Hierarchical Conjugated Polymer Systems Prepared by Controlled Chain-Growth Polymerization

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HIERARCHICAL CONJUGATED POLYMER SYSTEMS
PREPARED BY CONTROLLED CHAIN-GROWTH
POLYMERIZATION

A Dissertation
Submitted to the Graduate Faculty of the
Louisiana State University and
Agricultural and Mechanical College
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Doctor of Philosophy

in
The Department of Chemistry

by
Chunwa Peter Kei
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ABSTRACT

π-Conjugated polymer materials may have a significant economic impact on society by providing means for designing affordable, flexible, and portable organic electronic devices. Their successful commercialization will depend on major scientific advancements, which will challenge human society to seek out ever more detailed and fundamental processes to command. Controlled polymerization affords such a power; allowing for the design of meticulous and precisely defined systems granting detailed insight into structure-property relationships in the polymer materials and bettering understanding of novel physical phenomena.

This dissertation primarily focuses on development and preparation of well-defined hierarchically organized macromolecular systems. The novel chain-growth polymerization methodologies described herein depended on gaining new fundamental insights into transition metal catalyzed controlled polymerization reactions, including many intriguing aspects of their catalytic and self-assembly processes.

For example, a general approach towards deeper mechanistic understanding and improving the controlled polymerization reactions for preparing conjugated polymer is presented; subsequent applying this knowledge resulted in an approach for a general synthesis of various classes of π-conjugated polymers precisely incorporating specific structural units. Preparation of such materials led to the observation of novel and unusual photophysical phenomena that exclusively appear in these unique polymer systems.

In addition, the photophysical properties and energy transfer processes occurring in nano- and mesoscale conjugated polymer donor-acceptor materials were found to depend on the non-equilibrium conformations of supramolecular assemblies forming in kinetically controlled catalyst-transfer polymerization.
CHAPTER 1. CONTROLLED POLYMERIZATION FOR PREPARING \(\pi\)-CONJUGATED MATERIALS: AN INTRODUCTION

1.1. Synthesis of Conjugated Polymers by Controlled Polymerization

Conjugated polymers are an important subject of studies in macromolecular chemistry. Generally, they are characterized by a polymer backbone of \(sp^2\)-hybridized carbon (or hetero) atoms that each contain electrons in the \(\pi\) orbital. The electrons are bound to a delocalized \(\pi\)-orbital system which allows them to move along the polymer chain. In 1977, Heeger, MacDiarmid and Shirakawa demonstrated electrical conductivity in polyacetylene thin films. It was a revolutionary discovery because it demonstrated unprecedented transport of energy in an organic polymer, which is a highly tunable medium. Up until that time, the idea of conductive polymers had not existed, and this event rapidly evolved into rich field of conjugated polymer research today.

Conjugated polymer research contains two dominating areas. The first is synthetic methodology in which development of controlled polymerization techniques and the expansion of the monomer scope is a primary goal. The second is applied functional systems, where the chemical, electrical, magnetic and optical properties of conjugated polymers are engineered into nanostructured materials with a specific purpose. These areas are markedly separated by how their chemical model systems are prepared, either by chain-growth polymerization in the synthetic methodology community, or step-growth polymerization used for preparing functional conjugated polymer materials.

Controlled polymerization is a special type of chain-growth polymerization. It is notably advantageous because it produces polymers with a pre-determined and readily controlled molecular weight and allows for achieving narrow polydispersity. Some controlled
polymerization reactions lack irreversible termination pathways and the polymer chain remains catalytically active after the monomers have been consumed. This property gives powerful control over the polymer chain end functionality. Controlled polymerization grants access to sequence defined polymers such as block co-polymers, graft co-polymers, dendrimers and hierarchical star polymers (Figure 1.1).

![Image of dendrimer-like star polymers](image)

**Figure 1.1.** Simplified representation of a 5th generation dendrimer-like star polymer prepared by anionic living polymerization (left) and its block-terpolymer structure (right). Reproduced with permission from Ref. 43. Copyright © 2014 American Chemical Society.

For conjugated polymers, controlled polymerization can be achieved with the reaction Catalyst-Transfer-Polymerization (CTP). Developments in CTP techniques have led to precisely defined structural topologies which are often needed for studying important properties of π-conjugated polymer systems. But unlike classical coupling reactions used in step-growth polymerization (Heck, Kumada, Murahashi, Negishi, Stille or Suzuki-Miyaura carbon-carbon coupling reactions), their chain-growth analogs using CTP require the catalyst to stay bound to the conjugated backbone through various π-associated complexes (Figure 1.2). This additional mechanistic demand makes it difficult to find matching catalyst/ancillary ligand combinations that are also compatible for preparing various classes of conjugated polymers.
Figure 1.2. DFT calculated free energies of various Ni$^0$ π-complexes with 2-bromothiophene. The “ring-walking” process shown here allows the catalyst to migrate over the polymer backbone and it is a mechanistic requirement for Catalyst-Transfer Polymerization. R = methyl. Reproduced with permission from Ref. 51. Copyright © 2018 Springer Nature.

On the other hand, using step-growth for making of π-conjugated polymers is a more mature technique. It has a wider synthetic scope and is applied ubiquitously for preparing conjugated polymer materials that require an elaborate chemical structure of the repeating unit. Such designer materials are often needed for today’s high performance organic electronic devices (Figure 1.3). Both areas of research are constantly striving to fill this discrepancy in synthetic knowledge.

Conjugated polymer research on controlled polymerization reactions strive to achieve the monomer scope used in modern conjugated polymer materials and applied conjugated polymer material ventures aspire for the control and narrow polydispersity afforded by controlled polymerization. The polydispersity index (PDI) is important parameter for optimizing conjugated polymer devices. It describes the distribution of polymer chain lengths within a material. Although they may be chemically similar, individual conjugated polymer chains of
various lengths are still distinct macromolecules; showing different conformational behavior, solid state packing energies and electronic structures. These small differences in chain lengths become critically important when the molecule is electronically coupled into an overall supramolecular potential energy landscape. An energy carrier may visit many individual chromophores in a conjugated polymer material during its excited-state lifetime.

Figure 1.3. Polymer acceptor material in high performance all polymer solar cell with a power conversion efficiency of 8.3%. Prepared by step-growth Stille coupling. Reproduced with permission from Ref. 52. Copyright © 2019 American Chemical Society.

Introducing lower or higher energy molecular orbitals that are significantly different than the average electronic energy levels within the material can hinder energy migration even at long distances from the location of the primary excitation. Therefore, the uniformity of the polymer material affects fundamental properties of conjugated polymer systems, and the development of robust controlled polymerization methods affording controlled polymer molecular weight and narrow polydispersity is an important technological challenge.
1.2. Recent Developments in Controlled Catalyst-Transfer Polymerization

1.2.1. The Initial Discovery and Mechanism of Kumada Catalyst-Transfer Polymerization

The development of controlled polymerization via CTP is dominated by the preparation of poly(3-alkylthiophene) (P3AT), poly(9,9-dialkylfluorenes) (PF) and poly(p-2,5-dialkylphenylene) or poly(p-2,5-dialkylxyphenylene) (PPP) and their block co-polymers.\textsuperscript{11,14} P3AT’s are the most prevalent class of conjugated polymers prepared by CTP. Their electron rich $\pi$-system can form effective associative complexes with phosphine and diimine-based Ni\textsuperscript{0} and Pd\textsuperscript{0} catalysts and facilitating the intramolecular catalyst-transfer process.\textsuperscript{62} In 2004, McCullough\textsuperscript{63} and Yozawa\textsuperscript{64} independently discovered the chain-growth nature of Ni-catalyzed polymerization of 2-bromo-5-chloromagnesio-3-hexylthiophene supported by 1,3-(diphenylphosphino)propane (dppp) ligation. The Kumada type reaction set the first example of preparing conjugated polymers by CTP. The mechanism of Kumada Catalyst-Transfer Polymerization (KCTP) follows a typical transition metal mediated carbon-carbon coupling catalytic cycle (Figure 1.4).

Figure 1.4. Kumada Catalyst-Transfer Polymerization catalytic cycle showing oxidative addition, transmetalation and reductive elimination steps. In structure I, a simplified Ni\textsuperscript{0}(dppp) $\pi$-associative complex is shown. Reproduced with permission from Ref. 53. Copyright © 2013 American Chemical Society.
First, the catalytically active $L_2\text{ArNi}^{\text{II}}\text{Br}$ species undergoes transmetalation with the organometallic Grignard monomer, and then reductive elimination increases the polymer chain length by one repeating unit. Afterwards, the catalyst must be able to migrate intramolecularly (by forming a series of $\pi$-associative complexes) to the carbon-halogen bond at the chain end. An efficient intramolecular catalyst-transfer is the vital step distinguishing chain-growth from the step-growth mechanism.\textsuperscript{65} Subsequently, oxidative addition into the carbon-halogen bond regenerates the catalytically active $L_2\text{Ar}_{n+1}\text{Ni}^{\text{II}}\text{Br}$ species. The mechanism only permits chain-growth to occur on polymer species that are bound to the transition metal. This process allows for the polymer molecular weight to be controlled using the monomer to catalyst molar ratio defined at the beginning of the reaction (known as feed ratio). KCTP can be used to reliably prepare P3HT with controlled chain-growth up to approximately 120 repeating units, with narrow polydispersity (PDI $<$ 1.2) and in excellent yields ($>$90\%).\textsuperscript{66}

Kumada type reactions formed the foundation of CTP research. But beyond the controlled polymerization of P3AT, very few ancillary ligand/catalyst combinations have shown reliable controlled chain-growth for other classes of conjugated polymers.\textsuperscript{12,14,34,36,65-66}

1.2.2. State-of-the-Art: Suzuki-Miyaura Catalyst-Transfer Polymerization

Yokozawa was also the first to report Suzuki-Miyaura Catalyst-Transfer Polymerization (SCTP). Much less is known about the reaction in comparison to KCTP and the complete mechanism has not been elucidated. In his initial 2007 report, Yokozawa et al. prepared poly(9,9-dioctylfluorene) and poly($p$-2,5-dibutylxylophenylene) using aryl boronic acid and pinacol ester monomers, and using PhPd(P'Bu$_3$)Br as an external initiator (Figure 1.5).\textsuperscript{67}
The resulting PF showed molecular weight up to 19 kDa and produced a modest PDI = 1.33. Likewise, preparation of PPP by this method gave modest results of $M_n = 11$ kDa and PDI = 1.53.

Later in 2010, Yokozawa et al. conducted a more detailed optimization study on the controlled preparation of PPP using 4-iodo-2,5-dihexyloxyphenylboronic acid (Figure 1.6). In the optimized conditions, he and co-workers were able to achieve a $M_n$ up to 21 kDa and PDI = 1.44-1.89 for PPP with greater than 60 repeating units. In that work, they adopted several strategies...
to improve the polymerization reaction. First, the reaction temperature was lowered to -20°C in order to control the temperature-dependent intermolecular chain-transfer process (involving dissociation of the π-associative complex). Second, they switched from aqueous Na$_2$CO$_3$ to CsF as a soluble in organic solvents base. Third, they increased the THF/H$_2$O ratio from 10:1 to 17:1 (v/v) based on the claim that excess amount of water was detrimental to the final $M_n$ and solubility of PPP. Lastly, they switched from bromoarene monomers to iodoarene boronic acid monomers. Polymerization reaction yields were not reported in that work.

Following Yokozawa’s reports, Hu et al. were able to improve SCTP preparation of PF using in-situ generated PhPd(‘Bu$_3$I as the external initiator and aqueous K$_3$PO$_4$ as base. In optimized conditions, they reported the preparation of PF with $M_n$ up to 31 kDa (however this significantly exceeded a theoretical $M_n$ of only 8.3 kDa), consistently low PDI 1.13-1.35 and good yields (>70%). Obtaining polymers via SCTP with higher $M_n$ than the theoretical $M_n$ is generally considered an indication of poor initiator efficiency. Where only a fraction of the initiator can undergo polymerization with the monomers, the effective $[M]_0/[I]_0$ ratio is increased proportionally and leading to higher molecular weight polymers.

Wang demonstrated the use of a non-phosphine containing ligand for SCTP and utilized Pd(IPr)(OAc)$_2$ as the precatalyst for controlled preparation of PF and P3HT. Pd CTP catalysts with NHC ligands (Figure 1.7) have longer catalyst lifetimes than their phosphine containing analogs.
Figure 1.7. Pd/NHC complexes used in SCTP reaction for preparing poly(9,9-dialkylfluorenes).

Also, such catalysts are suggested to facilitate the intramolecular catalyst-transfer process due to lower activation energies for the ring walking process. Nevertheless, the resulting polymers showed uncontrolled molecular weight and broad polydispersity. As such, Wang obtained PF with $M_n$ 69 kDa (PDI 1.56-1.78), ~70% yield and P3HT with $M_n$ 64 kDa (PDI 1.40-1.81) and >95% yields.

In typical Suzuki-Miyaura conditions, aryl boronic acids readily undergo hydrodeboronation; such monomer degradation results in significant loss of polymerization control.\textsuperscript{71} Carrillo et al. attempted to address this issue by utilizing $N$-methyliminodiacetic acid (MIDA) protected aryl boronic acid monomers.\textsuperscript{72} MIDA hydrolysis rate depends on the pH, temperature and water concentration; releasing cross-coupling active aryl boronic acid monomers at small steady concentrations over an extended time.\textsuperscript{72} The slow hydrolysis of MIDA boronates limits the transient concentration of ArB(OH)$_2$ and therefore decreases the rate of side reactions (Figure 1.8).
Figure 1.8. Synthesis of thiophene cross-coupled products using boronic acid (top) versus N-methyliminodiacetic acid (MIDA) protected boronate esters (bottom). Slow hydrolysis of MIDA boronates into their corresponding aryl boronic acids reduces protodeboronation and increases reaction yields. Reproduced with permission from Ref. 72. Copyright © 2014 American Chemical Society.

Carrillo et al. chose SPhos/Pd(OAc)$_2$ as the pre-catalyst for the preparation of P3HT. They conducted the polymerization reaction at a high temperature of 55-60°C, resulting in P3HT with broad polydispersity (PDI 2.7 – 3.1) in optimized conditions. Carrillo’s main focus was on improving the regioregularity of P3HT in Suzuki-Miyaura chain-growth polymerization via novel preparation of 5-bromo-4-hexylthiophene boronic acid MIDA ester using direct electrophilic borylation. Along this route, they were able to achieve a P3HT regioregularity of >97%. These authors also investigated the effect of various H$_2$O : monomer molar ratios on the SPhos/Pd catalyzed polymerization of 5-bromo-4-hexylthiophene boronic acid MIDA ester. They implied that the polymerization rate was accelerated using increasing amounts of H$_2$O (80 vs 40 equiv H$_2$O : monomer); an effect that is not discussed in detail in SCTP literature. However, this reaction caused severe hydrodeboronation and broadening of the PDI relative to reactions using 40 equiv H$_2$O : monomer. Carrillo was also one of the first to imply on the nature of the Pd catalytic cycle involved in the polymerization. They presumed that Suzuki-Miyaura polymerization of 5-bromo-
4-hexylthiophene boronic acid MIDA ester using SPhos/Pd system occurred via the boronate pathway of the Suzuki-Miyaura reaction.

Shortly after Carrillo et al.’s reported finding of the water effect on the course of SCTP, Yokozawa further commented on the role of H2O in Pd SCTP of triolborate thiophene monomers in anhydrous conditions.73 Similar to Carrillo, he also assumed that controlled SCTP proceeded via the boronate pathway of Suzuki-Miyaura reaction and that water was unnecessary for the reaction. But instead, Yokozawa found that water was essential for producing P3HT with narrow polydispersity. In anhydrous conditions, undesired intermolecular catalyst-transfer process dominated resulting in low quality P3HT (Mn = 3.6 kDa, PDI = 1.78 – 2.15). Yokozawa convincingly demonstrated that water, when present in the reaction mixture, facilitates intramolecular catalyst-transfer at room temperature, but he admitted that he could not conclude on the reason for its remarkable effects on the SCTP process.

In 2015, Hong reported similar results as Hu in the preparation of PF using p-TsOArX (X = I, Br) as the external catalytic initiator precursor.74 Both research groups were able to achieve low polydispersity, high yield, and yet the final molecular weight of the produced polymers was much higher than the theoretical Mn. Hong’s data demonstrated that in-situ initiation using aryl iodides were more efficient than by aryl bromides, and this was likely related to a finding in Yokozawa’s report showing that aryl iodide monomers were superior to aryl bromides in the SCTP reaction for preparing PPP. Hong et al. did not discuss the halogen's role in external initiator efficiency or how it related to average molecular weight of the polymers. Also, neither of Yokozawa’s systems were ideal (exhibiting uncontrolled molecular weight and broad polydispersity of the resulting polymers), so it would be difficult to explicitly elucidate the effect of varying halogens in the monomer or initiator. Hong et al. carried out multiple experiments using aryl iodide precursors
that resulted in better initiation than using aryl bromides (as indirectly indicated by the observed $M_n$ vs theoretical $M_n$). They obtained exceptionally clean MALDI-TOF spectra (Figure 1.9) of the PF polymers exhibiting only a single set of polymer chain distributions containing the external initiator. This clearly demonstrated that undesired chain-transfer process was essentially non-occurring in these polymerization conditions.

Figure 1.9. MALDI-TOF spectrum of poly(9,9-dihexylfluorene) prepared by Suzuki-Miyaura Catalyst-Transfer Polymerization. MALDI-TOF spectrum demonstrates a single set of polymer chains therefore indicating insignificant presence of chain-transfer products when polymerization is carried out using Pd$_2$(dba)$_3$/t-Bu$_3$P as the catalyst. Reproduced with permission from Ref. 74. Copyright © 2015 Royal Society of Chemistry.

Despite these findings, they chose to continue their study using aryl bromide (rather than aryl iodide) precursors combined with Pd$_2$(dba)$_3$/t-Bu$_3$P as the in-situ initiating species due to better control over the polymer chain ends. The combination of poor initiator efficiency and narrow polydispersity generally indicates the existence of a chemoselective polymerization process. Hypothetically speaking, either a fraction of the Pd species are deactivated at the beginning of the reaction (e.g. through Pd black nanoparticles formation or phosphine ligand oxidation), or the Pd species that are not initiated in the beginning of the reaction are forbidden from entering the
catalytic cycle throughout the reaction progress. However, catalyst degradation is a less likely hypothesis, considering that multiple independent groups reported this effect, and the chemical pathways for ligand dissociation are reversible. From a synthetic standpoint, Hu and Hong’s results were interesting because they showed two unexplored phenomena related to SCTP reactions: halogens in then initiator affecting the polymerization efficiency, and possible monomer deactivation in SCTP which results in narrow polydispersity. Many reports on CTP have demonstrated these effects, but the underlying process causing them has not been elucidated.

More recently, Hong also reported an interesting SCTP resulting in “ortho”-connected poly(5-alkyl-2,3-thiophene) (P5AT) which self-assembles into a helical structure (Figure 1.10). The highly twisted morphology of P5AT caused a large blue-shift in the fluorescence emission relative to P3HT and the nanostructure’s rigidity is apparent by narrow emission bands with sharp vibronic structures. The blue shifted emission is a result of a shorter average conjugation path length caused by kinking and twisting in the unusually connected polymer chain.

