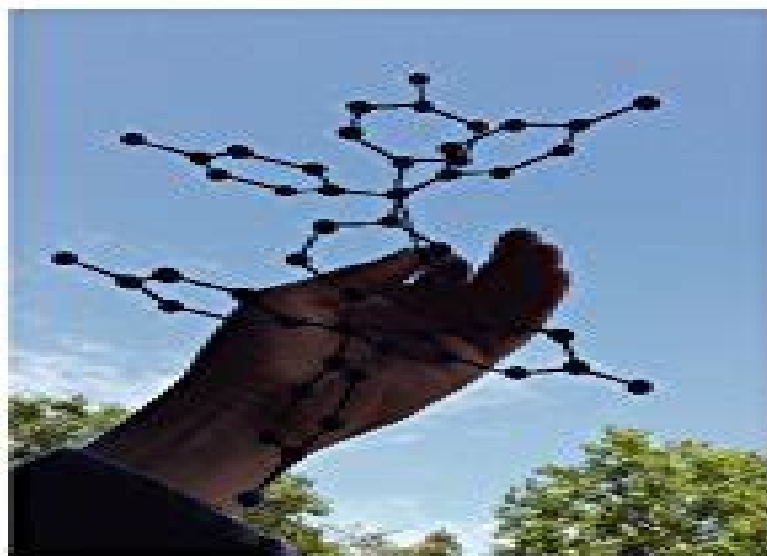


Alexandra Schade

Synthese *pseudo*-oktaedrischer
Hexaphenyl-*p*-xylol-Derivate als rigide Bausteine zur
Darstellung supramolekularer Netzwerke sowie deren
potentielle Anwendung als selektive Gasspeicher



Beiträge zur organischen Synthese
Hrsg.: Stefan Bräse

Λογος

Pseudo Oktaedrischer Hexaphenyl P Xylol Derivate Darstellung Supramolekularer Potentielle

Alexandra Maria Schade



Pseudo Oktaedrischer Hexaphenyl P Xylol Derivate Darstellung Supramolekularer Potentielle:

Synthese pseudo-oktaedrischer Hexaphenyl-p-xylol-Derivate als rigide Bausteine zur Darstellung supramolekularer Netzwerke sowie deren potentielle Anwendung als selektive Gasspeicher Alexandra Maria Schade, 2015-07-30 Der Bereich der Materialwissenschaften hat sich in den letzten zwei Dekaden insbesondere der Forschung und Entwicklung neuartiger poröser Materialien gewidmet. Dieses interdisziplinäre Fachgebiet der Nanotechnologie beschäftigt sich vor allem mit dem Design progressiver Materialien. Hierzu zählen neben kristallinen Netzwerken MOFs und COFs auch amorphe Polymere MOPs. Aufgrund ihrer herausragenden Eigenschaften wie der selektiven Gasspeicherung oder exzellenten Separationsfähigkeit finden solche funktionellen Nanomaterialien bereits erste industrielle Anwendung. Unter diesem Aspekt wurden im Rahmen der Arbeit zahlreiche neuartige Derivate des pseudo oktaedrischen Hexaphenyl p xylol HPX synthetisiert sowie optimiert und erstmals zum Aufbau poröser Netzwerke herangezogen. Die Vernetzung der dreidimensionalen HPX Bausteine zu Mikroporösen Organischen Polymeren erfolgte unter Zuhilfenahme etablierter Reaktionen wie z.B. der HUISGEN 1,3-dipolaren Cycloaddition, Click-Reaktion, verschiedener Kohlenstoff-Kohlenstoff-Kupplungsreaktionen sowie Reaktionen zur Trimerisierung von Nitrilen. Die amorphen Netzwerke gelten aufgrund ihrer Eigenschaften als potentielle Aspiranten im Bereich der selektiven Gasspeicherung.

