POLYCARBOXYLIC ACIDS

via

CATALYTIC HYDROCARBOXYLATION OF POLYBUTADIENES

a thesis presented by

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to the

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THE DEGREE OF DOCTOR OF PHILOSOPHY

St. Andrews September 1991
The BIT
DECLARATION

I, Narayanan Pondicherry, hereby certify that this thesis has been composed by myself, that it is a record of my own work, and that it has not been accepted in partial or complete fulfilment of any other degree or professional qualification.

St. Andrews Date 27.9.1991

I was admitted to the Faculty of Science of the University of St. Andrews under Ordinance General No.12 on 1st October 1988 and as a candidate for the degree of Ph.D on 1st September 1989.

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I hereby certify that the candidate has fulfilled the conditions of the Resolution and Regulations appropriate to the Degree of Ph.D.

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Dedicated with gratitude to my grandparents, parents, sister and brother
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ABBREVIATIONS

DIOP  2,3-o-Isopropylidene-2,3-dihydroxy-1,4-bis(diphenylphosphino)butane
DIPHOL  (S,S)-trans-4,5-bis[(5H-dibenzophospholyl)methyl]-2,2-dimethyl-1,3-dioxolan
dpb  1,4-bis(diphenylphosphino)butane
dpf  1,1'-bis(diphenylphosphino)butane
DMF  Dimethyformamide
MIBK  Methyl isobutyl ketone
PPh₃  Triphenylphosphine
PPh₃O  Triphenylphosphine oxide
rtm  Room temperature
THF  Tetrahydrofuran
ABSTRACT

Polybutadiene having varying degrees of 1,4 and 1,2 units (0 - 100% cis, 14 - 60% trans, 20-88% pendant carbon-carbon double bonds) have been reacted with carbon monoxide in the presence of water and palladium based catalysts to give new polymers in which the double bonds are hydrocarboxylated. When using [PdCl₂(PPh₃)₂]/SnCl₂ as catalysts, very high regiospecificities can be obtained with only the double bonds pendant from the chain being hydrocarboxylated. When using PdCl₂ as catalyst, in the presence of CuCl₂ and O₂, hydrocarboxylation takes place both at the pendant carbon-carbon double bonds and back bone carbon-carbon double bonds. Complete hydrocarboxylation of all the carbon-carbon double bonds in polybutadiene can be carried out by sequential catalytic reaction i.e., catalysis by [PdCl₂(PPh₃)₂]/SnCl₂ followed by (recatalysis) [PdCl₂(PPh₃)₂]/PPh₃. A new method based on ¹H and ¹³C n.m.r. has been developed to characterise polybutadienes and their hydrocarboxylated products.

The hydrocarboxylation reaction catalysed by [PdCl₂(PPh₃)₂]/SnCl₂ is truly homogeneous although catalytic decomposition is a problem because their exists a small temperature window between the onset of catalysis and the onset of catalytic decomposition. Unlike the starting materials the products, which are solids with melting points well above the room temperature are soluble in organic solvents or dilute base and have been shown to have corrosion inhibition and wood preservation properties.
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Chapter 1

Hydrocarboxylation of alkenes and polybutadienes
(background and literature review)

1.1 Introduction

This chapter reviews hydrocarboxylation reactions on alkenes in general and hydrocarboxylation modification reactions on polymers.

Water-soluble synthetic polymers are a family of materials that have been developed commercially and studied scientifically at an accelerating pace in recent years. Partly this is a reflection of the increasing diversity in the application of water-soluble polymers as a group in mineral processing, oil-recovery, etc.

Certain water-soluble synthetic polymers containing carboxylic acid groups have been found to exhibit antitumour and anti-viral effects. They are also used in the water treatment industries to prevent the deposition of inorganic salts on metal surfaces and to reduce metallic corrosion. These products have found major outlets in industrial cooling systems and desalination plants. Presently they are being widely used as surface coating materials.

The chemical modification of polymers is a post polymerisation process which is used in certain situations:

(i) to improve and optimise the chemical and mechanical properties of existing polymers, or

(ii) to introduce desirable functional groups into the polymer.

Many chemical reactions with polymers are possible, as reflected in the breadth and depth of the associated literature. The chemical
modification of unsaturated polymers via homogeneous catalysis offers a potentially useful method for the introduction of desirable functional groups to the polymer chains. The functionalisation involves hydroxylation, hydroformylation, hydrosilylation, hydrogenation, epoxidation, and various other chemical reactions involving synthetic and natural polymers.

The application of homogeneous catalysis can thus become a powerful tool in the synthesis of a wide range of polymeric compounds of industrial interest. One advantage of homogeneous catalysis over heterogeneous catalysis is its greater ability effectively to catalyse reactions on the polymeric substrates. The slow rates and low yields observed when heterogeneous catalysts are used in such reactions are generally ascribed to the difficulty of orientating a long polymer chain on the immobile active site. In contrast, a homogeneous catalyst offers catalytic mobility and circumvents such orientation problems, thus increasing their potential for diffusion on the polymer. (Homogeneous catalysis as used here implies that the catalytically active species is molecularly dispersed (solvated) in the polymer or a cosolvent. Heterogeneous catalysis implies that the catalytically active species and the substrates are in two different phases, e.g., solid catalyst, liquid substrates).

One of the ways to synthesise speciality polymers with certain desirable functional groups is by the polymerisation of monomers with the desirable functional groups. These polymers all have their functional groups directly attached to the main chain in the polymers.
However there are often a number of difficulties associated with polymerisation of the monomers such as

(i) difficulty in polymerisation
(ii) side reactions during polymerisation
(iii) difficulty in the preparation of desirable monomers
(iv) and, in the case of copolymerisation, unfavourable reactivity ratios.

Alternatively, catalytic chemical reactions on unsaturated polymers in which the desirable functional groups are introduced into available unsaturated sites in polymers, such as polybutadiene, can be employed to synthesise certain speciality polymers. Here, the functional group can be separated from the main chain by a flexible spacer group. Polybutadienes are of special interest in this context, since very regiospecific polybutadienes are available with desirable molecular weights and molecular weight distributions. The relative ratios of the three types of double bond (cis or trans in the backbone and pendant) can be controlled. The products thus obtained by the introduction of desirable functional groups have the same regiospecificity and chain length properties, provided that side reactions such as crosslinking and isomerisation, for example, are avoided.

Our interest in functionalised polymers is in the area of corrosion inhibition, calcium carbonate scale control, wood preservation etc., of which carboxylated polymers are finding most application. Thus, in this chapter we will review hydrocarboxylation reaction on alkenes in general and hydrocarboxylation modifications on polymers, particularly polybutadienes. Publications which are either important or relevant to our work have been discussed in detail.
Hydrocarboxylation refers to the reaction of alkenes with carbon monoxide and water in the presence of catalysts to give mixtures of straight chain and branched chain carboxylic acids (Equation 1.1).

\[ \text{Catalyst} \quad \text{CH}_2=\text{CH} + \text{CO} + \text{H}_2\text{O} \rightarrow \text{CH}_3\text{CH}_2\text{COOH} + \text{CH}_3\text{COOH} \]

Equation 1.1

If alcohols are used in place of water, then esters are produced. These hydrocarboxylation reactions are also known as Reppe reactions after the originator. In his original process Reppe used a nickel carbonyl catalyst for the synthesis of propanoic acid from ethene, at 270°C and carbon monoxide pressure of 200 atm (Equation 1.2).

\[ \text{CH}_2=\text{CH}_2 + \text{CO} + \text{H}_2\text{O} \xrightarrow{[\text{Ni(CO)}_4]} \text{CH}_3\text{CH}_2\text{COOH} \]

Equation 1.2

The reactions of alkenes with carbon monoxide and water (or alcohols) to form acids (or esters) was already known before Reppe's work. The earlier process proceeded only at high pressure (700-900 atm CO) and at high temperature (about 300°C) in the presence of catalysts such as phosphoric acid, heteropolyacids, boron trifluoride, and metal halides. Operation at such high temperature i.e., 250-300°C had certain inherent disadvantages. Carbonylation of alkenes, particularly above ca 150°C is always accompanied by side reactions such as the water-gas shift reaction (Equation 1.3).
CO + H_2O \rightarrow CO_2 + H_2

Equation 1.3

The hydrogen that is formed hydrogenates unreacted alkene and reacts with carbon monoxide and the alkene to form aldehydes, alcohols, and other products. Isomerisation, polymerisation, resinification and cleavage of the alkene also proceed readily at higher temperature. Hardy described experiments in which a mixture of ethene, carbon monoxide and steam were passed through H_3PO_4 at 290-300°C and 450 atmospheres, where, for every molecule of ethene converted into acid, approximately three were hydrated, and two were polymerised. Finally, the severe conditions required lead to the danger of corrosion of the walls of the reactor. This situation has led to a search for newer catalysts that would act at lower temperature than the conventional catalyst and would at the same time be relatively cheap, readily accessible, and offer the possibility of higher selectivity.

The development of homogeneous catalysis began over fifty years ago with the pioneering work of Reppe. Reppe and coworkers hypothesised that hydrocarboxylation of alkenes occurs as shown in Scheme 1.1. In this an alkene, RCH=CH_2, reacts with carbon monoxide to form intermediate [1]. This may be attacked either at (a) or (b), to yield straight chain and/or branched chain carboxylic acids respectively.
Depending on the amount of catalyst used (Reppe normally used the nickel complex \([\text{Ni(CO)}_4]\)), distinctions were made between a stoichiometric (Equation 1.4), and a catalytic conversion (Equation 1.5).

**Stoichiometric**

\[
4\text{CH}_2=\text{CH}_2 + \text{Ni(CO)}_4 + 4\text{C}_2\text{H}_5\text{OH} + 2\text{HCl} \rightarrow 4\text{CH}_2(\text{COOC}_2\text{H}_5)\text{CH}_3 + \text{NiCl}_2 + \text{H}_2
\]

Equation 1.4

**Catalytic**

\[
\text{CH}_2=\text{CH}_2 + \text{CO} + \text{C}_2\text{H}_5\text{OH} \xrightarrow{\text{Ni salt}} \text{CH}_2(\text{COOC}_2\text{H}_5)\text{CH}_3
\]

Equation 1.5
In each case, a compound possessing an active hydrogen atom, e.g. water or alcohol, is also present.

The stoichiometric method is carried out with nickel carbonyl and an acid. Less stringent conditions are necessary compared with the catalytic process, temperatures \( \text{ca} 160^\circ\text{C} \) at pressures of about 50 atmospheres of carbon monoxide are normally employed.

The catalytic method involves the formation of nickel carbonyl \textit{in situ} from carbon monoxide and a nickel salt such as \( \text{NiCl}_2 \), \( \text{NiI}_2 \), \( \text{[Ni(CO)]}_4 \), \( \text{[(C}_2\text{H}_5\text{CO})_2\text{Ni]} \), etc. The reaction temperatures are around 250\(^\circ\text{C} \), with carbon monoxide pressures of 200 atmospheres.

Once a moderately satisfactory technique for affecting the reaction had been evolved, Reppe applied the two methods to a variety of compounds with little variation in conditions. Over the years, Reppe and co-workers studied hydrocarboxylation reactions on unsaturated substrates quite extensively and published a number of papers and patents\(^4\). Since this discovery, the area of hydrocarboxylation has been one of frantic activity to become one of the most exciting and controversial new areas of chemistry. Many synthetically useful applications have been reported\(^2\).

Various classifications for the hydrocarboxylation reactions reviewed below are feasible. No method is pre-eminent as any single catalytic reaction can be accomplished by more than one transition metal complex and, not infrequently, a reaction may give rise to a multitude of products. For the purpose of this review, a classification in terms of the type of substrate, i.e., monoalkenes, di- and tri-alkenes, and functionalised alkenes has been adopted.

Most of the elements that have proved valuable in forming complexes suitable for catalytic homogeneous carbonylation appear in group VIII (now groups 8-10) of the periodic table. Salts and complexes
of Pd, Pt, Ru, Rh, Ir, Fe, Ni, and Co have been used as catalysts. For hydrocarboxylation reactions, complexes of Co, Ni and Pd have been most extensively studied, although strenuous efforts are being made to find catalysts from among the less expensive metals.

The catalytic activity of the transition metal salts and complexes is the result of a delicate balance of valence states and strengths of chemical bonds. Too strong a bond between alkene and the transition metal catalyst results in a stable compound showing no catalytic activity. Similarly, there is no catalytic activity if the reaction between alkene and the transition metal cannot occur. Not only must the alkene be accommodated by the transition metal, but also the carbon monoxide must be able to bind if transfer of carbon monoxide to the substrate is to occur. However, it must be noted that the products of catalytic hydrocarboxylation may themselves inhibit or show no catalytic activity. For example, [Pd(CO)(PPh₃)₃]₆, isolated from the reaction mixture during hydrocarboxylation of cyclohexene with carbon monoxide and ethanol, in the presence of [PdCl₂(PPh₃)₂] and a two molar excess of triphenylphosphine as the catalyst, showed no catalytic activity at all in the hydrocarboxylation of alkenes. Thus the catalytic activity depends on a delicate balance of energies with regard to the binding of the substrates.

The efficiency and the product distribution of the hydrocarboxylation reactions depends markedly on a number of factors: The nature of the nucleophile i.e., water or alcohol, the structure of the alkene, the catalyst, the solvent, and the physical parameters of the reaction such as variation of temperature and carbon monoxide pressure.

In the following section we will discuss the role of these various parameters and their importance in homogeneous hydrocarboxylation
reactions with respect to various substrates (i.e., monoalkenes, dialkenes, functionalised alkenes, etc.).

1.2 Hydrocarboxylation of 1-alkenes

Generally, 1-alkenes on hydrocarboxylation give rise to an isomeric mixture of carboxylic acids or their esters. As mentioned earlier the most active catalysts for hydrocarboxylation are found in the salts and complexes of Ni, Co, Pd, Pt and Rh complexes. A general observation on the characteristics of these metal catalysed hydrocarboxylation reactions of 1-alkenes is presented in Table 1.1.

In general, it has been found that nickel catalysed reactions with 1-alkenes give branched acids. Cobalt catalysts tend to cause more side reactions i.e. there is a greater tendency to form ketones. However, the presence of pyridine type bases markedly improves the selectivity to straight chain acids. In the case of palladium catalysts, regioselectivity very much depends on a number of factors such as ligands, the nature of R in R-CH=CH₂, solvent, hydrogen donor and physical parameters. When using rhodium catalysts, regioselectivity can be increased by using a promoter. Iridium catalysed reactions always tends to form terminal acids with 1-alkenes.
Table 1.1

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>Promoter</th>
<th>Conditions</th>
<th>Yield (linear)</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ni</td>
<td>X₂ or HX</td>
<td>250-300°C</td>
<td>80-90%</td>
<td>7,8</td>
</tr>
<tr>
<td></td>
<td></td>
<td>200-400 atm</td>
<td>(50%)</td>
<td></td>
</tr>
<tr>
<td>Co</td>
<td>HI, Py</td>
<td>180-220°C</td>
<td>70-95%</td>
<td>9,10,11</td>
</tr>
<tr>
<td></td>
<td></td>
<td>200 atm</td>
<td>(70%)</td>
<td></td>
</tr>
<tr>
<td>Pd</td>
<td>R₃P and HCl</td>
<td>80-100°C</td>
<td>95%</td>
<td>12,13,14,15</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30-700 atm</td>
<td>(70-90%)</td>
<td></td>
</tr>
<tr>
<td>Rh</td>
<td>HX</td>
<td>150-220°C</td>
<td>95%</td>
<td>16</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30 atm</td>
<td>(50-60%)</td>
<td></td>
</tr>
<tr>
<td>Ir</td>
<td>HX</td>
<td>150-220°C</td>
<td>95%</td>
<td>17</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>(70%)</td>
<td></td>
</tr>
</tbody>
</table>

1.2.1 Nickel catalysed reactions.

The hydrocarboxylation of 1-alkenes usually proceeds under aqueous acidic conditions in the presence of tetracarbonyl nickel or of a nickel salt such as nickel(II) iodide. Hydrocarboxylation of ethene using a three component catalytic system, namely [Ni(NH₃)₆Cl₂], NaI, and Raney nickel yielded, upon distillation, propanoic acid and tetracarbonyl nickel (Equation 1.6).

\[
\text{[Ni(NH₃)₆Cl₂], NaI, Raney Ni} \\
\text{CH₂=CH₂ + CO + H₂O} \xrightarrow{280°C, 250 \text{ atm}} \text{CH₃CH₂COOH + Ni(CO)₄}
\]

Equation 1.6
The presence of \([\text{Ni}(\text{CO})_4]\) proves that this species is formed under the reaction conditions. It has been suggested that the catalytic species in all the reactions catalysed by \([\text{Ni}(\text{CO})_4]\) is the hydrido nickel carbonyl complex (Equation 1.7). Nickel compounds must be used in the presence of promoters like hydrogen halides\(^8,19\).

\[
\text{[Ni(CO)\textsubscript{4}]} + \text{HX} \xrightleftharpoons[\text{CO}]{\text{CO}_\text{2}} \text{[NiHX(CO)\textsubscript{2}]} \\
\text{Equation 1.7}
\]

The hydrocarboxylation reaction mechanism\(^2\) is represented in Scheme 1.2.

The reaction catalysed by \([\text{Ni}(\text{CO})_4]\) is thought to be analogous to that catalysed by \([\text{HCo}(\text{CO})_4]\).

An alternative mechanism involving the formation of a nickel cyclobutanone intermediate (2) that reacts first with hydrogen halides has been proposed\(^2\) and is represented in Scheme 1.3.
Intermediate (2) is related to a ruthenium complex which has been formed by reaction of complexed carbon monoxide and alkyne, but (2) does not have the endocyclic double bond (Equation 1.8). No direct evidence for these proposed mechanisms has been established.

In nickel catalysed hydrocarboxylation reactions, ketone formation is sometimes observed. Ethene yields appreciable amounts of diethyl ketone in addition to propanoic acid\textsuperscript{22} (Equation 1.9).
\[
\text{CH}_2=\text{CH}_2 + \text{CO} + \text{H}_2\text{O} \xrightarrow{\text{[Ni(CO)]}_4, 300^\circ\text{C}, 450 \text{ atm}} \text{CH}_2(\text{COOH})\text{CH}_3 + \text{C}_2\text{H}_5\text{COC}_2\text{H}_5
\]

Equation 1.9

This is in accord with competition between water and alkene for reaction with an acyl-nickel intermediate (Scheme 1.4).

When [Ni(CO)]\(_4\) is used as the catalyst the reaction rate is dependent upon the pressure of carbon monoxide. An increase in carbon monoxide pressure increases the reaction rate; decrease in carbon monoxide pressure decreases the reaction rate.

Scheme 1.4
1.2.2 Cobalt catalysed reactions

Being good hydroformylation catalysts, cobalt complexes, in general, tend to cause more side reactions than nickel catalysts i.e., ketones are formed (Equation 1.10).

\[
\text{CH}_2=\text{CH}_2 + \text{CO} + \text{H}_2\text{O} \xrightarrow{234^\circ\text{C, 660 atm}} \text{Co propionate} \rightarrow \text{C}_2\text{H}_5\text{COOH} + \text{C}_2\text{H}_5\text{C}-\text{C}_2\text{H}_5 \\
(45\% \text{ yield}) \quad (14.5\% \text{ yield})
\]

Equation 1.10

With increasing temperature, the yields of acids decreases, and yields of ketone derivatives increases. With increase in carbon monoxide pressure the reaction rate is accelerated and the amounts of ketone derivatives decreases. The rate and selectivity to straight chain carboxylic acid derivatives is markedly improved by the presence of pyridine type bases. Thus hydrocarboxylation of 1-alkenes produces 80% of the linear acid.\textsuperscript{10,11}

The basicity and structure of the promoter are important. The effects\textsuperscript{17} are summarised in Table 1.2.

<table>
<thead>
<tr>
<th>Promoter</th>
<th>Relative rate of catalysis</th>
</tr>
</thead>
<tbody>
<tr>
<td>pyridine</td>
<td>10</td>
</tr>
<tr>
<td>α-picoline</td>
<td>1</td>
</tr>
<tr>
<td>β-picoline</td>
<td>10</td>
</tr>
<tr>
<td>2,6-Lutidine</td>
<td>0</td>
</tr>
<tr>
<td>Triethylamine</td>
<td>0</td>
</tr>
</tbody>
</table>

Table 1.2
It can be seen that a strong base which is sterically hindered completely stops the catalytic reaction. The most reasonable interpretation of the pyridine type bases is that they attack the intermediate acyl cobalt species, cleaving the cobalt-carbon bond, and giving rise to a pyridinium cation which is solvated to carboxylic acid or ester and the pyridinium cation.

Foster, Hershman and Morris\textsuperscript{17} suggested that pyridine acts as both an inhibitor and a promoter in the reaction. The Heck and Breslow\textsuperscript{23} mechanism (Scheme 1.5) agrees with the former if one postulates that the concentration of the acyl cobalt carbonyl intermediate is extremely small.

\[ \text{Scheme 1.5} \]
X-ray\textsuperscript{17} structural investigations carried out on a solid complex crystallised from a hydroesterification reaction mixture showed the compound to be $[\text{pyridine}_2\text{H}]^+[\text{Co(CO)}_4]$\textsuperscript{-}. The cation consists of a proton bonded to the nitrogen of the pyridines (Figure 1.1).

\begin{center}
\includegraphics[width=0.2\textwidth]{figure1.png}
\end{center}

\textbf{Figure 1.1}

It appears that the majority of the cobalt is in this catalytically inactive form (Equation 1.11).

\[
[\text{Py}_2\text{H}]^+ [\text{Co(CO)}_4]^+ \overset{\text{pyridine}}{\rightleftharpoons} [\text{HCo(CO)}_4]
\]

\textbf{Equation 1.11}

1.2.3 Palladium catalysed reactions

After the epoch making introduction in 1960 of the Wacker process\textsuperscript{24} by which acetaldehyde is produced from ethene using $\text{PdCl}_2$ as a catalyst (Equation 1.12), palladium complexes were recognised as
versatile catalysts for hydrocarboxylation reactions.

\[
CH_2=CH_2 + \frac{1}{2} O_2 \xrightarrow{\text{PdCl}_2/\text{CuCl}_2} CH_3CHO
\]

Equation 1.12

Alper and coworkers\(^{25}\) reported a modified PdCl\(_2\)-CuCl\(_2\) catalytic system that operated under mild conditions (room temperature and pressure), as an effective catalyst for hydrocarboxylating 1-alkenes with great regioselectivity to branched acids. Alper and coworkers in a patent\(^{26}\) extended this method for hydrocarboxylating polymers. Further detailed discussion is carried out in Chapter 4. A U.S. patent\(^{27}\) similarly reports hydrocarboxylation of cis- and trans-polybutadiene in the presence of PdCl\(_2\) as catalyst. However no details were given about the methods, or yields when using PdCl\(_2\) as catalyst.

The greatest advantage in using complex palladium catalysts is that they are very active even at low concentration and low temperatures. The activity of palladium catalysts are highly sensitive to the various factors governing the reaction. The efficiency and product distribution of the reaction depends markedly on the nature of the nucleophile i.e., water or alcohol, the structure of the alkene and the phosphine ligand, and other reaction variables such as solvent, temperature and carbon monoxide pressure.

For 1-octene hydrocarboxylation, \(n\) to iso ratios from 3.0 to 0.5 are observed depending on temperature, carbon monoxide pressure, nature and amount of added phosphines and additional reagents\(^{13}\). Consiglio and Marchelli reported\(^{28}\) that when triphenylphosphine was used as ligand attached to the palladium species, the branched chain carboxylic acid was obtained on hydrocarboxylation of styrene, Whilst
with DIOP as ligand the straight chain carboxylic acid was the major product (Figure 1.2).

\[
\begin{align*}
\text{CH}=\text{CH}_2 & \xrightarrow{[\text{PdCl}_2(\text{PPh}_3)_2]} \text{CH}_3\text{CHCOOH} + \text{CH}_2\text{CH}_2\text{COOH} \\
& \xrightarrow{[\text{PdCl}_2(\text{DIOP})]} \text{CH}_3\text{CHCOOH} + \text{CH}_2\text{CH}_2\text{COOH}
\end{align*}
\]

Figure 1.2

Fuchikami and coworkers\textsuperscript{29} studied hydrocarboxylation and hydroesterification of trifluoropropene (TFP) and pentafluorostyrene (PFS) in the presence of palladium complexes and report that hydroesterification of TFP in the presence of \([\text{PdCl}_2(\text{PPh}_3)_2]\) as the catalyst gave different ratios of branched to straight chain in different solvents under the same reaction conditions of temperature and pressure (100°C, 110 atm) (Table 1.3).

<table>
<thead>
<tr>
<th>Solvent</th>
<th>Yield</th>
<th>Branched/Unbranched</th>
</tr>
</thead>
<tbody>
<tr>
<td>(\text{C}_2\text{H}_5\text{OH})</td>
<td>74%</td>
<td>49 / 51</td>
</tr>
<tr>
<td>THF</td>
<td>93%</td>
<td>71 / 29</td>
</tr>
<tr>
<td>(\text{C}_6\text{H}_6)</td>
<td>95%</td>
<td>79 / 21</td>
</tr>
</tbody>
</table>

Table 1.3
By using the same solvent, acetone, for hydroesterification of TFP with ethanol as the hydrogen donor, different catalysts gave different yields and selectivity (Table 1.4).

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>Conditions</th>
<th>Yield</th>
<th>Branched/Unbr - anched</th>
</tr>
</thead>
<tbody>
<tr>
<td>PdCl$_2$(PPh$_3$)$_2$</td>
<td>100°C, 110 atm</td>
<td>82%</td>
<td>52 / 48</td>
</tr>
<tr>
<td>PdCl$_2$(dppb)</td>
<td>100°C, 110 atm</td>
<td>12%</td>
<td>&lt;1 / 99</td>
</tr>
<tr>
<td>PdCl$_2$(dppf)</td>
<td>120°C, 110 atm</td>
<td>16%</td>
<td>11 / 89</td>
</tr>
</tbody>
</table>

Table 1.4

They observed that during the hydrocarboxylation of PFS, when the source of hydrogen was changed from methanol to water, the selectivity of the catalyst [PdCl$_2$(PPh$_3$)$_2$] changed considerably from branched to unbranched product (Table 1.5).