Figure 1.10. Helical poly(5-alkyl-2,3-thiophene) prepared by Suzuki-Miyaura Catalyst-Transfer Polymerization using Pd/NHC catalyst and K₃PO₄ as base at room temperature. Reproduced with permission from Ref. 75. Copyright © 2016 American Chemical Society.

In 2017, reports of SCTP using Buchwald ligands/pre-catalysts began to surface in literature. Exploring a large set of Buchwald ligands, Yokozawa identified Pd/AmPhos as a modestly suitable
system for preparing P3HT (Mn = 16.6 kDa, PDI = 1.35, however theoretical Mn = 3.3 kDa).\textsuperscript{76} As in previous cases, the polymers showed uncontrolled molecular weight likely due to an inefficient initiation process similar to the situation described in Hong and Hu’s reports. To mitigate this problem, Yokozawa tried utilizing an isolated ArPd(AmPhos)Br catalyst to achieve uniform initiation but found that it resulted in an even broader polydispersity relative to the case of the initiators he generated in-situ (where the catalyst was incubated with aqueous base prior to polymerization). The data implied that ArPd(AmPhos)Br was actually the pre-catalytic species; a potentially valuable finding. Yokozawa mentioned that the odd observation might have been due to the slow formation of palladium hydroxo complex (a key intermediate in the oxo-palladium pathway of the Suzuki-Miyaura reaction) and all but conducted the necessary experiments proving this. He supported his statement by showing that the in-situ generated catalysts provided a more uniform initiation, yielding a narrower polydispersity versus the isolated Pd halide complex.

Seo \textit{et al.} authored the second report utilizing MIDA boronates to control hydrodeboronation in SCTP reactions.\textsuperscript{77} They found that SPhos and RuPhos supported Pd SCTP produced P3HT and P3EHT polymers with controllable molecular weight, consistently low polydispersity and excellent yields (Mn up to 17.6 kDa, PDI >1.08, >90% yield). This report is remarkable because it represented the first robust and completely controlled chain-growth polymerization for the preparation of a conjugated polymer by SCTP. They also showed that the initiation time could be reduced by using electron deficient initiator precursors, and they controlled the initiator efficiency by utilizing commercially available 3\textsuperscript{rd} generation Buchwald pre-catalysts and aryl iodide precursors. Using this strategy, they successfully prepared all-conjugated P3HT-\textit{block}-P3EHT with molecular weight up to 9.2 kDa and polydispersity index of 1.18.
Recently, Yokozawa reported studies of the intramolecular catalyst-transfer of $t$Bu$_3$P/Pd catalysts over various functional groups using 2nd generation Buchwald pre-catalysts. Poly($p$-phenylenevinylene)s (PPVs) are an important class of conjugated polymers used in organic electronic devices which have historically been difficult to synthesize using KCTP because of irreversible binding of Ni based catalysts onto the carbon-carbon double bonds. Using Pd($t$Bu$_3$) based 2nd generation Buchwald precatalyst, Yokozawa demonstrated that these can efficiently undergo intramolecular catalyst transfer through stilbene when the phenylene moiety is functionalized with alkyloxy substituents in the ortho position. He used CsF/18-crown-6 as a base in 18:1 (v/v) THF/H$_2$O. Using the same catalytic system, he also demonstrated intramolecular catalyst-transfer over carbon-carbon triple bond, and nitrogen-nitrogen double bonds. Most interestingly, Yokozawa et al. reported the intramolecular catalyst-transfer over methylene, carbonyl, tertiary nitrogen, oxygen and sulfoxide units. Intramolecular transfer over methylene unit was particularly odd because it is a non-conjugated functional group lacking the ability to form $\pi$-associative complexes.
KCTP and SCTP are the main and most promising synthetic methods for preparation of conjugated polymers using controlled chain-growth polymerization. The reports discussed above comprise the most recent and impactful developments in SCTP research. From analysis of these reports (and corresponding discussion) it is apparent that many important questions are still left unanswered. What is the unequivocal role of water in SCTP and how does it enhance the polymerization rate and polydispersity as noticed in Carrillo and Yokozawa’s reports? Does SCTP proceed via the oxo-palladium or boronate pathway of the Suzuki-Miyaura cross-coupling reaction? What are the processes enabling halogens to play such a critical role in the initiating catalytic cycle of Pd SCTP as reported in Hu and Hong’s report? And is monomer deactivation responsible for the narrow polydispersity in reactions that have inefficient initiation? Chapter 2 of this dissertation will include a detailed discussion of the research project we undertook to address the underlying process at the center of these phenomena.

1.2.3. Other Notable Developments in CTP Controlled Chain-Growth Polymerization

Besides these and other avenues of research that are reviewed above, some unconventional techniques have been recently developed for expanding the possible monomer scope and π-conjugated material properties achieved by CTP. There were also some reports revealing neat mechanistic insights of the transmetalation step in SCTP reactions (still remaining a controversial topic).

The dihedral angle between repeating units is an uncommon but effective element for controlling the electronic structure of π-conjugated polymer systems. Sugiyasu provided\textsuperscript{81} proof-of-concept of a cyclopolymerization towards P3HT using KCTP to control the twisting of the conjugated polymer backbone. Controlling the dihedral angle between chromophores allows for
tuning excited state electronic energy levels and is valuable for the field of thermally activated delayed fluorescence.\textsuperscript{82}

Koeckelberghs’ group has been spearheading the alternative preferential oxidative addition mechanism for the controlled chain-growth of conjugated polymers without an associative complex and demonstrated a broader monomer scope than traditional CTP.\textsuperscript{83-85} Jagadesan \textit{et al.} reported on preparation of poly(\textit{p}-phenyleneethynylene) (PPE) by Sonagashira chain-growth polymerization using ArPd(\textsuperscript{t}Bu\textsubscript{3})Br as an external initiator with efficient initiation, resulting in excellent yields (>85\%) and narrow polydispersity (PDI = 1.13-1.21).\textsuperscript{86} Grisorio and Suranna developed Buchwald-Hartwig CTP of aryl amines yielding polymers with good polydispersity (PDI = 1.11.1-79) and excellent yields (>80\%).\textsuperscript{87}

On the mechanistic side, Li \textit{et al.} demonstrated base-free Suzuki-Miyaura coupling of aryl bromides with aryl boronic acids in water. They found that the irreversible catalytic cycle drives the reaction towards gradually increasingly acidity of the reaction medium in the absence of base. Using JohnPhos supported Pd(OAc)\textsubscript{2} as their catalytic system, they conducted Suzuki-Miyaura coupling with near quantitative yields and measured the changing pH as the reaction progressed. They found that Suzuki-Miyaura coupling occurred down to pH 3.0 (the acidity they found at the completion of the reaction in their conditions, but might possibly operate at lower pH), with yields above 90\%. But below pH = 6.0, the phosphine ligand became protonated and dissociated from the catalyst. This is particularly interesting because it implies the existence of an unconventional palladium catalytic pathway that proceeds without base. This essentially rules out both the traditional oxo-palladium and boronate pathways. Li \textit{et al.} proposed an unconventional oxo-palladium pathway, generating the palladium hydroxo intermediate through intramolecular deprotonation of a Pd aqua complex by lone pairs existing on a 2-heteroarylene reactive ligand.\textsuperscript{88,89}
In our own lab, we tried the SCTP of 2-bromo-3-hexylthiophene boronic acid MIDA ester (with RuPhos Pd G3 as precatalyst) using THF/acetate buffer (pH = 4.75) and found obtained a moderate polydispersity polymer (PDI = 1.32) with $M_n = 4.2$ kDa (theoretical $M_n = 6.1$ kDa) and yield = 49%. Acidity control in mediation of SCTP has never been demonstrated. It is possible that utilizing acidic conditions may be an alternative method for deterring hydrodeboronation due to the pH dependent Kuivila process. Obliterating the slow MIDA hydrolysis rate while reducing monomer degradation could allow to control the speed of polymerization reactions to a tremendous degree (Buchwald has shown that quantitative conversion between $L_2ArPdOH$ and $ArB(OH)_2$ can occur in under two minutes in similar conditions as used in SCTP reactions). A fast polymerization rate for making conjugated polymer materials is useful for generating kinetically driven nanostructures, a topic which is discussed in Chapter 3.

1.3. Energy Transfer in Conjugated Polymer Materials

1.3.1. Conformational Subunits Within Conjugated Polymer Chains

For a planar and infinitely long conjugated polymer chain isolated in vacuum, the electronic structure of the macromolecule may be relatively homogenous. But in real-life disordered conjugated polymer materials (such as in solution, thin-films and nanoparticles), the dynamic nature of structural conformations produces segmented chromophores within the polymer chain.\(^{58}\) These chromophores having different electronic structures (referred to as conformational subunits) combine into a statistical distribution of energy levels within the material which govern the thermodynamics of energy migration.\(^{90}\)

1.3.2. Excitons Quasiparticles

Photoexcitation spatially separates the electron from its nuclei and produces an electric field polarization around the local atomic lattice. Concomitant, the altered electron density in the excited
state molecular orbitals causes nearby structural changes to occur (such as altered bond lengths).\textsuperscript{58} These perturbations and the associated electron/nuclei pair are together referred to as an exciton. Singlet intrachain excitons have strongly bound excited states and quickly recombine to emit a photon. This species is mainly responsible for the fluorescence observed in conjugated polymer materials.\textsuperscript{91} There are also non-emissive excitons where the electron/nuclei polarons (simply termed “electron” and “hole”) are deterred from recombination by existing on different polymer chains.\textsuperscript{92} Excitons dissipate energy as they migrate from higher energy chromophores in the conjugated polymer system to lower energy chromophores, eventually being trapped in the low energy states where they can fluoresce as a singlet intrachain exciton, exist as non-emissive excited states, or relax non-radiatively.\textsuperscript{93}

1.3.3. Energy Transfer Pathways in $\pi$-Conjugated Materials

The energy transfer of photoinduced excited states through $\pi$-conjugated materials occur through two main processes. In systems where the energy donor and acceptor have an efficient molecular orbital overlap, an excited state electron can physically transfer from the higher energy chromophore to the lower energy one (exchange, or Dexter mechanism). This process is often associated with intrachain energy migration through covalent bonds.\textsuperscript{94} Energy can also be transferred through dipole – induced dipole interactions that do not require direct molecular orbital overlap and thus can occur through space (Förster mechanism).\textsuperscript{95}

Dexter and Förster pathways are often intertwined in energy transfer processes.\textsuperscript{96} Their individual contributions can only be deciphered in exceptionally ordered or specially designed conjugated polymer systems. Swager has demonstrated the relative dynamics of the Dexter and Förster energy migration in a rigid poly($p$-phenyleneethynylene) containing a low energy gap acridine orange (AO) chromophore.\textsuperscript{97} The linear framework of $p$-phenyleneethynylene (PPE)
conjugated backbone limits some intrachain Förster energy pathways (as the rigid rod-like structure of PPE would prevent them from folding and therefore bringing intrachain chromophores in close proximity). Swager et al. synthesized highly aligned PPE Langmuir-Blodgett films with low energy AO traps within the film with well-defined number of layers. The meticulously controlled anisotropic energy transfer from the multiple PPE layers showed that photogenerated excitons undergo very efficient interchain energy transfer perpendicular to the molecular axis. The energy migration kinetics were modeled mathematically to show efficient Förster mechanism taking place.

Energy migration by the Dexter mechanism is generally considered slower and less efficient due to poor electronic coupling between intrachain chromophores and the presence of defects. Schwartz et al. showed that intrachain energy transfer within PPV chains isolated in mesoporous silica composites was orders of magnitudes slower than the interchain process. In some of the early reports of theoretical treatment of the energy migration process, Dexter energy transfer was completely neglected due to its perceived slower energy transfer rate relative to Förster energy transfer. However, there were also some reports showing essential role of the through-bond energy transfer in conjugated polymer systems, and it is still a critical element to consider in designing conjugated polymer systems with certain optical and optoelectronic properties.

1.3.4. Other Excited State Species Affecting Energy Migration

Photoexcitation energy can be diverted into other species that interact strongly with the initially photogenerated excited state and can also affect energy migration processes. For example, singlet excitons in polythiophenes (PTs) can undergo a fast intersystem crossing (ISC) to form relatively stable triplet states due to the large energy difference between the T1 and S1 energy levels. This stabilization effect may lower the intrinsic photoluminescence of PT, but also can be utilized to
facilitate charge transport in electronic devices. Alternatively, an excited state can be stabilized by interacting with a proximate chromophore in its ground state. The resulting excited state complex (exciplex) and the stabilizing effect manifest themselves as a bathochromic shift in the emission band (relative to the emission band of an isolated excited state) in the fluorescence spectra. Charge-transfer complexes are also common in donor-acceptor systems and also produce low lying energy levels, which can trap and non-radiatively deactivate excited states often lowering quantum yields.

The difficulty of controlling energy migration is further complicated by the fact that the formation of conjugated polymer nanostructures is strongly influenced by synthetic methodology. Schwartz has demonstrated that MEH-PPV chains can aggregate showing both a concentration and solvent dependency. In relatively poor solvents for MEH-PPV, aggregates are pre-formed in solution and translate their low energy electronic states into the thin-film, making up a significant fraction of the total excited states and reducing the photoluminescence quantum yield.\textsuperscript{100}

1.3.5. Regioregularity in P3HT by KCTP and Its Impact On Energy Migration

Structural abnormalities affecting the electronic levels of conjugated polymer materials can also come from chemical defects on the polymer chain.\textsuperscript{101} In the case of P3HT prepared by KCTP, the organometallic Grignard monomer is generated in-situ via magnesium-halogen exchange (metathesis) reaction which can produce one of two possible regioisomers of 2,5-dibromo-3-hexylthiophene precursors. Although the “wrong” isomer of the Grignard monomer has been shown to be less reactive and not actively participating in polymerization process, it still has the ability to incorporate in the resulting conjugated polymer chain. In a case of such incorporation, asymmetric segments of 3-hexylthiophene repeating units with alkyl side chains would be facing each other (tail-to-tail) experiencing increased steric bulkiness between repeating units and causing torsion strain along the backbone.\textsuperscript{102} The kinking due to these torsional defects in P3HT chains
can cause changes in electronic energy levels and reduce the conjugation pathlength and exciton diffusion length, thus inhibiting charge and energy transfer in the π-conjugated system.¹⁰³

1.4. Practical Significance of Conjugated Polymer Systems – Recent Examples

Conjugated polymers possess a number of unique fundamental properties that make this class of materials capable to be used for addressing some modern economic and technological problems. The previous sections discussed the controlled chain-growth polymerization for the preparation of conjugated polymers and how their careful preparation affects energy migration processes. With meticulous care, these fundamental properties can be harnessed to do useful work. This section of the dissertation will highlight more recent technologically and economically relevant advances in conjugated polymer devices based on their unique properties. Conjugated polymers are traditionally integrated into organic photovoltaic cells, organic light emitting diodes and organic field effect transistors. These traditional applications have been reviewed comprehensively elsewhere.⁹,3⁰,3¹,3⁷,3⁸,5² The focus of this section is to demonstrate broad utility and novel applications of modern conjugated polymer material systems.

1.4.1. Flexible Energy Harvesting Organic Electronic Devices

Organic solar cells incorporating thin-films of conjugated polymers experience a tremendous rate of development. The physical performance including power conversion efficiencies being reported are increasing exponentially with time (Figure 1.12).³⁷ All-polymer solar cells (all-PSCs) have a particular niche for wearable electronics. Yan et al. recently reported a flexible ternary heterojunction solar cells reaching an incredible 14.06% power conversion efficiency (PCE), a record increase from 12.5% reported just earlier in 2019.¹⁰³ They utilized conjugated polymers as both donor and acceptor materials and also incorporated PC₇₁BM as additive to facilitate exciton dissociation and charge carrier generation at the donor-acceptor interface. Ternary devices rely on
recently developed charge transfer models\textsuperscript{104} to affect the open-circuit voltage using an energy cascade mechanism.

Figure 1.12. A plot of the maximum power conversion efficiency and number of publications reported each year for research into all-polymer solar cells showing an exponential increase up to 2018. Reproduced with permission from Ref. 37. Copyright © 2019 American Chemical Society.

In an interesting development, Allison and Andrew created a wearable thermoelectric generator on cotton substrate by vapor deposition of poly(3,4-ethylenedioxythiophene).\textsuperscript{105} When wore on the hand, it generates 20 mV. The unique material has a Seebeck coefficient of 16 µV K\textsuperscript{-1} and thermopower factor of 0.48 µWm\textsuperscript{-1}K\textsuperscript{-2}. Important research is directed towards decreasing overall cost of conjugated polymer devices. For example, Sun \textit{et al.} reported a low-cost polymer solar cell with 12.70% PCE using an efficient two step synthesis (overall yield 87\%) involving step growth polymerization based on Stille coupling reaction.
1.4.2. Electroluminescent Devices

Lin et al. prepared a deep-blue OLED device based on poly(9,9-diphenylfluorene) which self-assembles into a linear rod structure. The solubilizing groups prevent interchain exciton transfer and also serve as self-encapsulating layer, preventing water and oxygen from degrading the electroluminescent material. The flexible thin-film devices demonstrated a power efficiency of 1.73 cd A\(^{-1}\) and a turn on voltage of 4.55 V (Figure 1.13).

![Diagram of electroluminescent devices](image)

Figure 1.13. A self-encapsulating polyfluorene-based polymer light-emitting diode. PF was prepared by Yamamoto coupling polycondensation.

Conjugated polymers possess inherently high large extinction coefficients and low cytotoxicity which are beneficial attributes for biological analysis applications. Recently, Tang et al. developed an optofluidic laser using PPV as gain medium which showed a 70% quantum yield in THF solution.
1.4.3. Electro-Mechanical Devices: Conjugated Polymer Actuators

Polypyrroles (PPy) have excellent mechanical and electrical properties that make them suitable for generating physical force while maintaining robust stability in various media. Almost two decades ago, Jager developed micro-robotic arm using PPy thin films to move objects over a small distance.\(^\text{108}\) The mechanical action was driven by mass transport of anions in and out of the polymer network in aqueous media. Oxidation and reduction of PPy causes anions to flood into the polymer network in order to achieve a neutral charge density and thus creates swelling and shrinking that affects the length of the film parallel to the plane of the material. Recently, 3D printing techniques have allowed for creating PPy materials with complex geometries, enabling multidirectional force to be applied at precise positions and amplifying the anisotropic swelling or shrinking mechanism.\(^\text{109}\)

1.4.4. Other Notable Developments: Energy Storage and Spintronics Materials

Recently, London \textit{et al.} reported a donor-acceptor conjugated polymer material utilizing cyclopentadithiophene as a donor and thiadiazoloquinoxaline as acceptor and having an optical bandgap of 0.30 eV. It possesses a narrow transition between the $S_1$ and $T_1$ states with a difference of 0.0093 kcal/mol and causes the polymer to exist in high-spin ground state. The polymer can be dissolved in organic solvents and the spin-state is preserved upon forming amorphous thin-films.\(^\text{110}\) High-spin polymers have enormous potential as energy storage, quantum computing, and magnetic materials to be used in various fields. A supercapacitor retaining high capacitance and coulombic efficiencies over 6000 cycles was developed by Liao \textit{et al.} based on 3D aminoanthraquinone oligomeric networks. This represents an exceptional system for $\pi$-conjugated materials for energy storage.\(^\text{111}\) Lately, conjugated polymer materials have also made considerable advances as electrode materials for rechargeable batteries.\(^\text{112,113}\)
Figure 1.14. High spin conjugated polymer synthesized by step-growth Stille coupling. ref. 108. Reprinted with permission from AAAS.

