RESEARCH IN RUBBER CHEMISTRY AND TECHNOLOGY: A BIBLIOMETRIC STUDY AND TREND REPORT 1994-1998

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CERTIFICATE

This is to certify that the dissertation entitled IN "RESEARCH RUBBER CHEMISTRY AND TECHNOLOGY: BIBLIOMETRIC Α STUDY AND TREND REPORT 1994-1998" presented by Ajitha A.S in partial fulfilment of the requirement for the award of the Degree of Master of Library and Information Science is an original work carried out by her under my guidance and supervision. No part of this work has been submitted earlier for the award of any other degree.

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CONTENTS

| | | Page No |
|--------|--|------------|
| СНАРТЕ | CR-I INTRODUCTION | 1-8 |
| 1.1 | Title of the study | |
| 1.2 | Definition of the Key Terms | |
| 1.3 | Objectives | |
| 1.4 | Methodology | |
| 1.5 | Implications and use | |
| 1.6 | Limitations | |
| 1.7 | Organization of the study | |
| СНАРТЕ | R-II RUBBER CHEMISTRY AND TECHNOLOGY : GENESIS, DEVELOPMENT AND DIMENSION | 9-22 NS |
| 2.0 | Introduction | |
| 2.1 | History | |
| 2.2 | The Chemistry of Rubber | |
| 2.3 | Discovery of Accelerators | |
| 2.4 | | |
| 2.5 | Shaping | |
| 2.6 | | |
| 2.7 | Rubber Industry | |
| 2.8 | Conclusion | |
| СНАРТЕ | R – III LITERATURE ON RUBBER CHEMISTRY AND TECHNOLOGY: A BIBLIOMETRIC ANALYSIS | 23-32 |
| 3.0 | Introduction | |
| 3.1 | Year-Wise Distribution of Materials | |
| 3.2 | Distribution of articles in periodicals | |
| 3.3 | Subject-wise Distribution | |
| 3.4 | Authorship pattern | |
| 3.5 | Conclusion | |
| | | |

CHAPTER – IV RESEARCH ON RUBBER 33-94 CHEMISTRY AND TECHNOLOGY: A TREND REPORT

4.1 Rubber Physics and Rubber Engineering

4.11 Rubber Rheology

| | 4.12 | Properties of rubber | |
|--------|----------|--|--------|
| | 4.13 | Rubber in engineering applications | |
| 4.2 | Dubba | r Chemistry | |
| 4.2 | | Elastomers | |
| | | | |
| | 4.22 | | |
| | 4.23 | | |
| | 4.24 | | |
| | 4.25 | | |
| | 4.26 | Latex | |
| 4.3 | Rubbe | Technology | |
| | 4.31 | | |
| | 4.32 | Rubber products | |
| | 4.33 | Vulcanization | |
| | 4.34 | | |
| | 4.35 | Application of compounding ingredients | |
| | 4.36 | Reclaimed rubber | |
| | 4.37 | Rubber blends and composites | |
| | 4.38 | Physical testing | |
| | 4.39 | Dynamic mechanical properties | 2 |
| | | 1 1 | |
| СНАРТЕ | R V | FINDINGS AND CONCLUSION | 95-106 |
| | | | |
| | Introdu | | |
| 5.1 | | gs Based on bibliometric analysis | |
| | | Year-wise Distribution | |
| | 5.12 | | |
| | 5.13 | 3 | |
| | 5.14 | Authorship Pattern | |
| 5.2 | Research | ch trends in Rubber Chemistry and Techno | ology. |
| 5.3 | Conclu | sion | |

LIST OF TABLES

| | | Pa | ge No. |
|----|--|----|--------|
| 1. | YEAR-WISE DISTRIBUTION OF PUBLICATIONS | ٠ | 21 |
| 2. | DISTRIBUTION OF PERIODICAL ARTICLES | | 26-28 |
| 3. | SUBJECT-WISE DISTRIBUTION OF ARTICLES | | 30 |
| 4. | AUTHORSHIP PATTERN | | 32 |

CHAPTER - I

INTRODUCTION

Rubber is one of our most interesting and most important raw material. This material has multifarious uses and there is hardly any segment of life which does not make use of rubber based materials. It is also a vital raw material having immense strategic importance. Rubber is the most fascinating material known to mankind both on account of its range of applications in everyday life, defence and civilian purposes and its behaviour under the most diverse conditions of applications.

During the 19th century, several epoch-making discoveries were made in the processing, compounding and use of natural rubber, which resulted in tremendous increase in the demand for this product. Research in rubber is directed mainly toward making a better synthetic rubber to provide improved rubber products for home and industrial use. In addition, many unusual types of rubber are required in the age of nuclear energy and space travel.

Researchers are also trying to develop rubber that can be processed with a minimum of energy.

India now occupies the fourth position in the world in terms of rubber production. More significantly, the present national average productivity of rubber is one of the highest among the rubber producing countries.

A number of studies have been conducted in the field of Rubber Chemistry and Technology. Considering all the above factors, an attempt is made to investigate research in Rubber Chemistry and Technology: A Bibliometric study and Trend Report.

1.1 TITLE OF THE STUDY

The title of the study is Research in Rubber Chemistry and Technology: A Bibliometric study and Trend Report, 1994-98.

1.2 DEFINITION OF THE KEY TERMS

The meaning of the terms used in the title are given by the following definitions.

RESEARCH

Research is repeated search or repeated inquiry.

Oxford dictionary defines research as a 'studious inquiry usually critical and exhaustive investigation of experimentation having for its aim, the revision of accepted conclusions in the light of newly discovered facts'.

RUBBER

It was originally named coutchous; literally means tears of weeping wood. English chemist Priestly found that this could erase pencil marks and hence got the name "Rubber".2

RUBBER CHEMISTRY AND TECHNOLOGY

It includes primarly processing, chemical modification, rubber technology and product development.

BIBLIOMETRICS

Harrod's Librarians' Glossary defines

Bibliometrics as the application of mathematical and statistical methods to the study of the use made of books and other media within and between library systems.3

TREND REPORT

It is a type of review literature of a subject intended to study the developments achieved during a particular period of time and the possible achievements in near future.

