appropriate thioethers in the presence of $(CF_3COO)_2Pd/CH_3COOAg$ at elevated temperatures. Thus, isomeric *t*-butyl peripherally functionalized 5,10,15-trithiatruxenes were obtained. Unfortunately, the reaction yields were much lower than that observed in the case of their oxygen counterparts.³⁰⁹

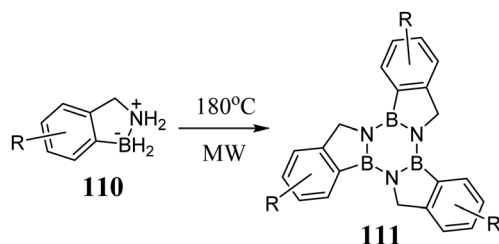
Similar to other symmetrical systems, 5,10,15-trithiatruxene reveals the first two electronic transitions forbidden by symmetry. The presence of sulfur atoms increases the intersystem crossing, resulting in phosphorescence in the solid state at room temperature.³⁰⁹ Miura *et al.* measured the phosphorescence spectra of two *tert*-butylated trithiatruxenes. The results are eye-catching. The phosphorescence of trithiatruxenes in PMMA films was only observed under vacuum. The authors measured the phosphorescence at low temperatures in methylcyclohexane. They observed almost pure phosphorescence (with minimal fluorescence signal) under these conditions for trithiatruxenes.

Hexa-substituted 5,10,15-trithiatruxenes, like their oxygen analogs, are characterized by rich polymorphism and columnar ordering in the solid state.^{311,313} This property has been exploited in epoxy resins, encapsulation, and electronic and structural laminates.³¹⁴ Currently, due to the dynamic development of optoelectronics, some 5,10,15-trithiatruxene derivatives can be used in OLEDs,^{315,316} OFET,³¹⁷ and OPVs.³¹⁸ The expanded, liquid-crystalline 5,10,15-trithiatruxene derivatives **107–109** developed by Aida *et al.* tend to self-assemble, increasing their semiconductive properties and leading to balanced, ambipolar conductivity of $0.18 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ (Scheme 39).³¹⁹

2.9. Borazatruxene

Truxenes in which the central benzene ring is replaced by borazine are structurally interesting systems, formally leading to hexaheterotruxenes. In 2019, Limberti *et al.* described a three-step synthesis in which **110** is converted to the corresponding borazatruxenes **111** in the presence of microwave radiation at 180°C (Scheme 40).³²⁰ The substrates enabling the synthesis of **110** are the appropriate nitriles or aldehydes. Their availability significantly increases the variety of borazatruxenes, allowing the synthesis of halogen derivatives and π -expanded systems. It should also be mentioned that the use





Scheme 40 Borazatruxene synthesis.

of appropriate aromatic ketones enables the synthesis of derivatives functionalized at the 5, 10, and 15 positions.

In the case of **111**, a clear blueshift can be observed in its absorption and emission spectra,³²⁰ while its fluorescence quantum yield does not exceed 10%. Borazatruxenes are air and moisture stable and exhibit better solubility, especially in moderately polar solvents, than the practically insoluble **4**. The flat structure of heterotruxene core and strong intramolecular interactions drive columnar ordering of the molecules in the solid state, creating potential organic semiconductors.

3. Non-symmetrical truxenes

The non-symmetrical truxenes described in this chapter are molecules of lower symmetry (without the triple-axis). Currently, the known non-symmetrical heterotruxenes possess at least one heteroatom. When working with polycyclic condensed aromatic systems, a remarkable problem is their poor solubility. The same issue also applies to truxenes. Truxene and 5,10,15-triazatruxene can be alkylated within 5-membered rings to improve their solubility. However, this procedure cannot be used for systems containing oxygen or sulfur atoms. This obstacle is partially solved by the peripheral functionalization of the π -electron system with alkyl substituents. Nevertheless, this procedure prevents their further modification. Another solution to overcome their low solubility, while maintaining their unique heteroatom properties, is the concept of 5-monohetero and 5,10-diheterotruxenes (Fig. 9).