1.5. Research Focus

The main focus of this dissertation is the development of precisely controlled conjugated polymer structures and nano- and mesoscale architectures through transition metal catalyzed controlled polymerizations. A review of recent developments in controlled chain-growth polymerization mechanisms for making conjugated polymers and the energy migration processes in \( \pi \)-conjugated materials underlines their importance in modern organic electronic devices.

In Chapter 2, rational, mechanism-based development of a universal system for externally initiated Suzuki-Miyaura Catalyst-Transfer controlled polymerization suitable for preparing poly(2,5-dialkylphenylenes), poly(9,9-dialkylfluorenes) and poly(3-alkylthiophenes) is described. An extensive discussion on the oxo-palladium and boronate pathways, as well as other mechanistic aspects of the polymerization catalytic cycle is supported by experimental evidence. Mechanism-based optimization studies on the polymerization reaction leading to PPP yielded discovery of the underlying role of the halide anion in the kinetics of the polymerization reaction. Using this mechanistic knowledge, the polymerization rate, \( M_n \), yield and polydispersity of the resulting conjugated polymers were substantially improved through the use of Ag(I) salts. Ag\(^+\) cations react
with Br\(^-\) to form AgBr during the polymerization and eliminate the anion’s antagonistic influence on the polymerization outcomes. The described method represents a novel and universally applicable approach to refining CTP reaction conditions and optimizing polymerization outcomes. By this way, CTP polymerization reaction can be rationally optimized synergistically or independently from any catalyst/ancillary ligand combination. In this work, Ag(I) mediated SCTP resulted in P3HT, PPP and PF with controlled molecular weight and consistently narrow polydispersity and excellent yields.

Chapter 3 describes the preparation of NIR fluorescent conjugated polymer nano- and mesoscale structures by kinetically driven self-assembly process using Kumada Catalyst-Transfer Polymerization. Perylenecarboxybisimide based external initiator was utilized to incorporate low energy chromophores into the structures, and the structurally dependent photophysical properties of these kinetically self-assembled systems were controlled by adjusting the reaction temperature and other reaction parameters. The photophysical properties of these kinetically stabilized nanostructures displayed a strong dependence on these parameters and their fluorescent emission could be tuned between 450-950 nm. Adjusting the reaction conditions also led to conjugated polymer nanostructures with varying conjugation path length, which allowed to attenuate the energy transfer related characteristics of the resulting materials. We also demonstrated that upon heating above the glass transition temperature, the metastable nano- and mesoscale structures rearranged into a common structural state as was observed from UV/vis absorbance and fluorescence spectroscopy. Detailed structural characterization using X-ray and neutron scattering experiments revealed a complex hierarchical structure of the systems.
1.6. References


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CHAPTER 2. SILVER(I) MEDIATED SUZUKI-MIYaura
CATALYST TRANSFER POLYMERIZATION: EVIDENCE OF THE
OXO-PALLADIUM PATHWAY

2.1. Introduction

Precision synthesis of complex macromolecular structures represents one of the major synthetic challenges still to be addressed in the polymer field. The ability to prepare macromolecular systems with structural exactness typical of the small-molecule synthesis is stimulated by the advanced practically useful properties as well as unique fundamental characteristics that the precise polymer systems may possess. Although scientists cannot yet rival Nature in the ability to prepare structurally precise polymers, recent developments in various living chain-growth polymerization methods started bringing such a possibility within reach. The field of conjugated polymers has recently experienced an impressive upswing towards the development of efficient controlled catalyst-transfer polymerization (CTP) protocols for the preparation of various classes of conjugated polymers (as described in Chapter 1). Conjugated polymers are materials of choice for a breadth of potential practical applications in electronic, optoelectronic and sensing devices due to tunability of their optical and electronic properties through manipulations of the polymers’ chemical structure. Whereas some classes of conjugated polymers (and block copolymers) can now be readily prepared via CTP methods, the real challenge remains in applying CTP for the preparation of more complex well-defined structures (e.g. where a specific structural unit would be precisely incorporated in the conjugated polymer chain – something that Nature routinely does for the case of biopolymers). For example, incorporation of a single electron deficient highly fluorescent perylenedicarboximide (PDCI) unit in a conjugated polymer chain has been previously demonstrated to dramatically alter the polymer’s optical properties, even when it was done in a non-precise, step-growth polymerization.\textsuperscript{2-3} One can argue that using an advanced CTP approach
to incorporate PDCI unit/s in conjugated polymer in a structurally precise fashion could deliver materials with novel and yet not well understood properties.\textsuperscript{4-5} Can such a process be implemented through a rational design of a conjugated polymer coupled with a precise synthesis? In this chapter, we focus on developing externally-initiated Pd-catalyzed Suzuki-Miyaura CTP as a functional group tolerant method of choice for precision polymerization to furnish various classes of conjugated polymers. Specifically, we describe how improving understanding of various mechanistic features related to the steps involved in Suzuki-Miyaura coupling can be used to enhance the efficiency of this CTP process and bring about the previously unavailable macromolecular synthetic capabilities, as well as conjugated polymer materials with novel properties that can be rationally fine-tuned through the precision polymer synthesis.

2.2. Results and Discussion

2.2.1. Design and Preparation of the Perylenedicarboximide External Initiator

Recently, Buchwald’s pre-catalysts have been shown to result in better initiation versus Pd('Bu\textsubscript{3}) based catalytic systems (which are the predominant Pd CTP catalytic system, making 40\% of the reports from 2008-2018).\textsuperscript{6-8} For example, Seo \textit{et al.} demonstrated that 3\textsuperscript{rd} generation SPhos and RuPhos Pd pre-catalysts are capable of controlled polymerization of 2-bromo-3-hexylthiophene boronic acid MIDA esters, leading to controlled molecular weight, excellent yields and consistently low polydispersity for poly(3-alkylthiophenes).\textsuperscript{9} The SCTP reaction using MIDA boronates is a remarkably complex chemical process involving more than a dozen chemical species, and many of them exist in dynamic equilibrium with each other. This requires the catalyst to be highly chemoselective in the presence of a variety of functional groups and conditions. The extraordinary performance of RuPhosPd\textsuperscript{0} in the SCTP reaction for preparing P3HT demonstrated that it is a robust system for all of these chemical environments. To efficiently obtain a chain-
growth polymerization method for preparing alternative conjugated polymers such as poly(p-phenylene) (PPP) and polyfluorene (PF), we chose to focus our efforts to optimize the well-established catalytic system of RuPhos supported Pd rather than to search for a new catalyst/ancillary ligand combination and having to optimize all the other parameters from scratch. Therefore, RuPhos supported Pd was chosen as the catalytic species installed on the external initiator. Also, utilizing PDCI as an initiator precursor allowed for incorporating an electron deficient chemical unit into the polymer backbone. Intramolecular catalyst-transfer over electron deficient monomers is still an obscure problem in CTP synthesis and this was a simple solution for overcoming this barrier. This strategy allowed to control the relative quantity of the acceptor molecule within the polymer chain and would allow us to study the structure dependent donor-acceptor photophysical properties in a precise manner. We also expected that the initiation process can be refined by isolating the external initiator as a solid, allowing for chemical purification and gravimetric measurements. External initiator 1 was prepared in one step by mixing a di-iodo derivative of PDCI (Figure 2.1) and stoichiometric ratio of RuPhos Pd G3 precatalyst in typical Suzuki-Miyaura conditions. After 1h at 50°C, the THF phase was precipitated into 1:5 (v/v) THF:hexanes, and then further purified by another reprecipitation step in the same solvent. Unreacted RuPhosPd⁰, carbazole (produced from the Buchwald pre-catalyst activation step) and the excess free RuPhos ligand are soluble in the reprecipitation solvent, whereas the external initiator is not. The resulting brown powder is stable in air-free environments, can be easily weighted and is soluble in THF. External initiator 1 displayed typical absorption and fluorescence spectra of PDCI with bithienyl functional groups attached to the 1,7 positions. The short wavelength absorption at $\lambda_{\text{max}} = 338$ nm derives from thiophene centered electronic transition.
Figure 2.1. Reaction scheme for the synthesis of the external initiator 1 (top). UV-Vis absorbance (black) and emission spectra (red) of the electron deficient PDCI molecule (bottom).

The long wavelength absorption at $\lambda_{\text{max}} = 613$ nm is attributed to $S_0$ to $S_1$ transition and the absorption peak at $\lambda_{\text{max}} = 460$ nm comes from $S_0$ to $S_2$ transition taking place along the perylene long molecular axis.\textsuperscript{11} Having bithiophene-substituted functionality in the 1,7 positions is
particularly advantageous for a functional Förster energy acceptor since both PF and PPP are expected to emit in the same UV-Vis region as the bithiophene absorbance band.\textsuperscript{12,13}

A strong overlap of the donor molecule fluorescence bands with the absorbance bands of the energy acceptor is a critical requirement for Förster energy transfer.\textsuperscript{14} Ideally, the design would allow for efficient long-range energy transfer from the excited state of PPP and PF to the low energy PDCI chromophore.

**2.2.2. SCTP Monomer Screening Studies Using Suzuki-Miyaura Coupling Experiments**

A robust intramolecular catalyst-transfer process is vital for CTP and its success is sensitive to the specific combination of the catalyst, ancillary ligand and monomer identity. As such, we explored the potential SCTP monomer scope of RuPhos Pd G3 by studying the Suzuki-Miyaura coupling of phenylboronic acid with 1,4-dibromo-2,5-dialkylbenzenes, 9,9-dialkyl-2,7-dibromofluorenes, 2,5-dibromo-3-alkylthiophenes and several related dihaloarenes.

Following the oxidative addition of $L_2\text{Pd}(0)$ onto the carbon-bromine bond and subsequent exchange of bromide ligand by hydroxide, $\text{BrArL}_2\text{PdOH}$ is expected to react with phenylboronic acid to produce a mono-substituted product $\text{M1}$ (Figure 2.2). If intramolecular catalyst-transfer occurs after, the subsequent oxidative addition, transmetalation and reductive elimination cycle will produce a disubstituted arene $\text{M2}$. An absolute selectivity for $\text{M2}$ indicates an ideal candidate for the controlled-polymerization studies. However, if $L_2\text{Pd}^0$ is not able to undergo intramolecular catalyst-transfer, only $\text{M1}$ is expected. Because the intermolecular catalyst-transfer process (leading to chain-transfer and loss of controlled-chain growth character) is accelerated at higher reaction temperatures, the monomer screening reactions are conducted at room temperature and only dihaloarenes that can exclusively form $\text{M2}$ under SCTP conditions are considered for further polymerization studies. Semi-quantitative analysis using liquid-chromatography/total ion count...
mass spectrometry (LC-MS) allowed for fast screening of 13 dihaloarenes. Ten of them demonstrated exclusive conversion to M2 with no M1 detected by mass spectrometry (Figure 2.2).

Figure 2.2. Summary of monomer screening studies showing dihaloarenes exclusively forming the intramolecular catalyst-transfer product M2 in Suzuki-Miyaura Catalyst-Transfer Polymerization reaction conditions using RuPhos Pd G3 as the pre-catalyst.

Along with several 1,4-dibromo-2,5-dialkylbenzenes (Figures 2.6 and 2.7), 9,9-dialkyl-2,7-dibromofluorenes (Figures 2.3 and 2.4) and 2,5-dibromo-3-alkythiophenes (Figures 2.10, 2.11 and 2.12), 3,9-dibromoperylene (Figure 2.9), 2,5-di(5'-bromo-2-thienyl)-3,4-ethylenedioxythiophene
(Figure 2.13) and 4-iodo-1-bromobenzene (Figure 2.14) showed exclusive formation of M2 and were identified to be promising candidates for Suzuki-Miyaura controlled-chain growth polymerization studies using RuPhos Pd G3 as the pre-catalyst. Expectedly, 2,5-divinyl-substituted 1,4-dibromobenzene (Figure 2.8) included in this experiment did not show strong intramolecular catalyst-transfer capabilities even at room temperature and thus was excluded from controlled chain-growth polymerization studies. Vinyl-substituents can form irreversible \( \pi \)- associative complexes with Ni and Pd based catalysts and inhibit controlled chain-growth. The exact mass of the M1 product resulting from coupling between 3,9-dibromoperylene and phenylboronic acid coincided with the mass spectrometer noise ions but no M1 products were detected above this background (Figure 2.9). 2,7-dibromo-9,9-di(6’-methoxyhexyl)fluorene produced both M1 and M2 products (Figure 2.5) and implies that either increased steric bulk of the solubilizing groups or the ether moiety interferes with the chain-walking process (relative to the excellent result of 2,7-dibromo-9,9-dialkylfluorene in Figure 2.4).

In Pd catalyzed SCTP reactions, oxidative addition onto aryl iodides are generally faster and more efficient than oxidative addition onto aryl bromides, and 2,5-methoxy-1,4-diiodobenzene was expected to undergo efficient intramolecular transfer with RuPhos supported Pd.\(^{15}\) Although 2,5-dimethoxy-1,4-diiodobenzene showed good selectivity (implying an efficient intramolecular catalyst transfer process), the low ion intensities of the M2 product relative to the starting material suggested an inefficient coupling reaction (Figure 2.15) in comparison to 2,5-dialkyl-1,4-dibromobenzene (Figure 2.7).
Figure 2.3. Reaction scheme of the Suzuki-Miyaura coupling between 2,7-dibromofluorene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.4. Reaction scheme of the Suzuki-Miyaura coupling between 9,9-dihexyl-2,7-dibromofluorene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.5. Reaction scheme of the Suzuki-Miyaura coupling between 9,9-di(6’-methoxyhexyl)-2,7-dibromofluorene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.6. Reaction scheme of the Suzuki-Miyaura coupling between 2,5-dihexyl-1,4-dibromobenzene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.7. Reaction scheme of the Suzuki-Miyaura coupling between 2,5-dimethyl-1,4-dibromobenzene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.8. Reaction scheme of the Suzuki-Miyaura coupling between 2,5-divinyl-1,4-dibromobenzene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.9. Reaction scheme of the Suzuki-Miyaura coupling between 3,9-dibromoperylene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.10. Reaction scheme of the Suzuki-Miyaura coupling between 2,5-dibromothiophene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.11. Reaction scheme of the Suzuki-Miyaura coupling between 3-methyl-2,5-dibromothiophene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.12. Reaction scheme of the Suzuki-Miyaura coupling between 3-hexyl-2,5-dibromothiophene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.13. Reaction scheme of the Suzuki-Miyaura coupling between 2,5-di(5’-bromo-2’-thienyl)-3,4-ethylenedioxythiophene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.14. Reaction scheme of the Suzuki-Miyaura coupling between 1-bromo-4-iodobenzene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Figure 2.15. Reaction scheme of the Suzuki-Miyaura coupling between 2,5-methyloxy-1,4-diiodobenzene and phenylboronic acid using RuPhos Pd G3 as pre-catalyst (top). Extracted ion chromatogram of the reaction mixture after 72h at rt (bottom).
Even though 2,5-dialkylxylophenylboronic acid monomers are typically used in SCTP reactions, the unexpected poor performance of 2,7-dibromo-9,9-di(6’methoxyhexyl)fluorene and 2,5-dimethyloxy-1,4-diiodobenzene (relative to their non-oxygenated analogs) indicated that oxygen atoms that are directly bonded to the π system or oxygen atoms installed on the solubilizing side chain have negative impacts on intramolecular catalyst-transfer mechanism.

2.2.3. Controlled Preparation of Poly(p-phenylene): Possible Theoretical Catalytic Pathways

The small molecule model experiments were conducted at room temperature to observe the highest potential for intramolecular catalyst-transfer. However, the initial SCTP polymerization studies were conducted at 45°C. This was necessary to accommodate for the slow hydrolysis rate of N-methyliminodiacetic acid (MIDA) boronates at ambient temperatures.

Yokozawa and co-workers first reported the preparation of poly(p-phenylenes) (PPPs) through SCTP of 4-iodo-2,5-dialkoxyphenylboronic acid, and using Pd(bb3)2 as the pre-catalyst.16 In this study, we also found that 1,4-diiodo-2,5-dialkoxybenzene could undergo intramolecular catalyst-transfer at room temperature using RuPhos Pd G3 as the precatalyst. However, it’s MIDA boronate analog did not produce an efficient SCTP polymerization reaction at 45°C (Figure 2.16).

![Figure 2.16](image)

Figure 2.16. Suzuki-Miyaura catalyst-transfer polymerization of 4-iodo-2,5-dialkxyoxyphenylboronic acid MIDA ester at 45°C. aPolymerization reaction conditions: 1 (3.3 µmol, 1 equiv), 4-iodo-2,5-dihexoxyphenylboronic acid MIDA ester (89 µmol, 27 equiv), K3PO4 (0.77 mmol, 3M), THF/H2O (30:1). bAbsolute molecular weight was measured by NMR. cMn/Mw was determined by GPC calibrated using polystyrene standard and with THF as eluent at 40°C.
The inability of 4-iodo-2,5-dihexyloxyphenylboronic MIDA ester to undergo efficient controlled chain-growth at 45°C is also in agreement with Yokozawa’s report, where they used 4-iodo-2,5-dihexyloxyphenylboronic acid pinacol ester and were compelled to lower the reaction temperature to -20°C in order to optimize the reaction. These results confirm that alkyloxy substituted phenylenes are not suitable for SCTP in these conditions.

All of the remaining phenylene monomers identified from the model reaction studies as potential candidates for externally initiated SCTP contain alkyl substituents as solubilizing groups. 4-bromo-2,5-dihexylphenylboronic acid MIDA ester 2 was chosen as a representative monomer for preparing poly(p-2,5-dialkylphenylenes) due to its expected solubility in THF/H₂O and ease of synthesis. Longer alkyl chain 1,4-dibromo-2,5-dialkylbenzenes are prone to precipitate during the lithium halogen exchange reaction at -78°C and were found to produce lower yields than 1,4-dibromo-2,5-dihexylbenzene.

