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Synthesis, processing and material properties of conjugated polymers

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Recent progress in the synthesis and characterization of well-defined conjugated polymers and oligomers is presented. The different synthetic schemes for most of the important conjugated polymers will be discussed using a subdivision into three main sections, covering hydrocarbon polymers, heterocyclic polymers and the oligomers, respectively. Although the materials properties will be referred to, the main emphasis of the review is directed to the different strategies that are now available to produce conjugated polymers and well-defined oligomers in a reproducible way and with full control over most of the important macromolecular properties. Copyright © 1996 Elsevier Science Ltd.

(Keywords: conjugated polymers; synthesis; processing; properties)

INTRODUCTION

The current interest in conjugated polymers is due to the substantial \( \pi \)-electron delocalization along their backbones which gives rise to interesting optical\(^1\) and nonlinear optical properties\(^2-4\) and allows them to become good electrical conductors typically when oxidized or reduced\(^5,6\). These properties may lead to a variety of practical applications such as information storage and optical signal processing, substitutes for batteries\(^1\) and materials for solar energy conversion\(^7\). Furthermore, electroluminescence from conjugated polymers is a rapidly expanding field of interest since the reports of poly(\( p \)-phenylene vinylene) based devices\(^9,10\). These possible applications are to some degree limited by the low stability towards atmospheric oxidation and the lack of processability often characterizing these materials. As a result, considerable research effort has been directed towards the preparation of well-defined conjugated polymers with improved processability and stability characteristics. The aim of this review is to discuss the recent progress in the synthesis and materials properties of polyconjugated polymers. The fundamental physics and applications of these materials in devices will be referred to, but for the detailed discussions of these matters the reader is referred to specialized reviews.

For the purpose of this review the field of conjugated polymer synthesis is divided into three main headings: namely, hydrocarbon conjugated polymers, conjugated polymers with heteroatoms in the main chain and well-defined oligomers. Each main class is subdivided into different subheadings, referring to the major classes of conjugated polymers. Within the subheadings recent progress for a particular polymer is discussed from a synthetic point of view.

HYDROCARBON POLYMERS

Polyacetylenes

Polyacetylene (Fig. 1) is the simplest polyconjugated organic polymer. The first reports on acetylene polymers date back to the last century. Cuprene, a highly crosslinked and extremely irregular product\(^11\) of acetylene polymerization in the presence of copper-containing catalysts, being a typical example.

Polycetylene was first prepared as a linear, high molecular weight, polyconjugated polymer of high crystallinity and regular structure in 1958 when Natta et al.\(^12\) polymerized acetylene in hexane using \( \text{Al}(\text{Et})_3/\text{Ti}(\text{OPr})_4 \) as the initiator system. However, for a long time polycetylene was considered of little interest because, irrespective of the method of preparation, it was obtained as an air sensitive, infusible and insoluble black powder.

A new phase in the study of polycetylene began in the 1970s when Shirakawa and co-workers\(^13,14\) developed a simple method for preparing film samples which on
Properties of conjugated polymers: W. J. Feast et al.

Figure 1 The structure of cis-polyacetylene

*Scheme 1* Shirakawa's route to polyacetylene

The synthetic procedure described above yields polyacetylene which is mainly cis. Baker et al. prepared almost pure trans polymer by reducing the Al/Ti ratio to unity and omitting the catalyst aging step. In another modification of the Shirakawa method, Naarmann and co-workers used silicone oil instead of toluene and prepared a polyacetylene sample which contained an approximately equal proportion of cis and trans vinylenes and a much lower concentration of sp²-hybridized carbon atoms than is normally present in these films. A year earlier, the same group developed the, so called, adding reducing agent method. This method involves the introduction of a reducing agent (such as n-butyl lithium) to the catalyst solution prior to the polymerization reaction. The polyacetylene samples thus produced are stretchable and on iodine doping conductivities of the order of 10⁵ S cm⁻¹ have been claimed. A further improvement in the observed conductivity was achieved by Tsukamoto. In this procedure, a solvent with high boiling point, such as decalin, is used and the catalyst is heat treated at around 200°C. The polyacetylene obtained by this procedure can be stretched by a factor of eight and yields a reported conductivity value of more than 10⁵ S cm⁻¹ after iodine doping.

Luttinger route. In 1980 Enkelmann et al. noted that powdered polyacetylene specimens prepared by traditional methods possessed properties which were essentially the same as those of the material produced by Shirakawa's method. Such powders could be suspended in an inert solvent and application of the suspension to a suitable substrate followed by evaporation of the solvent gave thin films of polyacetylene. They also showed that solid specimens of almost any desired size could be prepared by moulding polyacetylene powders. Later, Enkelmann et al. discovered that the best results are obtained using Co(NO₃)₂/NaBH₄ as the initiator system, systems of this type containing salts and complexes of group VIII metals together with a hydride reductant were first used by Luttinger twenty years earlier. In a typical experimental procedure ethanol is saturated with gaseous acetylene, and sodium borohydride followed by cobalt nitrate are added. After several hours at room temperature black flakes of polyacetylene precipitate. By lowering the reaction temperature to −30°C the formation of a predominantly cis structure is favoured (Scheme 2).

The main advantage of Luttinger's catalyst over that developed by Shirakawa is that it allows the polymerization reaction to be carried out even in the presence of water and oxygen. Frohner and Wucke studied the kinetics of the reaction and found that the initial polymerization rate is high and that the polyacetylene yield is proportional to the concentration of the cobalt species. Furthermore, the same workers observed that...
decreasing the reaction temperature results in an increase in polymer yield and crystallinity.

Precursor-polymer route. The inherent insolubility and infusibility of polyacetylene coupled with its sensitivity to air imposes a barrier to the processability of the polymer. As a result considerable research efforts have been directed towards obtaining polyacetylene from workable, easily processable polymers by means of polymer analogous transformations. In particular, the synthesis of polyacetylene from the dehydrohalogenation of poly(vinyl chloride)\(^{36}\) has attracted a lot of attention (presence of solvent) the morphology of the final product involving an exothermic symmetry allowed elimination of structural defects and cross-links. Nevertheless, materials produced in this way may be of practical use for EMR screening purposes\(^{35}\).

Another approach involves the use of prepolymers which can be thermally converted to polyacetylene. This approach was first described by Edwards and Feast\(^{19}\) (Scheme 3) and has been further refined by Feast and co-workers\(^{36-40}\).

The initial monomer, 7,8-bis(trifluoromethyl)tricyclo[4.2.2.0\(_2\)5]deca-3,7,9-triene is readily prepared by the thermal cycloaddition reaction between hexafluorobut-2-yne and cyclooctatetraene. The polymerization reaction is carried out in the presence of ring opening metathesis polymerization initiators, such as WCl\(_6\)/SnMe\(_4\), which act only on the strained four-membered ring to form a high molecular weight precursor polymer which is soluble in common organic solvents and can be purified, characterized and processed by conventional methods. On heating, the prepolymer can be easily and conveniently converted to polyacetylene with the evolution of 1,2-bis(trifluoromethyl)benzene. The conversion of the precursor polymer to polyacetylene is a complex process involving an exothermic symmetry allowed elimination with the formation of a new cis double bond and hexafluoro orthoxyylene, the cis double bond isomerizes to trans and the xylene migrates through the film and evaporates giving a large loss in mass and contraction in volume. By controlling the conversion reaction protocol (temperature, time, pressure, mechanical stress and presence of solvent) the morphology of the final product can be regulated, expanded amorphous foams through to fully dense oriented films are accessible.

The prepolymer shown in Scheme 3 decomposes spontaneously to the polyacetylene if stored at room temperature (half life: ca. 20 h). In order to produce a precursor polymer which was stable at room temperature a large number of prepolymers were examined but the majority of these possessed either insufficient or excessive thermal stability\(^{37}\). The synthesis of a precursor polymer with good room temperature stability\(^{40}\) was achieved by the isomerization of the original monomer, 7,8-bis(trifluoromethyl)tricyclo[4.2.2.0\(_2\)5]deca-3,7,9-triene, to 7,8-bis(trifluoromethyl)-pentacyclo[6.2.0.0\(_3\)4.0\(_7\)6.0\(_5\)2]deca-9-ene by ultraviolet irradiation in dilute pentene solution. The polymerization of this quadracycloane analogue (Scheme 4) gives a room temperature stable precursor polymer; however, the exothermicity of the conversion reaction presents a hazard and the polyacetylene obtained contains residual trifluoromethyl groups.

Schrock and co-workers\(^{41}\) have developed a series of well defined ROMP initiators of the type M(CH-t-Bu)\(_2\)(NAr)(O-t-Bu)\(_2\) in which M = W or Mo and Ar = 2,6-C\(_6\)H\(_5\)-i-Pr\(_2\) which allows better control over this polymerization reaction. Using these systems polyacetylene prepolymers of controlled molecular weight and polydispersities close to unity can be prepared. Furthermore, the use of these systems allows the controlled introduction of end-groups to the polymer chain\(^{42}\).

The main difference between the polyacetylene products obtained by the Shirakawa and precursor polymer routes lies in their morphology and relative order. Generally Shirakawa’s polymer is obtained in a fibrillar morphology with low bulk density and variable crystallinity whereas the precursor polymer route gives access to materials of more controlled morphology. Thus, continuous solid films with densities in the range 1.05–1.1 g cm\(^{-3}\) may be routinely obtained, which can be essentially amorphous, semicrystalline or highly crystalline and may be oriented by stretching during or prior to conversion.

More recently Swager et al.\(^{43}\) demonstrated an alternative precursor route to polyacetylene in which the production of volatile by-products is avoided. The method involves the ring opening polymerization of benzvalene and the subsequent catalytic isomerisation to polyacetylene of low crystallinity (Scheme 5).

Soluble polyacetylenes. Irrespective of the method adopted for its synthesis polyacetylene itself is an insolu-
Properties of conjugated polymers: W. J. Feast et al.