An unquestionable advantage of non-symmetrical truxenes is their solubility, resulting from *C*-alkylation. It is also worth mentioning that symmetry breaking of a molecule affects its physicochemical properties. Thus, targeted functionalization for mono-substitution is an undisputed advantage in low-symmetrical cores. However, this procedure in the case of C_3 -symmetric truxenes is a challenging task.

3.1. 5-Oxatruxene

5-Oxatruxene was synthesized by Górski *et al.* in 2018 during the photocyclization of the corresponding triene **113** (Scheme 41).³²¹ 5-Oxatruxene, similar to other unsubstituted truxenes, is characterized by relatively low solubility. However, the alkylation of its cyclopentadienyl rings in a basic environment significantly increases its solubility. For example, the obtained tetraethyl derivative **114** exhibits good solubility in most organic solvents. 5-Oxatruxene undergoes electrophilic substitution and

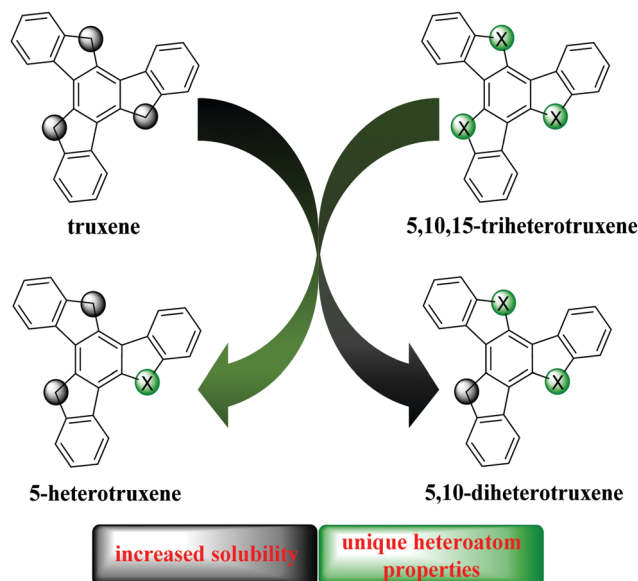
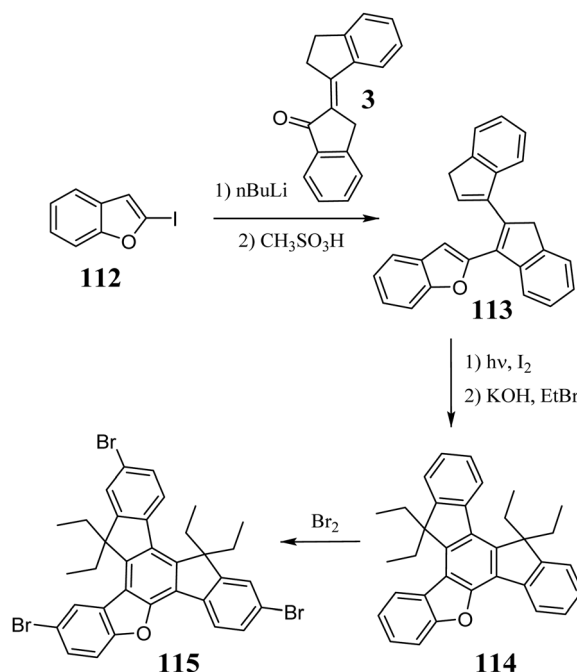


Fig. 9 Mono and diheterotruxenes as hybrids of symmetrical systems.

in excess Br_2 , can be converted to the corresponding tribromo derivative **115**.¹²⁵