1.3 OBJECTIVES

The study aims at the following

- 1. To examine the growth of research studies in Rubber Chemistry and Technology during 1994-98;
- 2. To identify the thrust areas of research;
- 3. To identify the year-wise distribution of publications;
- 4. To identify the most productive journal in the field;
- 5. To examine the past, the present and the anticipated near future developments in the field of Rubber Chemistry and Technology; and
- 6. To prepare a trend report on research in Rubber Chemistry and Technology.

1.4 METHODOLOGY

The methodologies applied in this study for data collection are 1. Literature survey; and 2. Bibliometrics. For this, primary journals in the field of Rubber Chemistry and Technology published during 1994-98 were examined and articles selected. Secondary Journals such as RAPRA abstracts, AEDR were also used for data collection. Bibliometric technique was also used for the quantitative analysis of data.

1.5 IMPLICATIONS AND USE

The study is aimed to serve the following.

- 1. To use this study as a reference tool by the research scholars, students and scientists in the field;
- 2. As a helpful tool for the compilation of bibliography in the field;
- 3. To enable the specialist in the area to become familiar with developments in the field so as to cope with it.

1.6 LIMITATIONS

The scope of the study is confined to literature published in the field of study during a period of five years from 1994-1998.

1.7 ORGANIZATION OF THE STUDY

The study is organized under the following chapters.

CHAPTER - I : INTRODUCTION

This chapter forms the introductory chapter where the subject is introduced, key terms are defined, objectives, methodology and limitations are provided.

CHAPTER - II : RUBBER CHEMISTRY AND TECHNOLOGY: GENESIS, DEVELOPMENT AND DIMENSIONS.

In this chapter, an overall view of the subject field, Rubber chemistry and Technology is included. Origin and development of the subject, its divisions and other related aspects are also explained in detail.

CHAPTER - III : LITERATURE ON RUBBER CHEMISTRY AND TECHNOLOGY: A BIBLIOMETRIC ANALYSIS

A quantitative analysis of the collected data are given in this chapter.

CHAPTER - IV : RESEARCH ON RUBBER CHEMISTRY AND TECHNOLOGY: A TREND REPORT.

A Trend Report showing the developments in the subject under study during 1994-1998 is given in this chapter.

CHAPTER - V : FINDINGS AND CONCLUSION

This chapter constitutes the major findings of the study. Conclusions are drawn and recorded.

REFERENCES

The studies used for the preparation of the Trend report are arranged systematically.

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CHAPTER - II

RUBBER CHEMISTRY AND TECHNOLOGY: GENESIS, DEVELOPMENT AND DIMENSIONS

2.0 INTRODUCTION

Natural rubber is a plantation crop cultivated in large areas as a monocrop, with an economic life cycle of around 32 years. Sustainable farming should essentially be adopted in rubber plantations as well. Natural rubber is nature's most versatile vegetable product. This material has multifarious uses and there is hardly any segment of life which does not make use of rubber based materials. Rubber is the most fascinating material known to mankind both on account of its range of applications in eveyday life, defence and civilian purposes.

2.1 HISTORY

Christopher Columbus is believed to have first found rubber in tropical South America. On return from his second voyage, 1493-1496, he brought a few balls made out

of the gum of a tree and used by the inhabitants of Haiti for playing games. There is also documentary evidence that in 1615 the Spanish troops in Mexico wore rubber-coated clothes to guard themselves from rain.

Guayule shrub was probably the first known source of rubber which was discovered in North America. Castilla tree from which rubber or ule was obtained and made into play-balls at Mexico was found to be the next one. Charles de la Condamine, a Frenchman, first described Hevea brasiliensis tree in 1751 as the source of rubber. He found it in the Amazon forests in 1736 during his visit to South America as a member of the party sent by the Academie des Sciences on scientific expedition. Condamine explain the crude methods adopted by the natives for extracting a milky fluid by cutting the bark of the trees and in using the fluid for coating fabric, making shoes and moulding water proof containers.

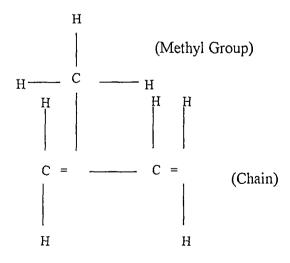
In India, rubber had been in use from time immemorial. In 1810, Roxburgh found that rubber was derived from Ficus elastica grown in Assam and he described that in his 'Flora Indica!' A rubber – yielding vine, Urceola elastica, was discovered in the straits settlements by

Howison in 1798. An inferior quality rubber was obtained from Landolphia vine in the west coast of Africa. Though rubber from various sources was available for centuries, the wild trees of Heve a brasiliensis in the South American forests have been the principal source of the world's supply of natural rubber in the early years. Since this was not sufficient to meet the increasing demand, it became an urgent necessity to increase the production by extending rubber cultivation to the Eastern hemisphere. The first suggestion to introduce rubber to this part was made by Thomas Hancock in 1857.

Hevea brasiliens's is a native of Brazil, which was introduced to tropical Asia in 1876, through kew, with the seeds brought from Brazil by sir Henry Wickham. The tree is now grown in the tropical regions of Asia, Africa and America. The rubber tree is sturdy, quick growing and tall and grows in many types of soils provided the soils are deep and drained. A warm humid, equable climate and a fairly distributed annual rainfall of not less than 200cm are necessary for the optimum growth of this plant. The tree, however, grows successfully under slightly varying conditions also.

2.2 THE CHEMISTRY OF RUBBER

The Chemistry of natural rubber has presented a challenge to scientists. In 1826, Michael Faraday, an English physicist and chemist, discovered that rubber is a That is, it belongs to a large family of hydrocarbon. substances that are composed of hydrogen and carbon. In 1860, another Englishman, Greville Williams, heated some rubber and obtained a colourless liquid that he called Each isoprene molecule contains five carbon atoms and eight hydrogen atoms (C5H8). The atoms in the isoprene molecule always form a definite pattern. Four of the carbon atoms form a chain. The fifth carbon atom branches off from one of the carbons in the chain. Three hydrogen atoms surrounded the fifth carbon atom to form a methyl group. The following chemical symbols show the arrangement of the five carbon and eight hydrogen atoms in the isoprene molecule.