<table>
<thead>
<tr>
<th>Source of hydrogen</th>
<th>Conditions</th>
<th>Conversion</th>
<th>Yield</th>
<th>Branched Unbranched</th>
</tr>
</thead>
<tbody>
<tr>
<td>CH$_3$OH</td>
<td>100°C, 70 atm</td>
<td>88%</td>
<td>71%</td>
<td>93 / 7</td>
</tr>
<tr>
<td>H$_2$O</td>
<td>125°C, 110 atm</td>
<td>100%</td>
<td>90%</td>
<td>27 / 73</td>
</tr>
</tbody>
</table>

Table 1.5

Knifton$^{30}$ reported hydrocarboxylation of 1-alkenes to give exclusively linear carboxylic esters in the presence of a variety of
palladium complexes such as [PdCl$_2$(PPh$_3$)$_2$]-SnCl$_2$, [PdCl$_2$(p-CH$_3$C$_6$H$_4$)$_3$P)$_2$]-SnCl$_2$, [PdCl$_2$(p-CH$_3$OC$_6$H$_4$)$_3$P)$_2$]-SnCl$_2$, and [PdCl$_2$(PPh$_3$)$_2$]-GeCl$_2$. For example, in the hydroesterification of 1-heptene the highest selectivity to methyl octanoate was obtained using [PdCl$_2$(p-ClC$_6$H$_5$)$_3$P)$_2$]-SnCl$_2$, and the highest yield of methyl octanoate (78%) was obtained using [PdCl$_2$(p-CH$_3$C$_6$H$_4$)$_3$P)$_2$]-SnCl$_2$. Correlation between the catalytic performance and either the size or electronic character of the coordinated ligands was not proposed.

Knifton found that [PdCl$_2$(PPh$_3$)$_2$]-SnCl$_2$ was the best catalytic system. This gave a yield of 76%, and selectivity of 87% to methyl octanoate on hydroesterification of 1-heptene, as shown in equation 1.13.

$$
\text{C}_7\text{H}_{16} + \text{CO} + \text{H}_2\text{O} \xrightarrow{[\text{PdCl}_2(\text{PPh}_3)_2]-10\text{SnCl}_2, 80^\circ\text{C}, 240\text{atm}} \text{C}_8\text{H}_{16}\text{O}_2
$$

Equation 1.13

The high regioselectivity is observed to be relatively insensitive to parameters such as carbon monoxide pressure, solvent, and the nature of the coreactant, but is significantly influenced by the structure of the alkenes and the composition of the active catalyst. Thus in the absence of cocatalyst SnCl$_2$, no regioselectivity was observed, i.e., equal amounts of linear and branched ester were obtained. Knifton observed that the catalyst [PdCl$_2$(PPh$_3$)$_2$]-SnCl$_2$ also carbonylates linear and branched 1-alkenes, internal alkenes, and cyclic alkenes. Sterically hindered branched 1-alkenes in which the alkyl substituent is on the $\beta$ or $\gamma$ carbon atoms also undergo regioselective hydroesterification, with selectivity
greater than 95%.

Internal, disubstituted alkenes undergo hydroesterification more slowly than 1-alkenes. With these compounds the catalyst is no longer selective and gives both straight and branched chain products.

Knifton also observed that under identical reaction conditions used for hydroesterification of 1-heptene, when methanol was replaced by water as the nucleophile coreactant 1-heptene, conversion was greater than 95% and selectivity to octanoic acid was 86%.

The mechanism of the Pd(II)-catalysed hydroesterification of 1-alkenes as proposed by Knifton (Scheme 1.6), is based on both electronic and steric factors. According to the author the high \( \pi \)-acceptor strength of the phosphorus and of SnCl\(_3\) lowers the electron density of the palladium metal. This favours the formation of a palladium hydride species, as well as coordination of nucleophiles such as carbon monoxide and the alkene. To explain the observed selectivity in his studies Knifton proposed that bulky ligands like PPh\(_3\) and SnCl\(_3^-\) give rise to the sterically hindered palladium complex (A) which favours both anti-Markownikov Pd-H addition (step 3) and high equilibrium concentrations of the less sterically hindered straight-chain \( \sigma \)-alkyl and \( \sigma \)-acyl platinum complexes such as D and F. Other evidence for the Pd-H mechanism was given by performing blank experiments i.e., in the absence of the substrate alkene, Knifton isolated the intermediate \([\text{PdH(SnCl}_3\text{)}(\text{P(C}_6\text{H}_5\text{)}_3\text{)})_2\] responsible for hydroesterification. Kingston and Scollary\(^{31}\) have similarly reported the isolation of the palladium hydride complex.
Similarly, Terekhora and coworkers\textsuperscript{32} reported high yield and high regioselectivity for linear acids upon hydrocarboxylation of alkenes in the presence of [PdCl\textsubscript{2}(PPh\textsubscript{3})\textsubscript{2}]. Addition of excess triphenylphosphine improved further the yield and selectivity to straight chain acids\textsuperscript{14}. Increase in selectivity, in some cases >99\%, and yield was also
observed upon addition of a cocatalyst for example, SnCl$_2$ to the catalytic system. With the same catalytic system it was observed that hydrocarboxylation of internal alkenes for example, 2-pentene and cyclic alkenes for example, cyclohexene is slower than for 1-alkenes. For straight chain alkenes, replacing carbon monoxide with synthesis gas resulted in higher yield and regioselectivity. However in the case of styrene this finding is reversed. Lapidus and coworkers reported that hydrocarboxylation of polybutadiene in the presence of [PdCl$_2$(PPh$_3$)$_2$] and excess triphenylphosphine occurred mainly at the pendant carbon-carbon double bond position. Further discussion about this work is carried out along with our work in Chapter 3.

1.2.4. Rhodium catalysed reactions

Hydrocarboxylation of alkenes in the presence of a rhodium catalyst and a halogen promoter also gives rise to both straight chain and branched chain carboxylic acids as do the other metals. Compared to other metals such as cobalt or nickel, rhodium catalysts operate at much lower pressures i.e., below 100 atmospheres. Reactions selective for linear carboxylic acids can be obtained under certain conditions. For example, Burke reported that 1-hexene on hydrocarboxylation in the presence of [RhCl$_3$].3H$_2$O, and the halide promoter CH$_3$I gave 68% straight chain carboxylic acid; 1-heptanoic acid and 9.5 % 2-methylhexanoic acid (Equation 1.14).
However the selectivity is not very high compared to palladium catalysts. The rate of hydrocarboxylation of terminal or internal alkenes can be increased by adding minor amounts of compounds of iron, cobalt or manganese.

A British patent describes hydrocarboxylation of 1-Hexene in high conversion (80%) and selectivity (99%) to C7 carboxylic acid using the rhodium complex RhCl₃.3H₂O as catalyst in the presence of a halide promoter in acetic acid solvent. The selectivity to straight chain heptanoic acid was less than 25% (Equation 1.15). In the absence of halide promoter no reaction occurred.
1.2.5. Reactions catalysed by other metal complexes

1.2.5.1. Platinum catalysed reactions

Platinum compounds, e.g. $\text{H}_2[\text{PtCl}_6]$ in the presence of $\text{SnCl}_2$, only catalyse the hydrocarboxylation of 1-alkenes, internal alkenes being relatively inert to hydrocarboxylation. Under drastic conditions\(^{40}\) they gave low conversions and mixtures of products. Alkene isomerisation becomes the almost exclusive reaction.

Ligand stabilised platinum (II) along with group 14 metal halide complexes have been reported\(^{41}\) to catalyse the homogeneous carbonylation of 1-alkenes to carboxylic acids or esters. For example, 1-heptene on hydroesterification in the presence of $[\text{PtCl}_2(\text{PPh}_3\text{O})]_2-\text{SnCl}_2$ gave 86% yield and 98% selectivity to methyl octanoate (Equation 1.16). In the absence of cocatalyst no reaction took place.

\[
\text{\begin{align*}
\text{C}_7\text{H}_{14} + \text{CO} + \text{CH}_3\text{OH} & \xrightarrow{[\text{PtCl}_2(\text{PPh}_3\text{O})]_2/\text{SnCl}_2} \text{C}_7\text{H}_{14}\text{COOCH}_3 \\
& \text{80°C, 240 atm} \\
> 93\% \text{ yield}
\end{align*}}
\]

Equation 1.16

1.2.5.2. Iridium catalysed reactions

Iridium complexes function as catalysts for hydrocarboxylation of alkenes under reaction conditions very similar to those employed
with rhodium complexes. The alkenes can be either terminal or internal and even with linear internal alkenes, a high proportion of terminal carboxylic acids are obtained. The iridium is most conveniently used in the form of an iridium trihalide. The iodide promoter can be used either as aqueous hydrogen iodide or as an alkyl iodide. Foster and coworkers obtained saturated acids on hydrocarboxylation of alkenes in the presence of the iridium complex catalyst \([\text{HIr(CO)}_2\text{I}_2\text{(H}_2\text{O})]\). Studies on the progress of the reaction showed that the catalyst composition changed, with conversion of part of it to the less reactive species \([\text{Ir(CO)}_3\text{I}]\), \([\text{Ir(CO)}_3\text{I}_3]\), \([\text{Ir(CO)}_2\text{I}_4]\)^+ etc., species which lowered the reaction rate significantly.

1.2.5.3. Copper catalysed reactions

In concentrated sulphuric acid, salts of copper(I) are active catalysts for the hydrocarboxylation of alkenes under very mild conditions (50°C, 1 atm). The final products are tertiary carboxylic acids\(^{42}\) (Equation 1.17).

\[
\begin{align*}
\text{R}^1_{\text{alkyl}} + \text{CO} \xrightleftharpoons{\text{[Cu}_2\text{O}], > 80\% \text{H}_2\text{SO}_4, 1 \text{ bar}} & \rightarrow \text{R}^1_{\text{alkyl}}\text{CH}_3\text{COOH} \\
\text{Where R}^1_{\text{alkyl}} \text{ and R}^2_{\text{alkyl}} &= \text{H, alkyl groups}
\end{align*}
\]

Equation 1.17

It has also been observed that, in the presence of saturated hydrocarbons containing a tertiary hydrogen, e.g. methylcyclohexane, which was used
as the solvent, the reaction occurred both on this hydrocarbon and on the alkene (Equation 1.18).

$$
\text{Cyclohexane} + \text{alkene} + \text{CO} \xrightarrow{[\text{Cu}_2\text{O}], > 80\% \text{H}_2\text{SO}_4, 1 \text{ bar}} \text{COOH} + \text{t-C}_7\text{ acids}
$$

Equation 1.18

The copper-catalysed hydrocarboxylation of alkenes involves first the formation of a carbonium ion. This carbonium ion rapidly rearranges to more stable carbonium ions which are trapped by carbon monoxide. The role of the copper(I) compound, presumably $[\text{Cu(CO)}_3]^+$, is to deliver carbon monoxide to the carbonium ion (Scheme 1.7). This species only exists at sulphuric acid concentrations greater than 80%.

Scheme 1.7

Souma and Sano\textsuperscript{43} obtained similar results on hydrocarboxylation
Souma and Sano\textsuperscript{43} obtained similar results on hydrocarboxylation of alkenes using copper carbonyl \([\text{Cu(CO)}_{n}^{+}]\) and silver carbonyl \([\text{Ag(CO)}_{2}^{+}]\) catalysts at room temperature and at atmospheric carbon monoxide pressure in BF\textsubscript{3}-H\textsubscript{2}O solutions. For example, 1-hexene gave a mixture of 2,2-dimethylpentanoic acid and 2-methyl-2-ethylbutanoic acid. Dienes also reacted with carbon monoxide to give mixtures of monocarboxylic acids, and dicarboxylic acids. The yield of terminal carboxylic acid and the reaction rate decrease with a decrease in BF\textsubscript{3} concentration. Saturated hydrocarbons also react with carbon monoxide in the presence of alkenes. In this case the hydride ion is abstracted from the saturated hydrocarbon by the carbonium ion formed from the alkene. Then the carbonium ion forms the tertiary carboxylic acid. Saturated hydrocarbons which have no tertiary hydrogens do not react with carbon monoxide. Hydride transfer takes place only in concentrated BF\textsubscript{3}-H\textsubscript{2}O solutions.

1.3 Hydrocarboxylation of dienes

Non-conjugated dienes

Non-conjugated dienes exhibit a similar behaviour to that of 1-alkenes. Depending upon the catalyst, non-conjugated dienes can give rise to dicarboxylic acids and/or unsaturated monocarboxylic acids. The catalysts most employed for hydrocarboxylation of non-conjugated dienes contain cobalt. However nickel and palladium also show some catalytic activity.
1.3.1. Cobalt catalysed reactions.

With the cobalt catalyst, $[\text{Co}_2(\text{CO})_8]$ non conjugated dienes give dicarboxylic and saturated monocarboxylic acids$^{44}$ on hydrocarboxylation (see Table 1.6).

1.3.2. Nickel catalysed reactions

Non conjugated dienes react with stoichiometric amounts of the nickel complex $[\text{Ni}(\text{CO})_4]$ to give mono carboxylic acids. However the yields are not very high$^{45}$ (Table 1.6).

<table>
<thead>
<tr>
<th>Diene</th>
<th>Catalyst</th>
<th>Conditions</th>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>1,5-Hexadiene</td>
<td>$\text{Co}_2(\text{CO})_8$</td>
<td>210°C, 250 atm</td>
<td>Octanedioic acid Saturated C$_7$-monocarboxylic acids</td>
<td>34%</td>
</tr>
<tr>
<td></td>
<td>$\text{Ni}(\text{CO})_4$</td>
<td>180°C</td>
<td>2-methylhex-5-enoic acid</td>
<td>20%</td>
</tr>
</tbody>
</table>

Table 1.6
1.3.3. Palladium catalysed reactions

If the double bonds are suitably juxtaposed intramolecular reactions are observed. For example, in the presence of the palladium complex, $[\text{PdI}_2(\text{PBu}_3)_2]$, a formal hydrocarboxylation followed by an intramolecular cyclisation takes place (Equation 1.19).

\[
\begin{align*}
\text{CH}_2\text{CH}=&\text{CH}_2 + \text{CO} + \text{CH}_3\text{OH} \quad \xrightarrow{\text{PdI}_2(\text{PBu}_3)_2} \\
&\text{CH}_2\text{CH}=&\text{CH}_2 \\
\end{align*}
\]

\[150^\circ\text{C}, 1000 \text{ atm}\]

Equation 1.19

1,5-Cyclooctadiene gave methyl cyclo-oct-4-ene-1-carboxylate in 45% yield and the dicarboxylic esters in 30% yield\(^\text{46}\) (Equation 1.20). It was observed that, as the reaction proceeded, the concentration of the unsaturated monoester increased rapidly to a maximum and then gradually decreased as the concentration of diesters increased.

\[
\begin{align*}
\text{C}_8\text{H}_{12} + \text{CO} + \text{CH}_3\text{OH} \quad \xrightarrow{\text{PdI}_2(\text{PBu}_3)_2} \\
&\text{C}_8\text{H}_{12} \quad \text{COOCH}_3 \\
&\text{H}_2\text{COOC} \\
\end{align*}
\]

\[150^\circ\text{C}, 1000 \text{ atm}\]

Equation 1.20
With tri-alkenes as substrates it is possible to target hydrocarboxylation at the required double bond position. For example, it was not previously possible to convert 1,5,9-cyclododecatriene into 4,8-cyclododecadiene-1-carboxylic acid, or its esters, or to the corresponding polycarboxylated compounds, since acenaphthene derivatives and polymerisation products were formed instead. Bittler and coworkers reported that in the presence of [PdCl$_2$(PPh$_3$)$_2$] as catalyst, it was possible to cause one, two or all three double bonds in cyclododecatriene to react depending on the reaction conditions to give mainly the mono, di-, and triester; the monoester could be obtained without any byproducts (Equation 1.21) at low temperatures (60°C).

\[
\begin{align*}
\text{+} & \quad x \text{CO} + x \text{C}_2\text{H}_5\text{OH} \\
& \quad \xrightarrow{\text{PdCl}_2(\text{PPh}_3)_2} \\
& \quad 105^\circ\text{C} \\
\text{x} &= 1, 2, 3 \\
\end{align*}
\]

\[
\begin{align*}
& \xrightarrow{60^\circ\text{C}} \\
& \quad \text{COOC}_2\text{H}_5 \\
& \xrightarrow{120^\circ\text{C}} \\
& \quad \text{COOC}_2\text{H}_5 \quad \text{COOC}_2\text{H}_5 \\
\end{align*}
\]

Equation 1.21
Conjugated Dienes

Under the usual high temperature conditions in the presence of nickel and cobalt catalysts, conjugated dienes fail to produce the expected hydrocarboxylation products. For example, conjugated 1,3-dienes undergo a formal 1,4-addition of the elements of formic acid to give $\beta, \gamma$-unsaturated acids. The most used catalysts for hydrocarboxylation of conjugated dienes are the cobalt complex, $[\text{Co}_2(\text{CO})_8]$, and complexes of palladium.

1.3.4. Cobalt catalysed reactions

Butadiene reacted in the presence of $[\text{Co}_2(\text{CO})_8]$ to give the rearranged unsaturated acid in high yield (Equation 1.22). By increasing the temperature to 210°C, reasonable yields of diacids, essentially adipic acid are obtained.

\[
\begin{align*}
\text{\text{CO}} + \text{H}_2\text{O} & \xrightarrow{[\text{Co}_2(\text{CO})_8] / \text{py}} \xrightarrow{160^\circ\text{C}, 250\text{ atm}} \text{\text{HOOC-----------------.COOH}} \\
& \xrightarrow{210^\circ\text{C}} \text{\text{HOOC---------COOH}}
\end{align*}
\]

Equation 1.22

Under hydrocarboxylation conditions, butadiene may also dimerise to vinyl cyclohexene. The solvent used has a great effect on the proportion of dicarboxylic derivatives formed. Pyridine seems to determine the preferential formation of these compounds, 49, 50, 51,
whereas with other solvents like dioxane, acetone, or isoquinoline, the main product is 3-pentenoic acid.

1.3.5. Palladium catalysed reactions

Palladium has been used as catalyst in the presence of methanol or ethanol at relatively low temperature (56-150°C) and very high carbon monoxide pressures (1000 atm). Unsaturated monocarboxylic esters were obtained.

<table>
<thead>
<tr>
<th>Diene</th>
<th>Catalyst</th>
<th>Conditions</th>
<th>Product (Yield)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Butadiene</td>
<td>PdCl₂(PPh₃)₂</td>
<td>120-140°C, 700 atm</td>
<td>3-pentenoic acid (70%)</td>
</tr>
<tr>
<td>Butadiene</td>
<td>Pd(PBu₃)₂</td>
<td>150°C, 1000 atm, CH₃OH</td>
<td>Methyl 3-pentenoate (73%)</td>
</tr>
<tr>
<td>1,3-Cyclohexadiene</td>
<td>PdCl₂</td>
<td>100°C, 1000 atm, C₂H₅OH</td>
<td>Methyl 4-pentenoate (1%)</td>
</tr>
<tr>
<td>1,3-Penta-diene</td>
<td>Na₂PdI₄</td>
<td>70°C, 1000 atm, CH₃OH</td>
<td>4-Vinylcyclohexene (1-2%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Ethyl 2-cyclohexene carboxylate (80%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Methyl-2-methylpentenoate (34%)</td>
</tr>
</tbody>
</table>

Table 1.7

With palladium catalysts, even small changes in the type of ligand bound...
to the palladium markedly influenced the composition of the reaction products (Table 1.7).

1.3.6. Nickel catalysed reactions

Stoichiometric hydrocarboxylation of butadiene\textsuperscript{4} first formed a vinyl cyclohexene which then gave complex mixtures of mono and dicarboxylic acids (Equation 1.23).

\[
\text{CH}_2=\text{CH}_2 + \text{CH}_3\text{COOH} + \text{H}_2\text{O} + [\text{Ni(CO)}_4] \xrightarrow{165^\circ\text{C}} \text{Complex mixtures of mono and dicarboxylic acids} \xrightarrow{\text{CH}_3\text{OH}} \text{C}_8\text{H}_{13}\text{COOCH}_3 + \text{C}_{12}\text{H}_{20}\text{O}_4
\]

Equation 1.23

The monocarboxylic acids on esterification with methanol yielded C\textsubscript{8}H\textsubscript{13}COOCH\textsubscript{3}, while a higher boiling fraction containing dicarboxylic acids gave on esterification the methyl ester of C\textsubscript{12}H\textsubscript{20}O\textsubscript{4}.

Catalytic hydrocarboxylation of butadiene\textsuperscript{55} in the presence of [Ni(CO)\textsubscript{4}] and a small amount of hydroquinone gave a mixture of 2-(3-cyclohexen-1-yl)propanoic acid and 2-(carboxycyclohexyl) propanoic acid (Equation 1.24).
1.3.7 Rhodium catalysed reactions

Garlaschelli and coworkers\textsuperscript{56} reported that by varying both the ligand attached to the catalyst and the physical parameters the yield and selectivity of isoprene hydrocarboxylation to give 4-methyl-3-pentenoic acid (pyroterebic acid) or its lactone derivative \( \gamma,\gamma\)-di-methyl-\( \gamma\)-butyrolactone can be altered (Equation 1.25).

When hydrocarboxylation of isoprene was carried out with [PPN][Rh(CO\textsubscript{2})X\textsubscript{2}] [PPN= bis( triphenylphosphine iminium; X=Cl, I) in the presence of aqueous HI as cocatalyst and tetrahydrofuran as the solvent, the products markedly depended on the HI/Rh ratio and
temperature. Thus the use of low HI/Rh ratio and lower temperature greatly depressed the formation of the lactone, and the pyroterebic acid could be obtained with an overall selectivity of ca. 85%. Conversely high HI/Rh ratio and higher temperature greatly favoured the formation of the γ,γ-di-methyl-γ-butyrolactone. Increase of either one or both of these parameters markedly lowered the selectivity towards oxygenated products owing to a steeper increase in the rate of the hydrogenation of isoprene to give an isomeric mixture of methyl butenes.

On the other hand with the [Rh₄(CO)₁₂]/HI catalytic system, lactone was always obtained as the major product with little pyroterebic acid. An increase in temperature and/or HI/Rh ratio lowered the selectivity. With a sub-stoichiometric isoprene/water ratio the system essentially catalysed the complete conversion of isoprene into γ,γ-dimethyl-γ-butyrolactone. When the isoprene/water molar ratio was increased so that water became the sub-stoichiometric reagent, the relative amount of pyroterebic acid present progressively increased at the expense of the lactones.

1.4 Hydrocarboxylation of functionalised alkenes

A fairly large number of unsaturated substrates containing various functional groups can be hydrocarboxylated in the presence of nickel, cobalt, and palladium catalysts. The functional groups in the unsaturated molecules are mainly halogens, carboxyl, carboalkoxy, and cyano groups. The products formed in these reactions are mixtures of isomeric esters or acids. Usually the reaction proceeds with the retention of the functional group, but in some cases, for example, when
the functional group in the unsaturated molecule is a hydroxyl group, hydrocarboxylation in the presence of cobalt catalysts \([\text{Co}_2(\text{CO})_8]\) does not give the expected products but gives lactones due to intramolecular cyclisation.

1.4.1 Cobalt catalysed reactions

Using \([\text{Co}_2(\text{CO})_8]\) in the presence of \((\text{C}_6\text{H}_5)_2\text{PCH}_2\text{CH}_2\text{P}(\text{C}_6\text{H}_5)_2\) as catalyst, acrylic acid upon hydrocarboxylation in ethanol gave 48% of \((\text{CH}_2\text{COOC}_2\text{H}_5)_2\) and 43% of \(\text{HOOCCH}_2\text{CH}_2\text{COOC}_2\text{H}_5\). Without \((\text{C}_6\text{H}_5)_2\text{PCH}_2\text{CH}_2\text{P}(\text{C}_6\text{H}_5)_2\) the highest yields were 25.5% and 73% respectively\(^{57}\) (Equation 1.26).

\[
\begin{align*}
\text{CH}_2=\text{CHCOOH} + \text{CO} + \text{H}_2\text{O} & \quad \text{[Co}_2(\text{CO})_8]\ / \ \text{Ph}_2\text{PCH}_2\text{CH}_2\text{PPh}_2 \quad \text{140°C, atm} \\
& \frac{\text{[(CH}_2\text{COOC}_2\text{H}_5)_2 + \text{HOOCCH}_2\text{CH}_2\text{COOC}_2\text{H}_5]}{\text{direct}}
\end{align*}
\]

Equation 1.26

In the presence of a \([\text{Co}_2(\text{CO})_8]\) / pyridine complex, isomeric hexenoic acids gave mixtures of \(\text{C}_7\)-dicarboxylic acid with an isomeric distribution of the hydrocarboxylation product. Only in the case of 5-hexenoic acid, was the unbranched \(\text{C}_7\)-dicarboxylic acid the main product. With other isomeric hexenoic acids, for example 2-hexenoic acid, propyl succinic acid was the main isomer in the resulting mixture of dicarboxylic acids. It was proposed\(^{58}\) that cobalt carbonyl forms a five membered ring complex with the unsaturated carboxylic acid
especially in the case of the pyridinium or alkali salt of the unsaturated acid. This stabilised isomeric cobalt intermediate would lead to the alkyl succinic acid or ester as the main isomeric product.

Pyridine and hydrogen gas improve conversions in the hydrocarboxylation of substrates like acrylonitrile\textsuperscript{59} and methyl acrylate\textsuperscript{60} in the presence of $[\text{Co}_2(\text{CO})_8]$.

### 1.4.2 Palladium catalysed reactions

Particularly good yields of esters are obtained by reacting vinyl chloride in the presence of $[\text{PdCl}_2(\text{PPh}_3)_2]$ and $\text{HCl}$\textsuperscript{47} in ethanol (Equation 1.27).

\[
\text{Cl} + \text{CO} + \text{C}_2\text{H}_5\text{OH} \xrightarrow{\text{PdCl}_2(\text{PPh}_3)_2 / \text{HCl}} \begin{align*}
\text{COOC}_2\text{H}_5 & \quad \text{Cl} \\
& \quad \text{Cl} \quad \text{COOC}_2\text{H}_5 (5\%)
\end{align*}
\]

Equation 1.27

Frankel and Thomas\textsuperscript{61} obtained 9(10) carboxystearic acids in high yields in the presence of a catalytic mixture of palladium (II) chloride and triphenylphosphine (Equation 1.28).
\[
\text{CH}_3(\text{CH}_2)_7\text{CH}=\text{CH}(\text{CH}_2)_7\text{COOH} \xrightarrow{\text{PdCl}_2/\text{PPh}_3} 150^\circ\text{C, 272 atm} \\
\]

\[
\begin{align*}
X & \quad Y \\
\text{CH}_3(\text{CH}_2)_7\text{CH}-\text{CH}(\text{CH}_2)_7\text{COOH} & \\
\text{When } X = \text{H}, \ Y = \text{COOH} & \\
X = \text{COOH}, \ Y = \text{H} & \\
\end{align*}
\]

Equation 1.28

Diels-Alder adducts such as the cyclohexene derivatives (Equation 1.29)

\[
\text{COOC}_2\text{H}_5 + \text{COOC}_2\text{H}_5 \xrightarrow{} \text{COOC}_2\text{H}_5 + \text{COOC}_2\text{H}_5
\]

Equation 1.29

and the corresponding norbornene derivative, which readily dissociate into their components at high temperatures, were effectively hydrocarboxylated with [PdCl\(_2\)(PPh\(_3\))\(_2\)] (Equation 1.30).
They report that a mixture of PdCl$_2$ and triphenylphosphine proved to be a more active and efficient catalytic system than preformed [PdCl$_2$(PPh$_3$)$_2$] in increasing the yield and reducing the time of the reaction.