**Scheme 4** Durham precursor route using a quadricyclane analogue as monomer

**Scheme 5** Swager's polyacetylene route starting from benzvalene

ble, infusible and generally intractable material. In order to obtain soluble polyacetylene analogues monosubstituted acetylenes\(^{44-46}\) have been polymerized to yield polymers with side groups attached to every second carbon atom. In most cases these materials are soluble, but the optical absorption maximum of the resulting polymer is higher in energy than that observed for polyacetylene indicating a lower effective conjugation length. The electrical conductivity of all substituted polyacetylene analogues reported to date is significantly lower than that of polyacetylene itself\(^{47}\). The reduction in conjugation length is attributed to steric repulsions between adjacent side groups which causes the polymer chains to twist. This loss in planarity is readily understood since the rotation barrier about the single bonds in polyacetylene has been calculated\(^{48}\) to be about 6 kcal mol\(^{-1}\). Recently, Grubbs and co-workers\(^{49}\) demonstrated that the ring opening metathesis polymerization of monosubstituted cyclooctatetraene derivatives leads to partially substituted polyacetylenes that are both soluble and highly conjugated (Scheme 6).

**Polydiacetylenes**

Polydiacetylenes are synthesized by the topochemical polymerization of diacetylenes\(^{50-73}\). Most work has focused on the polymerization of crystalline diacetylenes, while some investigations report polymerization in the liquid-crystalline state and as Langmuir–Blodgett multilayers. Before the different processes are discussed, a short review of monomer syntheses is given.

**Synthesis of diacetylenes.** Symmetrically disubstituted diacetylenes are usually obtained via Glaser\(^{74}\) coupling of terminal acetylenes. The reaction is carried out in the presence of a cuprous salt with air as oxidant. An alternative synthesis, the reaction of the terminal acetylene with cupric acetate in pyridine, was proposed in 1956 by Eglington and Galbraith\(^{75}\).

In 1957 Cadiot and Chodkiewicz reported\(^{76}\) coupling of a haloacetylene, usually obtained by reaction of the terminal acetylene with sodium hypobromite although alternative methods are also available\(^{80-82}\), with a terminal acetylene in the presence of cuprous salts (Scheme 7). This method is the most satisfactory procedure for the synthesis of unsymmetrically disubstituted diacetylenes.

Alternative approaches for the synthesis of diacetylenes include the iodine and cupric salt oxidation of Grignard acetylides\(^{77}\), the condensation of acetylenic Grignard reagents with 1-bromoacetylenes\(^{78}\) and the permanganate oxidation of sodium acetylides\(^{79}\). However, these methods are of limited use.

**Solid state polymerization.** The solid state polymerization of diacetylenes (thermally or by irradiation) can yield nearly perfect polymeric single crystals of large

**Scheme 6** Grubbs' soluble polyacetylene from monosubstituted cyclooctatetraene

**Scheme 7** Route to unsymmetrically disubstituted diacetylenes

\[
R^1\text{C}≡\text{C}≡\text{H} + \text{X}\text{C}≡\text{C}≡\text{R}^2 \xrightarrow{\text{CuCl \text{hv or } \Delta}} \ R^1\text{C}≡\text{C}≡\text{C}≡\text{C}≡\text{R}^2
\]
Properties of conjugated polymers: W. J. Feast et al.

The photoreactivity of diacetylenes was first noted as early as 1882 but it was not until 1969 that the current interest in the preparation and study of the macroscopic properties of single crystal diacetylene polymers was initiated by Wegner. The topochemical polymerization of an array of diacetylene monomers proceeds by 1,4-addition reactions of the rigid diacetylene units to yield a conjugated extended polymer chain (Scheme 8).

The diacetylene polymers are believed to exist in two mesomorphic forms (Scheme 8). The packing of the monomer units in the crystal is characterized by the separation (d) of the monomers in the direction of the array and the angle (γ) between the axis of the array and the diacetylenic rod. The perpendicular distance (s) between the diacetylene units seems to be critical for solid state polymerization. Correlation of the lattice packing of the diacetylene monomers with solid state reactivity have indicated that the reacting C1 and C4 carbon atoms must be closer than 4 Å for polymerization to occur. Bloor has related the effects of variation in d and γ to the reactivity of diacetylenes towards polymerization and found that, apart from the 4 Å criterion, the monomers should be parallel to one another, or, if non-parallel, the monomer molecules must be related by a glide plane or screw axis. Nevertheless, an example of a reactive non-parallel diacetylene monomer has been reported. The criteria for solid state reactivity are more easily satisfied by monomers with large end groups than by those with small end groups. The role of the end groups seems to be critical since the interactions of the end groups determine the monomer crystal lattice structure. The flexibility of the side groups attached to the diacetylene unit must also be taken into consideration when designing a reactive diacetylene molecule. There are examples known where a compact molecular structure is obtained in the crystal with the side groups impeding the reaction. Nevertheless considerable improvements in predicting expected crystal structures of diacetylenes from knowledge of molecular structures are required before the problem of obtaining a single crystal polymer from a specifically designed monomer molecule can be solved.

The nature of the propagating chain end has been the subject of much discussion but it is now generally accepted that the photoinitiation reaction step involves the production of diradicals from the interaction of two diacetylene molecules. The most widely used technique in the study of polymerization kinetics is differential scanning calorimetry. However, although a vast volume of literature has been published concerning 2,4-hexadiyne-1,6-diol, bis(p-toluene sulfonate), little work on other diacetylenes appears to have been done.

Diacetylene 2 may be polymerized thermally in the solid state. The time required for polymerization varies from 2 h at a temperature just below the melting point to two months at ambient temperature. The polymerization process is characterized by the occurrence of an induction period, during which time the polymerization rate increases by a factor of up to 200. Induction times during solid state reactions have been explained in terms of the incubation period necessary for the nucleation of a polymer crystal phase. This explanation, however, does not apply in the case of 2 where the polymer forms a solid-solution with the monomer. For polymerization of 2 the only structural change observed after polymerization is a 10% contraction of the crystal along the chain axis. The lattice strain generated by the mismatch of the polymer and monomer lattices is believed to be responsible for the autocatalytic effects observed during polymerization. On the basis of this model, Baughman developed a model which is consistent with the observed phenomena. According to this work, formation of the polymer chains is initiated during the induction period but the chain length is determined by the crystal strain field. Chain lengths of 10–20 units are estimated to be formed during the induction period. At about 10% conversion, irrespective
of the polymerization temperature, crystal strain is relieved as conversion to polymer takes place resulting in a strain free polymer crystal. Polymerization during the induction period and during the fast polymerization regime have been found to have the same activation energy. As a result of these observations, it was concluded that the rate determining step for this polymerization was chain initiation.

The determination of molecular weight distributions in polydiacetylenes by conventional methods has been difficult due to the tendency of polydiacetylenes to form aggregates or microgels in solution. However, methods making use of chain length dependent solid-state properties have proved useful. Wenz and Wegner have reported the molecular weight distributions of some γ-ray polymerized bis-toluenesulfonates. Chain lengths of about 1500 units were reported corresponding to weight average molecular weights of about 800,000 g mol⁻¹. Measurements were made using light scattering and confirmed by gel permeation chromatography. The variation in the molecular weight distribution with increasing degree of polymerization has also been studied. In general, at low polymer concentrations, short chains of approximately 60 repeat units are present. At higher conversion rates, the g.p.c. maximum rapidly shifts to higher values. A characteristic feature is that only the high molecular weight (1500 units) polymer is formed above about 20% conversion.

**Polymerization in Langmuir–Blodgett film structures.** The transfer of amphiphilic monolayers from the air–water interface to solid supports by the Langmuir–Blodgett film technique can yield ultra-thin, homogenous and well-ordered mono- and multi-layer films. The polymerization of amphiphilic diacetylene molecules organized by the Langmuir–Blodgett film technique can lead to polymer structures of controlled thickness and orientation. Langmuir–Blodgett films in which pentacosa-10,12-diynoic acid and henicosa-2,4-diynylamine were combined to form alternate-layer structures have been polymerized via irradiation with ultraviolet light and shown to possess pyroelectric properties.

**Polymerization in the liquid-crystalline state.** Meso-genic diacetylene monomers which form thermotropic liquid crystal phases can undergo polymerization on thermal annealing. In appropriately designed molecules the polymerization proceeds in a controlled manner to yield polydiacetylenes which also exhibit liquid crystalline behaviour.

**Polyphenylenes**

Four synthetic routes for the preparation of polyphenylenes have been employed fairly generally, namely: oxidative coupling, organometallic coupling, dehydrogenation of polycyclohexynes and cycloaddition reactions. These, together with the other methods available for the synthesis of polyphenylenes, are discussed in the sections below.

**Oxidative coupling.** The most commonly employed method for the preparation of polyphenylenes involves the oxidative coupling of substituted and unsubstituted benzenes via treatment with a Lewis acid catalyst/oxidant system. In 1963 Kovacic and Kyriakis synthesized poly(p-phenylene) by stirring benzene, anhydrous AlCl₃ and anhydrous CuCl₂ for 2 h at temperatures between 25°C and 35°C. The reaction is believed to involve an initial one-electron oxidation of benzene to its radical cation, followed by reaction of the radical cation with several benzene molecules to give an oligomeric radical cation. A second one-electron oxidation, followed by loss of two protons aromatizes the terminal rings, and oxidative rearomatization of the dihydro structures by CuCl₂ yields the polymer.

A number of other catalyst/oxidant systems have been employed for the conversion of benzenes to polyphenylenes. These include Cu²⁺ and Ru³⁺ ion-exchanged montmorillonite clays and AsF₅/AsF₃. The structural features and physical properties of the polymer prepared via oxidative polymerization depend to a large extent on the nature of the reagents. Thus, it has been claimed that polyphenylenes with ortho linkages can be obtained from the oxidative polymerization of monosubstituted benzenes (e.g. toluene, chlorobenzene) under conditions analogous to those employed for the preparation of poly(p-phenylene)⁶⁰. Polyphenylenes linked at the meta position are in general synthesized by treatment of m-terphenyl or mixtures of the m-terphenyl and biphenyl with anhydrous AlCl₃/CuCl₂ at elevated temperatures (85°C and 180°C). Polymers substituted in the para position are in general more difficult to process than the ortho and meta substituted analogues. Oxidations in liquid SO₂ at -75°C or concentrated...
sulfuric acid in the presence of AlCl₃ are claimed to result in the efficient formation of homogenous poly-(p-phenylene) which is almost completely amorphous. Highly crystalline films of this polymer can be deposited from benzene in concentrated sulfuric acid emulsion.