In natural rubber, thousands of tiny isoprene molecules link together in a giant, chainlike molecule, the rubber molecule. Chemists call such chainlike molecules polymers, meaning "many parts". They call single molecules, such as isoprene, monomers. The particular chainlike structure of the rubber polymer explains why rubber is elastic. Polymer molecules of unstretched rubber fold back on themselves somewhat like irregular coils. Stretching the rubber straightens the chain of folded molecules. Releasing the rubber lets the chain return to its coiled position.

Sulphur combines with the rubber during vulcanization to set up 'cross-links' between the rubber chains. The chains in unvulcanized rubber can slip, causing the rubber to be less elastic. During vulcanization, the cross links bind the chains together so they cannot slip past one another. This gives elasticity and strength to the vulcanized product. However, if the cross-linking is carried too far, the cross-links tend to stop the unfolding of the chains. This reduces the elasticity of rubber. The number of cross-links increases according to the amount of sulphur, added to the compound. With large amount of sulphur,

rubber becomes stiffer, tougher, and less stretchable, until it turns into hard rubber. Chemists believe that the characteristics of many rubberlike substances depend on the way their atoms are joined together. Natural rubber has many unsaturated carbon atoms. Oxygen atoms from the air gradually attach themselves to these carbon atoms. This breaks down the rubber polymers so that the rubber becomes brittle or soft and loses elasticity. The addition of antioxidants during compounding prevents this action.

2.3 DISCOVERY OF ACCELERATORS

The prime function of an accelerator, as the name implies, is to increase the rate of vulcanization. The first artificial accelerators were inorganic. Goodyear disclosed the use of basic lead carbonate in his original patent. It was largely replaced by litharge, which was the most widely used accelerator for over 50 years. It is considered as a good accelerator and is used in many compounds. Other inorganic materials which are, or have been widely used as accelerators are lime, magnesia, sodium hydroxide, and antimony sulfide. Ammonia was used as early as 1880.

Organic accelerators were first used Oenslager in 1906. About 1910, German chemists working on synthetic rubber discovered the accelerating activity of various organic bases. Among the first widely used organic accelerators were hexamethylenetetramine, various other aldehyde amines, DPG, and DOTG. Mercaptobenzothiazole, introduced around 1925, and its derivatives, especially MBTs, have continuously increased in popularity until today they are the most widely used of all accelerators. More or less concurrent has been the growth of thiurams and dithiocarbonates. Around 1920, it was discovered that zinc oxide improved the action of many accelerators and that the action of zinc oxide was in turn improved by the presence of fatty acids such as stearic acid. The recognition of this situation was a great stimulus to the use of organic accelerators, particularly of the thiazole and thiocarbomate types.

2.4 COMPOUNDING OF RUBBER

The term compounding in rubber technology is used for selection and incorporation of various additives into rubbers so as to give a homogeneous mix for processing.

Compounding helps to control the elasticity, strength, and other properties of the final product.

The materials used in compounding can be categorised as Elastomers, Fillers. Processing Acids, Vulcanizing agents, Accelerators, Activators, Antioxidants, Antiozonants and miscellaneous additives. Elastomers form the backbone and are the basic components. They can be natural or any synthetic rubber or thermoplastic elastomer and are selected to give specific physical properties to the final product. Fillers are used to reinforce or reduce the cost of the product. Carbon blacks are the most important fillers used for rubbers. Their incorporation brings about marked improvement in properties such as hardness, tensile strength, tear and abrasion resistance. The processing acids help during processing. Vulcanising agents bring about cross-linking resulting in a marked improvement in the physical properties of the rubber products. Accelerators reduce the time of vulcanization by accelerating the rate of vulcanization. Activators activate the accelerators to work more effectively and increase the cure rate. Antioxidants are added to slow down and improve the aging behaviour Antiozonants slow down the deterioration due to attack by

ozone. The miscellaneous additives are not normally incorporated in the rubber compounds but may be used for specific purposes.

2.5 SHAPING

Manufacturers use several methods to shape rubber into final products. These include Calendering, Extrusion, Moulding and Dipping.

Calendering means rolling rubber into sheets. It is done on a machine that has two to five rolls mounted one above the other. Rubber is passed between pairs of highly polished heated rolls under pressure. Proper calendering requires precise control of roll temperature, pressure and speed of rotation. High rubber mixes are difficult to calender and require higher temperature. Incorporation of fillers ease calendering. Calendering can be done single bank or double bank. Double bank calendering gives more uniform thickness as air blistering is reduced. The products include rubber flooring, toys, bedsheets, washers, rings etc.

In the extrusion process, rubber is propelled continuously along a screw through regions of high

temperature and pressure where it is melted and compacted and finally forced through a hole to give the final product. Extruded products include hoses, inner tubes and rubber strips. Extruded products are vulcanized after they have been formed.

Moulding produces shoe soles and heels, rubber tyres, hot-water bottles, industrial products etc. compression moulding is the most common method used in the industry. Many products are formed in moulds and vulcanized at the same time. During vulcanization, the rubber takes the exact shape of the mould in which it has been placed.

Dipping is used only to make products from liquid latex. Products made by dipping include rubber gloves and toy balloons.

2.6 SYNTHETIC RUBBER

Research in rubber is directed mainly toward making better synthetic rubbers to provide improved rubber products for home and industrial use. Rubber like materials made from chemicals are called synthetic rubbers because

they were intended as substitutes for natural rubber. Manufacturers group synthetic rubbers into two classes: general-purpose synthetic rubbers and special-purpose synthetic rubbers.

General-purpose synthetic rubbers have many uses. The most important general-purpose rubber is styrene-butadiene rubber (SBR). It usually consists of about three parts of butadiene and one part styrene. Butadiene, a gas, is made from petroleum. It must be compressed or condensed into liquid form for use in making rubber. Styrene is a liquid made from coal tar or petroleum.

Special-purpose rubbers have special properties such as resistance to oils and fuels, air and extream temperatures, that make them better than natural rubber for certain uses. Special purpose rubbers include Butyl rubber, Cis-polyisoprene rubber, Neoprene rubbers, Nitrile rubber, Polysulphide rubbers, Polyurethane rubbers, Silicone rubbers, Ethylene-propylene rubbers, Fluorocarbon rubbers and Thermoplastic rubbers.