### 1.4.3. Reactions catalysed by other metal complexes.

#### 1.4.3.1. Iron catalysed reactions

Recently, a fully catalytic regioselective hydrocarboxylation of acrylic acid to methyl malonic acid in the presence of [Fe(CO)$_5$] has been reported by Brunet and Passadargue$^{62}$ (Equation 1.31). This is the first reported use of iron in regioselective hydrocarboxylation.

$$
\text{CH}_2\text{=CHCOOH} + \text{CO} + \text{H}_2\text{O} \xrightarrow{[\text{Fe(CO)}_5]} \text{HOOC-CH-COOH}
$$

Equation 1.31
This reaction by [Fe(CO)_5] was carried out in the presence of a base Ca(OH)_2 in a H_2O-iPrOH mixture as solvent at 70°C under one atmosphere of carbon monoxide. Increase in carbon monoxide pressure inhibited the reaction suggesting that carbon monoxide dissociation did not take place at higher carbon monoxide pressure (Equation 1.32).

\[
[HFe(CO)_4]^\text{-} \longrightarrow [HFe(CO)_3]^\text{-} + \text{CO}
\]

Equation 1.32

The proposed mechanism is shown in Scheme 1.8.
1.5 Asymmetric hydrocarboxylation

Asymmetric hydrocarboxylation, in principle, could have enormous importance for synthesising important chemicals, as by making chiral catalysts through the use of chiral ligands, stereoselectivity can be obtained. There are certain problems associated with the synthesis however, due to the mechanistic complexity of the reactions, as itemised below:

1. the catalytic cycles of the reaction involve many steps.
2. the structures of the active catalyst species are strongly effected by reaction conditions, ratio of carbon monoxide to hydrogen, alcohol or water, and ratios of central transition metals to chiral ligands.
3. the regioselectivities of the reactions producing the desired optically active compounds (e.g., branched acids from 1-alkenes) are not always high.
4. very high carbon monoxide pressures (~300-700 atm) are usually necessary to obtain high optical yields.

For asymmetric hydrocarboxylation, palladium has been exclusively employed as the metal and DIOP (Figure 1.3) type chiral diphosphines the most common ligands.

![](image_url)

**DIOP**

*Figure 1.3*

Consiglio reported asymmetric hydrocarboxylation of alkenes
to give high optical yields of branched esters in the presence of a
[PdCl₂(-)DIOP] and PPh₃ catalytic system. The reaction took place at a
temperature of 100°C, but required 400 atmospheres of carbon
monoxide pressure. For example, α-methyl styrene catalysed by
[PdCl₂(-)DIOP] gave the ester of 3-phenylbutanoic acid with an
optical purity of ~ 60%. Some ethyl-2-methyl-2-phenylpropanoate was
also obtained. The ratio of (-)DIOP to palladium exerted a considerable
influence on stereoselectivity for example, a ratio of 0.5 gave a better
result than a ratio of 1. Also monophosphines lead to predominant
formation of the internal ester, while (-)-DIOP gave mainly the
terminal ester.

\[\text{DIPHOL}\]

Figure 1.4

When DIPHOL (Figure 1.4) was used in place of DIOP and the pressure
lowered, the maximum yield observed was ~47% e.e.

Alper and Hamel⁶⁵ have reported the first asymmetric
carbonylation of allyl alcohols to yield optically active lactones using poly α- amino acids as added chiral ligands. Thus treatment of but-2-en-1-ol (Equation 1.33) under extremely mild conditions with carbon monoxide, oxygen, copper(II) chloride, and hydrochloric acid in tetrahydrofuran in the presence of palladium chloride and poly-L-leucine afforded (R)-α-methyl-γ-butyrolactone in 61% enantiomeric excess.

\[
\text{CH}_3\text{CH} = \text{CHCH}_2\text{OH} + \text{CO} \xrightarrow{\text{O}_2, \text{THF}, L^*} \text{CH}_3 \text{CH} = \text{CHCH}_2\text{OH} + \text{CO}
\]

Equation 1.33

The chiral discrimination step in the asymmetric process with poly-L-leucine was the intramolecular addition of palladium hydride to the co-ordinated double bond of the five co-ordinate palladium (II) complex (Figure 1.5), to give a metallocyclopentane [carbonyl insertion followed by reductive elimination to give the products].

![Figure 1.5](image)

(L)-diethyl tartrate and (R)-and (S)-2,2'-bis (diphenylphosphino)-1,1'-
binapthyl (BINAP) also produced optically active lactones, but in lower optical purity.

Ibuprofen [2-(p-iso-butylphenyl)propanoic acid](Figure 1.7) and naproxen [2-(6-methoxy-2-napthyl)propanoic acid] (Figure 1.6) are two commercially important non-steroidal anti inflammatory agents.

Metal-catalysed approaches to obtain optically active ibuprofen or naproxen have been in vogue recently. Ohata and coworkers obtained a high degree of optical purity by using BINAP ruthenium(II)-catalysed hydrogenation of 2-(6-methoxynapthyl)-2-propenoic acid. A high pressure of hydrogen (135 atm) was required, and the acrylic acid derivative had to be synthesised via several steps. In 1987, Parrinello and Stille obtained naproxen from p-isobutylstyrene by hydroformylation in the presence of a Pt(II) complex of (2S,4S)-N-(tert-butoxycarbonyl)-4-diphenylphosphino-2-[(diphenylphosphino(methyl) prolidine[(-)-BPPM], with SnCl₂. The hydroformylation reaction required drastic conditions (165 atm) and a subsequent step to
produce acids. It was observed that while the enantiomeric excess of the formed aldehyde approached 80%, the regioselectivity broke down with an unfavourable branched/linear ratio (~0.5).

Alper and Hamel\textsuperscript{68} report the preparation of these two commercially important drugs under very mild conditions by carrying out hydrocarboxylation reactions (Equation 1.34) of p-isobutylstyrene and 2-vinyl-6-methoxynaphthalene in the presence of PdCl\textsubscript{2} and (R)-(−)-or(S)-(−)-1,1′-binaphthyl-2,2′-diyl hydrogen phosphate (BNPPA)

\[
\text{RCH}=\text{CH}_2 + \text{CO} + \text{H}_2\text{O} \xrightarrow{\text{O}_2, \text{THF}, L^*} \text{RCH(CH}_3\text{)COOH} \\
PdCl_2, \text{CuCl}_2, \text{HCl}, r.t, 1 \text{ atm CO}
\]

Equation 1.34

as the chiral ligand to yield (S)-(−)-and(R)-(−)-ibuprofen in 83-84% optical yield. Similarly, naproxen was obtained in good yield and in upto 91% optical yield. When other chiral ligands like L-menthol, D-menthol, (R)-1,1′-bi-2-naphthol, D-diethyl tartrate (DET), and (S)-2,2′-bis(diphenyl phosphino)-1,1′-binapthyl (BINAP) were used, only poor optical yields (<10%) were obtained.
Chapter 2

Analysis of Polybutadiene

2.1 Introduction

Since the bulk of this thesis is concerned with the synthesis of new polymers via hydrocarboxylation of polybutadiene it is important to be able to analyse the microstructure of the product and starting polymers. In this chapter, we discuss the various analytical methods that we have adopted.

A number of methods have been proposed for the analysis of polybutadienes or substituted polybutadienes. In this chapter a critical appraisal of the various methods of analysis, (i) Conductometric titration, (ii) $^1$H and $^{13}$C N.M.R., and (iii) Infrared spectroscopy, for analysing polybutadiene and its hydrocarboxylated products is provided, using samples of hydrocarboxylated polybutadiene obtained from (a) regiospecific hydrocarboxylation of pendant carbon-carbon double bonds, and (b) hydrocarboxylation of both internal and pendant carbon-carbon double bonds, so that direct comparison can be made between the various methods.

2.2 Method of analysis

2.2.1 Conductometric titration

The electrical conductivity of a solution depends upon the number and mobility of any ions it contains. This principle is used to find the end points of acid-base and other titrations in a conductometric titration.
Conductometric titration can offer several advantages over the use of indicators. For example, coloured solutions which cannot be titrated by ordinary volumetric methods with the help of indicators can be successfully titrated conductometrically. This method can also be employed in the case of very dilute solutions and also for weak acids and bases. Furthermore, no special care is necessary near the endpoint as it is determined graphically.

Four types of conductometric titrations are commonly known:

1. titration of a strong acid, like hydrochloric acid, with a strong base, like sodium hydroxide.
2. titration of a weak acid, like acetic acid, with a strong base, like sodium hydroxide.
3. titration of a strong acid, like hydrochloric acid, with a weak base, like ammonium hydroxide.
4. titration of a mixture of a strong and a weak acid, such as hydrochloric acid and acetic acid, with a strong base, like sodium hydroxide.

For example, in the titration of hydrochloric acid with sodium hydroxide, the conductance of hydrochloric acid is due to the presence of hydrogen and chloride ions. As base is added gradually, the hydrogen ions are replaced by slow moving sodium ions, causing the conductivity of the solution to decrease (Equation 2.1).

\[
\text{H}^+ + \text{Cl}^- + [\text{Na}^+ + \text{OH}^-] \rightarrow \text{Na}^+ + \text{Cl}^- + \text{H}_2\text{O}
\]

Equation 2.1
The conductivity continues to decrease until the acid has been completely neutralised. After this, the addition of base will result in an increase in the number of ions in the solution. The conductivity, therefore reaches a minimum value at the end point and then begins to increase. When conductivity is plotted against the volume of base added, two straight lines AB and CD are obtained (Figure 2.1). The point where AB and CD meet i.e., X gives the end point.

![Diagram of conductivity vs volume of HCl](image)

**Figure 2.1. Titration of a strong acid against a strong base.**

When a mixture of a strong and a weak base is to be titrated against a strong acid, a combination of curves is obtained as shown in Figure 2.2. Suppose a mixture of NaOH and NH₃(aq) is titrated against HCl, NaOH being a much stronger base, will react first. The titration of NH₃(aq) will commence only after NaOH has been completely
Figure 2.2. Titration of a mixture of a strong and a weak base against a strong acid

neutralised. Being a weak base, the number of ions increases slightly as more HCl is added so that a slight increase in conductivity is observed until all of the hydroxyl ions have been neutralised. The conductivity then rises sharply as more HCl is added on account of the presence of a larger number of ions, but also on account of the higher mobility of the H\(^+\) ions. Hence, a combination of curves are obtained. In Figure 2.2 point B corresponds to the neutralisation of NaOH and point C corresponds to the neutralisation of NH\(_3\)(aq).

Our aim was to apply the above principle to find out the amount of -COOH groups in the polymer chain. For this the following procedure was adopted.
The hydrocarboxylated polybutadiene was first converted to its corresponding sodium salt by dissolving the hydrocarboxylated polybutadiene (Figure 2.3) in excess of NaOH (0.01M) and conductometric titration with HCl (0.01M) was carried out.

![Figure 2.3](image)

It was observed that initially all the excess NaOH was neutralised by HCl and the conductivity dropped because the mobile OH⁻ ions were being replaced by the less mobile Cl⁻ ions (Equation 2.2).

\[
\text{HCl} + \text{NaOH} \rightarrow \text{NaCl} + \text{H}_2\text{O}
\]

Equation 2.2

After all the NaOH had been neutralised, HCl started to react with \(\text{COONa}^+\) ions (Equation 2.3).

\[
\text{\(\text{COONa}^+\)} + \text{HCl} \rightarrow \text{Na}^+\text{Cl}^- + \text{\(\text{COOH}^+\)}
\]

where, \(\text{\(\text{COO}^-\)} = \text{poly(butadiene)}\)

Equation 2.3
Being a weak acid, the number of ions increases slightly as more HCl is added so that a slight increase in conductivity should be observed until all of the carboxylates have been neutralised. The conductivity then rose sharply as more HCl was added on account of the presence of the larger number of ions, but also on account of the higher mobility of the H\(^+\) ions. For polycarboxylic acids of the type under study, there is, however, a further complication. The protonated form of carboxylic acid is insoluble in water and hence precipitates during the titration.

In principle this is not a problem since it should just level the middle portion of the curve as is observed. However, one cannot be certain at what point a given polymer molecule precipitates i.e., how many of the carboxylic acid groups need to be protonated before precipitation occurs. If this is not 100% and if the ionic carboxylates that precipitated with the polymer molecule are not available for subsequent protonation (with release of Na\(^+\)), the final end point will not be estimated correctly (For example, see Figure 2.14). Whether or not this is a genuine problem for the types of polymer we are analysing can be determined since the final end point should correspond to the end point obtained for titration of the same amount of NaOH as was used for the back titration. In general, this appears to be the case so that all of the carboxylic acid anions in the polymer are available for protonation. However elemental analysis (see section 2.2.4, Microanalysis) shows the presence of sodium ions in the hydrocarboxylated product. This means that some of the sodium ions are bound to the product polymer (as carboxylate) and not available for titration with HCl. In certain cases, especially for polymers where there are -COOH groups on the backbone, there is considerable evidence that precipitation of partially neutralised polymers does occur (see section 2.2.2.2.3).
For a specific example we select a polymer (initial composition = 53% pendant, 31% internal C=C and 16% saturation) prepared by hydrocarboxylation of polybutadiene where all the pendant carbon-carbon double bonds have been selectively hydrocarboxylated. From $^1$H n.m.r. calculations (see section 2.2.2. $^1$H and $^{13}$C n.m.r. for details) we have calculated that the overall conversion of the carbon-carbon double bonds in the hydrocarboxylated product to be 43%.

![Figure 2.4 Conductometric titration between hydrocarboxylated polybutadiene (in NaOH) and HCl](image_url)

**Figure 2.4** Conductometric titration between hydrocarboxylated polybutadiene (in NaOH) and HCl
Calculation

Figure 2.4 shows a graphic representation of the conductometric titration performed between hydrocarboxylated polybutadiene in NaOH versus HCl.

0.01g of the hydrocarboxylated product under study was dissolved in 10 cm³ of 0.01 M NaOH and titrated with 0.01 M HCl. Blank titration of NaOH with HCl gave 9.2 cm³ as the end point i.e., neutralisation point. First end point when all NaOH has reacted with HCl = 3.55 cm³

As can be seen from the graph the final end point could not be determined precisely as precipitation of the polymer took place. The end point calculated based on titration comes to 9.0 cm³. This means that some of the Na⁺ ions bound to the polymer chain are not available for titration with HCl. However, since this is a back titration and that the final end point should correspond to the end point obtained for titration of the same amount of NaOH used. We, have, therefore, assumed that the final end point should be at 9.2 cm³.

Second end point when all COONa bound to the polymer has reacted with HCl (based on back titration) = 9.2 cm³

Therefore 0.01g of hydrocarboxylated product is equivalent to 5.65 cm³ of 0.01 M HCl.

This in terms of moles will be

\[
0.01g = \frac{5.65 \times 0.01}{1000} = 5.65 \times 10^{-5} \text{ moles}
\]

Let 'M' be the mass in a.m.u. of the hydrocarboxylated product, which contains one proton,

\[
\text{then } \frac{0.01g}{M} = 5.65 \times 10^{-5} \text{ moles}
\]
Therefore, \( M = 177 \)

The product butadiene will contain butadiene units of mass 54, carboxylic acid units of mass 100, and hydrogenated butadiene units (saturated units) of mass 56, in this case the latter accounts for 16% of the initial butadiene units in the polymer.

The mass containing one proton will be

\[
\frac{a \times 100 + b \times 54 + c \times 56}{a},
\]

where

- \( a \) is the proportion of carboxylic acid units in the chain,
- \( b \) is the proportion of unreacted units, and
- \( c \) is the proportion of saturation in the starting material = 0.16

We then have

\[
\frac{a \times 100 + b \times 54 + 0.16 \times 56}{a} = 177
\]

\[
b \times 54 + 0.16 \times 56 = 77a
\]

We know that \( a + b = 0.84 \)
Therefore \( b = 0.84 - a \)

\[
(0.84 - a) \times 54 + 8.96 = 77a
\]

\[
45.36 - 54a + 8.96 = 77a
\]

\[
54.32 = 131a
\]

\[
a = \frac{54.32}{131} = 0.4146 = 41 \%
\]
Therefore \( b = 0.84 - 0.41 = 0.43 \)

The final composition of the hydrocarboxylated product will be;
- Remaining carbon-carbon double bond = 43%
- Overall conversion (Carboxylic acid) = 41%
- Saturated carbon-carbon units = 16%

As can be seen the value obtained by using conductometric titration is in agreement with the values obtained from \(^1\)H n.m.r. (43%). However, information about the selectivity or the microstructure of the product cannot be obtained.

Conductometric titration can thus only be used to obtain information about the overall conversion of carboxylic acid units in the hydrocarboxylated product. However, a combination of a conductometric titration and \(^1\)H n.m.r. spectroscopy can give complete information on the microstructure of most of the polymers under study (see Chapter 4).

2.2.2 \(^1\)H and \(^{13}\)C n.m.r. spectroscopy

2.2.2.1 \(^{13}\)C n.m.r.

The use of \(^{13}\)C n.m.r. as a tool for investigating the microstructure of polybutadienes was first reported by Duch and Grant\(^69\) in 1970. Soon after quite a few papers were published using \(^{13}\)C n.m.r techniques for determining the microstructure of polybutadienes\(^70,71\). Our assignment of the various peaks corresponding to cis, trans and pendant carbon-carbon double bonds of polybutadiene were carried out according to Clague and coworkers\(^72\).
A $^{13}$C n.m.r. spectrum of polybutadiene is shown in Figure 2.5. The peaks at 115 ppm and at 145 ppm are assigned to the carbon atoms of the pendant carbon-carbon double bonds, while the peaks around 130 ppm are assigned to carbon atoms of the back bone carbon-carbon double bonds i.e., cis and trans.

$^{13}$C n.m.r studies of the hydrocarboxylated product firstly give an insight as to whether the hydrocarboxylation reaction has taken place or not, and secondly about the regioselectivity of the reaction. Based on the hydrocarboxylation reaction carried out using Alper's method (see chapter 4), on 100% cis polybutadiene (see Chapter 4, Table 4.2 polymer number 7, page 169 ) we were able to assign the single peak at 176 ppm to the carboxylic acid carbon attached directly to the back bone of the polymer. Based on this observation, assignments were then made for carboxylic acids peaks at 175 ppm and 173 ppm to the terminal carbon position and the penultimate carbon of the pendant group. It is worth mentioning here that when hydrocarboxylation is carried out based on Knifton's method (see chapter 3), only a single carboxylic acid peak at 175 ppm arises. Given that this method is known to be highly specific to hydrocarboxylation at the terminal carbon atom of terminal double bonds, we can assign the peak at 175 ppm to such terminal carboxylic acid groups suggesting that the hydrocarboxylation reaction is very regiospecific. When using Alper's method for polybutadienes containing both back bone and pendant double bonds, three carboxylic acid peaks i.e., at 176 ppm, 175 ppm and 173 ppm are observed, suggesting that the hydrocarboxylation reaction is non-selective, i.e., with the backbone (176 ppm), and terminal (175 ppm) and penultimate (173 ppm) carbon atoms of pendant groups all being functionalised.
Figure 2.5 $^{13}$C n.m.r spectrum of polybutadiene (phenyl terminated)

Figure 2.6 $^1$H n.m.r. spectrum of polybutadiene (phenyl terminated)
2.2.2.2 \textsuperscript{1}H n.m.r. spectroscopy

2.2.2.2.1 Analysis of starting materials

\textsuperscript{1}H n.m.r. spectroscopy is the precise tool used to calculate quantitatively the extent of hydrocarboxylation. Assignment of various peaks from the \textsuperscript{1}H n.m.r. spectrum of polybutadiene were made according to Rempel and coworkers\textsuperscript{73} and Zymonas and coworkers\textsuperscript{74}. Our calculations are based on the calculations made by Rempel and coworkers\textsuperscript{74} for hydrosilylation of polybutadienes. The \textsuperscript{1}H n.m.r. spectrum of polybutadiene is shown in Figure 2.6.

The peaks in the range of 5.0-5.8 ppm are characteristic of olefinic protons and in the range 1-2.3 ppm are due to aliphatic protons. The polybutadienes used in our study were either phenyl terminated or terminated with other groups like proton, hydroxyl, acid etc. In order to calculate the various constituents i.e., \textit{cis}, \textit{trans}, pendant units in the starting material polybutadiene, we first calculate the percentages of pendant carbon-carbon double bonds among overall double bonds present in the polymer as follows.

Calculation of percentage of terminal double bonds among double bonds

\textsuperscript{1}H n.m.r. spectroscopy is used to calculate the percentage of pendant carbon-carbon double bonds among double bonds present in the polymer. The \textsuperscript{1}H n.m.r. spectrum is shown in Figure 2.6 and the assignment of the corresponding peaks is shown in Figure 2.7.
In the region ‘x’ there are two protons ‘a’ and ‘a’ arising from the back bone double bonds and one proton ‘b’ from the penultimate carbon atom of the pendant double bonds. Two protons ‘c’ arising exclusively in the region ‘y’ of the $^1$H n.m.r. spectrum corresponds to the terminal proton of the pendant carbon-carbon double bonds. The proportion of pendant double bonds as a percentage of the total number of double bonds is given by:

$$\frac{1 \text{ proton from pendant position}}{1 \text{ proton from the pendant dbs } + 1 \text{ proton from bb dbs}} \times 100\%$$

(eq1)

( where bb = back bone and dbs = double bonds).

$$1 \text{ proton from pendant double bond} = \frac{y}{2}$$

(eq 2)

(as region ‘y’ corresponds to 2 protons in the $^1$H n.m.r. spectra)

$$1 \text{ proton from back bone double bonds} = \frac{x-y/2}{2}$$

(eq 3)

Substituting equations (2) and (3) in equation (1) we get

$$= \frac{\frac{y}{2}}{\frac{y}{2}+\frac{x-y/2}{2}} \times 100$$

$$= \frac{y}{y+x-y/2}$$
\[ y = \frac{y}{x+y/2} \]

i.e., \[ \frac{m}{m+n} = \frac{2y}{2x+y} \times 100 \] (eq 4)

For calculation of the percentages of the various constituents i.e., cis, trans and pendant carbon-carbon double bonds present in the starting material we use both \(^1\text{H}\) and \(^{13}\text{C}\) n.m.r. and the calculation is done as follows:

\[
\left(\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2\right)_{n}\left(\text{CH}_2-\text{CH}-\text{CH}_2\right)_{m}\left(\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2\right)_{q}
\]

Figure 2.8

a, b, c, d, e, f, g, h, i, and j are ten types of protons in different chemical environments. If n, m, and q are the fractions of butadiene units which are backbone, pendant and saturated respectively (see Figure 2.8), then

\[ n + m + q = 1 \]  

(eq 5)

Where  
\[ n = \text{fraction of back bone double bonds} \]
\[ m = \text{fraction of terminal double bonds} \]
\[ q = \text{fraction of saturated units} \]
The percentage of terminal double bonds among overall double bonds has been shown previously to be given by (see equation 4)

$$\frac{m}{m+n} = \frac{2y}{2x+y} \quad \text{eq (6)}$$

The degree of unsaturation for the starting material i.e., polybutadiene is calculated as follows.

$$\frac{2n+3m}{4n+3m+8q} = \frac{\text{integration of olefinic protons}}{\text{integration of aliphatic protons}} \quad \text{eq (7)}$$

Substituting equations (5) and (6) in equation (7) will give the values of m, n, and p. The percentages of cis and trans double bonds can be estimated from the integration of $^{13}$C n.m.r. (cis peak at 32.7 ppm and trans peak at 27 ppm) and from the value of m.

In the following section an example of a polymer with a number average weight of 1000 is chosen and the calculation is shown below.

Figure 2.6 is the H n.m.r. spectrum of polybutadiene phenyl terminated. Since there is a certain degree of saturation present in the polymer the value of \( q \) in these equations will be taken into account.

$$n + m + q = 1 \quad \text{eq (8)}$$

From the \(^1\text{H}\) n.m.r. the integral value of \( 'y' = 12 \text{ mm} \) and the integral value of \( 'x' = 13 \text{ mm} \). Therefore,

$$\frac{m}{m+n} = \frac{2y}{2x+y} = 0.63 \quad \text{eq (9)}$$

\[ m (1-0.63) = 0.63n, \quad \text{Therefore,} \quad m = 1.7n \]
Substituting the value of $m$ from equation (9) in equation (8)

\[
p = 1 - 2.7n\text{ eq}(10)
\]

\[
\frac{2n + 3m}{4n + 3m + 8p} = \frac{25}{46} = 0.54\text{ eq}(11)
\]

Substituting the values of $m$ from equation (9) and $p$ from equation (10) in equation (11), we get

\[
\frac{2n + 3 \times 1.7n}{4n + 3 \times 1.7n + 8 - 8 \times 2.7n} = 0.54
\]

\[
\frac{7.1n}{8 - 12.5n} = 0.54
\]

\[n = 0.31\]

Substituting the value of $n$ in equations (9) and (10) we get

\[m = 0.53, \text{ and } q = 0.16\]

It is possible to estimate the relative amounts of cis and trans double bonds from the $^1$H n.m.r. spectrum. However, this is obtained from the $^{13}$C n.m.r. spectrum as follows.

Carbon atoms $\alpha$ to trans double bonds give rise to a signal at 32.7 ppm, whilst carbon atoms $\alpha$ to cis double bonds give rise to a signal at 27 ppm. Being in similar environments, their Overhauser enhancement are likely to be similar to one another so that integration can be used to compare the relative amounts of these species. The $^{13}$C n.m.r. spectrum of the same polymer as described above, is shown in Figure 2.5.
Measuring the height of the integration peak at 32.7 ppm gives $trans = 2$

Measuring the height of the integration peak at 27 ppm gives $cis = 1.5$

Therefore proportion of $trans = \frac{2}{3.5} = 0.57 \sim 0.6$

proportion of $cis = \frac{1.5}{3.5} = 0.42 \sim 0.4$

From the $^1$H n.m.r. spectrum we know the percentage of backbone double bonds as 31%

Therefore $31 \times 0.6 = 18.6 \sim 19\%$ $trans$

$31 \times 0.4 = 12.4 \sim 12\%$ $cis$.

The composition of polybutadiene is $cis = 12\%$

$trans = 19\%$

pendant = 53\%

saturation = 16\%

The reported value from Aldrich Chemical Company was 5\% $cis$, 10\% $trans$, 45\% pendant, and 40\% saturated units.