Previously, Carrillo and co-workers reported that the theoretical stoichiometric ratio of H₂O to K₃PO₄ to MIDA boronate monomer in the SCTP of 5-bromo-4-hexylthiophene boronic acid MIDA ester 3 was 3:3:1. They proposed that two equivalents of base are needed to hydrolyze MIDA protecting group, and the third was used to form the boronate anion under the assumption that the transmetalation proceeded through the reaction between L₂ArPdBr and BrArB(OH)₃K (the boronate pathway of Suzuki-Miyaura reaction). However, Burke later published comprehensive mechanistic studies demonstrating that the slow hydrolysis of MIDA boronates in THF and aqueous K₃PO₄ occurs predominately with H₂O and not hydroxide as the nucleophile in both hydrolysis steps of the reaction. Therefore, these stoichiometries are not likely to be correct. Since Carrillo’s report, there has also been strong evidence that the Suzuki-Miyaura coupling between L₂ArPdOH and neutral phenylboronic acid (the oxo-palladium pathway) is orders of
magnitude faster than the reaction between $L_2\text{ArPdBr}$ and anionic $\text{BrArB(OH)}_3\text{K}$, suggesting when the two species exist in equilibrium that the oxo-palladium pathway would be dominant in the Pd-catalyzed Suzuki Miyaura reaction between aryl halides and phenylboronic acid.\textsuperscript{19-23}

Based on these reports, the minimum theoretical stoichiometric ratio between $\text{H}_2\text{O} : \text{K}_3\text{PO}_4 : \text{BrArBMIDA}$ in the Pd catalyzed SCTP of $2$ should be $4:4:1$ via the oxo-palladium pathway. One stochiometric equivalent of base is needed to form the active palladium hydroxo complex $L_2\text{ArPdOH}$ in the transmetalation step, and three are required to neutralize the side products produced per CTP catalytic cycle with MIDA boronate monomers. The side product $N$-methyliminodiacetic acid has a $pK_{a1}$ of 2.81 and $pK_{a2}$ of 10.18. Boric acid has a $pK_a$ of 9.24, whereas the conjugate acid of $\text{K}_3\text{PO}_4$ has a $pK_a$ of 12.67. Based on $pK_a$ values, the neutralization of these side products occurs almost quantitatively in the presence of excess $\text{K}_3\text{PO}_4$.

Additionally, based on findings by Amatore and co-workers, $\text{OH}$ catalyzes the reductive elimination of $L_2\text{ArPdArBr}$ via a pentacoordinate $\text{Pd}^{II}$ complex, and thus a catalytic concentration of $\text{OH}$ should be implemented in the reaction design to for transmetalation to be rate limiting.\textsuperscript{21}

To differentiate between the two palladium pathways, $2$ was polymerized in typical SCTP conditions using 3.5 equivalents of base (entries 1-4, Table 2.1). A $3.5 : 1$ molar ratio of $\text{K}_3\text{PO}_4 : 2$ was chosen in consideration of the boronate pathway and its stoichiometry, an excess of 0.5 equivalents of $\text{K}_3\text{PO}_4$ was employed as a catalyst for reductive elimination as proposed by Amatore. Carrillo also found that a large excess of water was necessary for successful Pd catalyzed SCTP of 5-bromo-4-hexylthiophene boronic acid MIDA ester. In the following experiments (entries 1-4, Table 2.1), the molar ratio of $\text{H}_2\text{O}$ to $2$ was 66. This amount is consistent with the 40-80 molar ratio of $\text{H}_2\text{O}$ to monomer proposed by Carrillo and co-workers. Theoretically, if the boronate pathway was dominant, this stoichiometry should lead to an efficient reaction.
Alternatively, if the oxo-palladium pathway is the dominant catalytic pathway, then the expedient neutralization of K₃PO₄ is expected. The catalytic amount of –OH would also be exhausted, leading to a critical reduction in reaction rate before all the monomers are consumed. Ultimately, if the SCTP of 2 occurs via the oxo-palladium pathway, the sub-stoichiometric amount of base is expected to lead to a characteristic decrease of the Mₙ and yield via suppression of the polymerization rate relative to hydrodeboronation. Utilizing a 3.5 to 1 molar ratio of K₃PO₄ in typical SCTP conditions⁹ utilized for the polymerization of 3, the polymerization of 2 resulted in a low molecular weight, yield and moderate polydispersity. (Table 2.1, entry 1)

Table 2.1. Externally Initiated SCTP of 4-bromo-2,5-dihexylphenylboronic acid MIDA ester 2 using RuPhos supported Pd external initiator 1.

<table>
<thead>
<tr>
<th>entry</th>
<th>THF (mL)</th>
<th>H₂O (mL)</th>
<th>Mₙ⁺ (PDI)</th>
<th>yield¹ (%)</th>
<th>X_H₂O</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>8</td>
<td>0.15</td>
<td>1.2k (1.31)</td>
<td>5.6%</td>
<td>0.08</td>
</tr>
<tr>
<td>2</td>
<td>4</td>
<td>0.15</td>
<td>1.8k (1.30)</td>
<td>8.0%</td>
<td>0.14</td>
</tr>
<tr>
<td>3</td>
<td>2</td>
<td>0.15</td>
<td>3.0k (1.21)</td>
<td>15.6%</td>
<td>0.25</td>
</tr>
<tr>
<td>4</td>
<td>1</td>
<td>0.15</td>
<td>3.6k (1.19)</td>
<td>46.6%</td>
<td>0.40</td>
</tr>
</tbody>
</table>

⁹ Reaction conditions: 1 (3.3µmol, 1 equiv), K₃PO₄ (0.45mmol, 3.0M), [2]₀/[1]₀ = 38. ¹ Absolute molecular weight was determined by quantitative NMR. ² Mₚ/Mₙ was determined by GPC calibrated using polystyrene standard and with THF as eluent at 40°C. ³ Isolated yield after precipitation in acetone.
The utilization of MIDA boronates as protecting groups is beneficial because the slow release of BrArB(OH)₂ limits the overall rate of hydrodeboronation. In highly basic conditions, the buildup of transient unreacted BrArB(OH)₂ would readily result in hydrodeboronation via the Kuivila mechanism and would theoretically reduce both the Mₙ and yield in a correlated fashion via monomer degradation.¹⁹-²¹ Vice versa, an increase in polymerization rate relative to hydrodeboronation would improve Mₙ and yield in the same correlated fashion. The characteristic low yield and Mₙ observed for entry 1 (Table 2.1) implied a slow polymerization rate relative to MIDA hydrolysis. To test this theory and to obtain better results, we conducted the next series of reactions with increased concentration in the THF phase in order to accelerate the polymerization (entries 2-4, Table 2.1). This resulted in the expected concurrent improvement of the molecular weight of PDCI-PPP up to 3.6 kDa, a narrowing of the polydispersity index from 1.31 to 1.19 and large increase in yield from 5.6% to 46.6%.

In the externally initiated polymerization of 2, both oxidative addition and reductive elimination are assumed to be unimolecular processes. Since increasing the solute concentration in the THF phase of entries 1-4 resulted in accelerated reaction rate relative to hydrodeboronation (as indicated by higher molecular weight and yield), transmetalation was conjectured to be rate limiting.

Reducing the THF volume affects the kinetics of the reaction in two major ways. First, the concentrations of catalytic Pd centers and monomers are increased. This would theoretically increase the rate of polymerization up to 64 times from entry 1 to entry 4 (assuming that rate limiting transmetalation is an ideal bimolecular reaction between active catalytic species and monomer). Second, the lowering of THF volume also increases the water molar fraction which appears to be more impactful on the polymerization reaction. The water molar fraction (X_H₂O) dictates water activity in the THF phase and increasing the water activity accelerates the MIDA
hydrolysis reaction.\textsuperscript{18,22} Clearly, increasing the rate of MIDA hydrolysis relative to rate of polymerization would exacerbate hydrodeboronation, leading to lower yields and thus cannot be the reason for the improved results. Most critically, water activity strongly influences the equilibrium position of the ligand exchange reaction between $L_2$ArPdOH and $L_2$ArPdBr.\textsuperscript{23} In the oxo-palladium pathway, $L_2$ArPdOH is reactive and $L_2$ArPdBr is not. Because of this, their dynamic equilibrium concentration may be determining the average polymerization rate of Pd catalyzed SCTP reactions of boronic acid containing monomers. To understand the phenomenon observed in entry 1-4 (Table 2.1) and to conclude on the palladium catalytic pathway, a full understanding on how $X_{H2O}$ and water activity affects the $M_n$, yield and polydispersity of PDCI-PPP prepared by SCTP is required.

2.2.4. Two Opposing Roles of Water Molar Fraction on the Oxo-Palladium Pathway

Kinetic studies by Amatore, Hartwig and others of base mediated Pd-catalyzed Suzuki-Miyaura reactions between aryl bromides and phenylboronic acid employing THF/H$_2$O as the solvent (between 50:1 and 10:1 (v:v) THF/H$_2$O ratio) and utilizing a 1 : 1 molar ratio of $L_2$ArPdOH and Br$^-$, an increase of $X_{H2O}$ has been reported to drive $K_{eq}$ from 1 up to 9.3 favoring the formation of unreactive $L_2$PdArBr as the dominate palladium species in solution.\textsuperscript{23-28}

In the case of SCTP of bromo-substituted arene monomers, the solution concentration of Br$^-$ is directly proportional to the average degree of polymerization due to Br$^-$ being a side product of the catalytic cycle. At the beginning of the reaction (low DP) where the concentration of Br$^-$ is low, implementing a high $X_{H2O}$ in the reaction design is theoretically expected to increase the concentration of -OH in the THF phase and would drive the formation of the active $L_2$ArPdOH even with a relatively high $K_{eq}$ (determined by $X_{H2O}$ shown by Hartwig) between the ligand exchange reaction of $L_2$ArPdOH to $L_2$ArPdBr; thus increasing the polymerization rate at the
beginning of the reaction. However, increasing concentrations of Br\(^-\) due to catalytic turnover would cause a significant buildup of unreactive \(L_2\text{ArPdBr}\) and attenuating the average polymerization rate as the reaction progresses. This dynamic equilibrium concentrations of \(L_2\text{ArPdOH}\) and \(L_2\text{ArPdBr}\) throughout the reaction progress would also result in polymer chain ends with a distribution of reactivities and conceivably result in broadening or narrowing of polydispersity as a function of \(X_{\text{H2O}}\) in basic conditions such as observed in Table 2.1, entry 1-4. Water’s dual role in the ligand exchange reaction between the two palladium species may also be the reason for Carrillo and co-workers’ finding that an excess of water is necessary for efficient polymerization of 3 using Pd supported by SPhos. But, at the same time, they found that further increasing \(X_{\text{H2O}}\) has harmful effects on the SCTP reaction. Overall, water activity can enhance and inhibit the average rate of transmetalation in Pd catalyzed SCTP reactions. To test this theory and contrast the effects of THF concentration versus water molar fraction on the \(M_n\), yield and polydispersity, a series of reactions were conducted at varying water and THF volumes in the SCTP of 2 using 1 as the external initiator. (Table 2.2)

<table>
<thead>
<tr>
<th>entry</th>
<th>THF (mL)</th>
<th>H(_2)O (mL)</th>
<th>(M_n^{b}) (PDI(^c))</th>
<th>yield(^d)</th>
<th>(X_{\text{H2O}})</th>
<th>([\text{Pd}]) (µM)</th>
<th>([\text{K}_3\text{PO}_4]/[2])</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>8</td>
<td>0.25</td>
<td>3.6k (1.27)</td>
<td>8.0%</td>
<td>0.12</td>
<td>0.82</td>
<td>6.00</td>
</tr>
<tr>
<td>6</td>
<td>1</td>
<td>0.15</td>
<td>7.0k (1.19)</td>
<td>59.2%</td>
<td>0.40</td>
<td>6.56</td>
<td>7.00(^e)</td>
</tr>
<tr>
<td>7</td>
<td>8</td>
<td>0.42</td>
<td>3.1k (1.17)</td>
<td>20.0%</td>
<td>0.19</td>
<td>0.82</td>
<td>9.00</td>
</tr>
<tr>
<td>8</td>
<td>2</td>
<td>0.42</td>
<td>7.6k (1.16)</td>
<td>61.7%</td>
<td>0.48</td>
<td>3.28</td>
<td>9.00</td>
</tr>
<tr>
<td>9</td>
<td>4</td>
<td>0.63</td>
<td>6.8k (1.15)</td>
<td>70.1%</td>
<td>0.41</td>
<td>1.64</td>
<td>15.00</td>
</tr>
<tr>
<td>10</td>
<td>8</td>
<td>0.63</td>
<td>5.7k (1.16)</td>
<td>44.5%</td>
<td>0.26</td>
<td>0.82</td>
<td>15.00</td>
</tr>
<tr>
<td>11</td>
<td>16</td>
<td>0.63</td>
<td>3.7k (1.15)</td>
<td>27.1%</td>
<td>0.15</td>
<td>0.42</td>
<td>15.00</td>
</tr>
</tbody>
</table>

\(^e\)Reaction conditions: 1 (3.3µmol, 1 equiv), \(\text{K}_3\text{PO}_4\) (3.0 M), \([2]/[1]_0 = 38\). \(^b\)Absolute molecular weight was determined by quantitative NMR. \(^c\)\(M_w/M_n\) was determined by GPC calibrated using polystyrene standard and with THF as eluent at 40°C. \(^d\)Isolated yield after precipitation in acetone. \(^e\)6.0M \(\text{K}_3\text{PO}_4\).
Entry 5 is notably different than entry 1 because of the higher base stoichiometry (6:1 $\text{K}_3\text{PO}_4 : 2$ in entry 5 vs 3.5:1 in entry 1) and slightly higher $X_{\text{H}_2\text{O}}$ (0.12 in entry 5 vs 0.08 in entry 1) which resulted in small improvements in molecular weight, polydispersity and yield after reprecipitation in acetone (3.6 kDa, PDI = 1.27, 8.0% yield). In this case, the improvements of entry 5 vs entry 1 are not likely due to increased $X_{\text{H}_2\text{O}}$ because entry 2 (with higher $X_{\text{H}_2\text{O}}$ of 0.14 vs 0.12 in entry 5) did not show these improvements. We repeated the most optimized reaction (entry 4) in the series of entry 1-4 with identical conditions except using 7:1 equivalents of $\text{K}_3\text{PO}_4 : 2$ instead of 3.5:1 (Table 2.2, entry 6). This resulted in doubling of the molecular weight up to 7.0 kDa and a significant increase of yield from 46.6% to 59.2%. These improvement in yield and molecular weight of entry 5 and 6 vs entry 1 and 4 respectively demonstrate that 3.5 molar equivalents of base relative to monomer is not optimal for the Pd catalyzed SCTP of 2; rebutting the theoretical stoichiometry for the boronate pathway and supporting the oxo-palladium mechanism where at least 4 equivalents of $\text{K}_3\text{PO}_4$ relative to 2 is necessary for an efficient polymerization reaction. In conditions where the stoichiometric amount of base is insufficient, the low $M_n$ and yield are hypothetically attributed to the formation of $\text{L}_2\text{ArPdBr}$ due to low $\text{–OH}$ concentration.

Using at least 6 : 1 ratio of $\text{K}_3\text{PO}_4 : 2$, increasing $X_{\text{H}_2\text{O}}$ from 0.12 to 0.26 by increasing the water volume and keeping THF volume constant (entry 5, 7 and 10) lead to characteristic improvements of $M_n$ from 3.6 kDa to 5.7 kDa, polydispersity from 1.27 to 1.16 and yield from 8.0% to 44.5%. Likewise, increasing the $X_{\text{H}_2\text{O}}$ from 0.19 to 0.48 (entry 7 and entry 8) by changing the THF concentration while keeping volume of water constant also showed correlated improvements of the $M_n$ from 3.1 kDa up to 7.6kDa, polydispersity from 1.17 to 1.16 and yield from 20.0% to 61.7%. These results show that the improved polymerization reaction from entries 1-4 is not due to the increased concentration of solutes in the THF phase, but from increasing the water molar
fraction. This refutes the hypothesis that reaction between $L_2\text{ArPdOH}$ and $\text{BrArB(OH)}_2$ due to low concentration in the THF phase is limiting the overall rate of polymerization via transmetalation. It should be noted that the molar quantities of $K_3\text{PO}_4$ in entries 5-11 must be increased as the volume of water increases to maintain the concentration of 3.0 M $K_3\text{PO}_4$ in the aqueous phase. Additional to its role as a base, $K_3\text{PO}_4$ also acts as a strong electrolyte which lowers the water activity in THF, slowing down the rate of MIDA hydrolysis that is vital for an efficient SCTP reaction.

To gain further insight into the behavior of rate limiting step, a series of SCTP polymerization reactions of 2 was conducted at relatively high $X_{\text{H}_2\text{O}}$ (between 0.15 and 0.41) and varying amounts of THF up to 16mL (entries 9-11). Even at this dilute concentration of 0.42 µM of palladium centers (16 mL THF, entry 11), the polymerization reactions produced remarkable results for the controlled synthesis of poly($p$-2,5-dihexylphenylene) ($M_n = 3.7\text{kDa}$, $\text{PDI} = 1.15$, yield = 27.1%) and adhering to the theory proposed herein. Entries 9-11 displayed a decrease in yield and $M_n$ with increasing THF volume (and decreasing $X_{\text{H}_2\text{O}}$) and again demonstrated how water molar fraction is fundamentally tied to the reagent stoichiometries and polymerization results. Entries 5-11 show that changing the quantities of THF and H$_2$O affects the reaction only through $X_{\text{H}_2\text{O}}$, and that increasing $X_{\text{H}_2\text{O}}$ in these conditions results in enhancement of the $M_n$, yield and polydispersity significantly.

From these initial studies, it is apparent that the minimum $X_{\text{H}_2\text{O}}$ observed for the 3.0M $K_3\text{PO}_4$ mediated externally initiated SCTP of 2 using 1 must be greater than 0.15 (entry 11) in order to produce a narrow polydispersity (PDI < 1.2) of polymer chains. Implementing even higher water molar fractions up to 0.48 dramatically improve $M_n$ and yield of poly($p$-phenylene) presumably deriving from more efficient formation of catalytically active $L_2\text{ArPdOH}$. This is contingent on a
sufficient molar amount of $K_3PO_4$ relative to the 2, which is proposed to be at least the theoretical minimum stoichiometry of 4:1. The concentration of Pd species in solution leading to successful externally initiated SCTP can be as low as 0.42 µM, and the low concentration of palladium centers does not appear to be limiting the polymerization rate. These observations are consistent with rate limiting process involving the equilibrium exchange reaction between $L_2ArPdOH$, $Br^-$, $L_2ArPdBr$ and $-OH$. The premises are integrated into the proposed catalytic cycle via the oxo-palladium pathway illustrated in Figure 2.17. Such a mechanism of SCTP polymerization of MIDA boronates and associated catalytic cycle has never been postulated.

![Proposed catalytic cycle for the Externally Initiated Suzuki-Miyaura Catalyst-Transfer Polymerization of 4-bromo-2,5-dihexylphenylboronic acid MIDA ester](image)

Figure 2.17. Proposed catalytic cycle for the Externally Initiated Suzuki-Miyaura Catalyst-Transfer Polymerization of 4-bromo-2,5-dihexylphenylboronic acid MIDA ester 2 initiated by 1 via the Oxo-Palladium Pathway.

In this proposed catalytic cycle, the external initiator 1 becomes activated through ligand exchange reaction of $-OH$ with $I^-$ centered on the Pd$^{II}$ precatalytic species to generate the oxo-
palladium intermediate. I\(^-\) is less nucleophilic than Br\(^-\) and has a smaller \(K_{eq}\) in the equilibrium reaction between the Oxo-Pd intermediate and ArL\(_2\)PdX, therefore iodine derived external initiators are theoretically expected to better form Oxo-Pd intermediate and result in more efficient initiation than initiators with aryl bromide precursors.\(^{23}\) 2 is converted to the free boronic acid via the slow mechanism of MIDA boronate hydrolysis, and the by-product \(N\)-methyliminodiacetic acid is deprotonated by 2 equivalents of K\(_3\)PO\(_4\). The neutral boronic acid undergoes transmetalation with the oxo-palladium intermediate and boric acid is produced as a side product, consuming another equivalent of K\(_3\)PO\(_4\). Following fast reductive elimination, intramolecular catalyst-transfer and oxidative addition steps, the subsequent conversion of ArL\(_2\)PdBr to ArL\(_2\)PdOH by hydroxide ligand exchange reaction produces Br\(^-\) as a side product and regenerates the oxo-palladium complex.