2.7 RUBBER INDUSTRY

Thomas Hancock made many remarkable observations and inventions from 1819 which established a real base for the rubber industry. It was he who performed for the first time the operation known as 'rubber compounding'. The 'masticator' fabricated by Hancock and the mixing mill and calender developed by Chaffee in 1836 are landmarks in the history of rubber industry. Charles Goodyear spent his whole life and money on improving the properties of rubber. His work and repeated failures from 1831 finally made him succeed in discovering 'vulcanization' in 1839 and this laid the foundation upon which the rubber industry is built.

The first pneumatic tyre was invented in 1845 by a Scotsman called Thomson but this was not taken seriously for about half a century. It was J.B. Dunlop, a veterinary surgeon of Belfast, who reinvented pneumatic tyre and he patended it in 1888. The invention of pneumatic tyre was a milestone in the history of rubber industry as the tyre sector today is the biggest consumer of rubber. Goodrich started his programme of manufacturing rubber goods in 1870 at Akron, and this place turned out to

be the leading rubber manufacturing centre in the world.

The continuous developments in the technology have developed a great deal in the commercial production of a large variety of rubber goods.

2.8 CONCLUSION

Over the past 100 years, the importance of rubber to civilized living has increased at a constantly accelerated pace. According to present estimate, the consumption of rubber in the world stands at about 16.66 million tonnes. For these reasons, tremendous interest is shown in research on rubber products. The future of natural rubber will also be bright in view of its special qualities and its competitiveness with synthetic rubber, the production of which is dependent on the high- priced and non-renewable petroleum resources.

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CHAPTER - III

LITERATURE ON RUBBER CHEMISTRY AND TECHNOLOGY: A BIBLIOMETRIC ANALYSIS

3.0 INTRODUCTION

Bibliometric distributions are used to <u>study</u> the frequency of occurrence of words in a text, productivity of authors in terms of scientific papers and scattering of articles over different journals.

A bibliometric analysis of literature published during 1994 – 1998 in the field of Rubber Chemistry and Technology is provided in this chapter. The variables used for analysis include year, subject, core journals and authorship pattern. The articles were collected from various primary periodicals in the field of Rubber Chemistry and Technology. Secondary journals such as RAPRA abstracts, AEDR were also examined. 500 articles published in the field of study during 1994 – 1998 were selected. The analysis of collected data are carried out under the following heads.

3.1 YEAR-WISE DISTRIBUTION OF MATERIALS

An attempt has been made to find out the year-wise distribution of the papers. The data given in Table I shows the year-wise distribution of publications from 1994 – 1998.

TABLE - I
YEAR-WISE DISTRIBUTION OF PUBLICATIONS

| Year | Number of Articles | Percentage |
|--------------------------------------|------------------------------|---|
| 1994 1995 1996 1997 1998 | 69 128 154 74 75 | 13.8% 25.6% 30.8% 14.8% 15.0% |
| Total | 500 | 100.0% |

From the data given in the table it is clear that maximum contribution, 154 (30.8%) published in the year 1996. It was followed by 1995 with 128 contributions (25.6%). The year 1998 gets the third position in the case of publications with 75 contributions (15%). Since much of the

papers of 1998 will appear in the issues of 1999, it is difficult to give the overall contribution of 1998.

3.2 DISTRIBUTION OF ARTICLES IN PERIODICALS

Journal productivity refers to the number of articles published by a journal over a particular period of time. It implies that the journals that publish more number of articles on a subject are more productive than others. Journals constitute the bulk of the literature in any discipline and serve as the forum where different intellectual contributions are made by different authors and carry current information.

Attempts have been made to identify the core journals that produce maximum number of articles in the field of Rubber Chemistry and Technology. 46 journals were identified and listed. They were ranked on the basis of the largest number of the articles in the field of Rubber Chemistry and Technology.

The data given in Table 2 shows the core journals in the field of Rubber Chemistry and Technology arranged in decreasing frequency of productivity.

TABLE - II

DISTRIBUTION OF PERIODICAL ARTICLES

| SL. No. | NAME OF JOURNAL | NUMBER OF ARTICLES | PERCE- NTAGE | RANK No. |
|------------|--|--------------------------|-----------------|-------------|
| 1 | Journal of Applied Polymer Science | 109 | 21.8 | I |
| 2 | Rubber Chemistry and Technology | 88 | 17.6 | II |
| 3 | Kautschuk Gummi Kunststoffe | 75 | 15.0 | III |
| 4 | Journal of Rubber Research (Formerly Journal of Natural Rubber Research) | 54 | 10.8 | IV |
| 5 | Rubber World | 24 | 4.8 | .V |
| 6 | Journal of Elastomers and Plastics | 23 | 4.6 | VI |
| 7 | Plastics Rubber and Composites: Processing and Applications | 15 | 3.0 | VII |
| 8 | Rubber Chem Review | 11 | 2.2 | VIII |
| 9 | Indonesian Journal of Natural Rubber Research | 9 | 1.8 | IX |
| 10 | International Polymer Science and Technology | 9 | 1.8 | IX |
| 11 | Rubber India | 9 | 1.8 | IX |

| 12 | Draggage in Dubban and | 6 | 1.2 | X |
|----|---|-----|-----|------|
| 12 | Progress in Rubber and Plastics Technology | 0 | 1.2 | |
| 13 | Rubber Developments | 6 | 1.2 | X |
| 14 | Polymer | . 5 | 1.0 | XI |
| 15 | Polymer International | 5 | 1.0 | XI |
| 16 | Journal of Adhesion Science and Technology | 4 | 0.8 | XII |
| 17 | Journal of the Rubber Research Institute of SriLanka | 4 | 0.8 | XII |
| 18 | Rubber Asia | 4 | 0.8 | XII |
| 19 | Bulletin of the Rubber Research Institute of SriLanka | 3 | 0.6 | XIII |
| 20 | Polymer Degradation and stability | 3 | 0.6 | XIII |
| 21 | European Polymer Journal | 2 | 0.4 | XIV |
| 22 | Indian Journal of Natural Rubber Research | 2 | 0.4 | XIV |
| 23 | Journal of Biological Chemistry | 2 | 0.4 | XIV |
| 24 | Journal of Polymer Materials | 2 | 0.4 | XIV |
| 25 | Journal of polymeric Materials | · 2 | 0.4 | XIV |
| 26 | Polymer Engineering and Science | 2" | 0.4 | XIV |
| 27 | Rubber and Plastics News | 2 | 0.4 | XIV |
| 28 | Rubber News | 2 | 0.4 | XIV |
| 29 | ANRPC News | 1 | 0.2 | XV |