An alternative means for the synthesis of polyphenylenes involves the anodic electrochemical oxidation of benzene or biphenyl in liquid sulfur dioxide on an appropriate electrode. The reaction yields passivating films if quaternary ammonium perchlorate is used as the electrolyte or conductive dendritic deposits if quaternary ammonium tetrafluoroborate is employed. In both cases the polymer formed is reported to be linked at the para position but contains variable amounts of oxygen as phenolic groups. The electrochemical polymerization of benzene in nitromethane with aluminum chloride and water or an amine as additive results in the deposition of black polyphenylene on a platinum electrode. Freestanding polyphenylene films containing ortho, para and meta links are obtained if the electrochemical oxidation of benzene is carried out in a two-phase HF/benzene system. The anodic oxidation of benzene in nitrobenzene solution with CuCl₂ and LiAsF₆ results in the formation of flexible, electrically conducting films of poly(p-phenylene) as does the anodic oxidation of benzene in the presence of BF₃ OEt₂. The yields of the polymers prepared by this method are restricted by the area of the anode irrespective of whether the reaction is carried out in nitrobenzene or in the bulk.

**Organometallic coupling.** Ullmann and Wurtz-Fittig type reactions have been utilized for the synthesis of polyphenylene oligomers but these methods cannot be applied to the synthesis of polymers since yields are low. The preparation of polyphenylenes via the coupling of Grignard reagents has proved more suitable. Thus the coupling of the mono-Grignard reagent of dihalobenzene in the presence of organometallic or organic promoters has been used for the synthesis of a range of para and meta substituted polyphenylenes. However, the combination of lithiated anthracene and anthraquinone species is reported to be the most efficient method for the synthesis of polyanthrylenes.

**Cycloaddition reactions.** 1,4-Cycloadditions of bis-cyclopentadienones with bisacetylenes can yield high molecular weight polymers which contain approximately equal proportions of meta and para substituted phenylene units. Low molecular weight polyphenylenes containing ethynyl branches have been prepared by polycyclotrimerization of diacetylene in the presence of Al(i-Bu)₃ and TiC₁₄ whereas polyphenylene copolymers with vinyl and alkyl branches can be obtained by the reaction of diacetylene with an appropriately substituted acetylene. The thermal eliminative ring closure of 1,4-diphenyl-3-(N,N-dimethylamino)-hex-5-en-1-yne has been shown to give terphenyl and by an analogous method the poly(N,N-dimethylaminohex-5-en-1-yne) derivative shown in Scheme 12 affords phenyl substituted poly(p-phenylene)s on pyrolysis.

**Other synthetic routes.** Poly(p-phenylene) contaminated with small amounts of bridging diazo groups has been prepared by heating the bisdiazonium salt of...
Properties of conjugated polymers: W. J. Feast et al.

Scheme 12 Poly(p-phenylene) by thermal ring closure of a poly(N,N-dimethylamino-hex-5-en-1-yne) derivative

\[
\text{\begin{align*}
\text{\(N(CH_3)_2\)} & \quad \text{C}_6\text{H}_5 \\
\text{\(-\)} & \quad \text{\(-\)} \\
\text{\(\text{C}_6\text{H}_5\)} & \quad \text{H}_5\text{C}_6 \\
\end{align*}}
\]

Scheme 13 Poly(p-phenylene) via coupling of a bisdiazonium salt

\[
\text{n N}_2^+ \quad \text{\(\text{C}_6\text{H}_5\)} \\
\text{CuCl}_2/\text{NH}_3 \quad \text{n} \\
\text{\(\text{C}_6\text{H}_5\)} \\
\]

p-diaminobenzene in ammoniacal Cu\(^{2+}\) or Fe\(^{2+}\). In this case the polymerization mechanism is thought to involve homolytic cleavage of the carbon to diazonium group bonds followed by C–C coupling (Scheme 13).

The electrochemical reduction of 1,4-dibromobenzene or 4,4′-dibromobiphenyl in THF/HMPA on a mercury pool electrode with Ni(0) complex catalysts and lithium perchlorate as the electrolyte gives good yields of insoluble poly(p-phenylene) as a pale yellow powder. By analogy, using an acetonitrile solution of Ni(PPPh\(_3\))\(_2\), p-dibromobenzene has been reduced on a platinum or glassy carbon electrode with a tetrabutylammonium salt as electrolyte. The resulting electroactive polymer coating is of the poly(p-phenylene) type but it is contaminated with one nickel atom for every six or seven polymer repeat units. Homogeneous undoped films of poly(p-phenylene) can be deposited onto a solid cathode by electroreduction of 4,4′-dibromophenyl activated by a Ni(0) complex of 1,2-bis(diphenylphosphino-ethane) in an equimolar proportion. Electrodeposited thin layers of this polymer can be either oxidized or reduced in acetonitrile solution containing lithium salt as the supporting electrolyte. This method has been extended to the polycondensation of 2,7-dibromo-9,10-dihydrophenanthrene with isolated zero-valent nickel complexes and electrochemically generated zero-valent nickel complexes to afford poly(9,10-dihydrophenanthrene-2,7-diyl) which essentially possesses the same conjugation system as poly(p-phenylene).

Another route to substituted poly(p-phenylene) derivatives makes use of the Bergman cyclization of enediyynes (Scheme 14). Thermal treatment (T = 50–160°C) of the enediyne yields poly(p-phenylene) derivatives with \(M_n\) values of 1500–2500. Similarly poly(1,4-naphthalene)s can be prepared by thermolysis of ortho-dialkynylbenzenes.

Precursor routes to cross-linked polyphenylenes have been reported. Soluble substituted polyphenylenes with number average molecular weights of up to 6300 are obtained by the nickel-catalysed homocoupling reaction of bis[(trifluoromethyl)sulfonyl]oxy) derivatives of substituted hydroquinone and benzene derivatives having two bromine or chlorine leaving groups. Other classes of soluble poly(p-phenylene)s have been synthesized by the polymerization of 2,5-dibromo-1,4-di-n-alkylbenzenes and 2,5-dibromobiphenyl, via Yamamoto coupling of 4-bromo-2,5-di-n-hexylbenzenem-glycolic acid and (3,5-dibromophenyl)boronic acid, anionic polymerization of 2-phenyl-1,3-cyclohexadiene followed by aromatization and spontaneous polymerization of 1-bromo-4-lithiobenzene in hexamethylphosphoramide.

Scheme 14 Poly(dihydrophenanthrene) via a nickel catalysed polycondensation

\[
\text{n Br} \\
n \text{Ni}_0\text{H}_3 \\
\text{n} \\
\]

Scheme 15 Poly(p-phenylene)s via thermolysis of enediyynes

\[
\text{n} \\
\]

5024 POLYMER Volume 37 Number 22 1996
Poly(phenylene vinylene)s

Poly(phenylene vinylene)s (PPVs) possess a chemical structure which is intermediate between that of polyacetylene and polyphenylene. Oriented PPV is highly crystalline\textsuperscript{136,137}, mechanically strong and environmentally stable. A number of approaches have been developed for the synthesis of this polymer and its analogues\textsuperscript{138–140}. Early attempts utilized dehydrohalogenation or Wittig condensation reactions which resulted in the formation of intractable oligomeric powders, but since the development of the water-soluble precursor method dense, highly oriented free standing polymer films of high molecular weight are reproducibly prepared\textsuperscript{137'141–143}. The Wessling-Zimmermann precursor route is outlined in Scheme 16.

The precursor sulfonium polyelectrolyte is prepared in aqueous solution by the base induced polymerization of an appropriate bis-sulfonium monomer, see Scheme 16. The reaction is usually carried out at low temperatures in fairly dilute monomer solutions and in equimolar (or lower) base to monomer ratios in order to suppress the premature formation of unsaturated polymer segments by thermal or base induced elimination of solubilizing side chains. The polymerization reaction is terminated by the addition of dilute aqueous hydrochloric acid to the reaction mixture which is then dialysed against water in order to separate the high molecular weight fraction from the monomeric and oligomeric residues as well as the sodium and chloride ions. A study of the polymerization mechanism\textsuperscript{144} revealed the existence of an intermediate of quinoid structure during the early stages of polymerization. In order to shift the reaction equilibrium towards the side of the polymer the stabilization of the quinoid intermediate or the removal of the thioether byproduct is desirable. This is readily achieved by the introduction of hexane in the reaction mixture\textsuperscript{145}. The hexane/water medium creates a two-phase system which, with vigorous stirring, produces an inverse emulsion polymerization environment. By maintaining a fine dispersion good contact between the two phases and effective exchange of the reaction components is ensured. The yields achieved by the two-phase polymerization method are normally about twice as high as those obtained by other procedures\textsuperscript{145a}. In addition to the improved reaction yields, the inverse emulsion polymerization method produces precursor polymers with higher molecular weights.

Another group of interesting PPVs are the cyano substituted PPVs which are prepared by a Knoevenagel condensation reaction between a diacetonitrile and a dialdehyde (Scheme 17)\textsuperscript{145b}.

Because of the versatility of the Knoevenagel condensation reaction this route can also be used for the synthesis of hetero-aromatic cyanopolymers such as the

\begin{center}
\textbf{Scheme 16} Poly(phenylene vinylene) by sulfonium polyelectrolyte route\textsuperscript{141}
\end{center}

\begin{center}
\textbf{Scheme 17} Knoevenagel condensation synthesis of cyano substituted PPVs\textsuperscript{145b}
\end{center}
Properties of conjugated polymers: W. J. Feast et al.

![Figure 3](image1.png)

**Figure 3** Heteroaromatic cyanopolymers

![Figure 4](image2.png)

**Figure 4** Deuterated PPV

poly(thiénylene-p-phenylene vinylene) copolymer 3 and poly(thiénylene vinylene) 4 (Figure 3). Properties of PPV films. Homogeneous dense precursor films may be readily prepared as coatings on substrates or in free-standing form by slow evaporation of the solvent. Casting the solution on a PTFE block or a glass dish treated with dichlorodimethylsilane as release agent, facilitates the preparation of free-standing films from which those thicker than approximately 1 μm can easily be lifted free. The conversion to the conjugated polymer is carried out by placing the cast and oriented films under vacuum or in an inert gas atmosphere in a constant-temperature bath at a desired temperature (determined by the nature and number of substituents attached on the benzene ring). Control of the sample morphology is readily achieved by varying the casting and conversion protocol. The full conversion of the precursor monomer to PPV requires a temperature of at least 300°C but annealing at temperatures of 400°C or higher leads to degradation. In the absence of an inert gas atmosphere or vacuum a proportion of the vinylene carbons are oxidized to carbonyls during the elimination process.