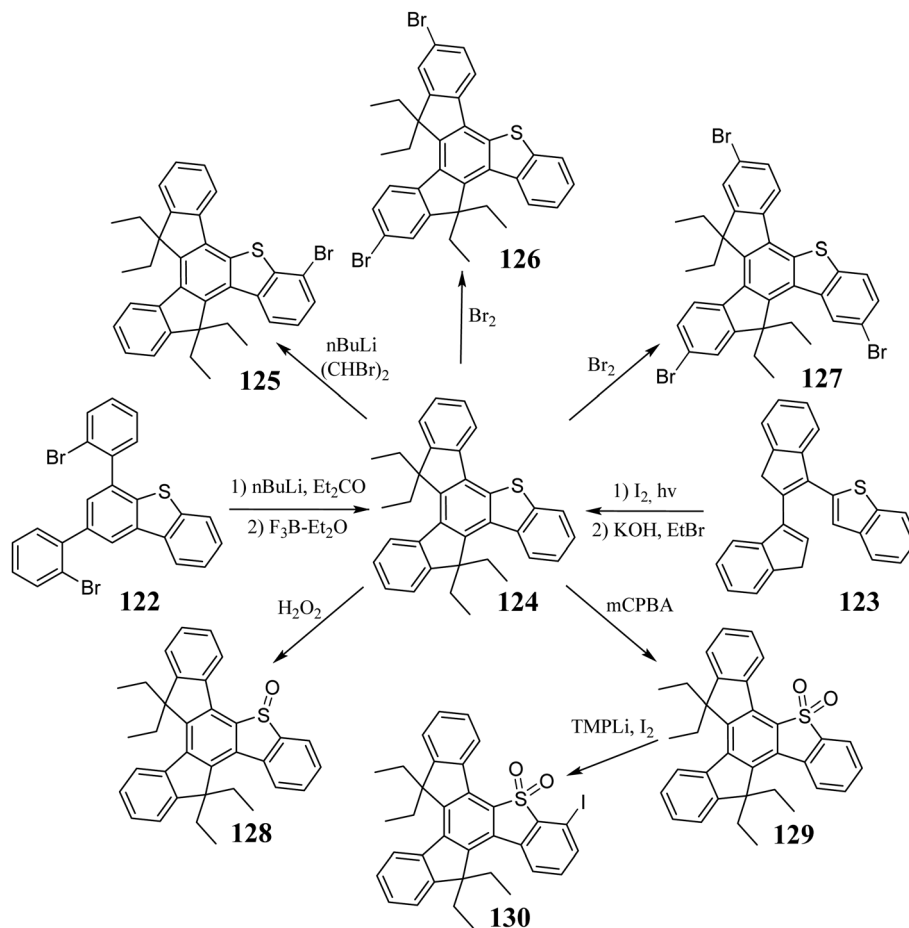
The presence of an oxygen atom in the molecule causes structural changes, which affect the electronic levels of 5-oxatruxene. Symmetry breaking lifts the HOMO and LUMO degeneration, which is visible in the redshift in its absorption and emission spectra. Moreover, a 4-fold increase in the fluorescence quantum yield is observed.³²¹ Unlike truxene, 5-oxatruxene exhibits better thermal properties, remarkable photostability, and a reversible oxidation process.¹²⁵ The presence of an oxygen



Scheme 41 Synthesis and reactivity of 5-oxatruxene.



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Scheme 43 Synthesis and reactivity of 5-thiatriuxene.

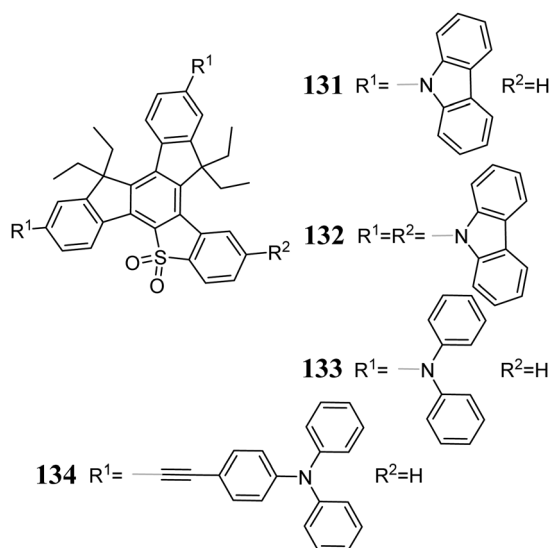
HOMO and stabilization of its LUMO compared to **4**. Thus, a redshift in its absorption and emission spectra is observed due to the decrease in HOMO–LUMO. The fluorescence quantum yield of **136** is 39.5%. The presence of oxygen atoms in the

5,10-dioxatriuxene core enables effective π -stacking, making 5,10-dioxatriuxenes promising materials for solution-processable organic semiconductors.