| 30 | Biomaterials | 1 | 0.2 | XV |
|----|--|-----|-----|----|
| 31 | European Journal of Biochemistry | 1 | 0.2 | XV |
| 32 | European Rubber Journal | 1 | 0.2 | XV |
| 33 | FEBS Letters | 1 | 0.2 | XV |
| 34 | Indian Rubber Journal | 1 | 0.2 | XV |
| 35 | International Journal of Polymeric Materials | 1 | 0.2 | XV |
| 36 | Journal of Adhesion | 1 | 0.2 | XV |
| 37 | Journal of Polymer Science: Polymer Chemistry | 1 | 0.2 | XV |
| 38 | Journal of Polymer Science: Polymer Physics | · 1 | 0.2 | XV |
| 39 | Macromolecular Reports | 1 | 0.2 | XV |
| 40 | Materials World | 1 | 0.2 | XV |
| 41 | Phytochemistry | 1 | 0.2 | XV |
| 42 | Plantations, Recherche, Development | 1 | 0.2 | χV |
| 43 | The Planters' Chronicle | 1 | 0.2 | XV |
| 44 | Plastics and Rubber Weekly | 1 | 0.2 | XV |
| 45 | Polymer Plastics Technology Engineering | 1 | 0.2 | XV |
| 46 | Trends in Polymer Science | 1 | 0.2 | XV |
| | TOTAL | | 100 | |

The data given in Table 2 shows that Journal of Applied polymer Science published 109 article (21.8%) in the field of Rubber Chemistry and Technology. So this is the most productive journal in this field. The second position goes to Rubber chemistry and Technology. This journal published 88 articles (17.6%). The third position goes to Kautschuk Gummi Kunststoffe, published 75 articles (15%). The fourth position goes to Journal of Rubber Research which published 54 articles (10.8%) out of the total articles published during the period of study.

3.3 SUBJECT-WISE DISTRIBUTION

Subject-wise distribution of publications in the field of study is subjected to detailed analysis. The publications are grouped under the following subjects.

TABLE - III
SUBJECT-WISE DISTRIBUTION OF ARTICLES

| S1. No. | SUBJECT | NUMBER OF ARTICLES | PERCE- NTAGE | |
|------------|--|--------------------------|-----------------|--|
| 1 | Latex | 70 | 14.00% | |
| 2 | Rubber Blends and Composites | 61 | 12.20% | |
| 3 | Synthesis of Elastomers | 46 | 9.20% | |
| 4 | Chemical Analysis | 41 | 8.20% | |
| 5 | Rheology of Rubber | 38 | 7.60% | |
| 6 | Techniques of Vulcanization | 35 | 7.00% | |
| 7 | Rubber compounding | 34 | 6.80% | |
| 8 | Chemical reactions and modifications | 28 | 5.60% | |
| 9 | Instrumental analysis | 25 | 5.00% | |
| 10 | Properties of Rubber | 25 | 5.00% | |
| 11 | Physical Testing of Rubber | 16 | 3.20% | |
| 12 | Dynamic mechanical properties of Rubber | 16 | 3.20% | |
| 13 | Rubber products | 15 | 3.00% | |
| 14 | Application and uses of compounding ingredients | 15 | 3.00% | |
| 15 | Latex Allergy | 12 | 2.40% | |
| 16 | Rubber In Engineering Applications | 9 | 1.80% | |
| 17 | Preparation and reactions of compounding ingredients | 7 | 1.40% | |
| 18 | Reclaimed Rubber | 4 | 0.80% | |
| 19 | Testing equipment | 3 | 0.60% | |

The analysis of the data shows that the major subjects under study which had contributed the maximum number of studies in the field of latex. It had 70 papers to its credit (14%) out of the total 500 papers. It is followed by Rubber blends and composites with 61 papers (12.2%). The third position goes to Synthesis of Elastomers with 46 papers (9.2%).

3.4 AUTHORSHIP PATTERN

Authorship is a desirable phenomenon reflecting the contemporary practices by clearly showing patterns of communication, productivity and influences among researchers. Scientific productivity is frequently measured in terms of the published output, mostly because the data on the number of publications by authors can be easily collected and are quite reliable. The contributions of authors in terms of number of papers published on the subject is an indicator of the productivity of the author. The higher the contribution, more the productivity of authors.

TABLE - IV
AUTHORSHIP PATTERN

| AUTHORSHIP | YEAR | | | | momaz | PERCE- | |
|---------------------|------|------|------|------|-------|--------|-------|
| PATTERN | 1994 | 1995 | 1996 | 1997 | 1998 | TOTAL | NTAGE |
| Single Author | 12 | 28 | 21 | 9 | 6 | 76 | 15.2 |
| Two Authors | 19 | 47 | 37 | 23 | 21 | 147 | 29.4 |
| Multiple Authors | 38 | 53 | 96 | 42 | 48 | 277 | 55.4 |
| TOTAL | 69 | 128 | 154 | 74 | 75 | 500 | 100 |

From the table it is clear that majority of the publications are brought out by multiple authors, 277 (55.4%). Publications contributed by two authors are 147 (29.4%). The data reveals that single author contributions are relatively less. This indicates that collaborative research is more in the field of study.

3.5 CONCLUSION

The bibliometric analysis helped in identifying the various characteristics of the contributions in Rubber Chemistry and Technology using various parameters.

CHAPTER - IV

RESEARCH ON RUBBER CHEMISTRY AND TECHNOLOGY: A TREND REPORT

This chapter provides a trend report on Rubber Chemistry and Technology. The trends have been traced through the analysis of abstracts of the publications published in the field. The period covered is from 1994 to 1998.

The analysis is carried out mainly under 3 broad classes and each of them is again divided on the basis of narrower concepts dealt within. The subject groupings followed in the study are listed below.