The molecular orientation in uniaxially drawn PPV films has been characterized by infrared dichroism and X-ray diffraction and very high values of the Hermans orientation function have been reported. Recently, Simpson et al. employed deuterium quadru-pole-echo n.m.r. experiments to study the chain orientation of drawn PPV films in which the phenylene ring was deuterated (Figure 4).

The films were prepared from water-cast films of the precursor polymer poly[(2,3,5,6-tetradeterio-p-xylylidene) tetrahydrothiophenium chloride] and were heated and stretched to effect simultaneous orientation and elimination. The results demonstrated that the average tilt of the phenylene ring relative to the chain axis was 7.7°, close but not exactly equal to the 9.2° predicted for a trans-stilbene-like structure. The difference was attributed to chain disorder within crystallites or at domain boundaries.

Highly oriented PPV shows high electrical conductivity upon vapour-phase doping with AsF5 but iodine doped films show only modest conductivities. The crystal structures of neutral pristine and doped forms have been studied by wide-angle X-ray diffraction. Poly-(p-phenylene vinylene) was found to undergo a first-order crystal-crystal phase transition when chemically doped with AsF5, SbS5, or sulfuric acid or when electrochemically oxidized with ClO4 as the counterion. The doping process does not disrupt the original orientation of the PPV crystallites and the crystalline phases obtained with all the above dopants are similar in character indicating a closely related family of electrically conductive structures all of which possess orthorhombic symmetry. On the basis of this work an electrically conductive phase consisting of layers of polymer chains separated by a layer of the chemical dopant was proposed. The influence of crystal defects and chain disorder on the electrical properties of PPV polymers have also been investigated. Highly drawn films of fully converted PPV exhibit a high degree of crystallinity and near-perfect orientation but, since distinctive paracrystalline diffraction patterns have been observed, there must be some structural disorder.

The one-dimensional band structures of PPV have been calculated. DaCosta et al. have extended the work by calculation of the three-dimensional band structure of PPV in order to study the effects of interchain coupling. The major effects are reported to be due to coupling of carbons in the ring to their closest neighbours (hydrogens or rings in adjacent chains). According to this study the existence of polaron in a perfect PPV crystal is not permitted but since these have been experimentally identified their existence is explained in terms of the presence of traces of residual precursor polymer within the matrix.

Arsenic pentafluoride doped PPV is not stable in air. In order to stabilize the p-type doped form of the polymer and decrease its ionization potential and band gap, the phenylene ring may be substituted with electron-donating substituents or replaced by heterocyclic aromatic molecules. By adopting this approach highly conducting iodine-doped films of poly(2,5-dimethoxy-p-phenylene vinylene) (Figure 5) have been prepared. The synthesis of the donor acceptor substituted poly(2-methoxy-5-nitro-p-phenylene vinylene) (10) and its copolymers with PPV has also been reported but these materials have a higher band gap than PPV itself. The interest in these materials is as potential...
Properties of conjugated polymers: W. J. Feast et al.

Figure 5 An anthology of poly(phenylene vinylene).

Figure 6 Linear polyacenes and polyarylenes.

electroluminescent components of light emitting diodes and there is a great amount of activity in the area of new poly(arylene vinylene) synthesis at present.

Two-dimensional ladder structures

Two-dimensional, ribbon-type conjugated polymer structures are known to exhibit better thermal stability and higher rigidity than their one-dimensional analogues\textsuperscript{166,167}. Furthermore, materials of this type are expected to exhibit low band gaps. The synthesis of defect free ladder or ribbon polymers is a demanding task which has attracted the attention of many research groups. The main synthetic strategies for the preparation of such materials is the utilization of repetitive cycloaddition reactions and, more recently, a two step approach in which a linear polymer is subjected to polymer-analogous ring-closure reactions with formation of ladder structures\textsuperscript{168,169}.

Linear polyacenes (Figure 6, 11) and polyarylenes (12) can be considered as two-dimensional subunits of graphite. Polyacenes are not stable if more than a few repeating units long. However, polyarylenes can be prepared from oligonaphthalene precursors by an electron-transfer induced electrocyclic process (oxidation or reduction) in which neighbouring naphthyl components are fused to perylene units\textsuperscript{170,171}.

The synthesis of polyacenes, high molecular weight analogues of biphenylene, has been explored\textsuperscript{172}. The most commonly employed procedure for the synthesis of this class of compounds involves the $\eta^2$-C\textsubscript{2}H\textsubscript{2}CoL\textsubscript{2} (L = CO, CH\textsubscript{2}CH\textsubscript{2}) catalysed cocyclization of orthotetraethynylarenes which can result in a series of multi[N]phenylenes\textsuperscript{173} (Scheme 18).

Because the alkynylarenes are readily prepared from the corresponding haloarenes by Pd-catalysed alkylation and since Si(CH\textsubscript{3})\textsubscript{3} can function as a masked halogen the method lends itself to the preparation of well-defined oligomers by an iterative procedure\textsuperscript{173-176}.

Another method to prepare polyarylenes makes use of a palladium-catalysed Suzuki coupling\textsuperscript{177} followed by a cyclization reaction (Scheme 19). After the cross-coupling reaction the resulting polyphenylene has to be

1) \((\text{CH}_3)_3\text{Si} - C\equiv C - H, \text{Pd}(\text{C}_6\text{H}_5\text{CN})_2\text{Cl}_2, \text{CuI}\)

2) KOH

3) \((\text{CH}_3)_3\text{Si} - C\equiv C - \text{Si}(\text{CH}_3)_3, \text{CpCo(CO)}_2, \text{hv}, \Delta\)

Scheme 18 Vollhardt’s route to multi[N]phenylenes\textsuperscript{172}
Properties of conjugated polymers. W. J. Feast et al.

Scheme 19 Polyarenes by the Suzuki/BF₃-cyclization sequence

Scheme 20 Polyacenes by thermal polymerization

Scheme 21 Thermochemically induced polyarylene formation

Planarized. This is performed by a BF₃ catalysed cyclization reaction which results in the double-stranded, fully soluble poly(fluoreneacene)s.

Other routes to polyacenes make use of Diels-Alder reactions, condensation-type cyclization reactions, a 'zipping-up' polymerization of vinyl substituted polymers, or polymerization of butadiynes as precursors. Unfortunately some of these methods give insoluble products from which no reliable structural information can be obtained.

A well-known route to polyarylenes is the pyrolysis of perilene-3,4,9,10-tetracarboxylic acid dianhydride in vacuo or under inert atmosphere giving black insoluble polymers which, from detailed structural analysis of films (X-ray, electron diffraction analysis, Raman spectroscopy), appear to be two-dimensional carbonaceous networks (Scheme 21). Under well-controlled conditions mirror like thin films can be formed on appropriate substrates.

Another route is shown in Scheme 22, the first step is the formation of poly(naphthalene) via a palladium (0) catalysed Suzuki-type condensation of naphthalenes containing bromo and boronic acid functionalities. In the next step this polymer is cyclized in a two-step procedure resulting in soluble macromolecules with ladder-type segments up to quaterarylene units. Unfortunately it is impossible to get complete cyclization via this approach.

Besides these hydrocarbon ladder polymers there are also a number of heteroatom containing species.

HETEROCYCLIC POLYMERS

Polythiophenes

The first polythiophene synthesis was described in 1883 when the purification of thiophene...
with sulfuric acid yielded a dark insoluble material\textsuperscript{201}. However, it was not until the early 1980s that any well-defined polymeric material was obtained. After the first reports of a controlled synthesis by Yamamoto\textsuperscript{202} and Lin\textsuperscript{203}, using the Grignard-type coupling of 2,5-dibromo-thiophene, a vast number of articles concerning the synthesis and properties of polythiophenes has been published.

Extended $\pi$-conjugation in polythiophenes is only possible in polymers with perfectly 2,5-linked repeating units; however, 2,4- and 2,3-couplings as well as hydrogenated thiophene units can also be found in the polymers\textsuperscript{204}. These structural defects interrupt the conjugation and, as a result, will impair the development of properties related to conductivity and nonlinear optics.

Polythiophenes, like many other linear polyaromatic compounds, are insoluble in organic solvents due to their rigid backbone. This lack of solubility and processability, as well as problems related to the characterization of polythiophenes, has been overcome by the introduction of flexible side chains at the 3- and/or 4-position. Appropriate solubility in common organic solvents has been achieved with an alkyl side chain of more than four carbon atoms at every repeating unit\textsuperscript{205}. Longer alkyl side chains are required in the case of copolymers with less than one side chain per repeating unit.

With the introduction of substituents at the 3-position of polythiophenes a number of different regioisomers are possible, namely head-to-tail (HT), head-to-head (HH) and random configurations (Scheme 23)\textsuperscript{206,207}.

Head-to-head coupling of alkyl groups is sterically unfavourable for coplanarity and hence causes a significant loss of conjugation; however, head-to-tail coupling does not limit conjugation. The significant difference in coplanarity between head-to-head and head-to-tail couplings shows the subtleties in the trade-off between resonance energy and steric hindrance in substituted polyheterocycles. Studies on regioregular polythiophenes, recently accessible via a number of elegant routes, have shown that the crystallinity increases with regioregularity and that the possibility of side-chain crystallinity is essential for the development of optimal properties. In the case of regiorandom polymers the optimal chain length for properties like conductivity and optical nonlinearities has been determined to be in the range of seven to nine carbon atoms\textsuperscript{208}; the conductivity of doped regioregular polythiophenes with an $n$-dodecyl side chain at the 3-position surpasses that of polymers with an $n$-octyl side chain\textsuperscript{209}. Even self-assembly in thin films of regioregular 3-substituted polythiophenes has been observed. In addition to improved solubility and crystallinity, the introduction of side chains in the 3- and/or 4-position also decreases the probability of $\alpha$-$\beta$ couplings, since at least one of the $\beta$ positions is already occupied\textsuperscript{210,211}.

3,4-Dialkyl substitution in monomers results in a significant loss of coplanarity\textsuperscript{212}, while in 3,4-dialkoxy and 3-alkyl-4-alkoxy polythiophenes the presence of an oxygen atom attached to the thiophene unit, is enough to decrease steric hindrance and limit the loss of conjugation\textsuperscript{213}. An unexpectedly high regioregularity was found in the polymerization of 3-methyl-4-alkoxythiophenes\textsuperscript{214}.

An interesting and intriguing property of substituted polythiophenes is the temperature dependence of the u.v. absorption; an increase in temperature causes the $\lambda_{\text{max}}$ to shift to shorter wavelengths. This thermochromism arises from a conformational change in the aromatic backbone as a result of an increased disorder of the alkyl
Properties of conjugated polymers: W. J. Feast et al.