An important question remains: which is the rate determining step? The reaction rate is seemingly affected by the equilibrium between ArL\(_2\)PdBr and the oxo-Pd intermediate; strongly suggesting that neither of the transmetalation, oxidative addition, reductive elimination nor intramolecular catalyst-transfer itself is limiting the average rate of polymerization. Practically (for controlling the overall polymerization rate), the ligand exchange reaction of \(\cdot\)OH and ArL\(_2\)PdBr to form ArL\(_2\)PdOH seems to be the rate limiting step. This hypothesis is supported by the observed influence of the Br\(^-\) anion on the polymerization rate, which is not participating in any of the elementary catalytic processes.

Alternatively, the rate of MIDA hydrolysis of 2 could be rate limiting. It’s slow formation certainly warrants consideration as the rate limiting step. The two mechanisms would show different rate kinetics. If the formation of the oxo-palladium intermediate from ArL\(_2\)PdBr was rate limiting, it should show a zero-order reaction in excess base because the concentration of Pd
centers and the pH of the THF phase can be assumed to be constant. If MIDA hydrolysis was rate limiting, then a first-order kinetics is expected proportional to the concentration of 2. It would progressively slow down as the monomer is depleted. Initial studies of the rate kinetics showed a zero-order reaction, showing a linear dependency of the molecular weight vs time, but more experiments were needed to confirm this result.

2.2.5. Capturing Br⁻ using Ag⁺: Enhancement of the Mₙ, Polydispersity and Yield

Halides play opposing roles in the boronate and oxo-palladium pathways. In the boronate pathway, the halide anion increases the reaction rate through L₂ArPdX species which undergoes transmetalation with the anionic boronate monomer. In the oxo-palladium pathway, halides drive the formation of inactive L₂ArPdX. The results obtained from the polymerization study strongly suggests that Br⁻ plays an antagonistic role, thus favoring the oxo-palladium pathway.

We hypothesized that the polymerization could be further enhanced by removing Br⁻ from solution. Ag⁺ readily reacts with Br⁻ to form AgBr and has been used in small molecule Suzuki coupling for accelerating reactions. Thus, SCTP of 2 was conducted in the presence of various silver(I) salts to enhance the polymerization reaction results (Table 2.3). The use of Ag⁺ to accelerate Suzuki-Miyaura reaction rate was pioneered by Kishi et al. Ag⁺ can react with Br⁻ and −OH to form AgBr and Ag₂O respectively. In basic conditions, Ag⁺ is expected to form Ag₂O almost quantitatively and reduce the availability of reactive ions in solution. As so, the equilibrium reaction between aqueous Ag⁺ and Ag₂O must be considered in order to integrate this strategy effectively. For the following reactions (entries 12 to 24), AgOAc and Ag₂SO₄ were chosen because of their limited solubility in H₂O, which was hypothesized to slowly form in-situ Ag₂O with a significant surface area.
Table 2.3. Externally Initiated SCTP of 4-bromo-2,5-dihexylphenylboronic acid MIDA ester 2 using AgOAc and Ag₂SO₄ as bromide capturing agents.

<table>
<thead>
<tr>
<th>entry</th>
<th>THF (mL)</th>
<th>H₂O (mL)</th>
<th>Mₙ (PDI)</th>
<th>yield (%)</th>
<th>Χ_H₂O</th>
<th>Ag⁺ (equiv)</th>
<th>[K₃PO₄]/[2]</th>
<th>temp (°C)</th>
<th>time (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>12ᵃ</td>
<td>8</td>
<td>0.25</td>
<td>8.8k (1.13)</td>
<td>31.2%</td>
<td>0.12</td>
<td>AgOAc (1)</td>
<td>12</td>
<td>45°C</td>
<td>48</td>
</tr>
<tr>
<td>13ᵃ</td>
<td>8</td>
<td>0.25</td>
<td>9.8k (1.14)</td>
<td>33.6%</td>
<td>0.12</td>
<td>Ag₂SO₄ (1)</td>
<td>12</td>
<td>45°C</td>
<td>48</td>
</tr>
<tr>
<td>14ᵃ</td>
<td>8</td>
<td>0.5</td>
<td>9.3k (1.15)</td>
<td>39.8%</td>
<td>0.22</td>
<td>AgOAc (1)</td>
<td>24</td>
<td>45°C</td>
<td>48</td>
</tr>
<tr>
<td>15ᵃ</td>
<td>8</td>
<td>0.5</td>
<td>9.8k (1.16)</td>
<td>34.8%</td>
<td>0.22</td>
<td>Ag₂SO₄ (1)</td>
<td>24</td>
<td>45°C</td>
<td>48</td>
</tr>
<tr>
<td>16ᵃ</td>
<td>8</td>
<td>0.25</td>
<td>6.8k (1.17)</td>
<td>50.1%</td>
<td>0.12</td>
<td>Ag₂SO₄ (1)</td>
<td>6</td>
<td>45°C</td>
<td>48</td>
</tr>
<tr>
<td>17</td>
<td>8</td>
<td>0.25</td>
<td>7.5k (1.21)</td>
<td>60.6%</td>
<td>0.12</td>
<td>Ag₂SO₄ (2)</td>
<td>11</td>
<td>rt</td>
<td>96</td>
</tr>
<tr>
<td>18</td>
<td>8</td>
<td>0.25</td>
<td>7.5k (1.18)</td>
<td>60.9%</td>
<td>0.12</td>
<td>Ag₂SO₄ (2)</td>
<td>13</td>
<td>rt</td>
<td>96</td>
</tr>
<tr>
<td>19</td>
<td>8</td>
<td>0.45</td>
<td>6.9k (1.16)</td>
<td>71.8%</td>
<td>0.20</td>
<td>Ag₂SO₄ (2)</td>
<td>12</td>
<td>rt</td>
<td>96</td>
</tr>
<tr>
<td>20</td>
<td>8</td>
<td>0.75</td>
<td>7.5k (1.14)</td>
<td>67.8%</td>
<td>0.30</td>
<td>Ag₂SO₄ (2)</td>
<td>20</td>
<td>rt</td>
<td>96</td>
</tr>
<tr>
<td>21</td>
<td>8</td>
<td>1.2</td>
<td>6.5k (1.12)</td>
<td>52.9%</td>
<td>0.40</td>
<td>Ag₂SO₄ (2)</td>
<td>32</td>
<td>rt</td>
<td>96</td>
</tr>
<tr>
<td>22</td>
<td>16</td>
<td>0.5</td>
<td>8.4k (1.16)</td>
<td>86.4%</td>
<td>0.12</td>
<td>Ag₂SO₄ (2)</td>
<td>13</td>
<td>rt</td>
<td>96</td>
</tr>
<tr>
<td>23</td>
<td>8</td>
<td>0.25</td>
<td>5.7k (1.16)</td>
<td>59.7%</td>
<td>0.12</td>
<td>Ag₂SO₄ (2)</td>
<td>13</td>
<td>rt</td>
<td>66</td>
</tr>
<tr>
<td>24</td>
<td>8</td>
<td>0.25</td>
<td>7.9k (1.17)</td>
<td>74.0%</td>
<td>0.12</td>
<td>Ag₂SO₄ (2)</td>
<td>13</td>
<td>rt</td>
<td>91</td>
</tr>
</tbody>
</table>

ᵃReaction conditions: [K₃PO₄] = 3.0M, [2]₀/[1]₀ = 29.ᵇAbsolute molecular weight was determined by quantitative NMR. ᶜMₙ/Mₚ was determined by GPC calibrated using polystyrene standard and -with THF as eluent at 40°C. ᵈIsolated yield after precipitation in acetone. ᵉ5.0M K₃PO₄. ᶠ6.0M K₃PO₄. ᵍ[2]₀/[1]₀ = 45.

Using 1 molar equivalent of AgOAc (entry 12, 14) or Ag₂SO₄ (entry 13, 15) relative to 2 resulted in narrow polydispersity at X_H₂O between 0.12 and 0.22; and significantly higher Mₙ (up to 9.8 kDa, theoretical Mₙ = 10.4 kDa) than previously obtained. This showed the compatibility of AgOAc and Ag₂SO₄ in SCTP conditions and also affirmed the innocent nature of acetate and sulfate counter-anions. Using 1 equivalent of silver salt relative to 2 and at X_H₂O = 0.12, Ag₂SO₄ (entry 15) produced higher Mₙ (9.8 kDa) PDCI-PPP than AgOAc (8.8 kDa, entry 14). Ag₂SO₄ contains double the amount of Ag⁺ per mole of reagent versus AgOAc and this demonstrated that silver(I) salts are not effective in stoichiometric amounts relative to the monomer. At X_H₂O = 0.22 however (entry 14), AgOAc produced a higher molecular weight polymer (9.3 kDa) than it did in conditions with X_H₂O = 0.12, and this implied that water volume affects the availability of Ag⁺.
significantly. Both silver salts demonstrated excellent compatibility and reaction enhancement capabilities. Due to its more uniform effects in varying $X_{\text{H}_2\text{O}}$, $\text{Ag}_2\text{SO}_4$ was selected for further studies.

Next, we repeated entry 5 with exact reaction conditions except adding 1 molar equivalent of $\text{Ag}_2\text{SO}_4$ relative to the monomer (entry 16). In previous conditions, narrow polydispersity PDI < 1.2 was only obtained at $X_{\text{H}_2\text{O}} > 0.15$. Along with results from entry 12 and 13, entry 16 confirmed silver’s polymerization rate enhancing abilities and produced PDCI-PPP with consistently better polydispersity (PDI = 1.13 - 1.17, $X_{\text{H}_2\text{O}} = 0.12$) relative to entry 5 (PDI = 1.27, $X_{\text{H}_2\text{O}} = 0.12$). The improvement can only be rationalized from the oxo-Pd pathway where the removal of $\text{Br}^-$ in the beginning of reaction would result in a more uniform initiation step, deriving from a higher concentration of $\text{L}_2\text{ArPdOH}$ vs $\text{L}_2\text{ArPdBr}$. Also, adding $\text{Ag}_2\text{SO}_4$ improved the $M_n$ and yield from 3.6 kDa in entry 5 to 6.8 kDa in entry 16, and yield from 8.0% in 50.1%.

The molecular weight of PDCI-PPP prepared from entry 16 using 6 : 1 molar ratio of $K_3\text{PO}_4$ to 2 was lower than entry 12 and 13 using 12 : 1 molar ratio of $K_3\text{PO}_4$ to 2. In this case, $\text{Ag}_2\text{SO}_4$ is likely acting as a Brønsted acid in the hydrolysis reaction to form $\text{Ag}_2\text{O}$, consuming $\text{-OH}$ and reducing the stoichiometry of the base below the necessary amount for the polymerization reaction. Therefore, additional stoichiometric amount of $K_3\text{PO}_4$ was added in following reactions to allow for efficient silver mediated SCTP reaction to occur. For PDCI-PPP polymers with $M_n > 9$ kDa, precipitation was observed during the extraction process using CHCl$_3$/brine as organic and aqueous phases and led to lower yields for entry 12-15.

A final reaction optimization series of SCTP of 2 was conducted using $\text{Ag}_2\text{SO}_4$, in various relative amounts of THF, $\text{H}_2\text{O}$ and $K_3\text{PO}_4$. Using the knowledge gained in previous studies, the objective of experiments in entries 17 – 25 is to obtain ideal silver mediated SCTP conditions.
Along this route, the reactions were conducted at room temperature to reduce intermolecular chain-transfer process, but this increased the reaction time dramatically. Two equivalents of Ag₂SO₄ was utilized, and therefore additional stoichiometric amount of K₃PO₄ was also increased. Due to precipitates observed at higher molecular weights (entries 12 – 15), the theoretical molecular weight for reactions 17 - 25 was lowered to 8.1 kDa. Two identical reactions were conducted to obtain the total reaction time at X_H₂O = 0.12 (entry 23, 24). At 66h, the entry 23 reached a yield of 59.7% and molecular weight 5.7 kDa (70% of the theoretical molecular weight). At 91h, the second identical reaction (entry 24) reached 74% yield and 7.9 kDa (97% theoretical molecular weight). Therefore, the reaction time was estimated to be 4 days at 0.12 water molar fraction and at room temperature.

Next, additional molar quantity of base was added (to accommodate for the acidic nature of Ag⁺) by increasing the aqueous concentration of K₃PO₄ to 5.0M (entry 17) and 6.0 M (entry 18). Using 5.0 M and 6.0 M K₃PO₄ produced similar results, producing PDCI-PPP with molecular weight up to 7.5 kDa (93% of the theoretical M_n), with similar yields of 61% and PDI 1.18-1.21. In ideal controlled polymerization systems, the % theoretical M_n (observed M_n / theoretical M_n × 100%) is an effective intrinsic indicator of reaction progress. The molar quantities of base can also be added by increasing the volume of H₂O while maintaining 3.0 M concentration of K₃PO₄. Concurrently, this increased the water molar fraction to 0.20 (entry 19), 0.30 (entry 20) and 0.40 (entry 21). Increasing X_H₂O again improved the polydispersity proportionally from 1.16 in entry 19 to 1.12 in entry 21, which is attributed to the increased formation of ArL₂PdOH vs ArL₂PdBr. Surprisingly, the yield decreased as the water molar fraction increased in stark contrast to entries 9-11 where the increasing X_H₂O resulted in higher yields.
In the theory proposed here, the increase in $X_{\text{H}_2\text{O}}$ plays a beneficial role in driving the formation of ArL$_2$PdOH versus ArL$_2$PdBr. But at the same time, increasing $X_{\text{H}_2\text{O}}$ raises the pH and results in monomer degradation (via formation of BrArB(OH)$_3$K, a precursor to hydrodeboronation via the Kuivila mechanism$^{19-21}$). When a significant amount of Br$^-$ is present (with no Ag$^+$ added), increasing $X_{\text{H}_2\text{O}}$ battles the negative effects of Br$^-$, thus enhances the polymerization rate, $M_n$, yield, and polydispersity. But in the absence of Br$^-$, the beneficial effects of increasing $X_{\text{H}_2\text{O}}$ are neutralized because the equilibrium between the oxo-palladium and palladium halide species is already being boosted by Ag$^+$. Then, only the negative effects of increasing $X_{\text{H}_2\text{O}}$ can be observed. Thus, the dual effects of increasing $X_{\text{H}_2\text{O}}$ materialize as two different results in silver and non-silver mediated SCTP. This process explains why the best polymerization reaction results in silver mediated SCTP are obtained at lower $X_{\text{H}_2\text{O}}$, because it reduces the rate of hydrodeboronation. PDCI-PPP polymers in entries 19-21 reached a maximum % theoretical $M_n$ of 92% (entry 20).

The molar amounts of base relative to 2 could also be added by scaling up the entire reaction (adding THF, H$_2$O and K$_3$PO$_4$) while maintaining the molar quantities of the initiator 2, and monomers 5 (entry 22); relative to entry 16. This allowed the reaction to maintain a low $X_{\text{H}_2\text{O}}$ of 0.12. These conditions produced the best results, reaching the highest yield of 86.4%, a quantitative % theoretical $M_n$ and narrow polydispersity of 1.16.

Applying the optimized silver mediated SCTP conditions just obtained, a series of PDCI-PPP were prepared with increasing degree of polymerization up to 50 (Table 2.4). A control reaction was conducted without silver to contrast its effects at room temperature (entry 31). The polymers were prepared with excellent yields and consistently low polydispersity. The number averaged molecular weight of the polymers were plotted to show a linear dependence versus feed ratio [2]/[1]$_0$ (Figure 2.18), demonstrating excellent controlled chain-growth behavior.
Ag\(^+\) does not seem to be totally innocent. Although the \(M_n\) and yield are dramatically enhanced versus the control in entry 31, the polydispersity is actually lower without silver. Albeit, the silver reactions still demonstrated narrow polydispersity indexes of PDI < 1.2 (entry 25-30). Amatore has described a similar effect where the cations of bases used in Suzuki coupling has been shown to coordinate to the oxygen lone pair on the oxo-palladium complex, making it inactive towards transmetalation.\(^{26}\) Ag\(^+\) is also an oxophilic Lewis acid, and it would be logical that it can behave in the same way. This is likely to be the reason for the increased polydispersity in entries 25-30. A constant concentration of silver would statistically deactivate a portion of the oxo-palladium complexes via Amatore’s proposed mechanism.

Table 2.4. Preparation of PDCI-PPP\(_n\) with various degrees of polymerization via silver mediated SCTP.

<table>
<thead>
<tr>
<th>entry</th>
<th>(M_n) (^b) (PDI(^c))</th>
<th>yield(^d) (%)</th>
<th>([2]_0/[1]_0)</th>
</tr>
</thead>
<tbody>
<tr>
<td>25</td>
<td>4.9k (1.19)</td>
<td>80.9%</td>
<td>17</td>
</tr>
<tr>
<td>26</td>
<td>5.7k (1.16)</td>
<td>71.5%</td>
<td>20</td>
</tr>
<tr>
<td>27</td>
<td>6.7k (1.19)</td>
<td>72.4%</td>
<td>25</td>
</tr>
<tr>
<td>28</td>
<td>7.8k (1.18)</td>
<td>71.7%</td>
<td>30</td>
</tr>
<tr>
<td>29</td>
<td>10.0k (1.17)</td>
<td>57.8%</td>
<td>40</td>
</tr>
<tr>
<td>30</td>
<td>12.5k (1.20)</td>
<td>71.6%</td>
<td>50</td>
</tr>
<tr>
<td>31(^e)</td>
<td>3.4k (1.09)</td>
<td>12.6%</td>
<td>25</td>
</tr>
</tbody>
</table>

\(^a\)Reaction conditions: \([K_3PO_4]\) = 3.0 M, rt, 96h. \(^b\)Absolute molecular weight was determined by quantitative NMR. \(^c\)\(M_n/M_w\) was determined by GPC calibrated using polystyrene standard and with THF as eluent at 40°C. \(^d\)Isolated yield after precipitation in acetone. \(^e\)No Ag\(_2\)SO\(_4\) was added.
And similar to how the dynamic equilibrium between ArL₂PdBr and ArL₂PdOH can effect polydispersity because of their relative reactivities, it’s probable that the equilibrium between the free oxo-palladium complex and the silver ligated oxo-palladium complex could also result in a statistical distribution of chain-end reactivities; giving rise to a polydispersity phenomenon observed in entries 25-31.