3.5. 5,10-Diazatriuxene

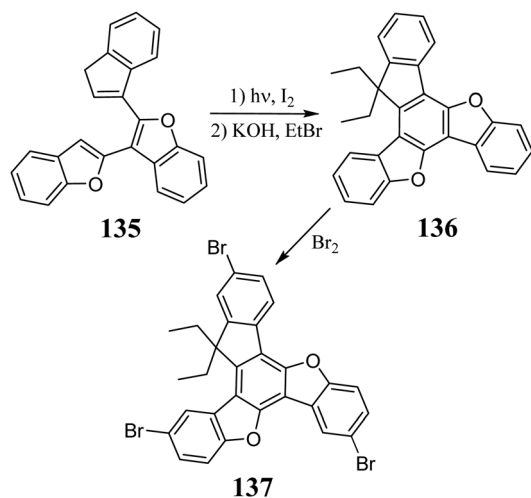
The first references to 5,10-diazatriuxene appeared in the work by El Sayed *et al.* in 2017.³²⁵ Consequently, the condensation of **56** was performed (Scheme 46). Subsequent studies also allowed condensation of 6-bromo³²⁶ and 5-nitroindoles,³²⁷ directly leading to doubly substituted 5,10-diazatriuxenes. Moreover, the N-substituted indoles also undergo this reaction, allowing the direct synthesis of N-protected 5,10-diazatriuxene-15-ones **140c**,³²⁴ and condensation using a stoichiometric amount of MeSO₃H acid in 1,2-dichloroethane.

In the presence of a base, **140a** and **b** undergo ionization, allowing reaction with various electrophiles, such as acyl chlorides,³²⁵ simple alkyl halides, and more complex derivatives containing a piperidine or oxirane ring.³²⁷ Alternatively, reducing the carbonyl group in **140c** by treatment with N₂H₄ in boiling ethylene glycol makes it possible to obtain the parent 5,10-diazatriuxene **142**,³²⁴ which after alkylation, is transformed to the highly soluble **143** (Scheme 47). Moreover, the presence of the carbonyl group enables the reaction between **140c** and organometallic substances, leading to the appropriate 3° alcohols



Scheme 44 5-Thiatriuxene-based D–A systems.

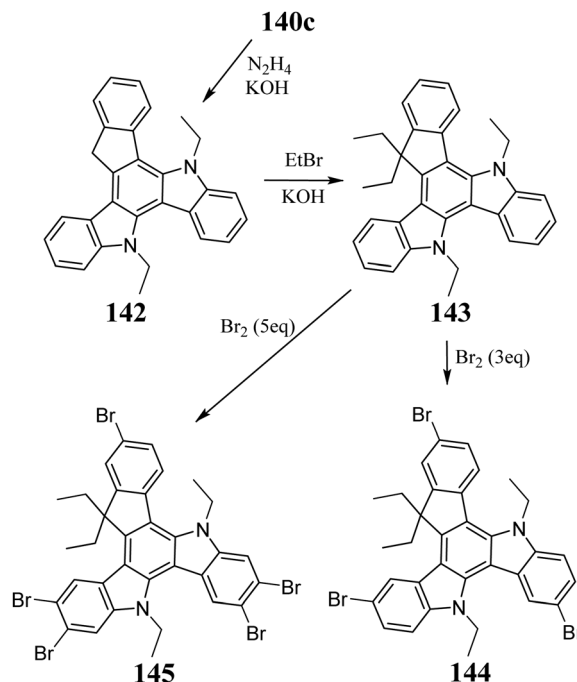




Scheme 45 Synthesis and reactivity of 5,10-dioxatruxene.