Scheme 23 Different regioisomers in polythiophene

Figure 8 Thermochromism in a 3-substituted polythiophene

Much controversy exists about the actual molecular weight of polythiophenes. G.p.c. determinations using polystyrene standards overestimate the molecular weight by a factor of ten compared to the determination relative to oligothiophene standards\(^{19}\). A similar phenomenon has been observed in the molecular weight determination of poly(\(\text{t-butyl} n\)-naphtalene) although in that case the deviation is only around 10\%\(^{20}\). By using multi-angle light scattering (MALLS) the calculated molecular weight values for polythiophenes are two to five times larger than those obtained by g.p.c. relative to polystyrene standards\(^{21}\).

For the preparation of polythiophenes three main synthetic strategies have been employed, namely: electrochemical coupling, oxidative coupling and organo-metallic coupling of 2,5-disubstituted monomers. These three routes will be discussed in the following sections.

**Electrochemical synthesis of polythiophenes.** Since the first reports by Diaz\(^{22}\), the synthesis of polythiophenes by electrochemical oxidation has been widely used. Although the mechanism is not fully understood, it is proposed that the polymerization proceeds via the coupling of two radical cations, formed by the oxidation of the monomer as outlined in Scheme 24\(^{23}\).

Aromatization of the bithiophene intermediate is the driving force for the transformation of the dihydro dimer. The dimer, having a lower oxidation potential than the monomer, is readily oxidized and undergoes further coupling. The polymer is deposited in its oxidized conducting form onto the electrode, allowing the polymerization to proceed. This method has the advantage that during the polymerization homogeneous, stable films are formed. These films can be characterized by optical and electrochemical methods. The electrochemical polymerization has been applied for the synthesis of unsubstituted polythiophene\(^{24,25}\) and 3-substituted polythiophenes including poly(3-methylthiophene)\(^{26}\), poly(3-ethylthiophene)\(^{27}\) and number of different soluble poly(3-alkylthiophene)s\(^{28,29}\). However, in all cases reported so far, the polymers obtained possess a regiorandom structure.

The electrochemical polymerization of substituted thiophenes has also been used to synthesize a variety of functional polymers. Oxygen containing substituents on the 3-position of thiophenes have been used, including methoxy\(^{30}\) and polyether substituents\(^{31}\), leading to materials with conductivities as high as 1000 S cm\(^{-1}\). A highly transparent conducting polymer has been obtained from the electrochemical polymerization of 3,4-ethylenedioxythiophene (Figure 9, 19)\(^{32}\).

Similarly polyfluoroalkyl substituted polythiophenes...
Properties of conjugated polymers: W. J. Feast et al.

Scheme 24  Electrochemical polymerization of thiophene

Figure 9  Poly(3,4-ethylenedioxythiophene)

Thiophene oligomers such as bithiophenes and terthiophenes possess lower oxidation potentials than the thiophene monomers (1.05 V, 1.31 V, and 2.07 V (vs. SCE) for terthiophene, bithiophene and thiophene, respectively) and are therefore more suitable for the electrochemical polymerization since this lower oxidation potential decreases the risk of over oxidation. Also 2,5-disilyl substituted thiophenes have been polymerized electrochemically with concomitant elimination of the silyl substituents.

Oxidative coupling to polythiophenes. Conducting polythiophenes are formed in the oxidation of thiophene or 2,2'-bithiophenes with arsenic(V) pentafluoride (AsF$_5$). Reduction of the insoluble material with ammonia clearly demonstrated that polymerization has occurred. However, due to the poisonous properties of arsenic(V) pentafluoride this method has not been widely adopted. A more convenient method was developed by Yoshino et al. who used iron(III) trichloride (FeCl$_3$) as the oxidizing agent and chloroform as the solvent under anhydrous conditions. Subsequent reduction with ammonia provided the neutral polymer in good yields. This coupling reaction is simple and has now been widely employed for the synthesis of polythiophenes from thiophene with alkyl, alkoxy and alkylsulfonic acid substituents and bithiophenes with one or two alkyl substituent. These bithiophene monomers can also be polymerized using milder oxidants.

Recently, scientists have been able to regioselectively polymerize 3-(4-octylphenyl)-thiophene using FeCl$_3$. By adding the FeCl$_3$ slowly and therefore maintaining a low Fe$^{3+}$/Fe$^{2+}$ ratio, this 'soft' polymerization leads to a polymer with up to 94% head-to-tail couplings (Scheme 25).

The oxidative coupling of thiophenes provides materials with higher molecular weights than the route described above. However, in a recent study a comparison has been made between the polymerization of 3-benzylthiophene and its deuterated analogue indicating that considerable crosslinking occurs through the alkyl side chain. The degree of crosslinking is reduced by performing the reaction at temperatures below -20°C, which also increases the yield of soluble 3-substituted polythiophenes.
Organometallic mediated syntheses of polythiophenes.
The nickel\textsuperscript{253} and palladium\textsuperscript{254} catalysed cross-coupling
of an aromatic organometallic compound with an aromatic organohalide is a well-known method for the
synthesis of oligoheterocycles\textsuperscript{255}. Organomagnesium
derivatives are most widely employed in polythiophene
chemistry, despite the availability of a variety of organo-
metallic species\textsuperscript{256}. The reaction of 2,5-dibromothio-
phene with magnesium and subsequent polymerization
using a nickel catalyst was described in 1980, with a
reported yield of 40-60\% based on 2,5-
dibromothiophene\textsuperscript{202,203}. The yields have been improved
up to 93\% by using 2,5-diodothiophene, magnesium
and Ni(dppp)Cl\textsubscript{2} as catalyst\textsuperscript{257}.

The synthesis of 3-alkylsubstituted polythiophenes,
with molecular weights varying from 2500 to 18 000, has
also been accomplished by this method, although the
structures obtained are regiorandom\textsuperscript{258,259}. This regio-
randomness has been explained in terms of the inhomoge-
neity of the Grignard iodide mixture, which consists of
diiodo-, bis-Grignard- and two monoiodo-mono-
Grignard compounds\textsuperscript{260,261}. Cross-coupling has also
been utilized for the synthesis of 3-substituted thiophene
precursors\textsuperscript{262,263}.

Highly regioregular head-to-tail 3-alkylpolythio-
phenes have been prepared by McCullough \textit{et al.} via
the polymerization of 2-bromomagnesio-5-bromo-3-
alkylthiophenes (Scheme 26)\textsuperscript{209,264--266}.

By this elegant route a variety of highly regioregular
(over 98\% head-to-tail) polymers has been obtained in
yields ranging from 33 to 69\%. These polymers exhibit
enhanced conductivity and optical properties when compared with the regiorandom materials. The con-
ductivity of poly(3-dodecylthiophene) reaches values
upto 10 000 S m\textsuperscript{-1} whereas for the regiorandom material
the conductivity is limited to 20 S cm\textsuperscript{-1}. The difference
in the position of the maximum absorption (450 vs.
434 nm in solution and 480 vs. 526 nm in the solid
state) are indicative of enhanced coplanarity in these
regioregular polythiophenes. As a result, the thermo-
chromism of these regioregular polythiophenes is much
more pronounced. X-ray analysis indicates a crystalline
and self-assembled structure and a well-defined dis-
tance between the polymer chains\textsuperscript{267}. Cyclic voltam-
metry exhibits two oxidation potentials vs. one broad
oxidation potential for the regiorandom poly(3-
alkylthiophene)\textsuperscript{268}.

Another approach to modulate the regioregularity is
by using zinc instead of magnesium in the nickel\textsuperscript{269,270}
or palladium\textsuperscript{271} catalysed polymerizations. The use of
Ni(dppe)Cl\textsubscript{2} affords regioregular samples (Scheme 27)\textsuperscript{72}
and Japanese workers had similar experience with
duzovalent nickel\textsuperscript{273,274}. In the latter case the 2,5-
dihalothiophene was coupled without being transformed
to an organometallic intermediate, and therefore, yields
a regiorandom polythiophene.

Due to the improved properties of the regioregular 3-
substituted polythiophenes, the polymerization based on
the organometallic coupling of thiophene monomers is
now by far the most valuable method of synthesis.
Recently, a number of polymers with functional groups
has been synthesized using the organometallic route,
giving rise to polythiophenes with unique sensor proper-
ties for ionic impurities\textsuperscript{275--277}.

\textbf{Polypyrrole}

The first synthesis of polypyrrole (Figure 10, 20),
which included a description of the conducting proper-
ties of the product, was published in 1968\textsuperscript{278}. The
electrochemical oxidation of pyrrole in 0.1 N sulfuric
acid yielded a black conducting film. Improvements by
using organic solvents and different electrolytes have

\begin{center}
\begin{tikzpicture}

\begin{scope}
  \node[above] at (-1.5,0) {R};
  \node[below] at (-1.5,-1) {Br};

  \node[above] at (1.5,0) {R};
  \node[below] at (1.5,-1) {Br};

  \draw[-latex] (-1.5,0) -- (1.5,0);
  \draw[-latex] (-1.5,-1) -- (1.5,-1);

  \node at (-1.5,-0.5) {1) LDA, THF, -78°C to -40°C}
  \node at (1.5,-0.5) {2) MgBr\textsubscript{2}, OEt\textsubscript{2}, -60°C to -5°C}
  \node at (-1.5,-1.5) {3) NiCl\textsubscript{2}(dppe), -5°C to 25°C}

  \node at (-0.2,-0.8) {head-to-tail}

  \begin{scope}[xshift=4cm]
    \node[above] at (-1.5,0) {R};
    \node[below] at (-1.5,-1) {Br};

    \node[above] at (1.5,0) {R};
    \node[below] at (1.5,-1) {Br};

    \draw[-latex] (-1.5,0) -- (1.5,0);
    \draw[-latex] (-1.5,-1) -- (1.5,-1);

    \node at (-1.5,-0.5) {1) LDA, THF, -78°C to -40°C}
    \node at (1.5,-0.5) {2) MgBr\textsubscript{2}, OEt\textsubscript{2}, -60°C to -5°C}
    \node at (-1.5,-1.5) {3) NiCl\textsubscript{2}(dppe), -5°C to 25°C}