In principle, a very similar method can be used for determining the micro-structure of a functionalised polybutadiene provided either that the functional group introduced has resonances in the $^1$H n.m.r. which are significantly shifted from the other polymer resonances (e.g., for introduced -Si(CH$_3$)$_2$Cl or -CHO groups) or that the protons originally on the double bonds of the polymer resonate in a different
but distinctive region of the $^1$H n.m.r. spectrum (epoxidised polymers are those containing added -OH groups). For hydrocarboxylated polymers, however, neither of these criteria holds so that direct analysis of the degree of hydrocarboxylation of the polymer is difficult. It should be noted that the position and intensity of the signal from the -OH protons vary considerably depending upon the presence of water in the polymer or the n.m.r. solvent (usually $d^6$-acetone or $d^4$-methanol) so this cannot be used for determining the degree of hydrocarboxylation of the polymer. Uncertainty about the position and intensity of the resonance from the -OH protons also causes difficulties in using the relative intensities of the aliphatic and olefinic regions of the spectrum for product analysis, these are further exacerbated by the presence of traces of other solvents (e.g. methyl isobutyl ketone) which are difficult to remove from the polymer in their entirely. We have, therefore, had to develop other methods for the analysis of the hydrocarboxylated polymers.

### 2.2.2.2.2. Method 1

This method is used when hydrocarboxylation takes place only at the pendant double bonds

![Figure 2.9](image)

where $n = \text{fraction of back bone double bonds}$
Values of \( n, m, \) and \( q \) (Figure 2.9) are known from the starting material. Since the hydrocarboxylation reaction is very regiospecific, we assume that after the reaction the percentage of the backbone double bonds remains unchanged i.e., the value of \( n \) remains the same. Therefore in the hydrocarboxylated product

Let, \( n = n_0 \) \hspace{1cm} (eq12)

(where \( n_0 \) is the backbone double bonds in the starting material).

\[
\frac{m}{m+n} = \frac{2y}{2x+y} = A \hspace{1cm} \text{(eq13)}
\]

where \( A \) is the ratio of pendant double bonds to all double bonds in the hydrocarboxylated polybutadiene.

On substituting equation (13) in equation (12) we get

\[
m = n_0A + mA
\]

\[
m(1-A) = n_0A
\]

\[
m = \frac{n_0A}{1-A}
\]

\[
m + p = m_0
\]

(where \( m_0 \) is the percentage of pendant double bonds in the starting material).
\[ p = m_0 - \frac{n_0 A}{1-A} \]

Example

Let us consider a specific example where hydrocarboxylation takes place only at the terminal position of the pendant group.

![Graph with peaks and annotations](image)

Figure 2.10 \(^1\)H n.m.r. spectrum of hydrocarboxylated polybutadiene

The composition of the starting polybutadiene chosen as an example is

- **cis** = 12%
- **trans** = 19%
- **pendant** = 53%
- **Saturation** = 16%

therefore, accordingly \( n_0 = 31\% \), \( m_0 = 53\% \), and \( q_0 = 16\% \).
Then \( n = n_0 = 31\% \) \hspace{1cm} \text{(eq14)}

Since hydrocarboxylation takes place only at the terminal carbon atom, the value of \( n_0 \) remains unchanged. The percentage of terminal double bonds among double bonds is calculated by measuring the heights of the integration of the regions ‘x’ and ‘y’ of the \(^1\)H n.m.r. spectrum (see Figure 2.10).

\[
\frac{m}{m+n} = \frac{2y}{2x+y} = \frac{10}{79} = 0.13 \hspace{1cm} \text{(eq15)}
\]

On substituting the values of equation (15) in equation (14), we get

\[
m = m \times 0.13 + 0.31 \times 0.13
\]
\[
m (1-0.13) = 0.31 \times 0.13
\]
\[
m = 0.05
\]

Substituting the value of \( m \) in the equation

\[
m + p = m_0
\]

we get \( p = 0.53 - 0.05 = 0.48 \)

Hence the final composition of the hydrocarboxylated product is:

- cis = 12\%
- trans = 19\%
- pendant = 5\%
- COOH = 48\%
- saturation = 16\%
This would correspond to an "overall conversion" of butadiene units to carboxylic acids of 48% and a conversion based on pendant carbon-carbon double bonds of 91%.

2.2.2.2.3. Method 2

This method is used when hydrocarboxylation takes place either at the backbone double bond or at the pendant double bond position or both. The polymers studied were phenyl terminated and hence integrations relative to the phenyl region, which does not change after reaction is used as standard for calculating the degree of unsaturation in the hydrocarboxylated polybutadiene as shown below (Figure 2.11).

where,  \( \bullet = \text{C}_6\text{H}_5 \)

\( n = \) fraction of backbone double bonds
\( r = \) fraction of carboxylic acid at the backbone
\( m = \) fraction of terminal double bonds
\( p = \) fraction of carboxylic acid at the terminal carbon position
\( q = \) fraction of saturated units

Figure 2.11
The percentage of saturated carbon-carbon double bonds remains unaffected.

Firstly, we calculate the ratio of pendant double bonds to all double bonds in the hydrocarboxylated product from the $^1$H n.m.r. spectra as before i.e.,

$$\frac{m}{n+m} = A \quad \text{eq}(16)$$

(where $A = \frac{2y}{2x+y}$)

Secondly,

$$\frac{2n+3m}{\phi} = B_1 \quad \text{eq}(17)$$

(i.e., $\frac{\text{integration of olefinic protons in the product polymer}}{\text{integration of phenyl protons in the product polymer}}$)

$$\frac{2n_0+3m_0}{\phi} = B_2 \quad \text{eq}(18)$$

(i.e., $\frac{\text{integration of olefinic protons in the starting material}}{\text{integration of phenyl protons in the starting material}}$)

(where $n_0$ and $m_0$ are percentages of backbone double bonds and terminal double bonds in the starting material respectively).

Dividing equation (17) by equation (18) we get

$$\frac{2n+3m}{2n_0+3m_0} = \frac{B_1}{B_2} \quad \text{eq}(19)$$
We know, \[ n + r = n_0 \] \hspace{1cm} \text{eq}(20)

Similarly \[ m + p = m_0 \]

Solving the four equations (16), (18), (19), and (20) simultaneously we get the composition of the hydrocarboxylated polymer.

\[
\begin{align*}
n &= \frac{C \cdot AC}{A+2} \\
r &= n_0 - n \\
m &= \frac{AC}{2+A} \\
p &= m_0 - m
\end{align*}
\] \hspace{1cm} \text{eq}(21) \hspace{1cm} \text{eq}(22) \hspace{1cm} \text{eq}(23) \hspace{1cm} \text{eq}(24)

where, \[ C = \frac{B_1}{B_2} (2n_0) + \frac{B_1}{B_2} (3m_0) \]

However before applying this method for calculating the percentage of conversion at the internal and pendant carbon-carbon double bond position, we can check this method by applying to the previous hydrocarboxylation reaction where conversion takes place only at the pendant position. In theory we should not get any conversion of the back bone double bonds at all in the hydrocarboxylated product.

Figure 2.12 (a) and (b) shows the \(^1\)H n.m.r. spectra of the hydrocarboxylated product and the starting polybutadiene. Firstly, we calculate the ratio of pendant double bonds to all the double bonds in the hydrocarboxylated product as shown previously according to equation (4).
Figure 2.12 (a) 

Figure 2.12 (b) 

Figure 2.12 (a) $^1$H n.m.r. spectrum of hydrocarboxylated polybutadiene
(b) $^1$H n.m.r. spectrum of polybutadiene(phenyl terminated)
\[
\frac{m}{m+n} = A = 0.13 \quad \text{eq}(25)
\]

Secondly according to equation 17

\[
\frac{2n+3m}{\phi} = \frac{42}{14} = 3 = B_1 \quad \text{eq}(26)
\]

Thirdly, according to equation 18

\[
\frac{2n_o+3m_o}{\phi} = \frac{25}{3} = 8.3 = B_2 \quad \text{eq}(27)
\]

Therefore

\[
\frac{B_1}{B_2} = \frac{3}{8.3} = 0.36 \quad \text{eq}(28)
\]

Substituting the value obtained in equation 28 in the following equation will give the value of C

\[
C = \frac{B_1}{B_2} (2n_o) + \frac{B_1}{B_2} (3p_o)
\]

\[
C = 0.8
\]

Substituting the values of \(A=0.13\) and \(C=0.8\) in equations 21, 22, 23, and 24 will give the value of \(n, r, m, \) and \(p\) as \(n = 32.7\%\), \(r = -1.7\), \(m = 5\%\) and \(p = 48\%\).

The negative value obtained clearly shows that none of the backbone double bonds have reacted and it is close enough to give confidence in the method. Hence the final composition of the hydrocarboxylated product will be
Back bone double bonds = 31%
Remaining pendant double bonds = 5%
Carboxylic acid at the pendant position = 48%
(i.e., Overall conversion)

This would correspond to an "overall conversion" of butadiene units to carboxylic acids of 48% and a conversion based on pendant carbon-carbon double bonds of 91%.

The values obtained on applying this method is clearly in agreement to those of the values obtained using Method 1 (Overall conversion = 48%).

We will now apply this method to a specific example where hydrocarboxylation reaction occurs both at the back bone and at the pendant carbon-carbon double bond position. Figure 2.6 shows the $^1$H n.m.r. spectrum of the starting material and Figure 2.13 the $^1$H n.m.r. spectrum of the hydrocarboxylated product.

![Figure 2.13 $^1$H n.m.r. spectrum of hydrocarboxylated polybutadiene (phenyl terminated)](image-url)
Firstly, we calculate the ratio of the pendant double bonds to all double bonds using equation (4) as mentioned earlier.

\[ \frac{m}{n+m} \frac{2y}{2x+y} = \frac{14}{31} = 0.45 = A \]  
\hspace{1cm} \text{eq}(29) \]

Secondly, according to equation 17;

\[ \frac{2n+3m}{\phi} = \frac{19}{36.5} = 0.51 = B_1 \]  
\hspace{1cm} \text{eq}(30) \]

Thirdly, according to equation 18;

\[ \frac{2n_0+3m_0}{\phi} = \frac{25}{3} = 8.33 = B_2 \]  
\hspace{1cm} \text{eq}(31) \]

Therefore \( \frac{B_1}{B_2} = 0.062 \)  
\hspace{1cm} \text{eq}(32) \]

Substituting the value of \( \frac{B_1}{B_2} \) obtained in equation 32 in the following equation will give the value of C

\[ C = \frac{B_1}{B_2} (2n_0) + \frac{B_1}{B_2} (3m_0) \]

\[ C = 0.14 \]

Substituting the values of \( A = 0.45 \) and \( C = 0.14 \) in equations 21, 22, 23, and 24 will give the values of n, r, m, and p. As n = 3\%, r = 28\%, m = 3\%, p = 50\%. Thus the composition of the final hydrocarboxylated product is
Remaining back bone double bonds = 3%
Carboxylic acid at the back bone = 28%
Remaining pendant double bonds = 3%
Carboxylic acid at the pendant position = 50%
Saturated carbon-carbon bonds = 16%
Total carboxylic acid = 78%

In order to check the accuracy of the method applied a conductometric titration was performed on the above hydrocarboxylated polymer. A graphical representation of the titration is shown in Figure 2.14. A few problems were encountered during the titration.

![Conductometric titration graph](image)

**Figure 2.14.** Conductometric titration between hydrocarboxylated polybutadiene (in NaOH) and HCl
In this polymer also the final end point (see Figure 2.14) could not be determined precisely as precipitation of the polymer took place. The end point based on titration was 15.5. This means that some of the Na\(^+\) ions bound to the polymer chain are not available for titration with HCl. However, since this is a back titration, the final end point should correspond to the end point obtained for titration of the same amount of NaOH used. We, therefore, assumed that the final end point should be at 19 cm\(^3\).

**Calculation**

0.01 gm of the hydrocarboxylated polymer was dissolved in 20 cm\(^3\) of 0.01 M NaOH and titrated with 0.01 M of HCl. Blank titration of NaOH with HCl gave 19.0 cm\(^3\) as the end point i.e., neutralisation point. 
First end point when all the NaOH has reacted with HCl = 10.5
Second end point when COONa bound to the polymer has reacted with HCl (based on blank titration) = 19.0
Therefore 0.01 gm of hydrocarboxylated product is equivalent to 8.5 cm\(^3\) of 0.01 cm\(^3\) M HCl.
This in terms of moles will be

\[
0.01 \text{ gm} = \frac{8.5 \times 0.01}{1000} = 8.5 \times 10^{-5} \text{ moles}
\]

Let 'M' be the mass in a.m.u. of the hydrocarboxylated product, which contains one proton, then

\[
\frac{0.01 \text{gm}}{M} = 8.5 \times 10^{-5} \text{ moles}
\]

Therefore \( M = 118 \)
Substituting this value of M in the equation

\[ \frac{a \times 100 + b \times 54 + c \times 56}{a} = M \]

where \(a\) is the proportion of carboxylic acid units in the chain,
\(b\) is the proportion of unreacted units
\(c\) is the proportion of saturation in the starting material = 0.16

\[ \frac{a \times 100 + b \times 54 + 0.16 \times 56}{a} = 118 \]

\[ b \times 54 + 0.16 \times 56 = 18a \]

We know \(a + b = 0.84\)
Therefore \(b = 0.84 - a\)

\(0.84-a)54 + 0.16 \times 56 = 18a\)
\(45.36 - 54a + 8.96 = 18a\)
\(54.32 = 72a\)
\(a = 0.75\)

Substituting the value of \(a = 0.75\) in the equation \(a + b = 0.84\)
we can arrive at the final composition of the hydrocarboxylated product as

Overall carboxylic acid = 75%
Remaining double bonds = 9%
Saturated C-C bonds = 16%

As can be seen the results obtained from conductometric titration is in reasonable agreement with those from the end group analysis method.
2.2.2.4 Method 3

In some cases there are no suitable protons from end groups to use as a standard and where hydrocarboxylation occurs at both the pendant and backbone double bonds, it is necessary to use a combination of both $^1H$ and $^{13}C$ n.m.r. spectroscopy. Due to different relaxation times for the different carbon atoms and different nuclear Overhauser effects mean that integration of $^{13}C$ resonances cannot be used without some kind of calibration, it is necessary to obtain some indication of the relative sensitivities of the n.m.r. experiment to the $^{13}C$ resonances from the carboxylic acid carbon atoms and from the olefinic carbon atoms. For this method to be successful, it is necessary for all $^{13}C$ spectra of the product polymers to be run under identical conditions so that relative sensitivities obtained from one spectrum can be assumed to be the same as those existing in another spectrum.

We define a factor, $f$, which is the ratio between the receptivities of the olefinic and carboxylic carbon atom both terminal and internal. If $'g'$ is the integral obtained from the carboxylic acid carbon atom and $'h'$ is the integral obtained from the resonance near 145 ppm ($CH_2$ of pendant double bond), then the factor $f$ is then given by

$$\frac{gf}{h} = \frac{p}{m}$$

(eq33)

(where $p$ corresponds to the fraction of carboxylic acid and $m$ corresponds to the remaining pendant carbon-carbon double bonds (see Figure 2.11).

Calculating this from a series of different phenyl terminated polymers for which the end group analysis has been used to obtain $p$
and m gives values of f as shown in Table 2.1. An average value of f = 4.4 is obtained. (see Table 4.2, page 169 for composition*)

<table>
<thead>
<tr>
<th>Polymer Number *</th>
<th>'F' factor</th>
</tr>
</thead>
<tbody>
<tr>
<td>2</td>
<td>4.2</td>
</tr>
<tr>
<td>4</td>
<td>4.4</td>
</tr>
<tr>
<td>5</td>
<td>4.6</td>
</tr>
</tbody>
</table>

Table 2.1

Using this value of f = 4.4 we can calculate the COOH conversion at the back bone and at the terminal carbon position of the unknown hydrocarboxylated product as shown below.

The relative amounts of back bone and pendant double bonds remaining in the polymer are available from the $^1$H n.m.r. as described previously according to equation 4.

If the ratio of the integrals of the $^{13}$C resonances from the terminal carbon atom of the remaining pendant double bonds to that from the carboxylic carbon atoms $^{13}$C is given by D, then

$$\frac{m}{p} = \frac{D}{f} \quad \text{eq}(34)$$

remembering (from equation 4 and also see Figure 2.11)

$$\frac{m}{n + m} = A,$$

$$n + r = n_o, \text{ and } m + p = p_o,$$

Therefore, $p = \frac{m_o}{D} \frac{1}{1 + \frac{f}{f}} \quad \text{eq}(35)$
\[ m = \frac{m_0}{f(1+D)} \quad \text{or} \quad m = p_0 - p \quad \text{eq}(36) \]

\[ n = \frac{m_0}{A(1+D)}(1-A) \quad \text{eq}(37) \]

\[ r = n_0 - n \]

The values of \( p, m, n \) and \( r \) will give both the amount of unreacted double bonds and the amount of conversion (both at the backbone and at the pendant position).

However before applying this method for calculating the percentage conversion (at the internal and pendant carbon-carbon position) of a non-phenyl terminated product, we can check this method by applying to the previous hydrocarboxylation reaction (phenyl terminated polymer).

**Example 1** (phenyl terminated polymer)

The \(^1\text{H}\) n.m.r. spectrum is shown in Figure 2.13 and the \(^{13}\text{C}\) spectrum is shown in Figure 2.15.

From \(^{13}\text{C}\) n.m.r. spectrum we calculate the value of \( D = \frac{8}{27} = 0.30 \)

We know the value of \( f = 4.4 \) and \( A = 0.45 \) (value of \( A \) from eq 29)

Applying the values of \( f \) and \( D \) in equation 35 we get,
Figure 2.15  $^{13}$C n.m.r spectrum of hydrocarboxylated polybutadiene (phenyl terminated)
\[ p = \frac{0.53}{1 + \left(\frac{0.30}{4.4}\right)} = \frac{0.53}{1.06} = 0.50 \]

i.e., 50% pendant COOH

Subtracting this value from 53% (initial starting pendant C=C %) will give the remaining pendant carbon-carbon double bonds
i.e., \( m = 53\% - 50\% = 3\% \).

Once again applying the values of \( f = 4.4 \) and \( A = 0.45 \) and \( D = 0.30 \) in equation 37

\[ n = \frac{0.53}{0.45(1+\frac{4.4}{0.30})} (1-0.45) \]

\[ n = 0.0412 \text{ i.e, 4\% back bone double bonds} \]

\[ r = 31\% - 4\% = 27 \% \text{ back bone COOH} \]

The composition of the hydrocarboxylated product will be as shown in Table 2.2. Values obtained based on end group (phenyl) analysis for the same polymers are included for comparison.
Various constituents of hydrocarboxylated polybutadiene

<table>
<thead>
<tr>
<th></th>
<th>Results obtained based on 'f' factor (%)</th>
<th>Results obtained based on end group analysis method (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Remaining bb dbs</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>COOH at bb position</td>
<td>27</td>
<td>28</td>
</tr>
<tr>
<td>Remaining pd dbs</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>COOH at pd position</td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>Saturated C-C bonds</td>
<td>16</td>
<td>16</td>
</tr>
<tr>
<td>Total COOH (bb + pd)</td>
<td>77</td>
<td>78</td>
</tr>
</tbody>
</table>

where bb = back bone, pd = pendant, dbs = double bonds

Table 2.2

Calculation of percentage of carboxylic acid at the terminal carbon position and penultimate carbon position.

Once the amount of carboxylic acid at the pendant position is known it is possible to estimate from $^{13}$C n.m.r. the relative amounts of carboxylic acid at the terminal and the penultimate carbon positions of the pendant carbon units. This is done by measuring the heights of the peaks corresponding to terminal carbon position (175 ppm) and penultimate carbon position (173 ppm). An example is given below (see Figure 2.14)

We know from our calculations, (based on f factor, see Table 2.2) that amount of carboxylic acid at the pendant position = 50 %

Height of the peak at 175 ppm corresponding to terminal carbon position = 18 mm

Height of the peak at 173 ppm corresponding to penultimate carbon position = 9 mm
Therefore 50% COOH = 27 mm
Therefore 18 mm = \( \frac{50}{27} \times 18 = 33\% \)

Hence the amount of carboxylic acid at the terminal carbon position is equal to 33%. From this value obtained we can calculate the amount of carboxylic acid at the penultimate carbon position i.e.,

\[ 50\% - 33\% = 17\% \text{ COOH} \]

Hence the final composition of the hydrocarboxylated product is shown below (Table 2.3)

<table>
<thead>
<tr>
<th>Remaining db bonds%</th>
<th>Carboxylic acid(%)</th>
<th>Total carboxylic acid (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cis Trans Pendant</td>
<td>Bb Tr Int</td>
<td></td>
</tr>
<tr>
<td>4 3 27 33 17</td>
<td></td>
<td>77</td>
</tr>
</tbody>
</table>

where db = double bonds, Bb = back bone, Tr = terminal and Int = Internal

Table 2.3

As can be seen from the table above the values obtained on using factor agree with those of the end group analysis and conductometric titration.

Example 2 (non-phenyl terminated polymer)

The composition of the initial starting material was (37% trans, 23% cis and 40% pendant carbon-carbon double bonds). The \(^1\)H n.m.r
spectrum and $^{13}$C n.m.r. spectrum of the hydrocarboxylated product is shown in Figure 2.16 (a) and (b).

From $^1$H n.m.r. spectrum of the hydrocarboxylated product the value of $A$ is calculated using equation 4 i.e.,

\[
\frac{m}{m + n} = \frac{2 \times 13}{2 \times 19 + 13} = 0.51 = A
\]

From $^{13}$C n.m.r. spectrum of the hydrocarboxylated product the value of

\[D = \frac{11}{7} = 1.6\]

On substituting the values of $A = 0.51$, $D = 1.6$ and $\gamma = 4.4$ in equations 35, 36, 37 and 38 we arrive at the composition of the final hydrocarboxylated product as

Remaining back bone double bonds = 10%
Carboxylic acid at the back bone = 50%
Remaining pendant double bonds = 11%
Carboxylic acid at the pendant position = 29%

By measuring the heights of the peaks corresponding to terminal carbon position (175 ppm) and penultimate carbon position (173 ppm) of the $^{13}$C n.m.r. spectrum we can calculate the amount of carboxylic acid at the terminal position and at the penultimate carbon position.

Height of the peak at 175 ppm corresponding to terminal carbon position = 5.5 mm
Height of the peak at 173 ppm corresponding to penultimate carbon position = 1.5 mm
We know 29% COOH = 7 mm
Figure 2.16 (a) 

$^{1}H$ n.m.r. spectrum of hydrocarboxylated polybutadiene (hydrogen terminated)

Figure 2.16 (b) 

$^{13}C$ n.m.r. spectrum of corresponding hydrocarboxylated polybutadiene
Therefore 5.5 mm = \( ? \) COOH

\[ \frac{29}{7} \times 5.5 = 23\% \]

Hence the amount of carboxylic acid at the terminal carbon position is equal to 33%. From this value obtained we can calculate the amount of carboxylic acid at the penultimate carbon position, i.e.,

\[ 29\% - 23\% = 6\% \text{ COOH} \]

The final composition of the hydrocarboxylated product is shown below (Table 2.4).

<table>
<thead>
<tr>
<th>Remaining db bonds</th>
<th>Carboxylic acid</th>
<th>Total carboxylic acid</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cis Trans Pendant</td>
<td>Bb Tr Int</td>
<td></td>
</tr>
<tr>
<td>10 11 50 23 6</td>
<td>79</td>
<td></td>
</tr>
</tbody>
</table>

where db = double bonds, Bb = back bone, Tr = terminal and Int = Internal

Table 2.4

2.2.3. Infrared spectroscopy

Various workers have developed IR methods for analysing polymers. However, the earlier attempts were seriously handicapped by the lack of suitable reference compounds for selecting the analytical bands for calculating the absorption coefficients.
By using the bands at 966, 909, 724 cm\(^{-1}\) obtained from the I.R. analysis of polybutadienes which were not rich in one particular kind of saturation, Hampton\(^75\) tried to improve the absorption coefficients by a successive series of approximations. Binder\(^76\) assigned the band at 680 cm\(^{-1}\) to \textit{cis} units, by a modification of the method due to Hampton. Richardson\(^77\) did not directly measure 1,4 \textit{cis} units. Instead he used the typical wavelength at 966, and 909 cm\(^{-1}\) for calculating percentages of \textit{trans} and pendant units assuming the rest to be \textit{cis} units.

Silas and coworkers\(^78\) in 1959 published a method for determining 1,4 \textit{cis} units, using an empirical function for the area of the absorption bands between 833 and 634 cm\(^{-1}\). This was used for the calculation of the absorption coefficients of polybutadiene with quite high contents of 1,4 \textit{trans}, 1,2 and 1,4 \textit{cis} units. However this method did not take into account the region between 833 and 634 cm\(^{-1}\) where absorption bands can also occur due to non 1,4 \textit{cis} units.

In the same year Morrero and coworkers\(^79\) came up with formulae for calculating the various units of polybutadiene. They measured the bands at 967, 913-910, 740-732 (cm\(^{-1}\)) generated by out of plane deformation of the =C-H bond in 1,4 \textit{trans}, 1,2 and 1,4 \textit{cis} units respectively and proposed formulae based on measurement of absorption coefficients for calculating the different percentages of 1,4 and 1,2 units in polybutadiene.

\[
D_{967} = (K_{1,4 \text{ trans}} \cdot P_{1,4 \text{ trans}} + K_{1,2} \cdot P_{1,2} + K_{1,4 \text{ cis}} \cdot P_{1,4 \text{ cis}}) \cdot S
\]

\[
D_{913-910} = (K_{1,2} \cdot P_{1,2} + K_{1,4 \text{ cis}} \cdot P_{1,4 \text{ cis}}) \cdot S
\]

\[
D_{740-732} = (K_{1,2} \cdot P_{1,2} + K_{1,4 \text{ cis}} \cdot P_{1,4 \text{ cis}}) \cdot S
\]
Where $D_{\lambda}$ is the optical density at wavelength $\lambda$.

$K_{x,\lambda}$ is the absorption coefficient of the component $x$ at the wavelength $\lambda$ expressed in mg$^{-1}$ cm$^{-1}$, 10 cm$^3$

$P_x$ is the weight of the component $x$ in 10 cm$^3$ of the solution

$S$ is the thickness in cm of the cell used.

The relative percentage is given by

$$C_{1,4 \text{ trans} \%} = P_{1,4 \text{ trans}} \cdot \frac{100}{P}$$

$$C_{1,2 \%} = P_{1,2} \cdot \frac{100}{P}$$

$$C_{1,4 \text{ cis} \%} = P_{1,4 \text{ trans}} \cdot \frac{100}{P}$$

$P$ being the weight in milligrams of polybutadiene dissolved in 10 cm$^3$ CS$2$.