### 2.2.6. Photophysical Properties of PDCI-PPP Polymers With Varying Molecular Weight

PDCI-PPP polymers having varying degree of polymerizations from 15 to 50 were characterized by UV-VIS absorbance and photoluminescence spectroscopy (Figure 2.19). The absorption spectra of all PDCI-PPP polymers in this series were essentially identical, with the bands significantly red-shifted relative to the free PDCI, which indicated increased π-conjugation due to phenylene being electronically coupled to PDCI. $\lambda_{\text{max}}$ of the short wavelength emission shifted
from 338 to 365 nm, $\lambda_{\text{max}}$ of the S$_0$ to S$_2$ transition shifted from 466 to 495 nm, and the lowest energy S$_0$ to S$_1$ transition was red shifted from 613 to 645 nm. The polymers were essentially nonfluorescent. Interestingly, the expected emission band around 400 nm expected of poly($p$-phenylene) was not observed in PDCI-PPP$_{15}$, but became progressively more intense as the degree of polymerization of the polymer chain increased up to 50. This indicated an efficient quenching process of the PPP excited state energy by PDCI charge-transfer complex. As the chain length increases, the energy transfer may become less efficient and some of the excited state energy would be fluorescently emitted by the more distant chromophores situated on PPP. In the emission spectra, the NIR band is also dramatically red-shifted from the 775 nm emission of the free PDCI core to 860 nm in the PDCI-PPP polymer series.

Figure 2.19. UV-Vis absorbance (left) and emission spectra (right) of PDCI-PPP$_n$

2.2.7. Silver Mediated Externally Initiated SCTP for Preparing Poly(9,9-dioctylfluorene)s

Poly(2,5-dialkylphenylene)s and poly(9,9-dialkylfluorene)s (PFs) are chemically very similar, having alkylated solubilizing groups installed on a conjugated backbone composed of benzene rings. Therefore, 7-bromo-9,9-dioctylfluorene boronic acid MIDA ester 4 was synthesized and a series of PDCI-PF polymers having various theoretical molecular weights were prepared to test if
the optimized silver mediated SCTP was transferrable to preparing poly(9,9-dioctylfluorene) (Table 2.5). A control reaction was also conducted without silver (entry 37). PDCI-PF prepared by silver mediated SCTP demonstrated controlled chain-growth character up to 14.9 kDa and resulted in a modest yield and narrow polydispersity (Figure 2.20). The photophysical properties of PDCI-PF series having various theoretical molecular weights were characterized by UV-VIS Absorbance and photoluminescence spectroscopy. (Figure 2.21)

Table 2.5. Preparation of PDCI-PPPₙ with various molecular weights by silver mediated SCTP.

<table>
<thead>
<tr>
<th>entry</th>
<th>Mₙ (PDI)</th>
<th>yield (%)</th>
<th>[4]₀/[1]₀</th>
<th>QY</th>
</tr>
</thead>
<tbody>
<tr>
<td>32</td>
<td>3.5k (1.41)</td>
<td>41.7%</td>
<td>8</td>
<td>24.0%</td>
</tr>
<tr>
<td>33</td>
<td>5.6k (1.40)</td>
<td>36.6%</td>
<td>16</td>
<td>71.7%</td>
</tr>
<tr>
<td>34</td>
<td>9.5k (1.57)</td>
<td>43.0%</td>
<td>24</td>
<td>110.0%</td>
</tr>
<tr>
<td>35</td>
<td>11.1k (1.53)</td>
<td>47.8%</td>
<td>32</td>
<td>156.6%</td>
</tr>
<tr>
<td>36</td>
<td>14.9k (1.49)</td>
<td>37.4%</td>
<td>40</td>
<td>161.8%</td>
</tr>
<tr>
<td>37</td>
<td>2.4k (1.13)</td>
<td>12.80%</td>
<td>16</td>
<td>-</td>
</tr>
</tbody>
</table>

aReaction conditions: [K₃PO₄] = 3.0 M, rt, 96h. bAbsolute molecular weight was determined by quantitative NMR. cMₙ/Mₙ was determined by GPC calibrated using polystyrene standard and with THF as eluent at 40°C. dIsolated yield after precipitation in acetone. eNo Ag₂SO₄ was added.

The control reaction also showed similar results in comparison to the control reaction of PDCI-PPP polymers where it had lower molecular weight, yield, but more narrow polydispersity (entry 37). The absorbance spectra of PDCI-PFₙ was heavily influenced by PF chain length. Unlike PDCI-PPP where the absorption is dominated by the PDCI core (or related charge-transfer
complex), the absorbance by polyfluorene becomes more intense as the degree of polymerization is increased. The PDCI-PF was much more fluorescent and demonstrated up to an unprecedented 160% photoluminescence quantum yield (last column in Table 2.5), where the short wavelength and NIR bands demonstrated an approximately 1:1 ratio of PL quantum yields.

Figure 2.20. $M_n$ vs feed ratio ([4]₀/[1]₀) of PDCI-PFₙ polymers prepared by silver mediated Suzuki-Miyaura Catalyst-Transfer Polymerization.

Figure 2.21. UV-Vis absorbance (left) and emission spectra (right) of PDCI-PFₙ.
The photoluminescence quantum yield varied proportionally from 11.9% in the case of PDCI-PF₈, 35.8% for PDCI-PF₁₆, 55.3% for PDCI-PF₂₄, 74.0% for PDCI-PF₃₂ to 81.2% for PDCI-PF₄₀. Strong intramolecular energy transfer is expected via the Förster mechanism since the emission $\lambda_{\text{max}}$ of PF overlaps with the absorption $\lambda_{\text{max}}$ of the PDCI charge transfer complex (Figure 2.1). PDCI is expected to significantly quench PF emission and the chain-length dependent fluorescence quantum yield shows a highly efficient energy transfer taking place.

2.2.8. Externally Initiated Controlled Polymerization for Preparing Poly(3-hexylthiophenes)

The successful preparation of PDCI-P3HTₙ did not require the use of silver. A series of PDCI-P3HT polymers were prepared with degree of polymerization varying from 20 to 100. External initiation using I resulted in PDCI-P3HT with controlled molecular weight, excellent yields and modest polydispersity (Table 2.6). The PDCI-P3HTₙ polymers were characterized by UV-Vis absorbance and photoluminescence spectroscopy (Figure 2.22). PDCI-P3HTₙ series displayed a strong correlation between the molecular weight of the polymers and their PL quantum yield (Table 2.6), where the quantum yield progressed from 4.9% to 24.8% going from PDCI-P3HT₂₀ to PDCI-P3HT₁₀₀. The observed trend further demonstrates the quenching effect of PDCI core (or likely an associated charge-transfer state) and also shows effective energy transfer from P3HT to low energy PDCI. PDCI-P3HTₙ displayed a broad absorbance, owing to the dynamic conformational subunits caused by twisting and kinking of the polymer chains. The PL spectra of PDCI-P3HT was far more blue-shifted vs PDCI-PPP and PDCI-PF with an emission band $\lambda_{\text{max}}$ of 575 nm. A slight vibronic structure is also observed with an emission band appearing around 630 nm.
Table 2.6. Controlled molecular weight series of PDCI-PT\textsubscript{n} prepared by SCTP without silver.

<table>
<thead>
<tr>
<th>entry</th>
<th>$M_n$ (PDI)</th>
<th>Yield (%)</th>
<th>$[4]/[1]$ \textsubscript{0}</th>
<th>QY</th>
</tr>
</thead>
<tbody>
<tr>
<td>38</td>
<td>4.0k (1.21)</td>
<td>65.0%</td>
<td>20</td>
<td>4.9%</td>
</tr>
<tr>
<td>39</td>
<td>7.7k (1.23)</td>
<td>82.7%</td>
<td>40</td>
<td>2.4%</td>
</tr>
<tr>
<td>40</td>
<td>11.4k (1.29)</td>
<td>88.1%</td>
<td>60</td>
<td>12.4%</td>
</tr>
<tr>
<td>41</td>
<td>14.0k (1.40)</td>
<td>92.3%</td>
<td>80</td>
<td>15.2%</td>
</tr>
<tr>
<td>42</td>
<td>17.0k (1.45)</td>
<td>83.8%</td>
<td>100</td>
<td>24.8%</td>
</tr>
</tbody>
</table>

$^a$Reaction conditions: $[K_3PO_4] = 3.0$ M, 45°C, 48h. $^b$Absolute molecular weight was determined by quantitative NMR. $^c$M\textsubscript{w}/M\textsubscript{n} was determined by GPC calibrated using polystyrene standard and with THF as eluent at 40°C. $^d$Isolated yield after precipitation in acetone.

Figure 2.22. $M_n$ vs feed ratio ($[3]/[1]$\textsubscript{0}) of PDCI-P3HT\textsubscript{n} polymers prepared by silver mediated Suzuki-Miyaura Catalyst-Transfer Polymerization.
The shortest chain-length PDCI-P3HT$_{20}$ had a blue-shifted absorbance and also emission bands versus PDCI-P3HT$_{40-100}$. This is expected because of the symmetry of PDCI-P3HT having two short 10 repeating unit P3HT on each side of the chain instead of a single 20-unit polymer chain and thus the conjugation path length for these units is much shorter than expected for 20 repeating unit of typical P3HT. The PDCI absorbance is also more pronounced in PDCI-P3HT$_{20}$ versus the other polymer samples, demonstrating that polythiophene has a significant molecular weight dependent extinction coefficient relative to the PDCI core (unlike PDCI-PPP where the absorption was dominated by PDCI). These experiments show that the donor-acceptor polymers prepared by externally initiated SCTP are well-defined, highly precise platforms for studying the structure dependent properties of conjugated polymers. PDCI was shown to be an effective energy acceptor and exhibits interesting and novel behavior when incorporated into conjugated polymer backbones.
2.3 Conclusion

Suzuki-Miyaura Catalyst Transfer Polymerization is still a technique at its infancy with few reports able to produce robust methodologies broadly applicable for preparing various classes of conjugated polymers. This evidence-based study on the SCTP mechanism revealed several novel fundamental processes that will likely have a lasting impact in conjugated polymer research. First, overwhelming evidence has been provided in support of the oxo-palladium (rather than boronate) pathway to be in operation for Pd SCTP. Second, the unequivocal role of water was elucidated to show two opposing effects on the \( M_n \), yield and polydispersity of conjugated polymers prepared by SCTP. Third, the influential role of \( \text{Br}^- \) was discovered to seriously hamper the rate SCTP reactions. Fourth, based on the proposed mechanism, a synthetic solution utilizing silver was developed and integrated into the optimized SCTP methodology. These principles were combined to demonstrate controlled chain-growth preparation of P3HT, PPP and PF.

2.4 References


CHAPTER 3. TUNABLE NANOSTRUCTURES OF DONOR-ACCEPTOR NANOPARTICLES BY EXTERNALLY INITIATED KUMADA CATALYST-TRANSFER DISPERSION POLYMERIZATION

3.1. Introduction

Recently, there has been a growing interest in utilizing surfactant-free donor-acceptor conjugated polymer nanoparticles (CPN) for organic electronic devices such as organic light emitting diodes, organic field effect transistors and organic photovoltaic cells.\textsuperscript{1-10} Integrating CPNs into the active layers of organic electronic devices allows for the optoelectronic properties to be engineered separately from the thin-film construction. Donor-acceptor CPNs are prepared by reprecipitation or miniemulsion methods\textsuperscript{11-12} and their material properties\textsuperscript{13-15} can be tailored by varying the polymer concentration during the preparation process; resulting in nanoparticles of different sizes and properties. Alternatively, the relative molar ratios of donor-acceptor blends can be adjusted prior to nanoparticle formation to modify the microstructure within (such as morphology and average distance between donor and acceptor domains).\textsuperscript{16-21} But these methods don’t allow for controlling the properties of primary nanostructures such as chain-conformation and average conjugation path lengths to any significant degree.

Using ultrasonication is a major drawback for both reprecipitation and miniemulsion techniques. The heat and cavitation bubbles caused by high energy ultrasound waves completely alter the polymer self-assembly process and disallow for using temperature as a synthetic parameter for tuning CPN properties.\textsuperscript{22-27} For example, Luscombe and Pozzo have recently demonstrated that ultrasonication triggers the self-assembly of poly(3-hexylthiophene) into nanofibers even in good solvents such as chlorobenzene or chloroform.\textsuperscript{28} Therefore, an ability to prepare CPN’s without sonication to control the temperature-dependent self-assembly process would be an important fundamental approach for making CPNs with precise and tunable properties.
Herein, donor-acceptor polythiophene nanoparticles with tunable photophysical properties are prepared by externally initiated Kumada Catalyst-Transfer dispersion polymerization. The CPNs are prepared without sonication or surfactants in THF. An electron deficient perylenedicarboximide (PDCI) was utilized as precursor to the Ni(dppp) based external initiator and was covalently integrated into the polymer; forming a donor-acceptor system. The reaction temperature was found to affect the average conjugation path length of polythiophene chain segments that are kinetically trapped within the supramolecular assembly. Subsequently, the polythiophene fluorescence $\lambda_{\text{max}}$ from these chromophores can be controlled and alters the Förster energy transfer efficiency towards PDCI. The kinetically trapped conformations revert back to a common structure upon heating at 310°C. In addition to providing a synthetic handle on conjugated polymer self-assembly process affording various tunable photophysical properties, PDCI-PTh particles were investigated by X-ray powder diffraction (XRD), small-angle X-ray scattering (SAXS), small-angle neutron scattering (SANS), high-resolution scanning electron microscopy (SEM) experiments to provide more details on the structure of donor-acceptor nanoparticles on multiple length-scales.

Polythiophene is an ideal model system to study nanoparticle structure formation by controlled polymerization because of the abundance of structural characterization data, predictable self-assembly behavior and compatibility with transition metal catalyzed polymerization methods. For polythiophene synthesis, Kumada Catalyst-Transfer Polymerization (CTP) is well-established for its quasi-living chain-growth nature and its ability to produce polythiophenes. The controlled chain-growth nature of Kumada CTP was discovered independently by McCullough and Yokozawa in 2004. Kumada type CTP reactions were used to polymerize thiophene Grignard monomers in this study which led to polythiophenes. We also employed an efficient preparation
of the external initiator using Ni(dppp)$_2$ as the pre-catalyst; a method previously reported by our group.$^{35}$

PDCI is an electron-deficient $\pi$-conjugated fluorophore which can form various supramolecular assemblies$^{36-38}$ and donor-acceptor complexes with electron-rich molecules$^{39}$ (e.g. thiophene). Thus, depending on the reaction conditions, externally initiated PDCI catalyst can result in various initial supramolecular assemblies which can direct CTP towards different PDCI-PTh macromolecular mesoscale systems.

3.2 Results and Discussion

3.2.1. Preparation of the Regioisomerically Pure External Initiator

Classical bromination of perylene tetracarboxylic dianhydride 1 (Scheme 3.1) produces a mixture of constitutional and regioisomers which are practically inseparable because of their low solubility.$^{40}$ Although some researchers have debated$^{41}$ that the difference in photophysical properties of these isomers are insignificant, its crystallinity and self-assembly properties of the donor-acceptor polymers will be affected by the regioregularity of the central initiating core. We therefore decided to pursue isomerically pure starting materials. In 2014, Sengupta et al. reported$^{42}$ a four-step preparation of regioisomerically pure 1,7-dibrominated PDCI via a tetrabutyl ester intermediate which could be readily purified in five days. Utilizing this method, the synthesis began with perylene tetracarboxylic dianhydride 1 (Scheme 3.1), which was converted into a perylene tetrabutyl ester 2 using a 1:1(v/v) mixture of butanol and bromobutane. DBU was utilized as a strong organic base. Bromination of 2 afforded a 3:2 ratio of 1,7- and 1,6-dibrominated regioisomers 3a and 3b, respectively. The 1,7-dibrominated product 3a was isolated with a 27% recovery by recrystallization in a mixture of acetonitrile and dichloromethane.
Scheme 3.1. Preparation of External Catalytic Initiator for Kumada Catalyst-Transfer Dispersion Polymerization.

Hydrolysis of the isomerically pure tetrabutyl ester 3a yielded the isomerically pure 4. Its structure was confirmed by NMR in D$_2$SO$_4$. The $^1$H NMR spectrum showed a single regioisomer and high purity of compound 4. Isomerically pure 4 was condensed with 4-bromo-2,6-diisopropylaniline to yield perylenedicarboximide 5. Following a typical Stille coupling and subsequent NBS bromination, the small molecule precursor 7 was obtained. A solution of the external initiator 8 is produced upon mixing of precursor 7 with an excess of Ni(dppp)$_2$ in THF. The external initiator 8 was freshly prepared prior to each polymerization reaction.

3.2.2. Morphology of PDCI-PTh Prepared at -10°C to 22°C: Characterization by SEM

Preparation of PDCI-PTh nanoparticles begins with a 50 mM solution of 5-bromo-2-thienylchloromagnesium monomer which was made by mixing 2,5-dibromothiophene with a sub-stoichiometric amount of $^1$PrMgCl. Because of its mono-functionality and relatively high reactivity versus the target Grignard monomer, excess $^1$PrMgCl will cause chain termination in this KCTP system. Nanoparticles are formed when a solution 8 (1 mol% relative to the monomer) is introduced to the Grignard solution under air-free conditions and at various temperatures. After
five minutes, the reaction was quenched using 6.0M HCl to neutralize the Grignard monomer and the resulting polymer suspension was isolated and washed repeatedly with chloroform and methanol via centrifugation. The obtained nanoparticles were dried in vacuum and could be resuspended in a variety of solvents including water, methanol, chloroform, and 1,2-dichlorobenzene.

CHCl₃ suspensions of PDCI-PTh produced at 22°C, 7°C, 0°C and -10°C (PDCI-PTh₂₂°C, PDCI-PTh₇°C, PDCI-PTh₀°C and PDCI-PTh⁻₁₀°C) were drop-casted onto a copper substrate for scanning electron microscopy (SEM). SEM reveals that PDCI-PTh₂₂°C particles have an irregular, and jagged shape. As the reaction temperature is lowered, KCTP produced larger PDCI-PTh particle diameters and the morphology became progressively disk-like (Figures 3.1-3.4). The number averaged diameter of PDCI-PTh₂₂°C is 234 ± 26 nm. The odd morphology of PDCI-PTh nanoparticles was unexpected because CPNs prepared by reprecipitation (a similar CPN forming process) generally form spherical particles which minimize the material’s surface area in contact with the solvent; and the solvophobic effect is the dominant driving force for particle self-assembly by reprecipitation.¹¹ The higher surface area morphology of disk-shaped particles versus forming spheres is entropically unfavorable in this regard. nanoparticle that are not single-crystals are typically made of smaller primary units, and the tendency to form disk-like structures at length scale >100 nm point to the existence of sheet-like primary particles that would be necessary to direct hierarchal self-assembly towards two-dimensional morphology. At -10°C, finer sheet structures are observed protruding at the surface of the particles, forming distinct edges and supporting the hypothesis that the sheet-like primary structures are directing hierarchal self-assembly at longer length-scales. Gravimetric analysis and study of the supernatant after centrifugation showed a percent conversion of 53%, while the remaining solute in the supernatant
contained quenched monomer (42% recovery) and unreacted dibromothiophene. 2,5-dibromothiophene is not expected to cause chain-transfer in CTP reactions.40

Figure 3.1. SEM images of PDCI-PTh nanoparticles prepared by Kumada Catalyst-Transfer Dispersion Polymerization at 22°C.