    \node at (-0.2,-0.8) {head-to-tail}
  \end{scope}
\end{scope}
\end{tikzpicture}
\end{center}

\begin{center}
\textbf{Scheme 26} McCullough's route to regioregular polythiophenes\textsuperscript{209,264--266}
\end{center}

\begin{center}
\begin{tikzpicture}

\begin{scope}
  \node[above] at (-1.5,0) {Br};
  \node[below] at (-1.5,-1) {Br};

  \node[above] at (1.5,0) {Br};
  \node[below] at (1.5,-1) {Br};

  \draw[-latex] (-1.5,0) -- (1.5,0);
  \draw[-latex] (-1.5,-1) -- (1.5,-1);

  \node at (-1.5,-0.5) {1) LDA, THF, -78°C to -40°C}
  \node at (1.5,-0.5) {2) MgBr\textsubscript{2}, OEt\textsubscript{2}, -60°C to -5°C}
  \node at (-1.5,-1.5) {3) NiCl\textsubscript{2}(dppe), -5°C to 25°C}

  \node at (-0.2,-0.8) {head-to-tail}

  \begin{scope}[xshift=4cm]
    \node[above] at (-1.5,0) {Br};
    \node[below] at (-1.5,-1) {Br};

    \node[above] at (1.5,0) {Br};
    \node[below] at (1.5,-1) {Br};

    \draw[-latex] (-1.5,0) -- (1.5,0);
    \draw[-latex] (-1.5,-1) -- (1.5,-1);

    \node at (-1.5,-0.5) {1) LDA, THF, -78°C to -40°C}
    \node at (1.5,-0.5) {2) MgBr\textsubscript{2}, OEt\textsubscript{2}, -60°C to -5°C}
    \node at (-1.5,-1.5) {3) NiCl\textsubscript{2}(dppe), -5°C to 25°C}

    \node at (-0.2,-0.8) {head-to-tail}
  \end{scope}
\end{scope}
\end{tikzpicture}
\end{center}

\begin{center}
\textbf{Scheme 27} Riecke's regioselective route to HT-polythiophene\textsuperscript{72}
\end{center}
made the electrochemical method the most commonly employed polymerization technique\textsuperscript{29,280}. The mechanism of the electrochemical polymerization is similar to that of thiophene as shown in Scheme 24.

Oxidized polypyrrole is stable under ambient conditions and up to temperatures exceeding 300°C. The neutral form of polypyrrole, on the other hand, has not been isolated and characterized, due to its extreme susceptibility to oxidation (\textasciitilde-0.02 V vs. SCE).

The electrochemical route to polypyrrole provides good quality films. The counter ion has a considerable influence on the conductivity and mechanical properties\textsuperscript{281}. Changing the counter ion from oxide to perchlorate increases the conductivity by a factor of 10\textsuperscript{205,274}. Commercially available polypyrrole films with tosylate as counterion exhibit a conductivity of 15 S cm\textsuperscript{-1} and the stability of the film under ambient conditions is extremely good; a decrease in conductivity of less than 15% per year has been reported\textsuperscript{282}. Alkylsulfonates and phosphates have also been used as electrolyte\textsuperscript{283}, while processable polymer blends have been formed using sulfonated polystyrenes as counter ions\textsuperscript{284}.

Oxidation with chemical oxidizing agents (sulfuric acid\textsuperscript{285}, bromine and iodine\textsuperscript{286}, copper(II) perchlorate\textsuperscript{287}, iron trichloride\textsuperscript{288}) of a neutral polypyrrole film increases the conductivity compared to that of the electrochemical oxidized material\textsuperscript{289}. 2,2'-Bipyrrrole has been used as monomer in the chemical oxidative polymerization, but the properties of the product are similar to the polymer prepared directly from pyrrole\textsuperscript{290}.

Soluble polypyrroles can be prepared by the introduction of flexible side chains\textsuperscript{229,291-293}. In contrast with the progress made in the synthesis of regioregular 3-substituted polythiophenes, all 3-substituted polypyrroles reported so far are synthesized in a regiorandom fashion. 3-Alkyl pyrrroles, the monomers in the electrochemical polymerization, have been synthesized via Friedel–Crafts acylation of N-protected pyrrole, followed by Clemmenson reduction (Scheme 28)\textsuperscript{294-296}.

Substitution at nitrogen also affords soluble polymers. However, the conductivity is reduced drastically due to the strong steric interactions of this substituent at nitrogen and the hydrogen rings at the 3- and 4-positions of the adjacent pyrrole ring. Both rings are forced out of plane resulting in a loss of conjugation and a reduced conductivity\textsuperscript{297}. At first sight the difference between a 3-substituted polypyrrole and a N-substituted polypyrrole is marginal with respect to steric hindrance, but the conductivities differ significantly, again illustrating how subtle structural differences influence the structure–property relationship. Chemical oxidative polymerization has also been applied to 3-alkyl pyrrroles\textsuperscript{285,286}, and 3,4-dimethoxy pyrrrole\textsuperscript{298}.

The introduction of sulfonic acid groups in the alkyl side chain afforded water-soluble self-doped polypyrroles\textsuperscript{299}. When the sodium salt of the 3-alkylsulfonic acid pyrrole is used as monomer it also acts as an electrolyte for the electrochemical synthesis. A second long alkyl chain on the 4-position of the pyrrole ring affords a highly ordered lamellar polymer which is also soluble in chloroform\textsuperscript{300}. Langmuir–Blodgett techniques have been used to improve the ordering of polypyrrole films\textsuperscript{301,302}.

Structural defects like α-β couplings are inherent to the oxidative polymerization approach and are always present in the materials synthesized according to the methods described above. This failure to produce perfectly 2,5-linked polypyrroles has been overcome by using organometallic polymerization techniques. Pyrroles with a tert-butoxycarbonyl (BOC) protecting group at nitrogen have been polymerized via a Stille\textsuperscript{254} coupling, yielding a soluble non-planar precursor polymer which was deprotected by thermal treatment (Scheme 29)\textsuperscript{303}.

This polypyrrole, although of relatively low molecular weight (approximately 16 pyrrole units), exhibits a perfect α-α-structure and is fully characterized.

Recently, the reductive Ullmann\textsuperscript{304} coupling of 2,5-dibromo-N-BOC-pyrrole has been reported to yield similar polymers to those described above\textsuperscript{305}. Using preparative h.p.l.c. it appeared to be possible to isolate the first 20 members of the N-BOC protected oligopyrroles, which were all characterized. Similarly, a self-doped analogue of polypyrrole has been prepared using N-buty1-3,4-pyrroledione in the Ullmann polymerization\textsuperscript{306}.
Polyaniline

Polyaniline (Figure 11, 21), also known as 'aniline black', was first prepared in 1834 and has been the subject of intensive research ever since\(^\text{307}\). In the 1980s the conducting properties of polyaniline were recognized and the number of papers dealing with this conducting polymer grew rapidly. The main reasons for this growth, besides the scientific interest, are the low cost of aniline, the relatively easy production process and the stability of the conducting forms.

**Synthesis of polyaniline.** Oxidation of aniline is the most widely employed synthetic route to polyaniline and can be performed either electrochemically or chemically. The reaction is usually carried out in acidic media. The electrochemical method was originally developed by Letheby as a test for the determination of small quantities of aniline\(^\text{308}\). The method has been improved ever since\(^\text{309-311}\) and has also been applied to alkyl\(^\text{312,313}\), alkoxy\(^\text{314}\) and dimethoxy\(^\text{315}\) substituted anilines, the latter exhibiting a conductivity comparable to that of unsubstituted polyaniline.

Chemical oxidation is usually carried out in acidic aqueous environments with an oxidizing agent such as ammonium persulfate\(^\text{316}\), but has also been performed in chloroform using tetrabutyl ammonium periodate\(^\text{317}\). Alkyl\(^\text{312}\) and alkoxy\(^\text{314}\) substituted polyanilines have also been prepared utilizing this chemical method.

It has been stated that in polyaniline crosslinking occurs during the electrochemical synthesis due to the potential applied\(^\text{318}\). However, non-oxidatively synthesized polyaniline, exhibiting the same properties, has a fully linear structure\(^\text{319}\). Crosslinking in polyaniline can also be accomplished when the emeraldine base is heated up to 300°C\(^\text{320}\).

**Properties of polyaniline.** Polyaniline can occur in a number of well-defined different oxidation states\(^\text{320}\), each with its own name as originally attributed by Green and Woodhead (Figure 12)\(^\text{321}\).

The different states range from the fully reduced leucoemeraldine via protoemeraldine, emeraldine and nigraniline to the fully oxidized pernigraniline. Unlike in most other polyaromatics, the fully oxidized state in polyaniline is not conducting. As a matter of fact none of the above-described states are conducting. Polyaniline becomes conducting when the moderately oxidized states, in particular the emeraldine base, are protonated and charge carriers are generated. It is this process, generally called 'protonic acid doping'\(^\text{312}\), which makes polyaniline so unique; no electrons have to be added to or removed from the insulating material to make it conducting. The different oxidation states of polyaniline can also be generated by doping with oxidants such as iodine, but the resulting conductivity is lower than that obtained via protonic acid doping\(^\text{314}\).

The conducting mechanism is believed to involve polaron carriers; the protonated emeraldine consists of a delocalized poly(semiquinone radical cation)\(^\text{317,322,323}\). The conductivity is affected by the water content; completely dry samples are five times less conductive.
than samples containing some water. The emeraldine base is soluble in N-methyl-pyrrolidone, but protonated polyaniline is insoluble in organic solvents and only soluble in aqueous acids. Substitution of the aniline monomer with alky1 or alkoxy groups improves the solubility in organic solvents but has a negative influence on the conductivity. The position of the substituent also has an influence on the polymerization. The ortho and meta isomers give the same polymer, but the reactivity of the meta isomer is considerably lower, resulting in a lower yield. Self-doped polyaniline, containing sulfonic acid substituents, has been synthesized by sulfonation of the emeraldine base.

A different approach to the production of soluble polyanilines is the use of n-alkyl sulfonic acids as proton donor. Polyaniline doped with n-4-dodecylphenyl-sulfonic acid does indeed give soluble and highly conducting materials. Stable thin films of polyaniline, that can be processed out of solution, are prepared from the emeraldine base with camphorsulfonic acid as dopant. m-Cresol as cosolvent is essential to obtain a chiral crystalline structure.

Although no data are available concerning the health risks of polyaniline, the possible presence of benzidine moieties, which are well-known carcinogens, suggests careful manipulation of both aniline and its polymers is advisable.