Oberflächenbeschichtungen von Silizium, HOPG und Graphitanoden mittels Thiol-En/In Click-Chemie, deren Charakterisierung mit XPS und ToF-SIMS sowie elektrochemische Untersuchung Dominique Moock, 2018-08-18 Eine der größten Herausforderungen des 21. Jahrhunderts ist die Umstellung von fossilen Brennstoffen auf erneuerbare Energien. Wegen den natürlich auftretenden Schwankungen sind geeignete Speichersysteme zum Ausgleich entscheidend. Lithium-Ionen-Batterien spielen hierbei eine wichtige Rolle, da sie hohe Energiedichten aufweisen. Jedoch sind noch nicht alle Abläufe in diesen Batterien vollständig verstanden, wie beispielsweise der irreversible Kapazitätsverlust in den ersten Zyklen sowie die Zyklenstabilität. Bekannt ist, dass die Ausbildung einer sog. SEI (Solid Electrolyte Interface) auf der Graphitelektrode die Verwendung von Graphit als Anodenmaterial erst ermöglicht, jedoch auch für den irreversiblen Kapazitätsverlust verantwortlich ist. Im Rahmen dieser Arbeit wurden sowohl auf Modelloberflächen HOPG und Silizium sowie realen Graphitelektroden Ankergruppen der Diazoniumsalzchemie angebracht. Hierzu wurden unterschiedlich funktionalisierte Diazoniumsalze synthetisiert, welche als Ankergruppen zum weiteren Schichtaufbau mittels Thiol-En/In Click-Chemie auf den Oberflächen dienten. Die so erhaltenen modifizierten Oberflächen wurden mit Hilfe von Röntgenphotoelektronenspektroskopie (XPS) und Flugzeitmassenspektrometrie (ToF-SIMS) analysiert sowie anschließend hinsichtlich ihrer Eignung als künstlich erzeugte SEI elektrochemisch in einer Lithium-Ionen-Batterie untersucht.

Synthese organischer Bausteine für kovalent gebundene, poröse Netzwerke über Nitroxid-Austausch-Reaktion Isabelle D. Wessely, 2018 Ein entscheidender Punkt bei der Umstellung von fossilen Brennstoffen auf erneuerbare Energien ist die Entwicklung neuer Speichersysteme für Gase und eine Weiterentwicklung im Bereich der Solarzellen/Brennstoffzellen.

oder Speichermedien für elektrische Energie. Poröse Materialien mit geringer Dichte, großer Oberfläche, hoher Stabilität und periodischer Struktur sind hierfür von großem Interesse, und sogenannte COFs könnten hierbei die Materialien der Wahl werden. Die Synthese kristalliner COFs erfordert einen reversiblen Reaktionsmechanismus, weswegen die Nitroxid-Austausch-Reaktion einen neuen Ansatz darstellen kann. Für diesen neuen Syntheseansatz gelang es im Rahmen dieser Arbeit, einige geeignete Bausteine darzustellen. Zum einen konnten mehrere Tri- bzw. Tetranitroxide über die Verknüpfung mehrfach funktionalisierter Tetraphenylmethan- bzw. Benzolderivate mit geeigneten Nitroxidreagenzien über Click-Chemie oder Kreuzkupplungsreaktionen dargestellt werden. Zum anderen konnten für die Reaktionspartner in der Nitroxid-Austausch-Reaktion ein Modellsystem an einem linearen Dialkoxyamin für die Synthese mehrfach substituierter Alkoxyamine entwickelt werden, welches auf verschiedene aromatische Grundgerüste übertragen werden konnte und gleichzeitig eine zusätzliche Funktionalisierung in α -Position zur Alkoxyaminfunktionalität erlaubt. Zusätzlich wurde in dieser Arbeit erstmals die Nitroxid-Austausch-Reaktion mittels EPR-Spektroskopie verfolgt, um Erkenntnisse über das Zusammenspiel von Temperatur, Reaktionszeit und molaren Verhältnissen der an dem Austausch beteiligten Komponenten zu gewinnen, um so ein besseres Verständnis des Netzwerkaufbaus zu entwickeln.

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