Nippon Soda Company$^{80}$ modified the above method by measuring the absorption of the trans double bonds at 967 cm$^{-1}$ and 1,2 double bonds at 912 cm$^{-1}$ in carbon disulphide solution using a liquid cell and arrived at the empirical formulae as shown below.

$$V(1,2- \text{ C-H}) = V_{\text{abs}} \times 0.03746$$

$$T(\text{trans} \text{ C-H}) + T_{\text{abs}} \times 0.04292 - V_{\text{abs}} \times 0.00129$$

$$1,2- \text{ C-H} (%) = \frac{V}{V+T} \times 100$$

$$\text{Trans} \text{ C-H} (%) = \frac{T}{V+T} \times 100$$

(where $V$= pendant and $T$= trans units).
However there are a few problems that we encountered in using the above mentioned methods of analysis. This is illustrated as follows.

Polybutadiene of the composition - 12% cis, 19% trans, 53% pendant, and 16% saturation and its hydrocarboxylated product (selective hydrocarboxylation of the pendant carbon-carbon double bonds) of the composition 12% cis, 19% trans, 10% pendant, 43% COOH (overall conversion) and 16% saturation previously determined by $^1$H n.m.r. spectroscopy was chosen to illustrate the method of determining the composition of the hydrocarboxylated product by I.R. spectroscopy.

Figure 2.17 (a) and (b) shows the I.R. spectra of polybutadiene and its corresponding hydrocarboxylated product in the region 1200 - 750 cm$^{-1}$. The peaks at 912 cm$^{-1}$ is associated with pendant double bonds, and the peak at 967 cm$^{-1}$ is associated with trans double bonds. The heights of the peaks are measured from the base line as shown in Figure 2.17 (a) and (b).

<table>
<thead>
<tr>
<th>Polymer</th>
<th>Pendant (912 cm$^{-1}$)</th>
<th>Trans (967 cm$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Starting material</td>
<td>78 mm</td>
<td>33 mm</td>
</tr>
<tr>
<td>Product material</td>
<td>55 mm</td>
<td>40 mm</td>
</tr>
</tbody>
</table>

Therefore if the product had the same composition as the starting material, the height of the pendant peak at 912 cm$^{-1}$ should be

$$\frac{78 \times 40}{33} = 95.54 \text{ mm}$$

But actually the height of pendant peak at 912 cm$^{-1} = 55 \text{ mm}$
Figure 2.17 (a) I.R. spectrum of polybutadiene
(b) I.R. spectrum of corresponding hydrocarboxylated polybutadiene
Therefore amount of double bonds remaining

\[
= \frac{55}{94.54} = 0.58
\]

From this the percentage of polymer units remaining as pendant double bonds can be calculated i.e.,

\[
= 0.58 \times 53 = 31%
\]

Percentage of carboxylic acid = 53 - 31 = 22%

Hence the final composition of the hydrocarboxylated product as determined by I.R. is shown in Table 2.4. The results from \(^1\)H n.m.r. analysis for the same polymer is shown for comparison.

<table>
<thead>
<tr>
<th>Various constituents of hydrocarboxylated polybutadiene</th>
<th>Results obtained from Infrared Spectroscopy (%)</th>
<th>Results obtained from (^1)H n.m.r. spectroscopy (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cis</td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>Trans</td>
<td>19</td>
<td>19</td>
</tr>
<tr>
<td>Pendant</td>
<td>31</td>
<td>10</td>
</tr>
<tr>
<td>COOH (Over all)</td>
<td>22</td>
<td>43</td>
</tr>
<tr>
<td>Saturation</td>
<td>16</td>
<td>16</td>
</tr>
</tbody>
</table>

Table 2.4

From the table it can be seen that the results obtained from I.R. analysis is not in agreement with the \(^1\)H n.m.r. results. This may be due to the following:

1. Extensive overlap of the relevant bands in the hydrocarboxylated
Figure 2.18 (a) I.R. spectrum of polybutadiene
(b) I.R. spectrum of corresponding hydrocarboxylated polybutadiene
product make it difficult to obtain measurements, as an accurate assessment of the baseline could not be made (see Figure 2.17 (a) and (b)).

2. Furthermore, the products are only sparingly soluble in organic solvents and hence only give weak solutions.

Therefore it can be concluded from the above that only qualitative and not quantitative information could be obtained about the extent of the hydrocarboxylation reaction. The I.R. spectra of both the starting material i.e., polybutadiene showing the various peaks is shown in Figure 2.18 (a) and the hydrocarboxylated product is shown in Figure 2.18 (b). The disappearance of the characteristic peaks of unsaturation at 967, 913-910, and 740-732 cm$^{-1}$ and the appearance of the characteristic peak for the acid group at 1727 cm$^{-1}$ and a broad peak at 3310 cm$^{-1}$, characteristic of the OH stretching vibration confirms a qualitative hydrocarboxylation reaction.

### 2.2.4. Microanalysis

Microanalysis of polybutadiene (initial composition of the starting material 2% saturated units, 14%trans, 84% pendant carbon-carbon double bonds) selectively hydrocarboxylated at terminal carbon position of the pendant group (>80% conversion) gave the following results on elemental analysis.

<table>
<thead>
<tr>
<th></th>
<th>Carbon</th>
<th>Hydrogen</th>
<th>Sodium</th>
</tr>
</thead>
<tbody>
<tr>
<td>Theoretical Composition</td>
<td>62.12</td>
<td>7.42</td>
<td>6.66</td>
</tr>
<tr>
<td>Result of Analysis</td>
<td>61.22</td>
<td>10.87</td>
<td>7.10</td>
</tr>
</tbody>
</table>
N.M.R. studies of the hydrocarboxylated product does show the presence of a trace of triphenylphosphine oxide. The discrepancy in the increase of the percentage of hydrogen observed is probably due to the presence of water in the product, hydrogen bonded to sodium ions or to the carboxylic acid group.

The amount of sodium in the hydrocarboxylated product is higher than expected from the titration results. A possible reason for this is that during titration, the sodium ions have enough time to leave the polymer and react with HCl. In the preparations, however, the polymer is precipitated very quickly, by addition of a large number of excess of 7M HCl. Under these conditions, it is evident that some of the sodium ions are trapped within the bulk polymer (see Chapter 3, section 3.4 Experimental Section).

2.3 Conclusion

We have examined four polymers using a variety of different methods. The results obtained by various methods are shown in Table 2.5. Examination of the table shows a good correlation between conductometric titration or methods using $^1$H n.m.r. and $^{13}$C n.m.r. for the selectively functionalised polymer, suggesting that these methods are relatively reliable, although $^1$H and $^{13}$C n.m.r. spectroscopy give more detailed information. For polymers which are phenyl terminated and in which hydrocarboxylation occurs at more than one site, methods based on end group analysis are applicable whilst for polymers that are not phenyl terminated a combination of $^1$H and $^{13}$C n.m.r data are required for analysis. The level of confidence of the results obtained for these polymers must be considered lower than that pertaining to the
selectively hydrocarboxylated polymers. Finally, there appears to be serious problems for the analysis of hydrocarboxylated polybutadienes using I.R. spectrosocopy.

<table>
<thead>
<tr>
<th>Polymer</th>
<th>Method of analysis</th>
<th>bb dbs (n) (%)</th>
<th>bb COOH (r) (%)</th>
<th>pd dbs (m) (%)</th>
<th>pd COOH (p) (%)</th>
<th>Saturated units (q) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>(i) Titration</td>
<td>31</td>
<td>-</td>
<td>12</td>
<td>41</td>
<td>16</td>
</tr>
<tr>
<td></td>
<td>(ii) $^1$H n.m.r.</td>
<td>31</td>
<td>-</td>
<td>10</td>
<td>43</td>
<td>16</td>
</tr>
<tr>
<td></td>
<td>(iii) I.R.</td>
<td>31</td>
<td>-</td>
<td>31</td>
<td>22</td>
<td>16</td>
</tr>
<tr>
<td>B</td>
<td>(i) $^1$H n.m.r.</td>
<td>31</td>
<td>-</td>
<td>5</td>
<td>48</td>
<td>16</td>
</tr>
<tr>
<td></td>
<td>(ii) End group analysis</td>
<td>31</td>
<td>-</td>
<td>5</td>
<td>48</td>
<td>16</td>
</tr>
<tr>
<td>C</td>
<td>(i) End group analysis</td>
<td>3</td>
<td>28</td>
<td>3</td>
<td>50</td>
<td>16</td>
</tr>
<tr>
<td></td>
<td>(ii) $^1$H and $^{13}$C n.m.r.*</td>
<td>4</td>
<td>27</td>
<td>3</td>
<td>50</td>
<td>16</td>
</tr>
</tbody>
</table>

* denotes calculations based on f factor, bb = back bone, dbs = double bonds, pd = pendant, bb COOH = back bone carboxylic acid, pd COOH = pendant carboxylic acid.

Table 2.5
Chapter 3

Hydrocarboxylation of polybutadienes catalysed by the palladium complex \([\text{PdCl}_2(\text{PPh}_3)_2]\)-SnCl₂.

3.1. Introduction

In this chapter we will discuss our results on the hydrocarboxylation of polybutadienes catalysed by the palladium complex \([\text{PdCl}_2(\text{PPh}_3)_2]\)-SnCl₂. However, it is pertinent to first review the various post polymeric modifications such as, hydrosilylation, hydrogenation, hydroformylation and epoxidation, on preformed polymers, particularly polybutadienes.

3.1.1 Hydrosilylation

The first report on hydrosilylation of a polymer was reported by Barry and coworkers. They used trichlorosilane to hydrosilylate natural hydrocarbons e.g. guttapercha and balata, as well as synthetic elastomeric butadiene-styrene copolymers (Equation 3.1). No catalyst was used in these reactions.

\[
\text{CH} = \text{CH}_2 + \text{HSiCl}_3 \xrightarrow{300^\circ C} \text{CH}_2\text{CH}_2\text{Si} \quad \text{where, } \text{CH} = \text{CH}_2 = \text{natural rubber or synthetic polymer}
\]

Equation 3.1.
The first report of a catalytic, post hydrosilylation of polybutadiene using a transition metal catalyst complex appeared in a Japanese patent by Kunio and Kimitaka. They were effective in hydrosilylating 1,2 polybutadiene with pentamethyldisiloxane to give hydrosilylated polybutadiene (Equation 3.2).

\[
\begin{align*}
\text{[1,2-Polybutadiene]} + (\text{CH}_3\text{Si-O-Si(}\text{CH}_3\text{)H}) \rightarrow \text{[Hydrosilylated Polybutadiene]}
\end{align*}
\]

Equation 3.2

Cameron and Qureshi have reported the exclusive hydrosilylation of polybutadiene containing both internal and pendant alkene bonds exclusively at the terminal position (Equation 3.3). These hydrosilylated polybutadienes were then coupled with salts of polystyrene and poly(a-methyl styrene).

Michel and coworkers in a French patent report hydrosilylation of hydroxy terminated polybutadiene, in the presence of chloroplatinic acid, with the hydrosilylating agent, \([(\text{CH}_3)_2\text{SiH(}\text{CH}_2\text{)}_3\text{OCH}_2\text{CH}_2\text{OC}_2\text{H}_5.\)] Information about the composition of the final hydrosilylated polybutadiene is not given.
Equation 3.3

Michel and coworkers\textsuperscript{[85]} in another European patent report hydrosilylation of the pendant alkene bonds of hydroxy terminated polybutadiene (1,2 units $\geq 90\%$) with $[\text{Bi(\(\text{CH}_2\))_5\text{SiH(\(\text{CH}_3\))}_2]}$, again in the presence of chloroplatinic acid. The amount of conversion of the pendant alkene bonds is not reported.

Guo, Farwarha and Rempel\textsuperscript{[74]} have reported hydrosilylating the pendant groups of polybutadienes (with 1,2 units ranging from
45-90%), with a variety of hydrosilylating agents in the presence of 
[RhCl(PPh$_3$)$_3$] (Equation 3.4).

\[
\begin{align*}
\text{RhCl(PPh$_3$)$_3$} & \xrightarrow{109^\circ C} \text{SiR$_2$R'} \\
\end{align*}
\]

where \( p = 0 \) for \( R = R' = \text{CH}_3, \text{C}_2\text{H}_5, \text{C}_4\text{H}_9, \text{C}_2\text{H}_5\text{O} \)
and, \( p = 0 \) for \( R = \text{CH}_3, R' = \text{C}_6\text{H}_5 \)

Equation 3.4

Hydrosilylation with triethylsilane, trimethylsilane, tri-n-butylsilane or 
tri-ethoxy silane results in the exclusive formation of the anti-
Markownikov addition product, while dimethylphenylsilane addition 
produces both anti-Markownikov and Markownikov addition products. 
The results suggest that the nature of the substituents in the silanes play 
an important role in selectivity.

Cole-Hamilton and coworkers\textsuperscript{86} have successfully hydrosilylated 
the \textit{cis} and pendant carbon-carbon double bonds of polybutadiene (40-
90% 1,2 units) with a variety of hydrosilylating agents in the presence 
of chloroplatinic acid (Equation 3.5).
They made several observations. The terminal double bonds are the easiest to hydrosilylate, followed by the cis internal carbon-carbon double bonds. The trans internal carbon-carbon double bonds proved difficult if not impossible to hydrosilylate. For example, a polybutadiene containing 47% pendant carbon-carbon double bonds, 17% cis and 36% trans gave a modified polymer with complete conversion of the pendant carbon-carbon double bonds followed by cis (conversion >95%) and no conversion of the trans groups. Further studies are currently being carried out in order to fully understand the mechanism of selectivity of this reaction system.
3.1.2 Hydrogenation

The first report on hydrogenation of polybutadiene relevant to this thesis using a heterogeneous catalyst was by Yakubehik and Gromova. They hydrogenated polybutadiene at room temperature and pressure over Pd/CaCO₃, Pd on Ni, Pt black, Pt-C activated with chloroplatinic acid, and Raney nickel. They observed that the external double bonds tend to be saturated more rapidly than do the internal ones in the main chain. The catalyst showed decline in activity as the process proceeded. Information about the composition of the starting material or the product is not reported.

Rachapudy and coworkers reported partial hydrogenation of polybutadiene in the presence of homogeneous catalysts prepared by the reaction of metal alkyls (n-butyl lithium and triethyl aluminium) with metal salts (cobalt, nickel 2-ethyl hexanoate, platinum and palladium acetyl-acetonates) at 700°C and ~1 atmosphere for approximately 4 hours. Information about the starting material or the product is not reported.

Doi and co-workers reported complete hydrogenation of 1,4-, atactic 1,2-, and syndiotactic 1,2-polybutadienes under conditions of 100°C and 50 atmospheres of hydrogen in the presence of Wilkinson's catalyst [RhCl(PPh₃)₃] (Equation 3.6).

\[
\begin{align*}
\text{Equation 3.6} \\
103
\end{align*}
\]
Using the same catalyst Rempel and Mohammadi\textsuperscript{90} reported hydrogenation of a acrylonitrile-butadiene copolymer under mild reaction conditions. The carbon-carbon unsaturated bonds undergo quantitative hydrogenation without any hydrogenation of the nitrile functionality. The selectivity of the catalyst for pendant versus internal carbon-carbon double bonds present in the copolymer is markedly influenced by the nature of the solvent employed, although the reasons for this are still under investigation. When the hydrogenation was carried out in 2-butanone, the catalyst showed no selectivity for hydrogenation of pendant over internal carbon-carbon double bonds. On the other hand, when chlorobenzene was employed as the solvent the temperature doubled, the catalyst showed greater selectivity for pendant versus internal carbon-carbon double bonds (Equation 3.7).

\[
\begin{align*}
\text{CN} \quad &\quad \text{CN} \\
\left[\begin{array}{c}
\text{H} \\
\text{H}
\end{array}\right] &\quad \text{H} \\
\text{R} &\quad \text{R}
\end{align*}
\]

\[
\begin{align*}
\text{H}_2 &\quad \text{RhCl}(\text{PPh}_3)_3 \\
\text{20-50}^\circ\text{C}, \text{< 1 atm} &\quad \text{CN} \\
\left[\begin{array}{c}
\text{H} \\
\text{H}
\end{array}\right] &\quad \text{H} \\
\text{R} &\quad \text{R}
\end{align*}
\]

\text{Equation 3.7}

Fairly recently Gilliom\textsuperscript{91} reported the first bulk catalytic hydrogenation of 1,2 polybutadiene and a block copolymer of polystyrene and polybutadiene (PS-PB-PS) in the presence of [RhCl(PPh\textsubscript{3})\textsubscript{3}] and [Ir(COD)(P(CH\textsubscript{3})\textsubscript{2}Ph\textsubscript{2})\textsubscript{2}]PF (COD=1,5-cyclooctadiene). Samples were prepared by evaporating the solution containing the catalysts and the reducible hydrocarbon rubber and then
subjecting them to hydrogenation at moderate temperature and pressure (~3 atm, 60°C). In the case of PS-PB-PS, 80% conversion was obtained within 72 hours using either of the catalysts. More detailed comparison of the catalytic system could not be made owing to the lack of more uniform systems i.e., uniformity of surface area, density, or thickness, local concentration of hydrogen and its variation in permeability through the sample materials.

3.1.3. Hydroformylation

The first hydroformylation reaction on diene-based polymers was reported by Ramp and coworkers92. They hydroformylated 1,4-polybutadiene in the presence of a cobalt catalyst. 1,1-diethoxyethane was used as a condensing agent, preventing gel (cross linking) formation by interacting with the aldehyde groups to form acetals. Some alcohols were also obtained (Equation 3.8).

\[
\text{[Co}_2\text{(CO)}_8\text{]} \quad \text{OC}_2\text{H}_5
\]

\[
\text{OC}_2\text{H}_5
\]

\[
\text{CH}_3\text{CH}_2\text{CHO} \quad \text{OC}_2\text{H}_5
\]

\[
\text{CO}_2\text{H} + \text{H}_2 + \text{CO}
\]

\[
\text{[Co}_2\text{(CO)}_8\text{]} \quad \text{OC}_2\text{H}_5
\]

\[
\text{OC}_2\text{H}_5
\]

\[
\text{CH}_3\text{CH}_2\text{CHO} \quad \text{OC}_2\text{H}_5
\]

\[
\text{OC}_2\text{H}_5
\]

\[
\text{CHO} \quad \text{C}_2\text{H}_5\text{O} \quad \text{OC}_2\text{H}_5
\]

\[
\text{CH}_2\text{OH}
\]

Equation 3.8
The same conditions were then applied to study other diene polymers like high-styrene-butadiene copolymers, ethyl acrylate-butadiene copolymer, methyl methacrylate-butadiene copolymer, isoprene-butadiene copolymer, natural rubber, balata. Glycol was added in these cases to prevent cross linking. All the above mentioned polymers, including polybutadiene gave polyaldehyde acetals with nearly quantitative double-bond conversion.

Hydroformylation and subsequent hydrogenation of polybutadiene at relatively high temperature and pressure in the presence of a mixed ligand cobalt catalyst is reported in a British patent\textsuperscript{93} (Equation 3.9). After hydroformylation the carbon monoxide pressure was decreased and hydrogenation was carried out under increased hydrogen pressure. No information is given on the composition of both the starting material and the final product.

\begin{equation}
\begin{align*}
PBD + CO + H_2 & \xrightarrow{[Co_2(CO)_{7}(PBu_3)_{2}]} PBD-Hyf \\
200^\circ C, 30-35 \text{ atm} \\

PBD-Hyf + H_2 & \xrightarrow{[Co_2(CO)_{7}(PBu_3)_{2}]} PBD-Hyox \\
120 \text{ atm}
\end{align*}
\end{equation}

where, PBD = polybutadiene

PBD-Hyf = hydroformylated polybutadiene

PBD-Hyox = hydroxylated polybutadiene

Equation 3.9

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Soon after this, quite a number of patents came out on hydroformylation of polybutadiene in the presence of cobalt complexes. The hydroformylated polybutadiene was then mixed with other materials to give commercially viable products. For example, the hydroformylated polybutadiene was stirred in the presence of an anionic emulsifier, i.e. a salt of higher fatty acids, to prepare latex films, blended with certain epoxidised polyolefin co-agents to produce baked coating materials or mixed with high vinyl content (1,2 units) polybutadiene and cured with SiO₂, vinyl triethoxy silane, dicumyl peroxide and calcium stearate to give polymers having improved resistance to thermoxidative degradation.

The first report on the use of rhodium complexes for hydroformylation of polydienes was by Sanui and co-workers (Equation 3.10). In the presence of \([\text{RhH(CO)}(\text{PPh₃})₃]\) hydroformylation of polypentenamer was carried out under mild conditions to yield 20 mol% of formyl side groups on the polypentenamer.

\[
\begin{align*}
\text{[\(\begin{array}{c}
\text{\(\cdot\text{CH}_2\text{-}\text{CH}\cdot\text{-}\text{CH}_2\text{-}\text{CH}\cdot\text{-}\text{CH}_2\text{-}\text{CH}\cdot\text{-}\text{-}\)}
\end{array}\}\text{]} + \text{CO} + \text{H}_2 & \xrightarrow{\text{RhH(CO)(PPh₃)₃}} \text{[\(\begin{array}{c}
\text{\(\cdot\text{CHO}\)}
\end{array}\}\text{]} }
\end{align*}
\]

Equation 3.10
Further reactions were carried out on the formyl group, but they are not relevant to our work. Azuma and coworkers have reported a two-step hydroxymethylation of polydiene via catalytic hydroformylation and subsequent hydrogenation in the presence of [RhH(CO)(PPh$_3$)$_3$] catalyst and reducing agents such as sodium borohydride.

Fairly recently, working at relatively lower temperature and pressure conditions, Rempel and co-workers have reported catalytic synthetic methods for the preparation of hydrogenated, hydroformylated and hydroxymethylated high vinyl content (90\% 1,2 units) polybutadiene in the presence of rhodium and ruthenium complexes. For example, they carried out hydrogenation of polybutadiene in the presence of Wilkinson’s catalyst [RhCl(PPh$_3$)$_3$] at 60°C and 2 atmospheres of hydrogen (Equation 3.11).

\[
\text{Equation 3.11}
\]
A ruthenium catalyst [RuClH(CO)(PPh$_3$)$_3$], at 40 atmospheres hydrogen and 120°C, was subsequently used to hydrogenate the above hydroformylated product. These reactions are also known as hydroxymethylation reactions (Equation 3.13).
However, quantitative results of the polybutadiene conversion and selectivities to the hydroformylated products are not reported. It is worth mentioning here that the above catalytic reactions were done without adding excess triphenylphosphine to the catalytic system.

Tremont, Remsen and Mills\textsuperscript{100} have improved upon the results reported by Rempel and coworkers. Hydroformylation of polybutadiene of varying composition using a hydridocarbonyltris(triphenylphosphine) rhodium(I) \([\text{RhH(CO)}(\text{PPh}_3)_3]\) catalyst in the presence of excess triphenylphosphine under mild conditions yielded polyaldehyde product at nearly 100\% selectivity. Thus hydroformylation of 1,4 \textit{cis} polybutadiene yielded 53.9\% internally branched aldehyde product (Equation 3.14).

\[
\begin{align*}
\text{[\includegraphics[width=0.4\textwidth]{polybutadiene.png}}] + \text{CO} + \text{H}_2 & \xrightarrow{\text{RhH(CO)(PPh}_3)_3, 75^\circ\text{C, 20 atm}} \text{[\includegraphics[width=0.4\textwidth]{polyaldehyde.png}}]
\end{align*}
\]

Equation 3.14

Hydroformylation of 1,2 syndiotactic polybutadiene under similar reaction conditions yields 22\% of the terminal branched aldehyde. A trace of \textit{iso}-branched aldehyde is also observed (Equation 3.15).
When a mixture of polybutadiene containing both 1,2-syndiotactic (12 wt%) and 1,4 cis and trans isomer (88 wt%) were hydroformylated under similar reaction conditions, the aldehyde products obtained correspond to those obtained for the pure constituents (Equation 3.16).

Comparison of kinetic data derived from gas uptake measurements showed that initial reaction rates for hydroformylation of 1,2
syndiotactic polybutadiene to be about six times greater than that obtained for hydroformylation of 1,4 cis-polybutadiene. This difference in rate can be attributed to the accessibility of the carbon-carbon double bond.

Similarly, Cole-Hamilton and coworkers\textsuperscript{101} working at a slightly higher pressure and in the presence of the same catalyst report both selective and non-selective hydroformylation of polybutadiene, containing a mixture of both internal and pendant carbon-carbon double bonds. For example, in the presence of only \([\text{RhH(CO)(PPh}_3)_3]\) hydroformylation occurs at both the internal and pendant carbon-carbon double bond position. A trace of hydroformylation occurring at the penultimate carbon atom of the pendant carbon-carbon double bond is also observed. However in the presence of excess triphenylphosphine the reaction becomes highly selective (selectivity > 90\%) towards the terminal carbon position of the pendant carbon-carbon double bonds (Equation 3.17).

\[
\begin{align*}
\text{[polybutadiene]} & \quad + \text{CO} + \text{H}_2 \\
& \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \quad \ Quad
3.1.4 Epoxidation

Epoxidation of butadiene by peracids to give the corresponding epoxide has been known for a long time. The commonly used epoxidising agent was peracetic acid and the product yield ranged from 45-100%.

Epoxidation of polymeric alkene hydrocarbons with a peracid, such as peracetic acid, was first reported in 1958. The polymeric hydrocarbons studied had molecular weights in the range of 150-250,000. The product contained 1-3% oxirane oxygen and was used for film forming materials. No additional information about the starting materials and their composition is given.

Epoxidation of polybutadiene containing a mixture of 1,4 cis, 1,4 trans, and 1,2 units, using peracids - hydrogen peroxide, monoperoxyphthalic acid and with tert -butyl hydroperoxide in the presence of a soluble molybdenum catalyst [MoO2(acac)2] have shown a preference for the backbone double bonds in the order cis -1,4 > trans -1,4 >> 1,2 units. The greater reactivity of the cis -1,4 units than trans -1,4 units is almost certainly due primarily to steric effects, although the energy difference between the cis and trans isomer may also be important. The pendant carbon-carbon double bonds start to epoxidise before all the backbone double bonds have reacted (i.e., some of the backbone double bonds are left unreacted). Therefore no high selectivity is observed.

The first highly regioselective epoxidation of polybutadiene using a soluble molybdenum complex was reported by Cole-Hamilton and co-workers. Polybutadiene containing a mixture of cis -1,4-, trans -1,4, and 1,2-units were subjected to epoxidation with tert butyl hydroperoxide in the presence of a molybdenum catalyst [MoO2Cl2[3-
Figure 3.1

The reaction shows very high selectivity to epoxidation of back bone double bonds whilst the pendant carbon-carbon double bonds are hardly reacted (Equation 3.18).

where, $m = 55\%$, $n = 45\%$

Equation 3.18
Since the system reported by Knifton\textsuperscript{30} in Chapter 1 (see page 19) afforded a very convenient method of regioselectively hydrocarboxylating alkenes with terminal carbon–carbon double bonds in the presence of palladium complexes, to give linear carboxylic esters, we were interested in applying the same methods for hydrocarboxylating polybutadienes (containing both a very high content of pendant carbon–carbon double bonds, and/or mixtures of internal and pendant carbon–carbon double bonds).