Other polymers

In the field of conducting and \( \pi \)-conjugated polymers most attention has been paid to the polymeric systems described above. However, a variety of other polymers containing heterocyclic rings has been synthesized and studied. In the following sections some of the most promising classes, with respect to improved material properties, will be discussed.

Small band gap polymers. The search for small band gap polymers is a special field of research within the general area of synthetic metals. Although many participants recognize the interest and importance of this class of polymers, the number of dedicated papers is limited. In addition to their high conductivity, these systems are expected to be transparent in their conducting form, and when the band gap approaches zero, these materials are predicted to exhibit metallic properties.

In 1984 Wudl et al. published the synthesis of poly(isothianaphthene) (PITN) with a band gap of 1 eV (for comparison: polythiophene has a band gap of 2 eV). Limited synthetic work has been performed on these kinds of system ever since, but recently the syntheses of a substituted PITN and of poly(2,3-dihexylthieno[3,4-b]pyrazine) were published.

The small band gap is believed to arise from the relatively large contribution of the quinoid structure in the polymers. Calculations predict that a copolymer of thiophene and isonaphthothiophene will exhibit a band gap of 0.5 eV. Polycyclopenta[2,1-b;3,4-b]dithiophen-4-one has a band gap of 1.2 eV due to its reduced aromatic character. The introduction of one fused thiadiazole unit in a terthiophene molecule resulted in a polymer with a band gap of 0.9 eV, arising to some extent from the nitrogen-sulfur interactions improving the intra-chain charge mobility. The synthesis of a polymer with a band gap as low as 0.5 eV based on the idea of bringing together positive and negative charges was published by Havinga et al., using polysquarenes and polycroconanes which were synthesized by using a condensation polymerization. Furthermore, Jenekhe has suggested that the introduction of methine groups between thiophene moieties would provide polymers with band gaps as low as 0.75 eV; however, the validity of this approach has been questioned.

None of the small band gap materials reported so far has conductivities surpassing the conventional polymers like doped polythiophene, polypyrrole and polyaniline, an observation which puts the supposed importance of the band gap in perspective.

Fused monomers. Fused heterocyclic compounds like thieno[3,2-b]pyrrole (Figure 14, 27), thieno[3,2-b]thio-
Properties of conjugated polymers: W. J. Feast et al.

Phene\textsuperscript{345} (28) and dithieno[3,2-b; 2',3'-d]thiophene\textsuperscript{346} (29) have been polymerized electrochemically. Other nitrogen containing polymers such as poly(quinoxaline-5,8-diyl)\textsuperscript{347} (30), poly(quinoline-5,8-diyl) (31) and poly(isoquinoline-1,4-diyl)\textsuperscript{348} (32) have been synthesized by the condensation polymerization of the dibromo compounds under the influence of a zerovalent nickel catalyst.

A completely fused ladder polymer with a quinoline moiety was synthesized by cyclization of the amine and ketone functionalities of a polyphenylene backbone\textsuperscript{349}, yielding a polymer with a high degree of coplanarity.

A special kind of fused polyaromatic is polyphenylene sulfide. The polymer has been synthesized by a number of methods including electrochemically from thiophene,\textsuperscript{350} oxidatively from phenylene disulfide\textsuperscript{351} and chemically from 4,4'-dibromo-phenylene disulfide with copper\textsuperscript{352}. The aromatic rings of polyphenylene sulfide are coupled upon oxidation with arsenic(V) pentafluoride in arsenic(III) trifluoride leading to fused benzothiophene moieties\textsuperscript{353}. Conductivities up to 200 S m\textsuperscript{-1} have been reported.

Finally, some results from studies on fused conjugated polymers based on dyes have been published\textsuperscript{354,355}; these materials are claimed to be intrinsically conducting.

Other homopolymers. Polypyridine has been synthesized through the homocoupling of 2,5-dibromopyridine\textsuperscript{356} or 5,5'-dibromo-2,2'-bipyridine\textsuperscript{357} with zerovalent nickel catalysts (Scheme 30).

The polymers are only soluble in formic acid as is also the case in methyl substituted polypyridine\textsuperscript{358}. However, the introduction of hexyl groups induces solubility in other organic solvents\textsuperscript{359}. Recently, the peculiar properties of these polypyridines have been attributed to self-assembly\textsuperscript{360}. Poly(pyrimidine-2,5-diyl)\textsuperscript{361} has been prepared utilizing the same method. Other examples of heterocyclic homopolymers are polypyrilidazine\textsuperscript{362} and polythiazole\textsuperscript{363}.

Copolymers

The synthesis of alternating benzenoid–heteroaryl copolymers enables, in principle, the combination of the properties of different homopolymers into one material, increasing the scope of synthesis and applications. The synthesis of these copolymers has, however, received relatively little attention. This will probably change with the increasing interest in small band gap polymers. The idea of using copolymers to introduce alternating structures of electron-rich and electron-poor moieties has emerged. Three synthetic routes have been developed to make copolymers: (a) the direct polymerization of the different aromatics by either oxidative or chemical coupling methods; (b) ring closure of precursor polymers; and (c) polymerization of oligomers which already contain the desired array of units. The different strategies are outlined in the sections below.

Direct polymerization. Oxidative copolymerization

Figure 14 Fused monomers and fused polymers

Scheme 30 Nickel catalysed polymerization of pyridine derivatives\textsuperscript{356,357}
Properties of conjugated polymers: W. J. Feast et al.

requires monomers having similar oxidation potentials, otherwise the material obtained is more likely to have a block rather than the required alternating structure. This places limits on the oxidative approach but it has, nevertheless, been used for a pyrrole-thiophene copolymer, prepared by the electrochemical polymerization of pyrrole and \( \alpha \)-terthiophene\(^{364} \), monomers having oxidation potentials of 1.2 V and 1.05 V, respectively.

The chemical route to alternating polymers is much more general; the reaction of bis-metallated and dihaloaromatic species of which either one can contain the heterocyclic ring, yields a completely alternating copolymer when palladium or nickel catalysts are used. Thus, heterocycles such as thiophene, selenophene and pyridine have been coupled with benzene and biphenyl using magnesium and a nickel catalyst\(^{365,366} \). Thiophene and furan have been coupled with benzene and substituted benzenes utilizing zinc and palladium catalysts\(^{367,368} \). While alkoxy-substituted benzenes and thiophenes have been transformed into an alternating copolymer via the Stille coupling\(^{369} \). The Suzuki coupling\(^{370} \) (arylhalide and boronic acid derivative) has been used in the coupling of alkyl-substituted benzenes with N-BOC protected pyrroles\(^{371} \) (Scheme 31).

When zerovalent nickel is used as a catalyst in the copolymerization of thiophene with benzene or pyridine\(^{372} \), materials with a random structure are formed due to the absence of organometallic monomers in this type of polymerization\(^{373,374} \). Reaction of dibromo thiophenes and \( p \)-phenylene diamines under 'Ullmann conditions' yields copolymers of aniline and thiophene\(^{383} \).

A final example of copolymers prepared by the direct polymerization route are the random head-to-tail poly(3-alkylthiophene) copolymers prepared by McCullough et al. using two different monomers (Scheme 32)\(^{376} \).

**Precursor route.** Thiophenes and pyrroles can be synthesized by the ring closure of diacetylenes with hydrogen sulfide\(^{377} \) and primary amines\(^{378} \), respectively, or from 1,4-diketones with Lawesson's reagent\(^{379} \), primary amines\(^{380} \) and ammonium acetate\(^{381,382} \), respectively (Scheme 33).

An alternating copolymer of phenylene and diacetylene or 1,4-diketone moieties offers the possibility of the synthesis of an alternating copolymer consisting of phenylene and thiophene or pyrrole rings. Thus, 1,4-dithylyn benzene was oxidatively coupled to give the phenylene diacetylene copolymer, containing approximately 10% of the undesired meta isomer. Upon ring closure with hydrogen sulfide and aniline the phenylene thiophene and the phenylene N-phenyl pyrrole polymers were obtained\(^{381} \).

The reaction of terephthaldehyde and the bis-Mannich base of 1,4-diacetyl benzene with sodium cyanide as a catalyst (Stetter reaction\(^{384} \)) yielded polyphenylene-1,4-butanedione; the alternating phenylene 1,4-diketone polymer. Ring closure with Lawesson's reagent (LR) or ammonia yielded the phenylene thiophene and phenylene pyrrole copolymers in good yield (Scheme 34)\(^{385} \).

**Oligomer polymerization.** In order to avoid the formation of block copolymers by the oxidative polymerization of different monomers, it is possible to use oligomers containing the desired sequence of units which can then be oxidatively polymerized. In this way a copolymer of thiophene and pyrrole has been prepared by the electrochemical polymerization of 2,2'-thienylpyrrole\(^{386} \). Similarly, a number of trimers containing thiophene,
Properties of conjugated polymers: W. J. Feast et al.

**Scheme 33** Ring closure reaction to thiophene and pyrrole derivatives

**Scheme 34** Copolymers via Stetter/ring closure route

**Scheme 35** Organometallic coupling of a thienylpyridine monomer

Furan and N-alkyl-pyrrole moieties have been oxidatively polymerized with NOPF₆, yielding the expected copolymers.

An alternating copolymer of benzene and bithiophene units has been synthesized by the polymerization of 1,4-dithienyl benzenes either electrochemically or oxidatively with iron(III) trichloride. Introduction of two alkyl or alkoxy substituents on the phenyl ring afforded regular and soluble polymers.

Two final examples are a polymer prepared by chemically oxidation of 1,4-dipyrrylbenzene, and one containing thiophene and pyridine units which was prepared from the dibromide using zerovalent nickel (Scheme 35).

**OLIGOMERS**

During the last few years there has been a growing interest in the synthesis and characteristics of well-defined oligomeric conjugated structures. There are a number of reasons for this interest, including the ability of such oligomers to serve as model compounds for the different polymers with respect to the synthesis, spectroscopic analysis and physical properties as well as their potential applications. Recently Garnier et al. reported on the first all-organic transistor using the unique properties of a self-assembled layer of a substituted sexithiophene. The high electron mobility observed in this oligomer, almost comparable to amorphous silicon, is of critical importance to the construction of this prototype device. Although the class of oligothiophenes has received by far the most attention of all π-conjugated oligomers, other classes of oligomers, as oligopyrroles and oligoanilines, will also be discussed in the next sections.