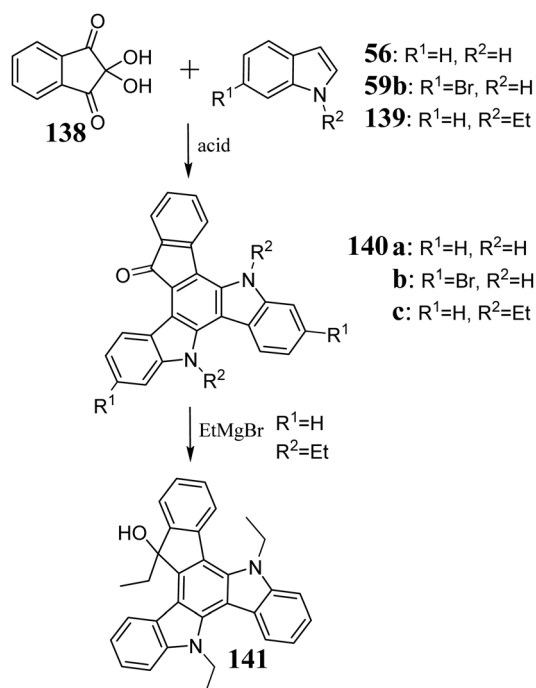
141 (Scheme 46), significantly extending the functionalization possibilities of the 5,10-diazatruxene core.³²⁴ Alternatively, **143** can be easily converted to the corresponding tri and pentabromo derivatives **144** and **145**, respectively, depending on the amount of Br_2 . **140b**, **144**, and **145** are crucial derivatives in further peripheral functionalization (Scheme 47).³²⁶

The two nitrogen atoms present in the aromatic system lift the electron level degeneration, destabilize the HOMO level significantly, and increase the fluorescence quantum yield (QY = 25.5%) compared to **4**.³²⁴ 5,10-Diazatruxene, like its dioxatruxene counterpart, tends to π -stacking interactions. Its high HOMO energy and aggregation in the solid state suggest the possibility of using this type of substance as an HTM.



Scheme 47 Synthesis and reactivity of 5,10-diazatruxene.

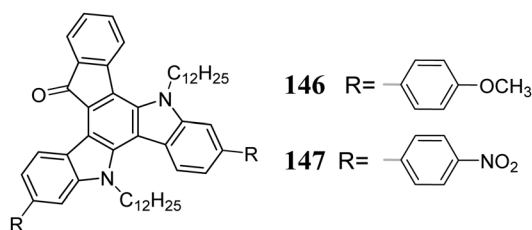
Peripherally functionalized 5,10-diazatruxene-15-ones with both donor **146** and acceptor **147** substituents were employed as promising semiconductors (Scheme 48). They exhibit reversible redox processes, which combined with their high π -stacking in the solid state, make them potential ambipolar semiconductors.³²⁶ Nevertheless, the interest in 5,10-diazatruxenes is not limited to optoelectronic applications. It has been found that the appropriate modification of the 5,10-diazatruxene core allows effective interaction with DNA, making it applicable as an anti-cancer agent.³²⁷



Scheme 46 Synthesis and reactivity of 5,10-diazatruxene-15-ones.

3.6. 5,10-Dithiatriuxene

In 2009, Suze *et al.* described the condensation between benzo[*b*]thiophene **148** and **138** in AcOH and the presence of H_2SO_4 , leading to 5,10-dithiatriuxene-15-one **149**, with a yield of 68% (Scheme 49).³²⁸ The same reaction can be performed using a stoichiometric amount of MeSO_3H in 1,2-dichloroethane, increasing the yield to 74%.³²⁴ However, due to the negligible solubility resulting from π -stacking, obtaining **149** and its further functionalization are significantly impeded. Nevertheless, the reaction

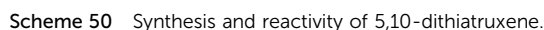


Scheme 48 5,10-Diazatruxene-15-one-based D-A ambipolar semiconductors.





The HOMO level of 5,10-dithiatruxene is destabilized compared to **4**.³²⁴ Due to the low symmetry of the molecule, the degeneration of its electron levels is lifted. However, despite the electron transitions being allowed, 5,10-dithiatruxene has a relatively low fluorescence quantum yield. The ability to exhibit strong π -stacking interactions in the solid state, characteristic for 5,10-diheterotruxenes, is also observed in the case of **149** and **152**. Therefore, 5,10-dithiatruxenes should be considered as potential organic semiconductors.



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