In theory, addition of carboxylic acid groups to polybutadiene can occur in three different ways (Equation 3.19). This would result in the formation of five types of hydrocarboxylated microstructures in the polymeric chain, as shown by species A–E.

\[
\begin{align*}
\left(\text{CH}_2\text{CH}=\text{CHCH}_2\right) + \text{CO} + \text{H}_2\text{O} & \rightarrow \left(\text{CH}_2\text{CHCH}_2\text{CH}_2\right)\text{COOH} \\
\text{1,4 units} & \\text{(A)}
\end{align*}
\]

\[
\begin{align*}
\left(\text{CH}_2\text{CH}\right)^{\text{CH}} + \text{CO} + \text{H}_2\text{O} & \rightarrow \left(\text{CH}_2\text{CH}\right)^{\text{CHCOOH}} + \left(\text{CH}_2\text{CH}\right)^{\text{CH}_3} \\
\text{1,2 units} & \\text{(B)} & \text{(C)}
\end{align*}
\]
Species A is formed when hydrocarboxylation occurs only on the 1,4 addition units, i.e., *cis* or *trans* only or a combination of both *cis* and *trans* structures. Products B and C are formed when the carboxylate group is introduced on the terminal or the internal carbon atom of the pendant carbon-carbon double bonds. Species B would thus be the result of an anti-Markownikov addition while species C would be a Markownikov addition product. Products D and E would be obtained when both the internal carbon-carbon double bonds and the pendant carbon-carbon double bonds are attacked by the -COOH acid groups.

One can expect that when using Knifton’s method, species A and B should be formed on hydrocarboxylation of polybutadiene.
3.2 Results and Discussion

There were three main purposes for this research. The first was to determine if Knifton's method of hydroesterification of 1-alkenes could be effectively applied for hydrocarboxylation of unsaturated polymers in general, and in particular polybutadiene. Secondly to see if the same method could be made regioselective in hydrocarboxylating polybutadienes. Finally, whether complete hydrocarboxylation of polybutadiene could be obtained by slightly altering the catalytic system \([\text{PdCl}_2(\text{PPh}_3)_2]\) (either by adding other cocatalysts or excess ligand, etc.)

For this purpose we treated polybutadienes with different microstructures, from 100% cis polybutadiene to polybutadienes containing large amounts of pendant carbon-carbon double bonds (1,2-units >80%) with the catalytic mixture \([\text{PdCl}_2(\text{PPh}_3)_2]-\text{SnCl}_2.2\text{H}_2\text{O}\) under high temperature and pressure. These polybutadienes were chosen so as to represent a wide range of olefinic structures. They were terminated with a variety of functional groups such as phenyl, hydroxyl, acid, etc. The polymers had some degree of saturation (see Table 3.1).

Much of Knifton's work was carried out at very high pressures (240 atmospheres of carbon monoxide). Technical limitations meant that we were limited to <120 atmospheres carbon monoxide pressure so initial studies were carried out at 80 atmospheres and 90°C. Initial studies were carried out on polymer no 7 (see Table 3.1). Hydrocarboxylation was observed and \(^{13}\text{C}\) n.m.r. studies indicated that the reaction was highly selective towards introduction of the carboxylic acid group on the terminal carbon atom of the pendant double bonds. A systematic survey of the reaction variables was then carried out in
<table>
<thead>
<tr>
<th>Polymer No</th>
<th>$M_n$</th>
<th>Cis double bonds (%)</th>
<th>Trans double bonds (%)</th>
<th>Pendant double bonds (%)</th>
<th>End group</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1000</td>
<td>12</td>
<td>19</td>
<td>53</td>
<td>Ph</td>
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<td>43</td>
<td>Ph</td>
</tr>
<tr>
<td>3</td>
<td>1480</td>
<td>-</td>
<td>16</td>
<td>81</td>
<td>COOH</td>
</tr>
<tr>
<td>4</td>
<td>1500</td>
<td>30</td>
<td>42</td>
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</tr>
<tr>
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<td>23</td>
<td>37</td>
<td>40</td>
<td>H</td>
</tr>
<tr>
<td>8</td>
<td>250000 $M_w$</td>
<td>100</td>
<td>-</td>
<td>-</td>
<td>H</td>
</tr>
<tr>
<td>9</td>
<td>-</td>
<td>20</td>
<td>60</td>
<td>20</td>
<td>Diol</td>
</tr>
<tr>
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<tr>
<td>11</td>
<td>3000</td>
<td>-</td>
<td>9</td>
<td>88</td>
<td>H*</td>
</tr>
</tbody>
</table>

* Narrow molecular weight distribution, $M_n$ = Number average molecular weight, $M_w$ = Molecular weight

% of different kinds of double bonds in starting polymer determined by $^1$H and $^{13}$C n.m.r. Where these do not sum to 100%, some butadiene units have been hydrogenated during the polymerisation process.

Table 3.1 Composition of polybutadienes investigated

118
order to obtain the optimum reaction conditions.

3.2.1 Effect of pressure

Pressures ranging from 70 -120 atmospheres carbon monoxide at 90°C for 6 hours were studied. At 70 atmospheres carbon monoxide pressure very little or no conversion took place. At 80 atmospheres complete conversion took place. Further increase in pressures did not cause any noticeable change in selectivity. The pressure was thus fixed at 80 atmospheres.

3.2.2 Effect of temperature

Temperatures ranging from 70°C-100°C at 80 atmospheres carbon monoxide for 6 hours were studied. At 70°C very little conversion took place. At 90°C complete conversion took place. The temperature was thus fixed at 90°C.

3.2.3 Reaction period

Since at 90°C and 80 atmospheres carbon monoxide pressure and 6 hours reaction period, complete conversion took place, reduced reaction periods were studied. It was observed that within 4 hours complete conversion took place. Further decrease in the reaction period decreased the yield. Further discussion of the progress of the reaction with time appears in the section '3.2.6. Catalyst stability'.
3.2.4 Amount of solvent

Initially experiments were carried out with 40 cm$^3$ of methyl isobutyl ketone. Experiments were then carried out using reduced amounts of solvent. It was found that the amount of solvent does not seem to affect the reaction greatly. The solvent was fixed at 10 cm$^3$, enough to dissolve the polybutadiene. The SnCl$_2$ dissolves in the aqueous phase.

3.2.4.1 Amount of polybutadiene

<table>
<thead>
<tr>
<th>Moles of polybutadiene</th>
<th>Overall conversion (%)</th>
<th>Conversion of pendant C=C bonds (%)</th>
<th>Catalyst turnovers</th>
</tr>
</thead>
<tbody>
<tr>
<td>3.697 x 10$^{-3}$</td>
<td>53.0</td>
<td>100</td>
<td>27.74</td>
</tr>
<tr>
<td>5.546 x 10$^{-3}$</td>
<td>41.2</td>
<td>77.39</td>
<td>32.35</td>
</tr>
<tr>
<td>7.395 x 10$^{-3}$</td>
<td>35.04</td>
<td>66.11</td>
<td>36.69</td>
</tr>
<tr>
<td>0.011</td>
<td>24.39</td>
<td>46.01</td>
<td>37.98</td>
</tr>
<tr>
<td>0.014</td>
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<td>40.36</td>
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<tr>
<td>0.018</td>
<td>18.34</td>
<td>34.60</td>
<td>46.75</td>
</tr>
</tbody>
</table>

Table 3.2

From Table 3.2 it is evident that increasing the moles of polybutadiene for the same amount of the catalyst decreases the percentage of conversion, although selectivity towards the terminal carbon atom of the pendant carbon-carbon double bond remains the same. This is due to the fact that there is not enough catalyst present in the reaction system to
react with the substrate. However, there is a small but probably significant increase in catalyst turnovers with increasing moles of substrate. This is in accord with the fact that at higher concentrations of the substrate the amount of unreacted substrate dispersed in the solvent is greater after 4 hours, thus increasing the chances for the interaction between the substrate and the catalyst (i.e., the rate will have fallen off more slowly for the more concentrated solutions).

Once the general conditions for specific hydrocarboxylation had been established, the method was applied to a variety of polybutadienes of varying composition under the optimum reaction conditions. The composition of the polybutadienes studied is shown in Table 3.1 and the analysis of the hydrocarboxylated polybutadiene is shown in Table 3.3.

3.2.5 Analysis of polybutadienes

Analysis of the hydrocarboxylated polybutadienes was carried out by various methods as follows. Polymer number 1 (see Table 3.3) is chosen as an example to show the various methods of analysis.

3.2.5.1 Infrared spectroscopy

The typical IR spectrum of polybutadiene and its corresponding hydrocarboxylated product is shown in parts (a) and (b) of Figure 3.2 respectively. The bands at 1640, 992, 969, 912, 740-732 cm\(^{-1}\) are due to carbon-carbon unsaturation. The IR spectra of the hydrocarboxylated polybutadiene lacks some of the characteristic peaks for unsaturation which indicates a degree of hydrocarboxylation. Besides the peaks for
<table>
<thead>
<tr>
<th>Starting polymer no</th>
<th>Cis double bonds (%)</th>
<th>Trans double bonds (%)</th>
<th>Pendant double bonds (%)</th>
<th>COOH (%)</th>
<th>Conversion of pendant C=C bonds (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>12</td>
<td>19</td>
<td>-</td>
<td>53</td>
<td>100</td>
</tr>
<tr>
<td>2</td>
<td>22</td>
<td>32</td>
<td>10</td>
<td>31</td>
<td>72</td>
</tr>
<tr>
<td>3</td>
<td>-</td>
<td>16</td>
<td>14</td>
<td>73</td>
<td>90</td>
</tr>
<tr>
<td>4</td>
<td>30</td>
<td>42</td>
<td>-</td>
<td>27</td>
<td>100</td>
</tr>
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<td>19</td>
<td>-</td>
<td>61</td>
<td>100</td>
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<td>77</td>
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<td>23</td>
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<td>40</td>
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<td>8</td>
<td>100</td>
<td>-</td>
<td>-</td>
<td>-</td>
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</tr>
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<td>20</td>
<td>60</td>
<td>12</td>
<td>8</td>
<td>40</td>
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<td>10</td>
<td>-</td>
<td>14</td>
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<td>84</td>
<td>100</td>
</tr>
<tr>
<td>11</td>
<td>-</td>
<td>9</td>
<td>9</td>
<td>78.3</td>
<td>89</td>
</tr>
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</table>

* Narrow molecular weight distribution,
  
  \([\text{Pbd}] = 3.7 \times 10^{-3} \text{ M} \); \([\text{Pd}]_{\text{total}} = 7.12 \times 10^{-5} \text{ M} \); \([\text{SnCl}_2] = 1.0 \times 10^{-3} \text{ M} \); Temperature = 90°C,
  
  CO pressure = 80 atm; \(\text{H}_2\text{O} = 0.5 \text{ cm}^3\); MIBK = 10 cm³

Table 3.3 Composition of product polymers obtained on hydrocarboxylation using \([\text{PdCl}_2(\text{PPh}_3)_2]-\text{SnCl}_2\)
Figure 3.2 (a) Infrared spectrum of polybutadiene
Figure 3.2 (b) Infrared spectrum of corresponding hydrocarboxylated (selectively) polybutadiene at the pendant carbon position.
unsaturation the characteristic band for the acid group C=O stretching at 1727 cm\(^{-1}\) is seen. The broad peak at 3310 cm\(^{-1}\) is characteristic of OH stretching.

3.2.5.2 \(^1\)H and \(^{13}\)C n.m.r. spectroscopy

3.2.5.2.1 \(^1\)H n.m.r.

The \(^1\)H n.m.r. spectra of polybutadiene and its hydrocarboxylated product is shown in Figure 3.3 (a) and (b). The conspicuous absence of a large peak at 5.0 ppm and the absence of peaks at 5.6-5.8 ppm shows almost complete conversion of the pendant carbon-carbon double bonds to hydrocarboxylated product.

3.2.5.2.2 \(^{13}\)C n.m.r.

The \(^{13}\)C n.m.r. spectra of both the starting material in CDCl\(_3\) and its hydrocarboxylated product in (CD\(_3\))\(_2\)CO is shown in Figure 3.4 (a) and (b) respectively. The absence of peaks at 115 ppm and 145 ppm (due to the carbon atoms of the pendant double bonds of polybutadiene) shows almost complete conversion of terminal carbon-carbon double bonds due to hydrocarboxylation. Consequently the presence of a new peak at 175 ppm due to the carboxylic acid group directly attached to the terminal carbon position is seen.

\(^{13}\)C n.m.r. spectroscopy of the crude product (see Figure 3.4 b) shows the presence of triphenyl phosphine oxide. This could be removed from the reaction mixture by precipitation of the polymer from the initial reaction solution with excess diethylether (see 3.3 Experimental section). \(^{13}\)C n.m.r. spectra of the hydrocarboxylated
Figure 3.3 (a) $^1$H n.m.r. spectrum of polybutadiene (phenyl terminated)
(b) $^1$H n.m.r. spectrum of corresponding hydrocarboxylated (selectively) polybutadiene at the terminal carbon position of the pendant group.
Figure 3.4 (a) $^{13}$C n.m.r. spectrum of polybutadiene (phenyl terminated)
(b) $^{13}$C n.m.r. spectrum of corresponding hydrocarboxylated (selectively) polybutadiene at the terminal carbon position of the pendant group.
Figure 3.5 (a) $^{13}$C n.m.r. spectrum of hydrocarboxylated product after reaction
(b) $^{13}$C n.m.r. spectrum of triphenylphosphine oxide
(c)$^{13}$C n.m.r. spectrum of hydrocarboxylated polybutadiene after treatment for removal of triphenylphosphine oxide.
product after reaction (Figure 3.5 a), triphenylphosphine oxide (Figure 3.5 b) and the hydrocarboxylated product (Figure 3.5 c) after treatment for removal of triphenylphosphine oxide are shown for comparison.

As can be seen from Table 3.3 no reaction took place when only cis polybutadiene (see polymer no 8) was subjected to hydrocarboxylation. Further proof for selective hydrocarboxylation comes from the observation that when polybutadienes containing a mixture of both, back bone carbon-carbon double bonds (cis or trans) and pendant carbon-carbon double bonds were subjected to hydrocarboxylation, reaction took place only at the terminal carbon atom of the pendant carbon-carbon double bonds resulting in the appearance of only one carboxylic acid peak at 175 ppm. It is evident therefore, that the catalytic system [PdCl2(PPh3)]-SnCl2 is highly regioselective to the introduction of carboxylic acid group on the terminal carbon atom of the pendant carbon-carbon double bonds of the polybutadienes under study (Equation 3.20).

![Chemical structure and reaction equation]

**Equation 3.20.**
3.2.5.3 Determination of weight average molecular weight ($M_w$) and number average molecular weight ($M_n$).

The hydrocarboxylated product of polymers number 1 and 5 (see Table 3.3) and their corresponding starting materials were chosen to study the change in weight average molecular weight ($M_w$) and number average molecular weight ($M_n$) upon hydrocarboxylation. Gel permeation chromatography (GPC) was carried out on our behalf at Ciba-Geigy. The samples were run in THF solvent using a polyethylene glycol standard. The results are shown in Table 3.4. and the GPC's of each polymer studied and their corresponding hydrocarboxylated products are shown in Figure 3.6 (a), (b) (polymer no. 1) and Figure 3.7 (a), (b) (polymer no.5) respectively.

<table>
<thead>
<tr>
<th>Polymer No</th>
<th>Cis db</th>
<th>Trans db</th>
<th>Pendant db</th>
<th>COOH</th>
<th>$M_w$</th>
<th>$M_n$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 SM</td>
<td>12</td>
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<td>53</td>
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<td>19</td>
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<td>49</td>
<td>8550</td>
<td>2170</td>
</tr>
</tbody>
</table>

Where SM = starting material, PR = product, db= double bonds

Table 3.4

The shapes of the gel permeation chromatographs for both the starting material and the hydrocarboxylated product is very similar. The molecular weight distribution also remains similar. There is perhaps a slight shifting of the curves for the hydrocarboxylated product to shorter elution times (higher molecular weight) and is
Figure 3.6 (a) GPC of polybutadiene (phenyl terminated, \( M_n = 1000 \))
(b) GPC of corresponding hydrocarboxylated polybutadiene
Figure 3.7 (a) GPC of polybutadiene (phenyl terminated, $M_n = 1800$)
(b) GPC of corresponding hydrocarboxylated polybutadiene
consistent with increase in molecular weights caused by the addition of HCOOH units in the formation of the carboxylic acid. Any further increase in molecular weight may be due to inter polymer hydrogen bonding as shown in Figure 3.8.

![Figure 3.8](image)

There is not any gross cross linking or chain scission. However the results obtained should not be taken too literally because of the fact that only polyethylene glycol was chosen as the standard and not polymers with closer structures to those of polycarboxylic acids.

3.2.5.4 Differential Scanning Calorimetry (DSC).

The effect of hydrocarboxylation of polybutadiene on the glass transition temperature (Tg) was studied by differential calorimetric
DSC studies on various polymers and their corresponding hydrocarboxylated products are shown in Table 3.5. The Tg of polymer number 10* is shown in Figure 3.9.

<table>
<thead>
<tr>
<th>Polymer number</th>
<th>Cis double bonds (%)</th>
<th>Trans double bonds (%)</th>
<th>Pendant double bonds (%)</th>
<th>Overall conversion (%)</th>
<th>Tg</th>
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<tbody>
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<td>19</td>
<td>53</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>PR</td>
<td>12</td>
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<td>-</td>
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<td>-</td>
</tr>
<tr>
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</tbody>
</table>

Where SM = starting polybutadiene, PR = Hydrocarboxylated polybutadiene, * = This represents a recatalysis reaction where all the carbon-carbon double bonds in polybutadiene have been carboxylated (see section 3.2.13 - Recatalysis).

Table 3.5

From the table it can be seen that Tg increases as the percentage of conversion of the carbon-carbon double bonds increases. The high Tg for the hydrocarboxylated product of polymer number 1 is due to a high percentage of saturated double bonds (16%) present in the starting
Figure 3.9 Tg of hydrocarboxylated polybutadiene (complete conversion of C=C, polymer no. 10)
Unlike the starting polymers, which are generally syrupy liquids with glass transition temperature (Tg) well below room temperature, the hydrocarboxylated product are solids which can be cast as films from solution. The products are soluble in polar solvents, e.g. acetone or methanol as well as in dilute NaOH, provided that the pH is kept ≥ 7.

3.2.6 Catalyst stability

The only problem with this catalytic system \([\text{PdCl}_2(\text{PPh}_3)_2]-\text{SnCl}_2\) is that the catalyst decomposed at the end of the reaction under these conditions. Time variation studies were carried out to find the onset of the reaction. From the $^{13}$C n.m.r. results obtained after various reaction periods it was evident that for the first 2 hours after introduction of the autoclave into the oven, there is very little conversion or no conversion at all. During this period there is no catalytic decomposition. After 2 hours reaction occurs rapidly and then levels off after 4 hours (see Figure 3.10), where by maximum conversion is obtained. After this time no further conversion was observed even after prolonged heating. As can be seen from Figure 3.10 it is during this short period when the reaction is occurring that the stability of the catalyst changes rapidly. The catalyst starts off with a slight decomposition and by the time the reaction is complete all the catalyst has decomposed. In other words there exists a small time window between the onset of the reaction and catalytic decomposition at which maximum conversion occurs.

Since recovery and reuse of catalyst to achieve maximum turn overs for a given amount of catalyst is of utmost importance in the
process of catalysis, the various factors affecting the stability of the catalyst such as effects of cocatalysts and its concentration, ligands, ligand concentration, percentage of water, solvent etc., were throughly
investigated. The reaction conditions; 80 atmospheres carbon monoxide pressure, 90°C and 4 hours were used throughout the reactions studied unless otherwise stated.

3.2.7 Effect of co-catalysts.

SnCl₂·2H₂O was found to have unique properties that make it the most effective co-catalyst for hydrocarboxylation. In the absence of SnCl₂·2H₂O no, or very little reaction takes place. That water in SnCl₂·2H₂O has no effect on the stability of the catalyst was shown by comparing the hydrocarboxylation with the dihydrate and anhydrous SnCl₂ as co-catalyst. The results were identical. For convenience, the dihydrate was therefore used throughout this work.

The results of varying the ratio of the cocatalyst to the catalyst was investigated (see Table 3.6).

<table>
<thead>
<tr>
<th>Mole ratio (C₆H₅)₃P : Pd : SnCl₂</th>
<th>Stability of catalyst</th>
<th>Conversion (%) of pendant C=C bonds</th>
</tr>
</thead>
<tbody>
<tr>
<td>2 : 1 : 1</td>
<td>C.D. observed</td>
<td>&lt;10%</td>
</tr>
<tr>
<td>2 : 1 : 5</td>
<td>C.D. observed</td>
<td>26%</td>
</tr>
<tr>
<td>2 : 1 : 10</td>
<td>C.D. observed</td>
<td>53%</td>
</tr>
<tr>
<td>2 : 1 : 30</td>
<td>C.D. observed</td>
<td>33%</td>
</tr>
</tbody>
</table>

Table 3.6

The results obtained revealed two things. Firstly, stability of the catalyst cannot be attained by changing the concentration of the cocatalyst and
secondly, conversion increasing with increasing tin-to-palladium ratio to a maximum ratio of 10 moles, then declining. Addition of excess SnCl₂ i.e., > 10 moles suppresses hydrocarboxylation reactions. This may be due to the fact that when a large excess of SnCl₂ is present, it may not only drive the above reaction towards the formation of a tin-rich complex, but may also block some of the sites on the palladium atom, which would otherwise be available for coordination with the alkene (polybutadiene).

3.2.8 Effect of adding excess triphenyl phosphine.

Although selectivity and conversion remained the same upon addition of excess triphenyl phosphine, stability of the catalyst did not show any improvement. Decomposition of catalyst was observed.

3.2.9 Effect of water

Water has a profound effect on catalyst stability. Several experiments were carried out and observations made to study the effect of water on catalyst stability.

Bittler and coworkers⁴⁷ observed that during hydroesterification of 1,5,9-cyclododecatriene in the presence of [PdCl₂(PPh₃)₂] at 75°C and 300 atmospheres carbon monoxide pressure, reduction of palladium (II) to metallic palladium became more pronounced when the "more water like" methanol is used instead of ethanol.

From our experiments we found that if water is replaced with an alcohol there was no decomposition of the catalyst but
hydroesterification also did not occur. It was observed that catalytic decomposition increases with increasing amounts of water, but the amount of conversion was also increased (see Table 3.7).

<table>
<thead>
<tr>
<th>Amount of water (cm³)</th>
<th>Stability of catalyst</th>
<th>Conversion (%) of the pendant C=C bonds</th>
</tr>
</thead>
<tbody>
<tr>
<td>No water</td>
<td>No C.D.</td>
<td>17.2</td>
</tr>
<tr>
<td>0.1</td>
<td>Slight C.D.</td>
<td>26.5</td>
</tr>
<tr>
<td>0.2</td>
<td>C.D. observed</td>
<td>30</td>
</tr>
<tr>
<td>0.3</td>
<td>C.D. observed</td>
<td>39</td>
</tr>
<tr>
<td>0.4</td>
<td>C.D. observed</td>
<td>42</td>
</tr>
<tr>
<td>0.5</td>
<td>C.D. observed</td>
<td>53</td>
</tr>
</tbody>
</table>

Table 3.7

We experimented on systems that would slowly release water into the reaction so that water can be used only as and when required. Several methods were tried to find an effective method of releasing water into the system. For example, molecular sieves (size 4A) were added along with water into the reaction vessel, with the aim that molecular sieves might absorb the water initially present in the reaction vessel and release it slowly as the reaction proceeded. In another experiment water was replaced with chloral hydrate to see if chloral hydrate would release the required amount of water during the reaction and hence avoid catalytic decomposition. Neither technique was successful.

To see if water kept in a separate compartment within the same reaction vessel might slowly distil into the reaction mixture and hence improve the catalyst stability, reactions were carried out with water
(0.5 cm³) being kept in a test tube inside the reaction vessel as shown in Figure 3.11.

![Diagram showing A: Glass liner, B: test tube]

**Figure 3.11**

<table>
<thead>
<tr>
<th>Height of test-tube</th>
<th>Stability of catalyst</th>
<th>Conversion (%) of pendant C=C bonds.</th>
</tr>
</thead>
<tbody>
<tr>
<td>8.5</td>
<td>No C.D.</td>
<td>18</td>
</tr>
<tr>
<td>7.5</td>
<td>No C.D.</td>
<td>22</td>
</tr>
<tr>
<td>2.5</td>
<td>No C.D*</td>
<td>18</td>
</tr>
</tbody>
</table>

*colour of the reaction solution changes from yellow to reddish brown

**Table 3.8**

After the reaction water was present in the test tubes in all cases. From Table 3.8 it is evident that the height of the test tube does not seem to matter. The conversion in all the cases resembles closely the results
obtained from the absence of water in the reaction (see Table 3.7., item 1). However when the height of the test tube was 2.5 cm the colour of the reaction solution changes from yellow to reddish brown. The reason for this observed result might be due to the fact that a very little amount of water enough to cause catalytic decomposition, but not enough to cause high conversion distilled into the reaction chamber during the reaction.

3.2.10 Effect of solvent

The solvent is an important factor governing the activity of a soluble catalyst in homogeneous reactions. The solvent may improve (a) the selectivity on varying the concentration of the reactants, the intermediates and final products in the reaction medium and (b) the mass transfer between the gas and liquid phases. The solvent can also act as a coolant or as a selective medium for the feed stock or product. Furthermore, certain solvents are not inert since they can react selectively with reaction intermediates or products or they can exert electronic effects even in the inner coordination sphere of the catalysts. Often it is important that there should be a single phase reaction i.e., solvent must dissolve both the reactants and the catalyst although in certain situations it is desirable to work in two phases.

Methyl isobutyl ketone was choosen as the solvent for the hydrocarboxylation reactions because of its convenience for dissolving the catalyst, co-catalyst, and polybutadiene. Since it forms an immiscible layer with water it has been very convenient for also treating polybutadiene to remove any peroxide contamination (see 3.4 Experimental section). Other solvents, such as tetrahydrofuran, acetone, methanol, etc., were studied, but it was found that methyl isobutyl
ketone gave the best results (see Table 3.9).