4.1 RUBBER PHYSICS AND RUBBER ENGINEERING

- 4.11 Rubber Rheology
- 4.12 Properties of rubber
- 4.13 Rubber in engineering applications

4.2 RUBBER CHEMISTRY

- 4.21 Elastomers
- 4.22 Polymerization
- 4.23 Reactions of compounding ingredients.
- 4.24 Chemical reactions and modifications
- 4.25 Chemical analysis
- 4.26 Latex

4.3 RUBBER TECHNOLOGY

- 4.31 Compounding
- 4.32 Rubber products
- 4.33 Vulcanization
- 4.34 Rubber machinery
- 4.35 Application of compounding ingredients
- 4.36 Reclaimed rubber
- 4.37 Rubber blends and composites
- 4.38 Physical testing
- 4.39 Dynamic mechanical properties

4.1 RUBBER PHYSICS AND RUBBER ENGINEERING

4.11 RUBBER RHEOLOGY

Aranguren and others studied the rheological properties of filled elastomers. They described about the rheology of silica-silicone rubber (1/1994). Ashalatha and others described about the rheological and mechanical properties of PP/EPDM/NR ternary blends (2/1994). Ashalatha et al conducted another study on the rheological and mechanical properties of compatible PP/CPE blends (3/1996).

Bhattacharya and Chakraborty conducted a study on rubber processing. They used peptiser for product performance (4/1997).

Claramma and Mathew described about the rheological behaviour of prevulcanized latex. They have studied about the viscosity modifiers (5/1998).

The evolution of rubber processing and its challenges have been studied by Freakley (6/1996).

Ganesan and others described about the carbon black dispersion in rubber by an image processing

technique. They have studied about the quantitative measurement of dispersion (7/1995).

Ganesan and others conducted another study about the quantitative fractography of rubber by image processing technique (8/1995). Gopalakrishnan and others outlines about the effect of blend rheology (9/1995).

Guriya and others described about the rheological properties of EDDM compounds. They have studied about the effect of blowing agent, curing agent and silica filler (10/1998).

Han and white explain about the rheological studies of dynamically vulcanized blends of polypropylene and ethylene-propylene rubber (11/1995).

Isayev and Wan conducted a study about the material characterization. They have studied about the injection molding of rubber compound with rheology affected by vulcanization (12/1996).

Ishikawa et al studied about the processing agents for silica-filled rubber (13/1998). Joseph and others described about the melt rheological behaviour (14/1994).

Karnika de Silva and others studied about the vulcanizable processing aid for natural rubber. They

described about the depolymerized natural rubber as a processing aid (15/1996).

Leblanc outlines about the rubber compound testing. He described about the discrepancies in testing rubber compounds with different type of commercial capillary rheometers (16/1994).

Leblanc and others described about the mixing behaviour and rheological properties of SBR compounds. They explained about the composite design experiments to study the relationship between the mixing behaviour and rheological properties (17/1994). Lee and others studied about the effect of network formation on the rheological properties of SBR modified asphalt binders (18/1997). Li and white described about the rheological behaviour of highly filled EPDM compounds with calcium carbonate, carbon black, silica and zinc oxide (19/1996).

Long explains about the current status and future prospects of rubber processing and products (20/1995).

In a study conducted by Marinelli and Magg described about the rheological properties of rubber compounds. They described about a new acrylonitrile-but adiene rubber for moulded articles (21/1995).

Mousa and others have studied about the mixing rheology (22/1997).

Mukhopadhyay described about the developments in rubber processing technology (23/1995).

Nair and others have studied about the viscosity modifier in nitrile rubber. They described about liquid natural rubber as a viscosity modifier (24/1997). Osanaiye et al outlines about the rheological properties of rubbercarbon black and carbon oil compounds (25/1995).

Ray and others have studied about the effect of zinc salt of unsaturated fatty acids as processing aid (26/1994). Reena Susan George and others described about the reheological and extrude behaviour of natural rubber (27/1995).

In a study by Schmidt and Kreipe described about the thermal utilization of waste rubber (28/1995).

Sambatsompop and others described about the rheological behaviour and flow vulcanization. They have studied about the flow analysis of natural rubber in a capillary rheometer (29/1997). Sombatsompop and wood have studied about the flow analysis of natural rubber in a

capillary rheometer. In this study the flow patterns and entrance velocity profiles are described (30/1997).

Soos and white described about the rheological characterization of NBR elastomers. The use of BF goodrich dynamic stress relaxometer is described in this study (31/1996).

Thomas and others studied about the processing properties of raw rubber. A comparative evaluation of ISNR 20 with conventional forms of natural rubber are used (32/1995).

Varkey and others studied about the effect of prevulcanizatgion on the rheological behaviour of natural rubber (33/1996). Wada outlines about the rubber friction test methods (34/1998). Wan and Isayev described about the injection molding of rubber compound with rheology affected by vulcanization (35/1996).

4.12 PROPERTIES OF RUBBER

Ahlbland et al outlines about the physical properties of rubber/cellulose fibre composites. They have studied about the effect of electron irradiation on physical properties (36/1996).

Anantharaman and others conducted an investigation on cure characteristics and the magnetic properties of rubber ferric composites (37/1998).

Azemi Bin Samsuri described about the properties of rubberised bitumen (38/1996). Bhattacharya and Gangopadhyay studied about the properties and processing techniques of butadiene rubber (39/1998). Gajewski and Prot described about the crosslink density and properties of rubber (40/1994). Galan and others outlines about the effect of adhesive composition on the properties (41/1996). Gibala and Hamed described about the cure behaviour (42/1994). Guyriya and Tripathy outlines about the dynamic viscoelastic properties of carbon black loaded closed-cell microcellular ethylene-propylene-diene rubber vulcanizates (43/1996).

Hagen and others studied the effects of the type of crosslink on viscoelastic properties of natural rubber (44/1996).

Heinrich and Dumler outlines about the wet skid properties of filled rubbers (45/1998).

Kakubo and others described about the origin of characteristic properties of natural rubber. They described

the effect of fatty acids on crystallization of cis-1, 4-polyisoprene (46/1998).

Karnika de Silva conducted study on some of the properties of blends of polyisoprene (47/1994). Li and others conducted study on variation of structure and properties of natural rubber (48/1998). Low et al described about the properties of rubber-modified cellulose-fibre-epoxy laminates (49/1994). Manas and others studied about the physical properties of NR/polystyrene blends. They studied the effects of dicumyl peroxide and liquid natural rubber (50/1998). Nishiyama and others described about the origin of characteristic properties of natural rubber (51/1996).