Figure 3.2. SEM images of PDCI-PTh nanoparticles prepared by Kumada Catalyst-Transfer Dispersion Polymerization at 7°C.
3.2.3. Effects of Polymerization Temperature on the Photophysical Properties of PDCI-PTh Nanoparticles

The polymerization reaction temperature had a dramatic effect on the photophysical properties of resulting PDCI-PTh nanoparticles. PDCI-PTh nanoparticles produced between -10°C and 22°C
displayed broad absorption spectra with $\lambda_{\text{max}}$ of 625nm (Figure 3.5). The disordered twisting of the polythiophene polymer backbone in the polymer supramolecular assemblies can result in a statistical distribution of chromophores (termed conformational subunits) with different electronic structures; resulting in a wide absorption spectra.\textsuperscript{41-43} PDCI-PTh$_{7^\circ\text{C}}$, PDCI-PTh$_{0^\circ\text{C}}$ and PDCI-PTh$_{10^\circ\text{C}}$ produced a more disordered absorption spectra versus the PDCI-PTh$_{22^\circ\text{C}}$, presumably due to the lower reaction temperature trapping polythiophene chains into non-equilibrium conformations.

The emission spectra of PDCI-PTh$_{22^\circ\text{C}}$, PDCI-PTh$_{7^\circ\text{C}}$, PDCI-PTh$_{0^\circ\text{C}}$ and PDCI-PTh$_{10^\circ\text{C}}$ displayed a broad emission from 400 – 950 nm with definitive dependence on the polymerization temperature (Figure 3.6). Albeit, these come from different excited states species responding to the change of polymerization reaction temperature in different ways. Notably, the short wavelength emission band ($\lambda_{\text{max}}$ between 450 nm and 500 nm) of the PDCI-PTh nanoparticle series (Figure 3.6) showed a dramatic red-shift with decreasing temperature. PDCI-PTh nanoparticles produced at lower reaction temperatures are expected to result in segments of polythiophene chains with more planar conformations, resulting in longer conjugation path length. This would explain red shift of PDCI-PTh particles emission ($\lambda_{\text{max}}$ between 450-500 nm) as reaction temperature decreases as observed in Figure 3.6.

There is also a reaction temperature dependent trend in the emission intensity of the PDCI core at $\lambda_{\text{max}} = 775$ nm relative to the short wavelength thiophene emission bands. The absorbance and emission spectra of the PDCI core is taken separately using the starting materials and confirmed the assignment of the PDCI-PTh nanoparticle emission at $\lambda_{\text{max}} = 775$ nm (Figure 3.7). An excitation spectrum of PDCI-PTh$_{22^\circ\text{C}}$ revealed that the origin of emission at $\lambda_{\text{max}} = 775$ nm (Figure 3.8) comes from absorption at 340 nm, and the shape of the excitation spectra is identical to the absorption of PDCI in Figure 3.7 (showing characteristic absorption maxima at both 340 nm and 92
460 nm). This further supports that the 775 nm emission band of the PDCI-PTh nanoparticle series originates from PDCI.

Unlike polythiophene, PDCI chromophore is not subject to conformation-dependent electronic structures and is expected to have a constant absorption wavelength $\lambda_{\text{max}} = 340$ nm and emission at $\lambda_{\text{max}} = 775$ nm (Figure 3.7). Indeed, the emission wavelength at $\lambda_{\text{max}} = 775$ nm of the PDCI-PTh remained constant for PDCI-PTh produced at various reaction temperatures. A significant overlap of the donor emission band and the acceptor absorbance band is a critical requirement for efficient dipole-induced dipole coupling needed for Förster energy transfer. Consequently, this characteristic of a constant absorbance wavelength in the acceptor but a variable emission wavelength of the donor affords a mechanism for tuning of the Förster energy transfer efficiency.

If Förster energy transfer is taking place from higher energy polythiophene conformational subunits to PDCI, intensity of the 775 nm emission band would be dramatically affected by the fluorescence wavelength of polythiophene. As the reaction temperature is decreased, the red shift of polythiophene conformational subunits may cause depletion of available energy donors that are able to undergo Förster energy transfer. Thus, for lower reaction temperatures, reduced energy transfer is expected to occur from polythiophene to the PDCI chromophore as it was observed in emission intensity of the 775 nm emission band in Figure 3.6.
Figure 3.5. UV-Vis absorbance spectra of PDCI-PTh particles prepared between -10°C and 22°C by Kumada Catalyst-Transfer Dispersion Polymerization.

Figure 3.6. Emission spectra of PDCI-PTh particles prepared between -10°C and 22°C by Kumada Catalyst-Transfer Dispersion Polymerization.
Figure 3.7. UV-Vis absorbance and emission spectra of PDCI precursor 5 and 7.

Figure 3.8. Excitation spectra acquired at $\lambda_{\text{max}} = 775$ nm emission bands of PDCI-PTh$_{22^\circ C}$. 
The conformation of polythiophene chains within the PDCI-PTh mesoscopic self-assemblies are shown herein to depend on reaction temperature, which affects the average polythiophene conformational subunit emission $\lambda_{\text{max}}$. The NIR emission bands at 775 nm was shown to correlate to the polythiophene emissions $\lambda_{\text{max}}$. Thus, the photophysical properties of PDCI-PTh nanoparticles have been demonstrated here to be highly dependent on reaction temperature, affecting the conjugation path length of polythiophene conformational subunits and subsequently altering the energy-transfer efficiency to PDCI via Förster mechanism.

**3.2.4. Kinetically Trapped Polythiophene Chains in PDCI-PTh$_{22^\circ C}$ and PDCI-PTh$_{0^\circ C}$**

The control over PDCI-PTh particles’ energy transfer behavior and photophysical properties rely on the hypothesis that the reaction temperature causes the formation of kinetically trapped conformational states. To test this idea, PDCI-PTh$_{22^\circ C}$ and PDCI-PTh$_{0^\circ C}$ were heated in an inert atmosphere at high temperature to allow for thermodynamically driven rearrangement of the polythiophene segments into a common supramolecular structural state. First, PDCI-PTh$_{22^\circ C}$ was characterized by differential scanning calorimetry (DSC) to identify possible thermal transitions (Figure 3.10).

![Figure 3.9. PDCI-PTh$_{22^\circ C}$ characterized by Differential Scanning Calorimetry.](image)

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DSC showed an endothermic transition around 310°C and thus this temperature was used for heating experiments. The absorbance and fluorescence of PDCI-PTh$_{22^\circ}$C and PDCI-PTh$_{0^\circ}$C before and after heating at 310°C is shown in Figure 3.11.

![Graphs showing absorbance and emission spectra](image)

Figure 3.10. UV-Vis absorbance (left) and emission spectra (right) of PDCI-PTh$_{22^\circ}$C and PDCI-PTh$_{0^\circ}$C before and after heating at 310°C.

Expectedly, heating transformed the absorbance and emission spectra of PDCI-PTh$_{22^\circ}$C and PDCI-PTh$_{0^\circ}$C towards a similar pattern, indicating a rearrangement of the polymer chain towards a common and more delocalized supramolecular structural state. In the absorbance spectra, the relative number of higher energy transition states (associated to more twisted polythiophene chains) decreased, while the longer wavelength absorbance increased (Figure 3.11, left), indicating the conversion of polythiophene segments with shorter conjugation path lengths into longer ones. This is expected as the DSC showed an endothermic transition at 310°C, implying that PDCI-PTh particles absorbed heat for a chemical process. The emission spectra also changed dramatically upon heating. After heating, the emission spectra of both PDCI-PTh$_{22^\circ}$C and PDCI-PTh$_{0^\circ}$C are perfectly superimposed within the range 600 – 800 nm (Figure 3.11, right), and the short wavelength emission $\lambda_{\text{max}}$ between 450 nm become redshifted in comparison to the pre-heated, kinetically trapped states. This also indicates that the short wavelength emission coming from $\lambda_{\text{max}}$
= 450nm for PDCI-PTh_{22^\circ C} and PDCI-PTh_{0^\circ C} are derived from segments of polythiophene conformational subunits which are converted into chains with longer conjugation length by heating. PDCI-PTh_{0^\circ C} had a characteristic decrease in the shortwave emission with concomitant increase in emission in the range 600 – 700 nm which are assigned to as the crystalline domains of semicrystalline polythiophene.

These experiments provide strong support that the lower temperature reactions produce segments of polythiophene chains with more planar and longer conjugation path lengths; resulting in tunable photophysical properties of primary nanostructures. At the same time, the lower reaction temperature does not allow polythiophene chains to rearrange into more crystalline supramolecular structures (involving more than one polythiophene chain) that are more delocalized.

3.2.5. Small Angle X-ray Scattering and X-ray Powder Diffraction

The internal structure of PDCI-PTh nanoparticles was characterized by X-ray and neutron scattering experiments. X-ray Powder Diffraction (XRD) describes the molecular packing of the polymer chains. PDCI-PTh_{22^\circ C} was chosen for comprehensive structural studies due to the more rich photophysical characteristics observed in its photoluminescence spectra. PDCI-PTh_{22^\circ C} showed typical XRD diffraction for crystalline polythiophenes and this type of pattern has been shown in literature to come from herringbone type packing of polythiophene.\textsuperscript{49-50} The broad peak at $2\Theta = 2.5-7.7^\circ$ is indicative of the amorphous regions of polythiophene while the sharp Bragg peaks are assigned to crystalline domains. Therefore, the PDCI-PTh_{22^\circ C} was shown to be a highly ordered semi-crystalline material. The semi-crystalline nature of PDCI-PTh_{22^\circ C} observed by XRD agrees with the distribution of chromophores assigned as segmented polythiophene chains and crystalline polythiophene observed by the emission spectra in Figure 3.6. XRD showed no other, alternative types of nanostructures or packing conformations.
Figure 3.11. X-ray Powder Diffraction Pattern of PDCI-PTh_{22°C}.

PDCI-PTh_{22°C} was also characterized by small angle X-ray scattering (SAXS). The crystalline domains of polythiophene are expected to be significantly denser than the amorphous components and would manifest as high contrast in the x-ray scattering length density (SLD). Thus, the SAXS pattern is expected to reveal the nanostructure of the crystalline polythiophene relative to the lesser dense amorphous polythiophene. At medium q range between 0.1 and 0.02 Å⁻¹, the slope of -2.67 is indicative of a planar sheet like structure such as to the one proposed upon analysis of the SEM data, and it is also expected because of the known sheet like self-assembled structures of polythiophene. Beyond q = 0.02 Å⁻¹, aggregation of the primary nanostructures is apparent by the increased slope to -3.7. The q = 0.02 Å⁻¹ is associated with the length of 31.4 nm and is assigned to a single size dimension of the primary nanostructure, presumably the sheet thickness.
3.2.6. Small-Angle Neutron Scattering Experiments

Neutrons are an ideal tool to differentiate and elucidate the structure of a multiple component nanostructure composed of light elements. Unlike X-rays which interact with the electron cloud (and can be used to differentiate between areas of different electron density), neutrons communicate with the atomic nuclei which results in greater sensitivity for elements such as carbon, hydrogen and nitrogen. Additionally, the scattering length density (SLD) of individual chemical components can be matched with the SLD of the solvent via isotopic labeling, absolving their contrast and scattering signal. In this way, the signal contributions from individual components can be highlighted while attenuating the signal from other chemical species (such as differentiating between PDCI and PTh nanostructures). PDCI-PTh$^{22^\circ \text{C}}$ and PDCI-PTh$^{0^\circ \text{C}}$ particles were suspended in either protonated (h)1,2-dichlorobenzene or deuterated (d)1,2-dichlorobenzene for SANS analysis.
Figure 3.13. Small-angle neutron scattering curves of PDCI-PTh synthesized at 22°C (left), PDCI-PTh at 0°C (right).

For these experiments, nanoparticles with deuterated and protonated PDCI and PTh units were synthesized, and referred to here as (d)PDCI, (h)PDCI, (d)PTh and (h)PTh. All the (d/h)PDCI-PTh particles exhibited a power law scattering at below \( q = 0.02 \) Å\(^{-1}\) and this derives from the aggregation of primary nanostructures with a characteristic size of 31.4 nm such as observed in SAXS studies. This is assigned to thickness of polythiophene sheet nanostructures. The intensity \( I(Q) \) is proportional to the square of the SLD contrast. Because of the low contrast between (d)PDCI (SLD = 4.43) (d)PTh (SLD=4.81) with the solvent (SLD = 4.59), (d)PDCI and (d)PTh is not expected to contribute significantly to scattering signal above the incoherent scattering background (green traces, Figure 3.14). The SLD of (d)PTh was measured experimentally using a control sample (d)PTh nanoparticles that do not contain PDCI component. Protonated components: (h)PDCI (SLD=1.79) and (h)PTh (SLD = 2.48) are expected to result in more than 4 times stronger signal to noise ratio in comparison to their deuterated counterparts. Therefore, only sub-structures containing (h)PDCI or (h)PTh are assumed to affect scattering data.
SAXS and SEM provided strong supporting evidence that polythiophenes form sheet-like primary structures that drive the self-assembly of higher order mesoscopic assemblies. The nanostructure of PTh was investigated using \((d)\text{PDCI-(}h\text{)PTh}\) nanoparticles prepared at 22°C and 0°C. Both samples showed power law scattering below \(q = 0.02\ \text{Å}^{-1}\) and did not show definite scattering slopes that can be measured quantitatively by SANS. This demonstrated that the polythiophene nanostructures had poor SLD contrast with its environment in the supramolecular assembly. Whereas in SAXS, the increased physical density of the crystalline PTh domains produced a stronger scattering signal (versus MeOH as solvent). The increased density of crystalline domains of PTh (vs amorphous domains) in SANS actually lowers the signal. Because the solvent’s SLD is 4.59 and the protonated PTh’s SLD is 2.48, higher PTh density will increase the SLD proportionally, lowering the contrast with the solvent and resulting in less signal to noise ratio versus the amorphous regions of polythiophene.

![Graphs showing Porod slope of (h)PDCI-(d)PTh prepared at 0°C and 22°C.](image)

Figure 3.14. Porod slope of (h)PDCI-(d)PTh prepared at 0°C and 22°C. \(I(Q) \sim q^{(-0.86)}\) for (h)PDCI-(d)PTh\(_{0\degree C}\) and \(I(Q) \sim q^{(-1.04)}\) for (h)PDCI-(d)PTh\(_{22\degree C}\), indicated a 1D structure.

However, protonated PDCI, deuterated polythiophene \((h)\text{PDCI-(}d\text{)PTh}\) prepared at 0°C and \((h)\text{PDCI-(}d\text{)PTh}\) prepared at 22°C showed distinct features by SANS. The \(I(Q)\) was proportional to \(q^{(-0.86)}\) for \((h)\text{PDCI-(}d\text{)PTh}_{100\degree C}\) and \(I(Q)\) was proportional to \(q^{(-1.0)}\) for \((h)\text{PDCI-}
(d)PTh$_{22}^\circ$C between q = 0.006 and 0.15 which strongly indicated a 1D object for PDCI sub-structure in the supramolecular structure (Figure 3.15). Perylene has often been reported to form 1D supramolecular structures in literature.$^{36-39}$ However, the photoluminescence spectra showed that PDCI-PTh nanoparticles exhibited similar NIR emission behaviors as the free PDCI precursor in Figure 3.7, which indicates that the PDCI core is electronically separated from other molecules. Therefore, PDCI is not expected to exist in strongly associated complexes with other PDCI molecules (such as H or J aggregates). More likely, the PDCI may be forming 1D structures due to the living character of KCTP reaction and the 2D symmetry of polythiophene self-assembled structures. As so, PDCI is expected to be randomly distributed in the PDCI-PTh nanoparticles except being forced into 1D structures by the supramolecular assembly of polythiophene 2D structures.

3.3 Conclusion

In conclusion, we described the first example of preparing donor-acceptor conjugated polymer nanoparticles by externally initiated Kumada Catalyst-Transfer dispersion polymerization. The approach allows for using reaction temperature to fine-tune the photophysical properties of conjugated polymer nanoparticles, which affects the conformations of polythiophene and their emission wavelength; also altering their ability to transfer energy efficiently to PDCI. Changing the reaction temperature allowed for tuning emission wavelength of PDCI-PTh nanoparticles between 400-600 nm, and also allows for adjusting NIR emission intensity bands at 775 nm. The obtained particles were dispersible in H$_2$O and a variety of organic solvents without dissolving. Extensive structural characterization studies revealed polythiophene sheet-like nanostructures, which can drive the hierarchical self-assembly at larger length scales, whereas PDCI chromophore formed loosely correlated 1D structures. This method represents a novel fundamental approach for
tuning the properties of donor-acceptor conjugated polymer nanoscale systems and affords a new surfactant- and sonication-free way to engineering materials for organic electronic devices.

3.4 References


CHAPTER 4. EXPERIMENTAL SECTION

4.1. General Considerations

All reactions were performed under an atmosphere of dry nitrogen (unless mentioned otherwise). Column chromatography was performed on silica gel (Sorbent Technologies, 60 Å, 40-63 µm) slurry packed into glass columns. Tetrahydrofuran (THF) and N,N-dimethylformamide (DMF) were dried by passing through activated alumina and molecular sieves, respectively, using a PS-400 Solvent Purification System from Innovative Technology, Inc. The water content of the solvents was periodically controlled by Karl Fischer titration (using a DL32 coulometric titrator from Mettler Toledo). Isopropylmagnesium chloride (2.0 M solution in THF) was purchased from Acros Organics, all other reagents and solvents were obtained from Sigma-Aldrich and Alfa Aesar and used without further purification. Organometallic reagents were titrated with salicylaldehyde phenylhydrazone prior to use. UV-Visible absorption spectra were recorded on Agilent Cary 5000 and Varian Cary 50 spectrophotometers. Fluorescence studies were carried out using a PTI QuantaMaster4/2006SE spectrofluorimeter. 1H NMR spectra were recorded at 400 MHz and are reported in ppm downfield from tetramethylsilane; 31P NMR spectra were obtained at 162 MHz and are reported in ppm relative to 80% aqueous H3PO4 as external standard. Ultrasonication was carried out using a Branson Sonifier 450 (400 W nominal power) operating at 70% power. GPC analysis of polymers was performed with Agilent 1100 chromatograph equipped with two PLgel 5 µm MIXED-C and one PLgel 5 µm 1000 Å columns connected in series, using THF as a mobile phase, and calibrated against polystyrene standards. SEM Images were obtained on a Quanta 3D DualBeam FEG-FIB-SEM instrument. (accelerating voltage of 20 kV). SEM samples were dropcasted from CHCl3 suspensions onto copper substrates. Samples for Total Ion Count Mass Spectrometry were analyzed on a Shimadzu LCMS-8045 system. The column used was a
Shimadzu C18 1.8 μm particle size of dimensions 50 x 4.6 mm. Mobil phases A and B were water and acetonitrile, respectively, each containing 0.1% formic acid. The analytical conditions consisted of: gradient, 95% B (20 min); flow rate, 0.20 mL/min; column temperature, 40°C; detection, APCI(+) and APCI(-); nebulizing gas, N2 at 2 L/min; drying gas, N2 at 10 L/min; heating gas, air at 10 L/min; probe voltage, 4.5 kV (+) and -3.5 kV (-); interface temperature, 220°C; DL temperature, 250°C; heating block temperature, 400°C.