<table>
<thead>
<tr>
<th>Solvent</th>
<th>Catalyst stability</th>
<th>Conversion (%) of pendant C=C bonds</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methyl isobutyl ketone</td>
<td>C.D. observed</td>
<td>53</td>
</tr>
<tr>
<td>THF</td>
<td>C.D. observed</td>
<td>0</td>
</tr>
<tr>
<td>Acetone</td>
<td>C.D. observed</td>
<td>36</td>
</tr>
<tr>
<td>Toluene</td>
<td>C.D. observed</td>
<td>0</td>
</tr>
<tr>
<td>Carbon tetrachloride</td>
<td>C.D. observed</td>
<td>0</td>
</tr>
</tbody>
</table>

Table 3.9

3.2.11 Studies on the relationship between stability and activity of the catalyst

From the above results, it is clear that high conversions to polycarboxylic acids are only obtained in cases where there is substantial catalyst decomposition. Particularly striking is the time profile of the reaction. Thus, data from reactions carried out for different times are shown in Figure 3.10. There appears to be a long initiation period (ca 2.5 hours). After the onset of reaction the conversion increases steadily for the next 1.5 hours until its completion. Examination of the recovered solutions indicates that no decomposition occurs for 2.5 hours but that essentially total decomposition has occurred after 4 hours. There are various possible explanations for these phenomenon;
(i) The reaction may in fact be catalysed by palladium metal formed in the decomposition,

(ii) Since triphenylphosphine oxide is a major product of the decomposition, it may be that triphenylphosphine oxide in some way activates the catalyst towards reactivity, or

(iii) Some physical reaction parameters may be responsible for these unusual findings.
We have attempted to distinguish between these possibilities by a variety of experiments.

3.2.11.1 Tests on the homogeneity of the reaction

Attempts to use the black solid obtained at the end of a typical catalytic reaction as catalyst in the presence of added SnCl₂, indicate that this is not itself active, but this does not rule out the possibility that colloidal palladium formed from the catalyst decomposition, is the active species and that it coagulates slowly enough to allow complete conversion of the pendant carbon-carbon double bonds on polybutadiene to carboxylic acids. Were this the case, the highly selective nature of the reaction would be surprising, but it stills needs to be definitively discounted.

Crabtree¹⁰⁸ has reported certain tests which allow the distinction between homogeneous and heterogeneous reactions. The simplest of these involves carrying out the reaction in the presence of metallic mercury which, in general, poisons heterogeneous (metal catalysed) reactions by amalgamating the metal.

We have, therefore, carried out a reaction in the presence of a stirred pool of mercury and found that, although catalyst decomposition still occurs, high conversion to polycarboxylic acid is observed. This
would appear to indicate that the reaction is genuinely homogeneous, but may not be definitive in this case since palladium does not readily form an amalgam with mercury.

3.2.11.2 Effect of triphenylphosphine oxide

The $^{13}$C n.m.r. of hydrocarboxylated polybutadienes always showed the presence of triphenylphosphine oxide. It was therefore thought that possibly triphenylphosphine oxide, generated *in situ* during the reaction, in conjunction with the catalytic mixture $[\text{PdCl}_2(\text{PPh}_3)_2] / \text{SnCl}_2$, generates the actual catalytic species for hydrocarboxylation, (analogous complexes, such as $[\text{PtCl}_2(\text{OPPh}_3)_2]$ are known). In order to test this hypothesis excess triphenyl phosphine oxide was added and the reaction carried out. It is evident from the

<table>
<thead>
<tr>
<th>Moles of PPh$_3$O:Pd</th>
<th>Catalyst stability</th>
<th>Conversion (%) of pendant C=C bonds</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 : 1</td>
<td>C.D.observed</td>
<td>53</td>
</tr>
<tr>
<td>10 : 1</td>
<td>C.D.observed</td>
<td>19</td>
</tr>
</tbody>
</table>

Table 3.10

Table 3.10 that addition of excess triphenyl phosphine did not improve the activity of the catalyst; the stability was not improved.
3.2.11.3 Possible physical effects on catalyst stability

Commercially, where reactions can be carried out at high temperatures and pressures, the catalytic solution is usually heated, with stirring under pressure in the reaction vessel. In another compartment, the substrate is kept at the required temperature and pressure. Once the required conditions in the reaction vessel are stabilised, the substrate is pumped into the reaction chamber and the contents stirred throughout the reaction period. The temperature and the uptake of the reactant gas eg., carbon monoxide could be monitored throughout the reaction period.

However, with our technique we encountered several problems. Firstly, since we typically place a cold autoclave into a hot oven, it seemed possible that the reaction temperature might be rather lower than we had imagined and that the unusual reaction profile shown in Figure 3.10 might arise because the autoclave only reaches temperatures at which the reaction is rapid after ca. 2.5 - 3 hours. Further increase in the temperature might then lead to catalyst decomposition, with only a rather narrow temperature window occurring between the onset of the reaction and the decomposition of the catalyst.

Accordingly, we modified an autoclave by introduction of a steel pocket dipping into the reaction solution (Figure 3.12) so that we could monitor the temperature of the solution. We found that the temperature of the autoclave reaches only 85° C in 4 hours although the oven was kept at 90° C (see Figure 3.13). Using a fan assisted oven, the temperature rose more quickly but still took an hour to reach the preset temperature.
Key to Figure 3.12

1. Screw cap for connecting the autoclave, through metal tubing, to the carbon monoxide cylinder
2. Valve for pressurising and depressurising the autoclave
3. Autoclave head/top
4. Autoclave screw top. This comes along with the autoclave head part 3 and fastens the autoclave screw top to the body/base of the autoclave.
5. Rubber gasket which goes between the head and the body of the autoclave
6. Steel autoclave body/base
7. Glass liner
8. A thin hollow steel tube (closed at the bottom) used for inserting the temperature probe to measure the temperature of the reaction solution. The temperature probe is connected to a digital thermometer.

The autoclave used for carrying out experiments was made of stainless steel.

Secondly in all the experiments mentioned above the contents of the reaction vessel were not stirred. It would also be desirable to stir the contents of the reaction mixture. Due to technical difficulties however, stirring of the contents could not be done in an oven so experiments were carried out in an oil bath with stirring. When the autoclave was placed in a preheated oil bath, and the temperature monitored throughout the reaction it was observed that the autoclave acts initially, like a large heat sink. The temperature of the oil bath falls to almost half and then starts to rise.
3.2.12 Effect of Heater design

The large heat capacity of the autoclave means that it is very difficult rapidly to achieve the desired reaction temperature. Examination of Figure 3.10 and 3.11 suggest that the onset of reaction
occurs at ca. 70°C or below. Experiments were performed by gradually lowering the temperature by 5°C each time. It was observed that between 50-55°C the catalyst did not decompose. However, the yield was comparatively very small < 50%. With the temperature between 50-55°C and the pressure at 80 atmospheres carbon monoxide, various reaction time periods were studied i.e., 4, 24, 48, and 100 hours. It was observed that after a reaction period of 100 hours complete conversion of polybutadiene takes place without any catalyst decomposition.

We conclude that the reaction is truly homogeneous but that there is only a small temperature window between the onset of catalysis and the onset of catalyst decomposition. Experiments were also run in which the autoclave was placed in a gas chromatograph oven. In one case the gas chromatograph oven was switched on after the autoclave had been placed in it, whilst in the other the gas chromatograph oven was kept at a preheated temperature and the autoclave introduced. In both cases catalyst decomposition was observed although complete conversion of polybutadiene was obtained. By heating the autoclave quickly, it is possible to obtain high conversions in a short time but catalyst decomposition also occurs. By using a band heater directly attached to the autoclave (see Figure 3.16 b), we have now managed to obtain a heating method which allows the temperature of the solution to reach the required temperature within 10 minutes as can be seen in Figure 3.13.

3.2.13 Recatalysis: Complete hydrocarboxylation of all the carbon-carbon double bonds in polybutadiene.

Lapidus and coworkers\textsuperscript{36} reported that hydrocarboxylation of
polybutadiene in the presence of \([\text{PdCl}_2(\text{PPh}_3)_2]\) and excess triphenylphosphine occurred mainly at the pendant carbon-carbon double bonds, with a yield of \(\text{ca} < 60\%\). No information is given about the catalyst stability. We carried out identical experiments and observed from \(^{13}\text{C}\) n.m.r. that hydrocarboxylation takes place at three positions namely at the internal double bond position (177 ppm), and at the terminal (175 ppm) and penultimate carbon atom position (173 ppm) of the pendant carbon-carbon double bonds (Figure 3.14). As can be seen from \(^{13}\text{C}\) n.m.r. the selectivity is greater towards the terminal carbon atoms of the pendant carbon-carbon double bonds, followed by the backbone carbon atoms. The final composition of the hydrocarboxylated product obtained (on using Lapidus and co-workers method) is shown in Table 3.11.

<table>
<thead>
<tr>
<th>Polymer Number</th>
<th>Double bonds</th>
<th>Carboxylic acid (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cis</td>
<td>Trans</td>
</tr>
<tr>
<td>1  SM</td>
<td>12</td>
<td>19</td>
</tr>
<tr>
<td>PR</td>
<td>4</td>
<td>2</td>
</tr>
</tbody>
</table>

Where SM = starting material, PR = product, Bb = backbone, Tr = terminal and Int = internal

Table 3.11

In order to see if we could effectively hydrocarboxylate all the carbon-carbon double bonds in polybutadiene, the hydrocarboxylated product obtained by applying \([\text{PdCl}_2(\text{PPh}_3)]-\text{SnCl}_2\) (Knifton's method) was subjected to further hydrocarboxylation using the same palladium catalyst \([\text{PdCl}_2(\text{PPh}_3)_2]\), but with excess \(\text{PPh}_3\) (Lapidus and coworkers
Figure 3.14 $^{13}$C n.m.r spectrum of hydrocarboxylated polybutadiene
(based on Lapidus and coworkers\textsuperscript{36} method)
Figure 3.15 (a) $^1$H n.m.r. spectrum of completely hydrocarboxylated polybutadiene (polymer no. 1)

(b) $^{13}$C n.m.r. spectrum of the corresponding hydrocarboxylated polybutadiene
method) rather than SnCl₂. The analysis of the hydrocarboxylated product by ¹H n.m.r. and ¹³C n.m.r. analysis shows (Figure 3.15 (a) and (b)) almost complete hydrocarboxylation of the remaining carbon-carbon double bonds i.e., internal double bonds in polybutadiene. The pendant double bonds are selectively functionalised at the terminal carbon atom. These reactions are outlined in Scheme 3.1.

![Scheme 3.1](image)

(i) \([\text{PdCl}_2(\text{PPh}_3)_2]\)-SnCl₂, MIBK, 90°C, 80 atm, 4 hrs

(ii) \([\text{PdCl}_2(\text{PPh}_3)_2]\)-PPh₃, benzene, 170°C, 80 atm, 4hrs

(Back bone double bonds can be cis or trans; only trans double bonds are shown for sake of clarity).

**Scheme 3.1**

### 3.2.14 Scale up

After having had initial success with selective hydrocarboxylation of polybutadienes at the terminal carbon position of the pendant carbon-carbon double bonds, we were interested in investigating whether the
same results could be reproduced on a larger scale.

Polymer number 1 (see Table 3.1) was initially chosen for our studies. The reactants were scaled up 50 times while the physical parameters such as temperature and pressure were unchanged. The reaction was carried out in a rocking autoclave (see Figure 3.16 a) of one litre capacity. Analysis of the product clearly demonstrated that the reaction is reproducible and the results were consistent with the results obtained on a small scale.

Reactions were then carried out with polybutadienes of different composition. We have obtained results for all these hydrocarboxylation reactions consistent with those obtained when the hydrocarboxylation are carried out on a small scale in a 100 cm³ capacity autoclave (Figure 3.16 b).

3.3 Conclusion

From the experiments carried out on hydrocarboxylation of polybutadiene using [PdCl₂(PPh₃)₂]-SnCl₂ the following conclusions can be made

1. Regioselective hydrocarboxylation of polybutadiene can be effectively carried out with complete conversion and selectivity to the introduction of the carboxylic acid group on the terminal carbon atom of the pendant carbon-carbon double bonds, at a comparatively low temperature and pressure.

2. The hydrocarboxylation reaction catalysed by [PdCl₂(PPh₃)₂]-SnCl₂ is truly homogeneous and there exists a small temperature window between the onset of catalysis and the onset of catalyst decomposition.
Figure 3.16 (a) Rocking high pressure autoclave of 1 Litre capacity with inbuilt thermostat

Figure 3.16 (b) High pressure autoclave of 100 cm³ capacity along with band heater, glass liner and thermostat.
3. Complete hydrocarboxylation of all the carbon-carbon double bonds in polybutadiene can also be carried out by sequential catalytic reaction, but at a slightly higher temperature.

In principle, the selectively functionalised i.e., hydrocarboxylated polybutadiene could be prepared by copolymerisation of butadiene with pent-4-enoic acid but this does not appear to have been attempted. The relative scarcity of pent-4-enoic acid does not make this copolymerisation reaction attractive. So hydrocarboxylation is an attractive and simple route to a wide range of new polycarboxylic acids.

3.4 Experimental Section

Treatment of polybutadiene

Polybutadiene was freed of peroxide before reaction by treatment with a saturated solution of freshly prepared ferrous ammonium sulphate. In a typical reaction polybutadiene (0.2 g, 3.7 x 10^{-3} moles) was dissolved in 10 cm^3 of methyl isobutyl ketone and to this was added an equal amount of freshly prepared saturated solution of ferrous ammonium sulphate in sulphuric acid (2 mol dm^{-3}). The contents were shaken and separated in a separating funnel. The organic phase was used directly, unless otherwise stated.

Reaction procedure

a. Knifton's method

Dichlorobis(triphenylphosphine) palladium (II) [PdCl_2(PPh_3)_2]
(0.05 g, 7.12 x 10⁻⁵ moles) and tin (II) chloride dihydrate (0.19 g, 8.42 x 10⁻³ moles) were added to a nitrogen saturated mixture of methyl isobutyl ketone (10 cm³) containing H₂O (0.5 cm³) and polybutadiene (0.2g, 3.7 x 10⁻³ moles) in a 100 cm³ capacity specially designed glass liner. The glass liners / inserts were used to exclude any effects due to foreign ions from the walls of the autoclave. The glass liner was placed in a specially designed autoclave which was deoxygenated with nitrogen, flushed 3-4 times with carbon monoxide at 20-30 atmospheres and finally pressurised at room temperature to 80 atmospheres of carbon monoxide and sealed. It was then placed in an oven or oil bath, as required (see section - Results and Discussion) and maintained at 90°C for 4 hours.

b. Lapidus and coworkers method

Dichlorobis(triphenylphosphine) palladium (II) [PdCl₂(PPh₃)₂] (0.056 g, 7.97 x 10⁻⁵ moles) and triphenylphosphine (0.23 g, 8.76 x10⁻⁴ moles) were added to a nitrogen saturated mixture of benzene (12 cm³) containing H₂O (0.45 cm³) and polybutadiene (0.75g, 0.0138 moles) in a 100 cm³ capacity specially designed glass liner. The glass liners / inserts were used to exclude any effects due to foreign ions from the walls of the autoclave. The glass liner was placed in a specially designed autoclave which was deoxygenated with nitrogen, flushed 3-4 times with carbon monoxide at 20-30 atmospheres and finally pressurised at room temperature to 80 atmospheres of carbon monoxide and sealed. It was then placed in an oven and maintained at 170°C for 4 hours.
Product work up

In both cases i.e., after reaction a or b as mentioned above, the autoclave was cooled to room temperature either by allowing it to stand at room temperature or by placing it in a water bath circulated with running water. The pressure was released gradually in a fume cupboard and the contents recovered. The solvent was evaporated using a rotary evaporator and the residue dissolved in NaOH (20 cm$^3$/2 mol dm$^{-3}$). The polymer was then precipitated by adding an equal amount of HCl (7 mol dm$^{-3}$) in portions. The polymer was separated, dissolved in acetone (20 cm$^3$) and MgSO$_4$ was added to remove any water present in the acetone solution. The solution was then filtered and evaporated to dryness under vacuum. The yield obtained in both cases was >85%.

c. Recatalysis

Dichlorobis(triphenylphosphine) palladium (II) [PdCl$_2$(PPh$_3$)$_2$] (0.125 g, 1.78 x 10$^{-4}$ moles) and tin(II) chloride dihydrate (0.475 g, 2.10 x 10$^{-3}$ moles) were added to a nitrogen saturated mixture of methyl isobutyl ketone (25 cm$^3$) containing H$_2$O (1.25 cm$^3$) and polybutadiene (0.5g, 9.25 x 10$^{-3}$ moles) in a 100 cm$^3$ capacity specially designed glass liner. The glass liners / inserts were used to exclude any effects due to foreign ions from the walls of the autoclave. The glass liner was placed in a specially designed autoclave which was deoxygenated with nitrogen, flushed 3-4 times with carbon monoxide at 20-30 atmospheres and finally pressurised at room temperature to 80 atmospheres of carbon monoxide and sealed. It was then placed in an oven and maintained at 90°C for 4 hours.

After the reaction the autoclave was cooled to room temperature
either by allowing it to stand at room temperature or by placing it in a water bath circulated with running water. The pressure was released gradually in a fume cupboard and the contents recovered. The solvent was evaporated using a rotary evaporator and the residue dissolved in NaOH (50 cm³/2 mol dm⁻³). The polymer was then precipitated by adding an equal amount of HCl (7 mol dm⁻³) in portions. The polymer was separated, dissolved in acetone (30 cm³) and MgSO₄ was added to remove any water present in the acetone solution. The solution was then filtered and evaporated to dryness under vacuum. The yield obtained was >85%.

Dichlorobis(triphenylphosphine) palladium (II) [PdCl₂(PPh₃)₂] (0.028 g, 3.98 x 10⁻⁵ moles) and triphenylphosphine (0.115 g, 4.38 x 10⁻⁴ moles) were added to a nitrogen saturated mixture of benzene (6.0 cm³) containing H₂O (0.225 cm³) and hydrocarboxylated polybutadiene (0.375 g, 4.76 x 10⁻³ moles) in a 100 cm³ capacity specially designed glass liner. The glass liners / inserts were used to exclude any effects due to foreign ions from the walls of the autoclave. The glass liner was placed in a specially designed autoclave which was deoxygenated with nitrogen, flushed 3-4 times with carbon monoxide at 20-30 atmospheres and finally pressurised at room temperature to 80 atmospheres of carbon monoxide and sealed. It was then placed in an oven and maintained at 170°C for 4 hours.

After the reaction the autoclave was cooled to room temperature either by allowing it to stand at room temperature or by placing it in a water bath circulated with running water. The pressure was released gradually in a fume cupboard and the contents recovered. The solvent was evaporated using a rotary evaporator and the residue dissolved in NaOH (20 cm³/2 mol dm⁻³). The polymer was then precipitated by adding an equal amount of HCl (7 mol dm⁻³) in portions. The polymer
was separated, dissolved in acetone (20 cm\(^3\)) and MgSO\(_4\) was added to remove any water present in the acetone solution. The solution was then filtered and evaporated to dryness under vacuum. The yield obtained was >75%.

**Procedure for removing excess triphenyl phosphine oxide.**

Immediately after the reaction, the solvent methyl isobutyl ketone was reduced to less than half the initial volume by evaporation *in vacuo*. The solution was then poured slowly and gently into a flask containing excess diethylether (100 cm\(^3\)) with constant stirring. The hydrocarboxylated polymer precipitated instantly. After a few minutes of continued stirring, the whole solution was left aside to settle. The product was finally separated by decantation and dried under vacuum. After this, the product was dissolved in NaOH and precipitated with hydrochloric acid, as detailed earlier.
Chapter 4

Hydrocarboxylation of polybutadiene catalysed by PdCl₂, in the presence of air and CuCl₂.

4.1. Introduction

This chapter discusses the results obtained on hydrocarboxylation of polybutadiene catalysed by PdCl₂, in the presence of air and CuCl₂.

The Wacker process

It has been known for a long time that an aqueous solution of palladium chloride can oxidise ethene to acetaldehyde with concomitant deposition of palladium\(^{109}\). The hydrolysis of Zeise's salt \(\{\text{K}[(\text{C}_2\text{H}_4)\text{PtCl}_3]\}\), produces ethene, acetaldehyde, and ethanol accompanied by deposition of platinum. The accidental rediscovery of these reactions by Smidt and co-workers in 1959 in the laboratories of the Consortium für Electrochemische Industrie and development by the parent company, Wacker Chemie, has resulted in the Wacker process for the oxidation of ethene to acetaldehyde\(^{24}\). This discovery became an important landmark in the development of organo-transition metal chemistry. The commercial success of this process provided an enormous stimulus for further studies on homogeneous catalysis.

The so called 'Wacker process' involves the stoichiometric oxidation of ethene to acetaldehyde, by Pd(II) in aqueous solution. In the presence of CuCl₂, palladium(0) is reoxidised to palladium(II) and
the CuCl$_2$ is in turn regenerated by dissolved oxygen. The sequence of reactions proposed is shown in Scheme 4.1.

\[
\begin{align*}
\text{C}_2\text{H}_4 + \text{PdCl}_2 + \text{H}_2\text{O} & \rightarrow \text{CH}_3\text{CHO} + \text{Pd} + 2\text{HCl} \\
\text{Pd} + 2\text{CuCl}_2 & \rightarrow \text{PdCl}_2 + 2\text{CuCl} \\
2\text{CuCl} + 2\text{HCl} + \frac{1}{2}\text{O}_2 & \rightarrow 2\text{CuCl}_2 + \text{H}_2\text{O} \\
\text{C}_2\text{H}_4 + \frac{1}{2}\text{O}_2 & \rightarrow \text{CH}_3\text{CHO}
\end{align*}
\]

Scheme 4.1

Aqueous solutions of other group VIII metal salts, such as salts of Pt(II), Ir(III), Ru(III) and Rh(III) were also found to oxidise alkenes, but Pd salts were found to be the most effective catalysts for alkene oxidation$^{110,111,112,113}$. Alkenes other than ethene were oxidised to ketones. For example propene was oxidised under similar conditions to acetone in 90% yield and 1-butene afforded methyl ethyl ketone in approximately 85% yield$^{114,115,116}$.

Alkenes branched at the double bond did not form ketone and underwent allylic oxidation. Terminal alkenes of higher molecular weight generally afforded low yields of ketones under the standard Wacker conditions, since mixtures of isomeric ketones are often formed as a result of the palladium-catalysed isomerisation of the alkenes. The extent of isomerisation was markedly dependent on the reaction temperature. For example, 1-octene which generally produced a mixture of 2-,3- and 4-octanones, gave a 97% yield of 2-octanone at 30°C, 85% at 60°C, but only 82% at 90°C.
Effective oxidation of higher terminal alkenes has been realised using alcohols\textsuperscript{117,118}, dimethylformamide\textsuperscript{119} or sulfolane\textsuperscript{120} as co-solvents. Thus, Lloyd and Luberooff\textsuperscript{118} found that most monosubstituted and 1,2-disubstituted alkenes readily oxidised to the corresponding ketones in aqueous methanol or ethanol solutions. Clement and Selwitz\textsuperscript{119} reported that in aqueous dimethylformamide 1-dodecene was oxidised to 2-dodecanone in greater than 80% yield, and 10-undecanoic acid afforded 10-ketoundecanoic acid in 83% yield (Equation 4.1).

\begin{equation}
\text{CH}_2=\text{CH(CH}_2\text{)}_8\text{COOH} + \text{O}_2 \xrightarrow{\text{PdCl}_2/\text{CuCl}_2, \text{DMF-H}_2\text{O}} \text{CH}_3\text{C(CH}_2\text{)}_8\text{COOH}
\end{equation}

Equation 4.1

Oxidation of cyclohexene with PdCl\textsubscript{2} and CuCl\textsubscript{2} in aqueous ethanol afforded cyclohexanone in 95% yield\textsuperscript{117} (Equation 4.2).

\begin{equation}
\text{C}_6\text{H}_{12} + \text{O}_2 \xrightarrow{\text{PdCl}_2/\text{CuCl}_2} \text{C}_6\text{H}_{12}\text{O}
\end{equation}

Equation 4.2

A wide range of oxidants has been employed including benzoquinones, ferric salts, potassium dichromate, potassium persulphate, potassium bromide and lead dioxide. Electrolytic oxidation
has also been used\textsuperscript{121,122}. However the most common oxidant employed is a cupric salt, either using a quantitative amount, or a lesser amount together with a supply of oxygen to reoxidise the cuprous ion. Direct aerial oxidation of palladium(0) to palladium(II), required to regenerate the catalyst is too slow. An additional advantage of using copper salts is that copper(I) forms alkene complexes thereby increasing the amount of alkene available in solution. As these complexes are less stable than the palladium ones, they do not compete with the reaction intermediate (see Scheme 4.2).

Anions can also influence the reaction. It has been found that\textsuperscript{123} the rate of the reaction is inversely proportional to the square of the chloride ion concentration. The chloride ion suppresses complex formation between the alkene and palladium(II). Anions such as sulphate, nitrate, phosphate, perchlorate and fluoride which have very limited coordinating ability have little effect on the reaction rate.

The mechanism of the Pd(II)-catalysed oxidation of alkenes as proposed by Bäckvall, Akermark and Ljunggren\textsuperscript{124} is shown in Scheme 4.2.

\[
PdCl_4^{2-} + C_2H_4 \overset{\text{PdCl}_3 \cdot C_2H_4]}{\longrightarrow} [PdCl_3 \cdot C_2H_4]^- + Cl^- \quad (1)
\]

\[
[PdCl_3 \cdot C_2H_4]^- + H_2O \overset{\text{[PdCl_2(H_2O) \cdot C_2H_4] + Cl^-}}{\longrightarrow} [PdCl_2(H_2O) \cdot C_2H_4] + Cl^- \quad (2)
\]

\[
[PdCl_2(H_2O) \cdot C_2H_4] + H_2O \overset{[HOCH_2CH_2PdCl_2(H_2O)]^- + H^+}{\longrightarrow} [HOCH_2CH_2PdCl_2(H_2O)]^- \quad (3)
\]

\[
[HOCH_2CH_2PdCl_2(H_2O)]^- \overset{HOCH_2CH_2PdCl(H_2O) + Cl^-}{\longrightarrow} \quad (4)
\]

\[
HOCH_2CH_2PdCl(H_2O) \overset{CH_3CHO + Pd + HCl + H_2O}{\longrightarrow} \quad (5)
\]

Scheme 4.2
The first step involves the formation of a π complex (eq 1), which reduces the electron density at the double bond rendering it susceptible to nucleophilic attack. In the second step (eq 2) aquation of the complex takes place in a reversible reaction by external attack of water, together with loss of chloride. Formation of the new C-O bond occurs via nucleophilic attack of external water on the coordinated alkene to give the anionic hydroxyethyl complex (eq 3). The dissociation of the chloride ligand in equation 4 would give a 14 electron complex which would be prone to undergo rapid β-elimination (eq 5) to give first vinyl alcohol which would rapidly tautomerise to acetaldehyde. The palladium hydride so formed is unstable and rapidly deposits palladium metal. The key features of this mechanism are the nucleophilic attack on the coordinated alkene in equation 3 and the β-hydride elimination of the hydroxyethylpalladium (II) intermediate in equation 5.