Okel and Waddell studied about the silica properties/rubber performance correlation (52/1994).

Patkar studied the viscoelastic properties. The effect of silica is described in this (53/1998). Pu and Ding described the properties of EVA preirradiation grafted NIPPAm. The preparation and characterization are also described (54/1996). Srilathkutty and others studied about the effect of fillers on the properties of NBR/PVC microcellular shoe soles (55/1998). Suwardin and others described about optimizing the quality control of SIR 20 crumb rubber (56/1995). To et al suggests sulfanimide

accelerators for improving the properties of rubber (57/1996).

Uchiyama described the non-slip properties of rubber (58/1998). Vink suggests the influence of filler surface treatment on properties of reinforced rubber products (59/1996). Wang and Wang suggest a new technique to quantitatively characterize crosslink density (60/1998).

Wang and others described the dynamic properties of filled rubber. They suggests the dynamic stress-softening of filled vulcanizates (61/1998).

Yang and Hsiue describes the physical and dielectric properties of acrylic acid grafted silicone rubber (62/1996).

4.13 RUBBER IN ENGINEERING APPLICATION

Bauman described high – value engineering materials from scrap rubber (63/1995).

described Bhuana and others about the earthquake rubber damper. They suggest the homogenization of temperature and reduction of vulcanization time by applying additional heat source (64/1996).

Chatterjee described about the rubber spring in earthquake protection (65/1995).

Davies and others outlines the stress relaxation under repeated stressing (66/1996). De Varies described about the heat-resistant conveyor belts (67/1995). Dierkes explain about the rubber recycling. The re-use of rubber waste is described in this (68/1995). Dutta gives an account of the engineering application of rubberişed fabrics (69/1994). Gopalakrishnan described about the rubberised roads (70/1995; 71/1994).

4.2 RUBBER CHEMISTRY

4.21 ELASOMERS

Besdo and Ihlemann describes the material behaviour of elastomers (72/1996).

Blume and others describes the kinetic aspects of nitrosamine formation during the production and storage of elastomers (73/1996).

Bowers outlines the new developments in improving the processibility of fluoelastomers (74/1997). Chaudhury and Bhowmic described the thermostable insulating thermoplastic elastomers (75/1996).

Class described the peroxide crosslinking of elastomers (76/1995).

Drake and others suggested the use of polybutadiene coagents in peroxide cured elastomers (77/1995).

Eisenbach described about thermoplastic elastomers and polyurethane elastomers. He described the morphology of filled structutre and rubber and thermoplastic elastomers (78/1996). Elleithy and others described about the fracture and fatigure failure behaviour fluoroelastomers (79/1996).Ellul suggests temperature resistant thermoplastic elastomers for speciality applications (80/1997).

Fukushima et al describes the synthesis of graff copolymers from highly deproteinised natural rubber (81/1998).

Furukawa and others described the preparation of functionally graded polyurethane elastomers. The method used is electrophoresis of monomer (82/1997).

Gerspacher and others proposed a mechanism for the reinforcement of elastomers (83/1994).

Hamed gives the characteristics and properties of crosslinked elastomers (84/1994). Hamed and Wu descibes the relationship between the cohesive strength and tack of elastomers (85/1995). Herd and Bobo give an account of the analysis and properties of carbon black filled elastomer blends (86/1995). Kelm and others described about the crosslink density of elastomers (87/1998). Kohler and O'Neill describe technology for а new the devulcanization of sulfur-cured scrap elastomers (88/1997). Kotz and Stevens described the improved proce3ssing variants of viton GFLT fluoroelastomers (89/1995). Mente described the antiozonant protection elastomers (90/1995). Liu and Li studied about the curing reaction kinetics and properties of elastomers (91/1996).

Martin et al described about the morphology and properties of polyurethane elastomers. They described the average soft segment length (92/1996). Meissner gives an account of bound rubber and elastomer – filler

interaction (93/1995). Mousa and others describes the oil-resistance study (94/1998). Nakauchi and others describes the crosslinking structure and properties of elastomers (95/1996). Nuyken and others studied about the reinforcing polymeric fillers. They suggests core-shell polymers as reinforcing polymeric fillers (96/1995). Parker et al suggests the applications of 'Dimide-Hydrogenated' styrene butadiene and polybutadiene thermoplastic elastomers (97/1994).

Pader and Agarwal studied about the thermoplastic elastomers for automotive applications (98/1998).

Roychoudhury and De describes about the elastomer-carbon black interaction (99/1995).

Song and others conducted studies on the relationships between network structure and mechanical properties of interpenetrating polymer networks (100/1995).

Steinbuchel describes about the synthesis and production of biodegradable elastomers (101/1996).

Suhara and others give an account of the stress relaxation of polyester fiber-polyurethane elastomer composite (102/1998).

Sykes describes about the structure and property relationships in polyurethane elastomers. (103/1996).

Thomas gives and account of the thermoplastic elastomers and their influence in future world elastomer consumption (104/1995). Tonelli et al describes about the new flurorinated thermoplastic elastomers (105/1996).

Van der pol describes about the fiber reinforcement of elastomers (106/1995). Vladkov and Amaudova describes the influence of oligoamidophosphates (107/1994). Xie and others describes about the synthesis and properties (108/1996).

4.22 POLYMERIZATION

Change Nemeth studied about the rubber particle agglomeration phenomena (109/1996). Datta and others describes about the electron beam initiated grafting and crosslinking (110/1996). Decker et al gives an account of the polymerization of multifunctional monomers. Photocrosslinking is used (111/1996; 112/1996;113/1996). El – Ghaffar describes about the emulsion polymerization of vinyl acetate. A semicontinuous polymerization technique is used (114/1995).

Fang and others described about the radiation-induced graft copolymerization (115/1996). He and others described about the formation mechanism of silicone rubber particles with core-shell structure by seeded emulsion polymerization (116/1996). Wood and Iroh describes the effect of electrolytes and process parameters on the electropolymerization (117/1996). Yang and Hsiue describes about the graft polymerization of acrylic acid. Preirradiation method is used (118/1995).

4.23 REACTIONS OF COMPOUNDING INGREDIENTS.

Ahagon describes about the evaluation of polymer chain during mixing of filled compounds (119/1996).