**Oligothiophenes**

Well-defined oligomers of thiophene have been isolated from Marigolds and have been synthesized by different routes including the cross-coupling reaction of a dihalo compound with organometallics of magnesium,
zinc or tin with a palladium or nickel catalyst (including the Kumada coupling\textsuperscript{399} to regioregular oligothiophenes, Scheme 36\textsuperscript{400–406}, the copper(II) chloride oxidation of a lithiated thiophene\textsuperscript{407–409} and the ring closure reaction of a 1,4-diketone with Lawesson’s reagent\textsuperscript{410–412}).

The conformational analysis of oligothiophenes has been studied in detail. The structures of oligothiophenes containing two\textsuperscript{413}, three\textsuperscript{414} and four\textsuperscript{415,416} thiophene rings have been resolved by X-ray analysis; all the compounds exhibit an all-trans conformation and are nearly coplanar. However, u.v. and n.m.r. analysis, together with calculations, have shown that in solution unsubstituted tetrathiophene exhibits a planar cis-trans-cis conformation. The introduction of methyl substituents induces an equilibrium between planar cis and twisted trans-conformations and when the substituents are head-to-head the twisted trans-conformation is favoured, resulting in a considerable loss of conjugation\textsuperscript{406, 417}.

The use of oligomers containing more than six thiophene rings is restricted to substituted compounds due to the insolubility of the unsubstituted oligomers. Most of the oligomers differ from the substituted polythiophenes in their substitution pattern; they do not contain one substituent on every ring as is the case in most soluble polythiophenes.

Upon oxidation some of the shorter oligomers, containing up to six thiophene rings, were found to polymerize\textsuperscript{418}, so the reported high conductivities of these oligomers are probably due to polymeric material\textsuperscript{419}. Oligomers containing more than six thiophene rings do not polymerize, while it has been found that oligomers containing 11 or 12 thiophene rings exhibit conductivities of 5–20 S cm\textsuperscript{−1}, being in the same range as that of polythiophene\textsuperscript{397,408}.

Optical measurements show absorption maxima exceeding that of polythiophene\textsuperscript{394,408,420}, but this is mainly due to a decrease of steric hindrance in the oligomers having less substituents. Extrapolation of these values results in a $\lambda_{\text{max}}$ for polythiophene of about 540 nm\textsuperscript{397,410}, implying that the perfect polythiophene has not yet been synthesized.

A number of oligothiophenes has been employed to investigate the nature of the different oxidation states in polythiophene. Oxidation of oligothiophenes proceeds in two steps. First a radical cation is obtained (the analogue of the polaron state in conjugated polymers), which exhibits a strong e.s.r. signal. Some authors have reported the dimerization of these radical cations to diamagnetic $\pi$-dimers\textsuperscript{421–425}. In the second step of the oxidation the radical cation is converted into the dication (the bipolaron analogue). It has been reported that the oligomers must consist of at least six thiophene rings to

![Scheme 36](image)

**Scheme 36** Kumada coupling to regioregular oligothiophenes\textsuperscript{406}

![Scheme 37](image)

**Scheme 37** Oligopyrroles by the Stille coupling reaction: (a) oligo(pyrrole-2,5-diyl)s; (b) phenyl blocked mixed oligomers based on pyrrole\textsuperscript{433–437}
Properties of conjugated polymers. W. J. Feast et al.

Scheme 38  H.p.l.c. analysis of the Ullmann polymerization of a dibrominated pyrrole monomer

undergo the second oxidation step, but this claim is at variance with other results.

Langmuir–Blodgett films of oligothiophenes and in particular 2,2',5',2''-terthiophenes derivatives exhibit large dielectric constants and high conductivities with or even without additional dopants. Oligothiophenes also display other characteristics that have been found in polythiophenes, such as thermochromism and hyperpolarizability.

Oligopyrroles

In sharp contrast to the oligothiophene series, relatively little work has been performed on the synthesis and characterization of oligopyrroles. This is mainly due to the complexity of these compounds. There are three main routes which result in oligomeric pyrrole systems. The first one consists of the oxidative coupling of o-lithiated N-methyl substituted oligopyrroles using NiCl₂ resulting in N-methyl substituted oligomers up to 16 pyrrole units. The second route makes use of the previously described Stille coupling. By performing a Pd catalysed cross-coupling reaction between an arylhalide and an arylstannyl derivative, both homo and co-oligomers based on pyrrole can be prepared (Scheme 37).

The crystal structures of the dimer (m = 0), trimer (m = 1) and pentamer (m = 3) of this oligopyrrole series have been resolved and clearly display coplanar structures for the unprotected oligomers and a 70 degree twist angle for the N-BOC-protected analogues. The unprotected oligomers have also been investigated by FT-i.r. and Raman spectroscopy, cyclic voltammetry and u.v.-vis-n.i.r. spectroscopy.

The Ullmann polymerization of 2,5-dibromo-N-BOC-pyrrole has been used to produce a mixture of oligomers up to 25 repeating pyrrole units (Scheme 38). Subsequent preparative h.p.l.c. separation yielded oligomers up to 20 repeating units which were all characterized by u.v. and n.m.r. spectroscopy. Upon doping (I₂) of the deprotected pentamer, a black material with a conductivity of 100 S cm⁻¹ was obtained.

Oligoanilines

Just like on oligopyrroles, little research has been performed on oligoanilines. However, some contributions are worth mentioning. The first members of the oligoaniline series were prepared by Yoffe et al. who, starting from p-aminodiphenylamine, were able to isolate tri- (Figure 15, m = 3) and tetraaniline (m = 4) by an oxidative coupling/reduction sequence.

Figure 15  Yoffe’s trimer and tetramer as prepared by the oxidative coupling/reduction sequence.

About twenty years later Honzl et al. synthesized a series of phenyl-blocked oligoanilines. The dimer and trimer were prepared by a Sandmeyer type of elimination of the amino groups of Yoffe’s tri- and tetraaniline while...

Scheme 39 Honzl's route to phenyl-blocked oligoanilines

Figure 16 Oligophenylene and oligo(phenylene vinylene)s

the tetramer and hexamer were prepared by modifying Liebermann's method of preparing derivatives of 2,5-diaminoterephthalic acid (Scheme 39).

Although this method appeared not to be successful for the preparation of the octamer \( (m = 8) \), Wudl et al. were able to isolate this oligomer by coupling two equivalent of Yoffe's tetramer with one equivalent of dihydroxydihydroterephthalic acid and subsequent reduction with phenylhydrazine. From physical characterization it appeared that the different oxidation states of this octamer fully represent those of polyaniline.

The last series of oligoanilines worth mentioning were prepared by Strohriegi et al. Starting from diphenylamine and \( N,N' \)-diphenyl-1,4-phenylenediamine, they managed to prepare the \( N \)-phenyl substituted and phenyl blocked dimer, trimer and tetramer using \( n \)-BuLi, iodosobenzen and 1,4-diiodobenzen in combination with CuI as catalyst.

Other oligomers

Unsubstituted phenylene oligomers (Figure 16, 32) have been prepared and their conducting behaviour has been studied as a function of chain length and structural order after oxidation with potassium. From some of these oligomers the crystal structures have also been resolved. Substituted phenylene oligomers have been doped and the formation of radical anions and dianions have been studied by optical measurements; this work highlights the influence of substituents on the aromatic backbone.

All-trans \( \beta \)-carotene has been studied as a model polyene. Oligomers containing two to four double bonds have been reduced and the different states were studied with e.s.r. (radical anions) and n.m.r. (dianions). Substituted phenylene vinylene oligomers (33) have been synthesized by a sequence of Wittig reactions and their radical anions and dianions studied by optical methods. Recently, these oligomers have attracted considerable interest due to the electroluminescent properties of the parent polymer.

CONCLUSIONS

In the different sections above, we have presented a review on the wide range of conjugated polymers and oligomers. We have attempted to highlight the considerable advances in chemistry and materials science which have taken place in recent years, but within the framework of a historical perspective. The synthetic routes now available for most of the important polymers are well-established and yield materials with full control over molecular weight, polydispersity and regioselectivity. This high degree of perfection at the molecular level allows material scientists to study the ultimate properties of conjugated polymers using advanced physical characterization techniques and prototype devices. Furthermore, self-assembly of polymers, oriented samples and well-defined oligomers are present to investigate the structure–property relationship in detail. It should be stressed that many of the earlier studies have been performed with materials that, by the standards of today, are ill-defined and the results are not indicative for the ultimate properties of the materials under investigation. There are still some polymeric materials, e.g. polypyrrole, and functionalized PPVs, that need further systematic studies before they can be reliably and reproducibly prepared. Hopefully, we have managed to demonstrate that if the synthetic developments progress at the same
rate, it will not be long before the vast array of potential practical applications associated with these materials begin to materialize. It is expected that in the next few years we will have full control over most of the important molecular parameters and that chemists are able to study the mesoscopic ordering of the well-defined conjugated polymers; the next hierarchy in the development of advanced materials.

REFERENCES

20 Bloor, D., Korski, L., Stevens, G. C., Preston, F. H. and Ando, J. D. J. Mater. Sci. 1975, 10, 1678
33 Bloor, D., Korski, L., Stevens, G. C., Preston, F. H. and Ando, J. D. J. Mater. Sci. 1975, 10, 1678
37 Enkelmann, V., Leyrer, R. J. and Wegner, G. Makromol. Chem. 1979, 180, 1787
39 Young, R. J., Bloor, D., Butcher, D. N. and Hubel, C. L. J. Mater. Sci. 1978, 15, 62
45 Glaser, C. Ber. 1869, 2, 422
46 Eglinton, G. and Galbraith, A. R. Chem. Ind. (London) 1956, 737
47 Chodkiewicz, W. Ann. Chem. 1957, 2, 819
49 Viehe, H. G. Ber. 1959, 92, 3064
51 Bredas, J. L. and Heeger, A. J. Macromolecules 1990, 23, 1150
<table>
<thead>
<tr>
<th>Reference</th>
<th>Authors</th>
<th>Journal</th>
<th>Year</th>
<th>Pages</th>
</tr>
</thead>
<tbody>
<tr>
<td>442</td>
<td>Liebermann, H.</td>
<td>Ann.</td>
<td>1914</td>
<td>404, 272</td>
</tr>
<tr>
<td>449</td>
<td>Schenk, R., Huber, W., Schade, P. and Müller, K.</td>
<td>Chem. Ber.</td>
<td>1988</td>
<td>121, 2201</td>
</tr>
</tbody>
</table>