Alper, Woell, Despeyroux and Smith\(^{25}\) successfully modified the Wacker process by bubbling carbon monoxide (1atm) through the reaction solution along with oxygen (Equation 4.3). This resulted in hydrocarboxylating straight chain alkenes, both terminal and internal, under exceedingly mild conditions. For terminal alkenes, the product are mainly α-methyl acids indicating stereospecific addition of the carboxyl group to the internal carbon atom of the double bond.

\[
\text{RCH=CH}_2 + \text{CO} + \text{H}_2\text{O} \xrightarrow{\text{PdCl}_2, \text{CuCl}_2, \text{HCl}, \text{O}_2, \text{rtm, 1atm}} \text{RCH(COOH)CH}_3
\]

Equation 4.3

Di-alkenes such as octa-1,7-diene and deca-1,9-diene exclusively formed the branched product.\( cis \) or \( trans \)-dec-2-ene gave only 2-methyl-
decanoic acid. Similarly, only 2,4-dimethylpentanoic acid was isolated using *cis*-4-methylpent-2-ene as the substrate. No isomeric acids were obtained in these reactions.

Concurrently in their patent, Alper and Smith\textsuperscript{26} had extended the above reaction to hydrocarboxylation of polymeric compounds containing residual carbon-carbon double bonds as starting materials. However there appears to be an anomaly in the patent as the authors have quoted polyisobutene as an example for the starting material. It may be noted that polymers of isobutene have two methyl groups on alternate carbon atoms along the chain and two hydrogen atoms on the other chain carbon atoms (Equation 4.4).

\[
\begin{align*}
\text{CH}_3 & \quad \text{CH}_3 \\
\text{CH}_3 & \quad \text{CH}_3 \\
\text{nH}_2\text{C} & \equiv \text{C} \\
\end{align*}
\]

\[\rightarrow \quad \left[ \begin{array}{c}
\text{CH}_3 \\
\text{-CH}_2 \\
\text{CH}_3 \\
\text{CH}_3 \\
\end{array} \right]_n\]

Equation 4.4

In other words polyisobutene is a saturated polymer with no double bonds available for hydrocarboxylation apart, perhaps from end groups of the chain. No information was given for the product yield or conversion.

Since the method by Alper and coworkers afforded a convenient method of hydrocarboxylating alkenes with internal double bonds, to give exclusively branched chain carboxylic acids, we were interested in using the same method for hydrocarboxylating polybutadiene (containing both internal double bonds only and/or mixtures of internal
and pendant carbon-carbon double bonds.

In theory application of Alper's method of hydrocarboxylation would result in the formation of products A, C, and E, as shown in Equation 3.19, Chapter 3, page 115.

4.2. Results and Discussion

The composition of the polymers employed in this study are shown in Table 4.1. The analysis of the product polymer is shown in Table 4.2. Polymer number 1 (see Table 4.2) was chosen to show the various methods of analysis.

4.2.1 Methods of analysis

4.2.1.1 I.R. analysis

The typical IR spectra of polybutadiene (polymer no. 1, as shown in Table 4.1) and its corresponding hydrocarboxylated product is shown in parts (a) and (b) of Figure 4.1 respectively. The bands at 1640, 992, 969, 912 and 740-732 cm\(^{-1}\) are due to carbon-carbon double bond unsaturation. IR spectra of the hydrocarboxylated polybutadiene lack some of the characteristic peaks for unsaturation which indicates a degree of hydrocarboxylation. Beside the peaks for unsaturation the characteristic peak for the acid group i.e., C=O stretching at 1727 cm\(^{-1}\) is seen. The broad peak at 3310 cm\(^{-1}\) is characteristic of the OH stretching vibration.
<table>
<thead>
<tr>
<th>Polymer number</th>
<th>$M_n$</th>
<th>Cis double bonds</th>
<th>Trans double bonds</th>
<th>Pendant double bonds</th>
<th>End group</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1000</td>
<td>12</td>
<td>19</td>
<td>53</td>
<td>Ph</td>
</tr>
<tr>
<td>2</td>
<td>1300</td>
<td>22</td>
<td>32</td>
<td>43</td>
<td>Ph</td>
</tr>
<tr>
<td>3</td>
<td>1480</td>
<td>-</td>
<td>16</td>
<td>81</td>
<td>COOH</td>
</tr>
<tr>
<td>4</td>
<td>1500</td>
<td>30</td>
<td>42</td>
<td>27</td>
<td>Ph</td>
</tr>
<tr>
<td>5</td>
<td>3400</td>
<td>22</td>
<td>50</td>
<td>26</td>
<td>Ph</td>
</tr>
<tr>
<td>6</td>
<td>4500</td>
<td>23</td>
<td>37</td>
<td>40</td>
<td>H</td>
</tr>
<tr>
<td>7</td>
<td>$250000M_w$</td>
<td>100</td>
<td>-</td>
<td>-</td>
<td>H</td>
</tr>
</tbody>
</table>

Where $M_n$ = Number average molecular weight, $M_w$ = weight average molecular weight.

% of different kinds of double bonds in starting polymer determined by $^1H$ and $^{13}C$ n.m.r. When these do not sum to 100%, some butadiene units have been hydrogenated during the polymerisation process.

Table 4.1 Composition of polybutadienes investigated
<table>
<thead>
<tr>
<th>Starting polymer number</th>
<th>Remaining Double bonds (%)</th>
<th>Carboxylic acids (%)</th>
<th>Overall Conversion of C=C bonds (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cis</td>
<td>Trans</td>
<td>Pendant</td>
</tr>
<tr>
<td>1</td>
<td>6</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>18</td>
<td>18</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>4</td>
<td>47</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>33</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>13</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>10</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>18</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* signal intensities from branched carboxylic acid is too small to measure

\[ [\text{Pbd} = 3.7 \times 10^{-3} \text{ M}; \text{PdCl}_2 = 7.8 \times 10^{-4} \text{ M}; \text{CuCl}_2 = 1.55 \times 10^{-3} \text{ M}; \text{H}_2\text{O} = 1\text{ cm}^3; \text{HCl} = 1\text{ cm}^3; 1\text{ atm CO, 1 atm O}_2] \]

Table 4.2 Composition of the product polymers on hydrocarboxylation using PdCl₂/CuCl₂/O₂
Figure 4.1 (a) I.R. spectrum of polybutadiene

(b) I.R. spectrum of corresponding hydrocarboxylated polybutadiene
4.2.1.2 $^1$H and $^{13}$C n.m.r analysis

4.2.1.2.1 $^1$H n.m.r.

The $^1$H n.m.r. spectrum of the hydrocarboxylated polybutadiene is shown in Figure 4.2. (for $^1$H n.m.r of the starting polybutadiene see chapter 3, page 125). The absence of peak at 5.7 ppm and the decrease in the height of the peak at 5.0 ppm indicates degree of hydrocarboxylation. Further evidence for this conversion comes from $^{13}$C n.m.r. spectrum.

4.2.1.2.2 $^{13}$C n.m.r.

The $^{13}$C n.m.r. spectrum of the hydrocarboxylated product is shown in Figure 4.3 (for $^{13}$C n.m.r of the starting polybutadiene see chapter 3, page 126). It can be seen that the peak intensity at 115 ppm and at 145 ppm are considerably reduced relative to the intensity of peaks at 130 ppm (arising due to back bone carbon-carbon double bonds). The presence of three new peaks at (δ173 ppm, δ175 ppm, δ177 ppm), characteristic of carboxylic acid is seen. From our studies using only cis polybutadiene (see Table 4.2, polymer number 7) assignment of the three peaks have been made as follows.

The peak at 177 ppm is due to hydrocarboxylation taking place at the internal double bond position i.e., either cis or trans. Hydrocarboxylation of pendant double bonds on the terminal carbon position gives rise to an acid peak at 175 ppm. The peak at 173 ppm thus results from hydrocarboxylation at the penultimate carbon atom of the pendant position.
Figure 4.2 $^1$H n.m.r. spectrum of hydrocarboxylated polybutadiene
Figure 4.3 $^{13}$C n.m.r. spectrum of hydrocarboxylated polybutadiene
4.2.2 Discussion

The results obtained i.e., COOH insertion at the *cis* or *trans* terminal and penultimate carbon-carbon double bond (pendant) position show that the catalytic system PdCl$_2$/CuCl$_2$/O$_2$ is more reactive towards the pendant carbon-carbon double bonds (Equation 4.5. Backbone double bonds can be *cis* or *trans*; only *trans* are shown for reasons of clarity).

\[
\text{[\begin{array}{c}
\text{[ \text{PdCl}_2,\text{CuCl}_2,\text{HCl} ]}
\end{array}]} + \text{CO} + \text{H}_2\text{O} \xrightarrow{\text{PdCl}_2,\text{CuCl}_2,\text{HCl}} \text{O}_2, \text{rtm, 1 atm}
\]

Equation 4.5

In Alpers work on simple alkenes, terminal alkenes have been shown to produce predominantly branched carboxylic acids. For polybutadienes, this is not the case and the reaction of the pendant double bonds show substantial, although not total, selectivity towards terminal carboxylic acid formation. This difference in selectivity presumably arises from steric problems associated with the presence of the main chain on the carbon atom α- to the pendant double bonds. A related effect can be noted$^{25}$ in the exclusive formation of 2,4-dimethylpentanoic acid on hydrocarboxylation of 4-methylpent-2-ene using the same system (Equation 4.6).
Here branching of the chain at the carbon α- to the double bond apparently directs the incoming carboxylate group to the end of the double bond remote from the branching point.

For the polybutadiene and reaction conditions considered in this study, the PdCl₂ catalyst adds the carboxylic acid functionality over the entire range of number average molecular weight. To find out if the reaction can be carried out regioselectively i.e., if the reaction can be regiospecific in hydrocarboxylating polybutadiene, other chloride complexes like SnCl₂, GeCl₂ were added to the existing PdCl₂/CuCl₂ system, but no reaction took place.

Increasing the temperature of the reaction caused loss of solvent tetrahydrofuran. This problem was overcome by attaching a water condenser to the reaction vessel and by passing CO and O₂ saturated with tetrahydrofuran before bubbling them through the reaction solution. On increasing the temperature to ~50°C, hydrocarboxylation takes place at the 3 carbon-carbon double bond position in polybutadiene i.e., at the back bone carbon-carbon double bond position, at the penultimate carbon-carbon double bond and at the terminal carbon-carbon double bond of the pendant carbon-carbon double bond position. Here selectivity appears to be greater towards the pendant carbon atoms relative to the back bone carbon atoms. The overall conversion is very high compared to a normal 24 hour reaction (see Table 4.4). ¹³C n.m.r. spectra (see Figure 4.4) of the hydrocarboxylated product shows a relatively higher conversion taking
place at the penultimate carbon position of the carbon-carbon double bonds in comparison with the $^{13}$C n.m.r. of a normal 24 hour reaction (Figure 4.3). However, a black coating of palladium metal coating occurred at the tip of the bubbling tubes with an hour after starting the reaction.

Varying the reaction time resulted in selectivity and varying degrees of hydrocarboxylation. For example, from $^{13}$C n.m.r. spectrum (see Figure 4.5) of a 12 hour reaction only one carboxylic acid peak corresponding to the terminal carbon position of the pendant carbon-carbon double bond can be seen, whilst for a 100 hour reaction time, reaction took place at two positions i.e., at the terminal carbon position (175 ppm) of the pendant carbon-carbon double bond and at the internal carbon position (177 ppm) of the back bone carbon-carbon double bond (see Figure 4.6). The corresponding conversions are shown in Table 4.4.

<table>
<thead>
<tr>
<th>Polymer number</th>
<th>Conditions</th>
<th>bb dbs (%)</th>
<th>bb COOH (%)</th>
<th>pendant dbs (%)</th>
<th>pendant COOH (%)</th>
<th>Overall conversion (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>12 hrs reaction time</td>
<td>10</td>
<td>21</td>
<td>14</td>
<td>39</td>
<td>71</td>
</tr>
<tr>
<td>1</td>
<td>24 hrs reaction time</td>
<td>6</td>
<td>25</td>
<td>5</td>
<td>48</td>
<td>87</td>
</tr>
<tr>
<td>1</td>
<td>100 hrs reaction time</td>
<td>3</td>
<td>28</td>
<td>5</td>
<td>48</td>
<td>90</td>
</tr>
<tr>
<td>1</td>
<td>Increase temperature (24 hours)</td>
<td>3</td>
<td>28</td>
<td>3</td>
<td>50</td>
<td>93</td>
</tr>
</tbody>
</table>

Table 4.4
Figure 4.4 $^{13}$C n.m.r. spectrum of hydrocarboxylated polybutadiene
(obtained by increasing reaction temperature)
Figure 4.5  $^{13}$C n.m.r. spectrum of hydrocarboxylated polybutadiene (12 hours reaction time)

Figure 4.6  $^{13}$C n.m.r. spectrum of hydrocarboxylated polybutadiene (100 hours reaction time)
As one would expect increasing the reaction time increases conversion. From the table it is evident that greater conversion can be achieved in a relatively shorter time by increasing the temperature i.e., increasing reaction rate.

The proportion of water used had a significant influence on the reaction. No reaction took place when water was used as the only solvent, as polybutadiene does not dissolve in water. Also when the amount of water was greater than 5 moles equivalent of the solvent tetrahydrofuran, the reaction failed to take place. No reaction took place in the absence of either oxygen or carbon monoxide. Also variation in the rate of bubbling carbon monoxide and/or oxygen through the solution had no observable effect on hydrocarboxylation.

One main drawback of this method is that the catalyst PdCl$_2$ could not be recovered for further catalysis. It was observed that if the reaction was left for a few days without bubbling oxygen or carbon monoxide, precipitation of palladium metal occurred as a thin coating on the sides of the reaction vessel. The recovered metal showed very little or no catalytic activity.

Application of Alper's method of hydrocarboxylation of alkenes to polymers gave reduced conversion. For example, according to Alper and coworkers hydrocarboxylation of alkenes such as deca-1,9-diene gave 100% conversion, but in the case of polybutadiene complete conversion could not be obtained (< 70%) on using normal reaction temperature and pressure conditions. This could account for the low reactivity of polybutadiene; due to the size and thus the steric effect of the bulky polymer chain.
4.3 Conclusion

From our experimental work we have made two main observations.
1. With *cis* polybutadiene, the hydrocarboxylation reaction does takes place and the conversion is > 60% i.e., species A is obtained according to Equation 3.19, Chapter 3, page 115.
2. When the polymer contains mixture of both internal and terminal double bonds, hydrocarboxylation takes place at three positions namely at the internal double bond position, terminal and at the penultimate carbon position of the pendant group. i.e., species D and E are obtained.

4.4 Experimental Section

Polybutadiene, obtained from Aldrich was used as received. Carbon monoxide was bubbled through a solution containing tetrahydrofuran (30 cm$^3$) and water (1 cm$^3$). PdCl$_2$ (0.14 g, 7.8 x 10$^{-4}$ moles) was added, followed by concentrated hydrochloric acid (1.0 cm$^3$) and the mixture was then stirred until PdCl$_2$ dissolved. CuCl$_2$.2H$_2$O (0.26 g, 1.55 x 10$^{-3}$ moles) was then added and O$_2$ bubbled through the solution. Polybutadiene (0.2 g, 3.7 x 10$^{-3}$ moles) dissolved in tetrahydrofuran (10 cm$^3$) was added and the reaction was stirred at room temperature for 16 hours or left to stir overnight. The gases were bubbled through the reaction mixture via gas inlet adaptors below the surface of the solvent as shown in Figure 4.1. The flow of
carbon monoxide and oxygen was maintained throughout the reaction period.

The solvent was evaporated in a rotary evaporator and the residue dissolved in NaOH (20 cm$^3$ / 2 mol dm$^{-3}$). The polymer was then precipitated by adding an equal amount of HCl (7 mol dm$^{-3}$) in portions, separated and dissolved in acetone (20 cm$^3$). MgSO$_4$ was added, to remove any water present in the acetone solution. The acetone solution was filtered and finally evaporated to dryness. The yield was < 60 %. 
Chapter 5

Future Work

5.1. Introduction

This chapter briefly discusses the current work being carried out using hydrocarboxylated polybutadienes. Based on the success of selectively hydrocarboxylating polybutadienes two patents have been filed, and currently, studies are being carried out to see if these hydrocarboxylated polymers could be used for

1. treatment of wood and,
2. corrosion inhibition.

The results obtained so far have been very encouraging. It is not intended to include a detailed discussion as studies are currently in progress and hence the description of the methods are only intended to be thumbnail sketches.

5.1.1 Wood treatment by polycarboxylic acids

The ready availability and unique physical properties of wood have made it the material of choice for mankind for thousands of years. Although wood is amongst the most resistant of all organic materials, as time passes, various chemical, morphological and physical changes take place resulting in its degradation. Wood exposed to natural weathering processes undergoes discolouration, and degradation by light and moisture, etc. Consequently wood has to be protected by paints, stains, and similar materials. Since the late 1960's and early 1970's
considerable research effort has gone into finding polymeric materials that would not only protect the surface of wood but would also penetrate the wood and strengthen it from inside by undergoing changes like cross linking, etc. Materials that penetrate deep inside the wood surface in a relatively short time and are also cheap are required and to this end polymers, particularly water soluble polymers are currently being investigated.

Our aim was to see if hydrocarboxylated polybutadiene which shows water solubility would fulfil the above requirements. Studies are being carried out at our laboratory. The initial studies and the results obtained are as follows.

As hydrocarboxylated polybutadiene in its acid form is sparingly soluble in water, it must be converted into the sodium salt to increase its solubility by the following method. Polybutadiene 84% unsaturation (53% pendant, 12% cis-1,4, 19% trans-1,4) was hydrocarboxylated so that all of the pendant carbon-carbon double bonds were converted into terminal carboxylic acid groups. A sample of this product polymer (0.5 g) was suspended in water (10 cm$^3$) and NaOH (2 mol dm$^{-3}$) was added dropwise until all the polymer dissolved, with the pH kept at 7.0.

Three pieces of dry balsa wood (30 cm x 0.3 cm x 0.3 cm) were soaked in water for 43 hours, during which time the water was shown by $^1$H n.m.r. imaging$^{125}$ to have penetrated (see Figure 5.1) ca 0.2 mm all round. In Figure 5.1 the white areas indicate the mobile protons (from water). The three pieces of wood were further allowed to stand in water for 14 days. After 14 days the wood pieces became water logged and sank. $^1$H n.m.r. imaging of such a piece of wood showed (see Figure 5.2) that water had penetrated uniformly into the wood.
Figure 5.1 $^1$H n.m.r transverse image of dry balsa wood after floating in water for 43 hours

Figure 5.2 $^1$H n.m.r transverse image of dry balsa wood after floating in water for 16 days

Figure 5.3 $^1$H n.m.r transverse image of dry balsa wood after floating in water for 52 hours followed by polymer solution for 14 days
Three pieces of balsa wood, which had been standing in water for 52 hours were transferred to the polymeric solution described above. After 14 days, the wood was still afloat. $^1$H n.m.r. imaging (see Figure 5.3) indicated that the polymer has prevented any further ingress of water into the wood. After 5 weeks, n.m.r. imaging again showed no further water had penetrated the wood. The pieces of wood are still floating at this point and were still floating after 18 months, suggesting that water had still not penetrated.

5.1.2. Corrosion inhibition

When pipes made from metals, such as iron or steel, are in contact with water for a prolonged period, chemical reactions take place between the ions present in water and the metallic surface resulting in corrosion. In industrial plants such as desalination plants, water cooling systems, steam-generating plants, bottle washing plants, closed circuit heating systems, waste treatment plants etc., the problem of corrosion is acute resulting in increased maintenance and/or replacement of the corroded pipes with costs running to billions of pounds. One way to circumvent this problem is to prevent the deposition of inorganic salts on metal surfaces and to reduce metallic corrosion. Water soluble polymers can offer an easy solution to this problem as they can be mixed with the water entering the system.

Our aim was to see if hydrocarboxylated polybutadiene would meet the above requirement. Tests are currently being carried out at Ciba-Geigy, and one of the tests, known as a rotating coupon test is described below. It is conducted using corrosive water of composition:
The test is conducted as follows:

In a 1 litre reservoir of the test water, two precleaned and preweighed mild steel coupons are rotated. The test is conducted over 48 hours in oxygenated water at 40°C using 20 ppm of the appropriate concentration of corrosion inhibitor.

The coupons are removed, scrubbed, immersed for one minute in hydrochloric acid inhibited by 1% by weight of hexamine and then rinsed, dried and weighed. A certain loss in weight will have occurred. A blank test i.e. immersion of mild steel specimen in the test water in the absence of any potential corrosion inhibitor, is carried out with each series of tests. The corrosion rate is calculated in milligrams of weight loss/square decimeter/day (m.d.d.). The results obtained in a series of tests are set out in the following Table 5.1.

<table>
<thead>
<tr>
<th>Example</th>
<th>Corrosion inhibitor</th>
<th>Corrosion rate (m.d.d.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Blank</td>
<td>197</td>
</tr>
<tr>
<td>2</td>
<td>polymer 5*</td>
<td>9.6</td>
</tr>
<tr>
<td>3</td>
<td>polymer 7*</td>
<td>32.9</td>
</tr>
</tbody>
</table>

* For composition of the hydrocarboxylated polymers see Table 3.3, Chapter 3, page 122.

Table 5.1
Traditionally zinc chromates have been used as corrosion inhibitors. As they are extremely toxic and harmful if released into the environment they have been replaced by less toxic corrosion inhibitors such as, polyphosphates and phosphoric acids. But these compounds do contain phosphorous in them which are harmful to aquatic life as they are the main cause of eutrophication in rivers, lakes, etc. Currently attention is being focussed on corrosion inhibitors, scale control reagents etc., that are not only environmentally friendly but also readily biodegradable.

The hydrocarboxylated polybutadiene synthesised by us and currently being tested for corrosion inhibition is amongst the most effective of compounds containing only C, H, and O, in them and also has the advantage of being readily biodegradable.

5.1.3. Hydrocarboxylated polybutadienes with other functional groups

Introduction of other functional group on to the hydrocarboxylated product might lead to unusual and interesting physical and chemical properties. For example, polybutadiene which has been selectively hydrocarboxylated at the pendant carbon-carbon double bond position can be epoxidised at the backbone carbon-carbon double bonds. The epoxidised product thus obtained might offer firstly, the advantage of being water soluble and secondly offer better cross linking than those of hydrocarboxylated product. The epoxidised hydrocarboxylated product might also interesting properties upon ring opening for example, form diols, which linkages between polymer chains giving a highly cross linked net work.
Hydrogenation of the hydrocarboxylated product might offer different physical properties which would be interesting to study. The resulting product would be less susceptible to cross linking than the hydrocarboxylated product and hence might find useful application as ion exchange resins.

5.2. Conclusion

Products of selective and non selective hydrocarboxylation of polybutadienes are being tested for possible commercial applications, particularly as corrosion inhibitors and in wood preservation. The principle of selectively hydrocarboxylating polybutadienes can be extended to other diene polymers. Introduction of more than one functional group on to a polymer will open scope for further research. The products thus obtained may have wide commercial application.
Chapter 6

Experimental techniques and Starting Materials

N.M.R. spectroscopy

$^1$H n.m.r. spectra were recorded on a Brucker AM300 spectrometer (300 MHz) or on a Brucker WP80 (80 MHz). $^{13}$C n.m.r. spectra were recorded on a Brucker AM300 instrument operating in the pulse Fourier Transform mode. The chemical shifts quoted in this thesis are relative to internal tetramethyl silane (TMS) for $^1$H and $^{13}$C spectra.

Infrared Spectroscopy

Infrared spectra of organic compounds were recorded on Perkin Elmer 1710 (Fourier Transform) spectrometer as Nujol mulls for solids and thin films for liquid between KBr plates.

Micro Analysis

Micro analysis were performed by the University of St.Andrews microanalytical service.

Differential Scanning Calorimetry (DSC)

All DSC scans were performed on a Perkin-Elmer DSC7 equipped with glove box and nitrogen purge. Samples were prepared using standard crimped sample pans. The heating rate was $10^\circ$C/min. Transition
temperatures have been corrected using Indium and Zinc standards. Tg's were taken as the midpoint in the transition curve.

**Solvents**

Methylisobutylketone (MIBK), tetrahydrofuran, diethylether, acetonitrile, acetone, benzene were obtained from Aldrich Chemical Company and, unless otherwise specified, were used as obtained.

**Vacuum Lines**

The experimental work previously described were carried out under nitrogen standard Schlenk line and catheter techniques. The nitrogen was further purified by passing through a Cr⁺ on silica column. The vacuum line used was a standard poly(tetrafluoroethylene) greaseless tap line with ball and socket compression joints.

**Starting materials**

1. Polybutadiene, *Cis* Average $M_w = 200,000$

2. Polybutadiene containing 45% vinyl, 55% *cis* and *trans*  
   Average $M_n = 4,500$

3. Polybutadiene diol containing 20% vinyl, 20% *cis*-1,4 and 60% *trans*-1,4

4. Polybutadiene, phenyl terminated containing 60% unsaturation  
   (45% vinyl, 10% *trans*-1,4 and 5% *cis*-1,4) Average $M_n = 1,000$
5. Polybutadiene, phenyl terminated containing 99% unsaturation (40% vinyl, 30% trans-1,4) 
   Average \( M_n = 1,300 \)

6. Polybutadiene, phenyl terminated containing 99% unsaturation (25% vinyl, 40% trans-1,4) 
   Average \( M_n = 1,500 \)

7. Polybutadiene, phenyl terminated containing 60% unsaturation (45% vinyl, 10% trans-1,4 and 5% cis-1,4) 
   Average \( M_n = 1,800 \)

8. Polybutadiene, phenyl terminated containing 99% unsaturation (25% vinyl, 40% trans-1,4) 
   Average \( M_n = 3,400 \)

9. Polybutadiene, hydrogen terminated containing 98% unsaturation (84% vinyl, 14% trans) 
   Average \( M_n = 3,000 \)

10. Polybutadiene, acid terminated containing 97% unsaturation (81% vinyl and 16% trans)

11. Polybutadiene, hydrogen terminated with narrow molecular weight distribution containing 97% unsaturation (88% vinyl, 9% trans)
Mₙ mentioned above means number average molecular weight and Mₘ means weight average molecular weight. Polybutadienes 1-8 were purchased from Aldrich Chemical Company. Polybutadienes 9-11 were obtained from Japanese Soda Company.

Tin(II)chloride dihydrate, Copper(II)chloride dihydrate, Triphenylphosphine, Triphenylphosphine Oxide, were purchased from Aldrich Chemical Company. PdCl₂ was purchased from Johnson Matthey and Company.

Preparation of Catalyst [PdCl₂(P(C₆H₅)₃)₂]

[PdCl₂(P(C₆H₅)₃)₂] was prepared according to standard procedures¹²⁶.
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