4.2. Synthetic Details

**Perylene-3,4,9,10-tetracarboxylic tetrabutylester (3-2).** 71.19mL BuOH and 1,8-diazabicyclo[5.4.0]undec-7-ene (23.69g, 155.6 mmol) was added to a suspension of commercially available perylene-3,4,9,10-tetracarboxylic dianhydride (15.26g, 38.9 mmol) in 500 mL acetonitrile. The reaction mixture was allowed to stir at 60°C for 1 hour before 71.0 mL of 1-bromobutane was added and brought up to reflux conditions to stir for 24 hours. After cooling to room temperature, the suspension was poured into 1 L ice cold deionized H2O. The precipitate was filtered and dried overnight at 60°C in vacuo to yield 23.07g (91%) orange solid. 1H NMR (CDCl3, 400MHz) δ8.31 (d, 4H), 8.05 (d, 4H), 4.33 (t, 8H), 1.81 - 1.74 (m, 8H), 1.55 - 1.43 (m, 8H), 0.99 ppm (t, J = 7.4 Hz, 12H)

**1,7-dibromo-Perylene-3,4,9,10-tetracarboxylic tetrabutylester (3-3a).** To a 1L round bottom flask was added 3-2 (23.07g, 35.34 mmol), K2CO3 (11.47g, 83.04 mmol) and 285 mL DCM. Br2 (23.04mL, 447.4 mmol) was added dropwise via addition funnel. After 22h, the reaction mixture was quenched with sat. 20% (w/v) Na2S2O3 pentahydrate in deionized H2O. The crude mixture was transferred to a separatory funnel, extracted with DCM and washed several times with deionized water. The organic layer was dried over Na2SO4 and concentrated to yield 27.45g red
viscous product which was dissolved in a minimal amount of dichloromethane (ca. 38mL) then subsequently 610 mL acetonitrile was slowly added. The solution was allowed to sit for 2-3 days to recrystallize. After 3 days, the crystals were filtered and recrystallized again according to the same procedure. After the 2nd recrystallization, the product was filtered, washed with ice cold acetonitrile then dried in vacuo at 60°C to yield 8.23g (28.8%) orange solid. ^1H NMR (CDCl₃, 400MHz)  δ8.95 (d, J = 8.0Hz, 2H), 8.29 (s, 2H), 8.10 (d, J = 8.0 Hz, 2H), 4.36-4.32 (m, 8H), 1.82-1.74 (m, 8H), 1.54-1.45 (m, 8H), 1.02-0.97 ppm (m, 12H).

1,7-dibromoperylene-3,4,9,10-tetracarboxylate dianhydride (3-4). p-Toluenesulfonic acid monohydrate (9.10g 47.9mmol) was added to a solution of regioisomerically pure 3a (7.76g, 9.57mmol) in 250 mL toluene. The reaction mixture was brought up to 100°C. After 48h, the reaction was allowed to cool to room temperature before it was filtered and dried overnight at 60°C in vacuo. The crude solid was then refluxed in CHCl₃ for 8 hours, filtered and washed with CHCl₃ to obtain the final product 4.93g (94%) red solid. ^1H NMR (D₂SO₄, 400MHz) δ9.70 (d, J = 8.32 Hz, 2H), 9.03 (s, 2H), 8.81 (d, J = 8.36 Hz).

N,N'-Di(2,6-diisopropylphenyl)-1,7-dibromoperylene-3,4,9,10-tetracarboxylic bisimide (3-5). 2,6-diisopropylaniline (3.55g, 20.02 mmol) was added to a suspension of compound 3-4 (1.0g 1.82mmol) in 25mL of propionic acid and brought to reflux. Let stir 3 days. The reaction was cooled slightly and filtered hot into a 2L filter flask containing 500 mL ice cold deionized H₂O. Saturated NaHCO₃ was added until the acid is neutralized. Then the product was extracted with EtOAc, washed several times with sat. NaHCO₃ and deionized water, then dried over Na₂SO₄ and concentrated in vacuo. The crude product was purified by column
chromatography on silica gel (eluent: CH$_2$Cl$_2$ to hexanes (5:1)) to afford 723mg (46%) of red powder. $^1$H NMR (CDCl$_3$, 400MHz) δ 9.56 (d, $J$ = 8.12 Hz, 2H), 9.02 (s, 2H), 8.80 (d, $J$ = 8.12 Hz, 2H), 7.52 (t, $J$ = 7.8 Hz, 2H), 7.36 (d, $J$=7.8 Hz, 4H), 2.77-2.70 (m, 4H), 1.19-1.18 (m, 24H). $R_f$ = 0.32.

**2-(2’-thienyl)-5-(trimethylstannyl)thiophene.** Bithiophene (3.33, 20.02 mmol) was dissolved in 35mL dry THF under an inert atmosphere and cooled to -78°C. Subsequently, 13.55 mL of 1.6M $n$-BuLi in hexanes (21.67 mmol) was added dropwise. The solidified reaction mixture was warmed to room temperature slowly over 30 minutes, and then cooled again to -78°C. Afterwards, trimethyltinchloride (4.38g, 22.02 mmol) was added and the reaction mixture was allowed to stir overnight warming up to room temperature. The reaction was quenched with deionized H$_2$O, then THF was removed in vacuo. The crude product was extracted using DCM and washed several times with deionized H$_2$O and brine. The organic layer was dried over Na$_2$SO$_4$ and concentrated in vacuo to yield 5.55 g (84%) of a green oil and used without further purification. $^1$H NMR (CDCl$_3$, 400MHz) δ 7.29-7.27 (m, 1H), 7.22-7.20 (m, 1H), 7.19-7.16 (m, 1H), 7.10-7.08 (m, 1H), 7.03-7.00 (m, 1H), 0.45-0.35 (m, 9H).

$N,N'$-Di(2,6-diisopropylphenyl)-1,7-di(2,2’bithien-5-yl)perylene-3,4,9,10-tetracarboxylate bisimide (3-6). Compound 3-5 (670mg, 0.77 mmol), 2-(2’-thienyl)-5-(trimethylstannyl)thiophene (955mg, 2.90 mmol), Pd(PPh$_3$)$_4$ (15mg, 0.013 mmol), and 90 mL of dry toluene was added to a dry Shlenk flask in air-free conditions and heated to 100°C. Stir overnight. Subsequently, the solvent was removed in vacuo and the crude product was separated by column chromatography using a short plug of silica gel (eluent: CH$_2$Cl$_2$ – hexanes (3:2)) to
yield 554mg (65%) of a black solid. $^1$H NMR (CDCl$_3$, 400MHz) δ8.76 (d, $J = 8.0$ Hz, 2H), 8.48 (t, $J = 9.7$ Hz, 2H), 8.36 (d, $J = 8.0$ Hz, 2H), 7.65 – 6.94 (m, 16H), 2.88 – 2.64 (m, 4H), 1.40 – 0.96 ppm (m, 24H). $R_f$ = 0.29.

$N,N'-$Di(2,6-diisopropylphenyl)-1,7-di(5'-bromo-2,2'-bithien-5-yl)perylene-3,4,9,10-tetracarboxylate bisimide (3-7). To a solution of 3-6 (550mg, 0.53 mmol) in 50mL of anhydrous DMF was added N-bromosuccinimide (188mg, 1.06mmol). The reaction was allowed to stir for 6 hours. Subsequently, the reaction mixture was transferred to a separatory funnel and the product was extracted with ethyl acetate. The organic layer was washed several times with deionized H$_2$O, brine, dried over Na$_2$SO$_4$ and concentrated in vacuo. The crude product was purified by recrystallization in boiling heptane (ca. 20mL) until dissolved and cooling to room temperature. The product was isolated by filtration to yield 386mg black solid. (61%) $^1$H NMR (CDCl$_3$, 400MHz) δ8.77 (s, 2H), 8.43 (d, $J = 8.12$Hz, 2H), 8.38 (d, $J = 8.12$ Hz, 2H), 7.50 (t, $J = 7.88$ Hz, 2H), 7.34 (d, $J = 7.8$ Hz, 4H), 7.31 (d, $J = 3.72$ Hz, 2H), 7.19 (d, $J = 3.72$, 2H), 7.01 (d, $J = 3.84$ Hz, 2H), 6.98 (d, $J = 3.84$ Hz, 2H), 2.78-2.71 (m, 4H), 1.19-1.16 (m, 24H).

Catalytic Initiator (3-8). Compound 3-7 (50mg, 0.04 mmol) and Ni(dppp)$_2$ (110 mg, 0.12 mmol) was dissolved in dry THF (10 mL) and stirred in glovebox for 3 hours. The 4.18mM solution was used as is.

PDCI-PTh nanoparticles. Isopropylmagnesium chloride (0.21 mL of 1.62M solution in THF) was added dropwise to a solution of 2,5-dibromothiophene (100 mg, 0.41 mmol) in 8 mL dry THF at 0°C and in air-free conditions to generate the Grignard monomer solution. After 90
minutes, 1 mL of 4.18mM catalytic initiator solution is injected promptly under ultrasonication and allowed to shear for 10 minutes. Subsequently the reaction mixture was quenched in 24mL of 0.1M solution of HCl in MeOH. The solid suspension was isolated by centrifugation at 3500 RPM for 15 minutes. The resulting particles were washed repeatedly by re-suspending in 25%(v/v) CHCl₃ in MeOH and centrifugation until the supernatant is clear. The resulting solids were dried in vacuo to yield 22.38 mg (58%) maroon powder.

2,6-diisopropylaniline-d₁₉. To a high pressure Parr reactor was added 56 g 2,6-diisopropylaniline, 5.6g 10% wt. palladium on carbon, 5.6g 10% wt. platinum on carbon, and 650mL of D₂O. The Parr reactor was flushed with hydrogen gas for 15 minutes before it was sealed and heated to 150°C, reaching 180 PSI and allowing to stir for 72 hours. After cooling to room temperature, the crude slurry was filtered and the product was extracted from the filtrate using dichloromethane. The organic layer was dried over MgSO₄ and concentrated in vacuo to yield 29g (46%) of a light brown oil.

N,N'-Di(2,6-diisopropylphenyl-d₁₇)-1,7-dibromoperylene-3,4,9,10-tetracarboxylic bisimide. 2,6-diisopropylaniline-d₁₉ (7.76g, 39.52 mmol) was added to a suspension of regioisomerically pure 3-4 (4.35g, 7.9 mmol) in propionic acid (150 ml) and brought to reflux to stir for 3 days. After cooling to room temperature, the reaction was poured into 1L of ice cold deionized H₂O and neutralized with sat. NaHCO₃. Then, the product was extracted using EtOAc, dried over MgSO₄, and the solvent was removed in vacuo to yield a crude solid which was purified by column chromatography (eluent: Hx – DCM (3:4)) to yield 1.57g red solid. The
percent deuteration of alkyl protons in 2,6-diisopropylaniline 92%, 50% in the meta- position and 10% in the para- position due to hydrodebromination. \( R_f = 0.15 \)

**\( N,N'-\text{Di}(2,6\text{-diisopropylphenyl-d}17\)-1,7-di(2,2’bithien-5-yl)perylene-3,4,9,10-tetracarboxylate bisimide}.** In a dry Schlenk flask, \( N,N'-\text{Di}(2,6\text{-diisopropylphenyl-d}17)-1,7\)-dibromoperylene-3,4,9,10-tetracarboxylic bisimide (1.57g, 1.74 mmol), 2-(2’-thienyl)-5-(trimethylstannyl)thiophene (1.70g, 5.16 mmol) and Pd(PPh\(_3\))\(_2\)Cl\(_2\) (80mg, 0.11 mmol) were dissolved in anhydrous toluene (175 mL) under air-free conditions, sealed, heated to 100°C, and allowed to stir 18h. Subsequently, the solvent was removed in vacuo and the product was purified by column chromatography on silica gel (eluent DCM – Hx (3:2)) to afford 733 mg (39.3%) black solid. \( R_f = 0.3 \)

**\( N,N'-\text{Di}(2,6\text{-diisopropylphenyl-d}17)-1,7\)-di(5’-bromo-2,2’-bithien-5-yl)perylene-3,4,9,10-tetracarboxylate bisimide}.** N-bromosuccinimide (243mg, 1.366 mmol) was added stepwise over 10 minutes to a solution of \( N,N'-\text{Di}(2,6\text{-diisopropylphenyl-d}17)-1,7\)-di(2,2’bithien-5-yl)perylene-3,4,9,10-tetracarboxylate bisimide (733 mg, 0.68 mmol) in DMF. After 3 hours, the reaction mixture was transferred to a separatory funnel and extracted using EtOAc. The organic layer was washed several times with deionized H\(_2\)O, brine, and dried over MgSO\(_4\) before the solvent was removed in vacuo. The crude solid was recrystallized by boiling in heptane until dissolved and slowly cooling back to room temperature. The suspension was filtered and dried in vacuo to yield 310 mg (36%) black powder.
**(d)PDCI catalytic initiator.** *N,N’-Di(2,6-diisopropylphenyl-d17)-1,7-di(5’-bromo-2,2’-bithien-5-yl)perylene-3,4,9,10-tetracarboxylate bisimide* (130 mg, 0.11 mmol), Ni(dppp)$_2$ (286 mg, 0.32 mmol) and 26 mL of dry THF was mixed together and allowed to stir 3 hours. The 4.18 mM solution was used as prepared.

**(d)PDCI-(h)PTh$_{100}$ nanoparticles.** 0.69 mL of 1.62M Isopropylmagnesium chloride (1.12 mmol) was added dropwise to a solution of 2,5-dibromothiophene (300 mg, 1.24 mmol) in 24 mL dry THF at 0°. After 90 minutes, 3 mL of the 4.18 mM (d)PDCI dinickel catalytic initiator was injected promptly at the decided reaction temperature. Subsequently, the reaction was quenched in 1 mL of 6.0M HCl solution in MeOH and concentrated by centrifugation at 5000 RPM for 15 minutes. The supernatant was removed and the resulting pellet was washed repeatedly by re-suspending the particles in 25%(v/v)CHCl$_3$ solution in MeOH and concentrating by centrifugation to yield 28 mg (24%) maroon powder.

**(d)PDCI-(d)PTh$_{100}$ nanoparticles.** (d)PDCI-(d)PTh$_{100}$ particles were prepared in the same way as (d)PDCI-(h)PTh$_{100}$ except using 300 mg (1.23 mmol) of 2,5-dibromothiophene-d$_2$ yielding 25 mg (22%) maroon powder.

**PDCI(PdRuPhosI)$_2$ External Initiator (2-1).** RuPhos (244mg, 0.624 mmol, 3 eq), RuPhos Pd G3 95% (307.9mg, 0.367mmol, 2.10 eq), K$_3$PO$_4$ (380mg, 1.83 mmol), 3-7 (225.8mg, 0.175mmol, 1 eq) was added to a 20 mL scintillation vial with 0.8 mL H$_2$O and 8 mL dry THF. The reaction was heated to 50°C, for 1.5h. After, the reaction was allowed to cool to room temperature, and the organic layer was precipitated into 5:1 hexanes:THF. The solids were
collected by centrifugation in inert atmosphere. The crude powder was dissolved in dry THF and reprecipitated again in 5:1 hexanes:THF to result in a brown powder.

**General procedure for preparing conjugated polymers using 2-1 as the external initiator.** To a 20 mL scintillation vial was added 2-1 (9.64mg, 0.0039mmol), K$_3$PO$_4$ (320mg, 1.5mmol), silver sulfate (78mg, 0.25 mmol), 14mL THF and 0.5 mL H$_2$O. Mix until K$_3$PO$_4$ dissolves. After, add the monomer as a solution in 2 mL THF). Stir for 96 hr at room temperature.

**PDCI-PPP.** Following the general procedure, 55mg of 4-bromo-2,5-dihexylphenylboronic acid MIDA ester (0.114 mmol, 29 eq) was added to the solution of external initiator. After 96h, the reaction was quenched with 3 mL 6N HCl and extracted using CHCl$_3$. The organic layer was washed with brine and deionized H$_2$O repeatedly. The organic layer was dried over sodium sulfate and concentrated in vacuo. The crude polymer was dissolved in minimal amount of CHCl$_3$ and reprecipitated in acetone. The precipitates were collected by centrifugation and washed repeatedly with acetone and MeOH to yield a green solid (27.8mg, 86% yield).

**PDCI-PF.** Following the general procedure, 32.2 mg of 7-bromo-9,9-dioctylfluorene-2-lboronic acid MIDA ester (0.0515 mmol, 13.2 eq) was added to the solution of external initiator. After 96h, the reaction was quenched with 3 mL 6N HCl and extracted using CHCl$_3$. The organic layer was washed with brine and deionized H$_2$O repeatedly. The organic layer was dried over sodium sulfate and concentrated in vacuo. The crude polymer was dissolved in minimal amount of CHCl$_3$ and reprecipitated in acetone. The precipitates were collected by centrifugation and washed repeatedly with acetone and MeOH to yield a green solid (11.2mg, 46.4% yield).
**PDCI-PT.** Silver sulfate was not used for preparing PDCI-PT. Otherwise, following the general procedure, 50 mg of 2-bromo-3-hexylthiophene-5-boronic acid MIDA ester (0.124 mmol, 31.4 eq) was added to the solution of external initiator. After 96h, the reaction was quenched with 3 mL 6N HCl and extracted using CHCl₃. The organic layer was washed with brine and deionized H₂O repeatedly. The organic layer was dried over sodium sulfate and concentrated in vacuo. The crude polymer was dissolved in minimal amount of CHCl₃ and reprecipitated in acetone. The precipitates were collected by centrifugation and washed repeatedly with acetone and MeOH to yield a green solid (21.4mg, 86.5% yield).

4.3. References

APPENDIX A: PERMISSIONS

Title: Advances in Living Anionic Polymerization: From Functional Monomers, Polymerization Systems, to Macromolecular Architectures
Author: Akira Hirao, Raita Goseki, Takashi Ishizone
Publication: Macromolecules
Publisher: American Chemical Society
Date: Mar 1, 2014
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Title: Identifying the missing link in catalyst transfer polymerization

Author: Weiying He et al

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APPENDIX B: Nuclear Magnetic Resonance Spectroscopy
LIST OF REFERENCES


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VITA

Peter Kei was born in 1988 in Hong Kong. He moved to the U.S. in 1996 and grew up in Quincy, MA. After graduating from North Quincy High School, he obtained his Associates in Science Degree at Quincy College. In 2010, he enrolled in Bridgewater State University as a Biochemistry major. There, he began his research career in the laboratory of Prof. Stephen A. Waratuke, conducting multistep synthesis of asymmetric carbamate ligands and investigated titanium catalyzed hydroamination reactions using them. After graduating with a Bachelor of Science, Peter moved to Baton Rouge in 2014 to join the laboratory of Prof. Evgueni Nesterov. Soon after, Peter received an Honorable Mention by the National Science Foundation Graduate Research Fellowship Program. He also received U.S. Department of Energy Office of Science Graduate Student Research Award, which supported his studies at Oak Ridge National Laboratory in 2017. In 2018, Peter moved along with the laboratory of Prof. Evgueni Nesterov to Northern Illinois University. His research contributions include development of controlled polymerizations for the preparation of well-defined conjugated polymer architectures. Peter is a candidate for the Doctor of Philosophy in Chemistry. His dissertation is entitled: “Hierarchical Conjugated Polymer Systems Prepared by Controlled Chain-Growth Polymerization”. 