Arnaudova and others describes about the influence of oligoamidephosphates (120/1996).

Ghosh and others suggest a plant product modified carbon black as a reinforcing filler (121/1995).

Compounds of natural rubber containing different fillers were evaulated for their air permeability. The results indicated that incorporation of filler generally reduce the permeability. Certain fillers such as mica, graphite and

clay produced substantial reduction which was attributed to the particular morphology of these fillers as evidenced from photomicrographs (122/1997).

Pramanik describes about LLDPE composites (123/1995). Pushpa and Billingham describes about the solubility of antioxidants in rubber (124/1996).

Yusof and others describes about the purification and characterisation of an inhibitor of rubber biosyntesis (125/1998).

4.24 CHEMICAL REACTIONS AND MODIFICATIONS

Abbate and others give an account of a new reactive liquid rubber with maleimide end groups (126/1996).

Abdel-Aziz describes about the influence of polyfuctional monomers (127/1995).

Avirah and Joseph conducted studies on natural rubber bound para-phenylenediamine antioxidants (128/1995).

Bielinski and others describes about the influence of chemical modification on tribological properties of elastomers (129/1995).

Cho et al have found the effect of interfacial adhesion on toughening behaviour of rubber modified poly(methyl methacrylate) (130/1997). Debnath and Basu conducted studies on the effect of thiuram disulphide on naturtal rubber vulcanization (131/1996). Derouet and others described about the phosphorus modification of expoxidized natural rubber. This can improve flame resistance (132/1994).

De Sarkar and others describes about the optimization of reaction conditions (133/1997). Ellul et al outlines the crosslink densities and phase morphologies (134/1995). Evans and Waddell describes about the ultrahigh reinforcing precipitated silica (135/1995). Ghosh et al outlines the effect of modified carbon black (136/1997). Huang and Tsiang describes the effect of tetrahydrofuran as a structure modifier (137/1996). Isayev and others describes the elastic recovery of rubber vulcanizates (138/1996). Ishiaku et al describes the effect of di-2-ethylhexyl phosphate (139/1996). Kali Ray explains the chemical modification of EPDM rubber (140/1997). Lee and

others explains the method for estimating the chemical crosslink densities of cured natural rubber (141/1994).

Myhre and Mackillop outlines the modification of crumb rubber (142/1996).

Nasar and Goman describes about the influence of physical ageing (143/1995).

O' conner and ohters describes about the role of rubber particles (144/1997). Okieimen and Urhoghide conducted studies on miscibility of poly(vinyl chloride) (145/1996). Ou and others describes the effects of alkylation of silica filler on rubber reinforcement (146/1994).

Puskas and Wilds describes about the kinetics of the epoxidation of butyl rubber. They describes the development of high precision analytical method (147/1997).

Roychoudhury and others shows the chemical interaction between carbon black and elastomers (148/1994).

Tanka and others gives the structural characterisation of natural rubber (149/1997). Tangpakdee and Tanka describes about the branching of natural rubber (150/1998). Thammathadanukul and others shows a

comparison of rubber reinforcement using various surfacemodified precipitated silica (151/1996).

4.25 CHEMICAL ANALYSIS

Anachkov and others described about the DSC study of thermal decomposition (152/1996). Ansarifar and Lake explains the mechanics of rubber-to-metal bon failure (153/1995). Bandyopadhyay and others described about the surface-oxidized carbon black. 3-Aminopropyltriethoxysilane is used as a promoter in the crosslinking of carboxylated nitrile rubber (154/1996; 155/1996). Bayers described about the silane-coupling agents for enhanced silica performance (156/1998). Bhaumic and Datta describes about the thermal analysis (157/1995). Blume and others explains the analysis of N-Nitrosamines (158/1996).

Bohin and others describes the using of a dispersion model for single agglomerates. They describes the analysis of power requirements and dispersion quality in batch compounding (159/1996).

Bonfils and others explains the chromatography study of natural rubber films (160/1995).

FT-NMR is used in the analysis of rubber blends crosslink distribution in carbon black filled blends of NR and cis-BR have studied. Sulphenamide accelerators are

used in this study. (161/1995)

n.

others described Buding and about influence of peptizer on rubber-to-steel cord adhesion Burbank and others outlines about the (162/1998).processing continuous of high quality compounds (163/1995).Campbell et al describes about characterisation of the chemical structure of sulfur vulcanizates of natural rubber. They used the 1H NMR spectroscopy method for the analysis (164/1994). Campbell and Farley describes about the gel formation during latex epoxidation. Acid-catalysed hydrolysis is used (165/195).

In a study chandra and others describes about the scanning electron microscopy, energy dispersive X-ray studies of the influence of a cobalt adhesion promoter on the interface between rubber skim compounds and tyre steel cord (166/1996).

¹³C NMR spectroscopy has been used to study vulcanised blends of natural rubber. Effect of transpolyoctenylene on crosslink structure of NR and SBR are explained (167/1997).

Chetri and others outlines about the homogeneous acetalization and esterification of poly (vinyl) alcohol) in non-aqueous medium Ethyl nitrate dimethyl sulfoxide is used (168/1998). Sulfur vulcanization of NR, BR, SBN and their blends have been studied. This study is conducted by using rheometer and DSC (169/1996). Effective processability measurements of acrylonitrile butadiene rubber have been described by Cousins and Dick (170/1998).

The experimental results of the crack growth in twisted rubber disks have been studied by De and Gent (171/1998). Dehnicke studied about the acqueous adhesives as an alternative to conventional rubber-to-metal adhesives (172/1998).

An investigation on the distribution of fillers in natural rubber by element spectroscopic transmission electron microscopy is described (173/1994). Ellul and Hazelton studied about the chemical surface treatments. The effects on friction and adhesion are explained (174/1994).

The reaction of moist ozone with natural rubber have been studied using attenuated total reflectance spectroscopy (175/1998). ¹³C NMR study of oxidative

degraded rubbers is conducted by Eng and others (176/1998). A study is conducted by Fong and others about the mathematical model of diffusion of solvents in rubber (177/1998).

Hydrogenation of natural rubber using nickel 2-ethylhexanate catalyst in combination with trisobutylaluminium is described by Gan and others (178/1996). Ganghoffer and Schultz described about the cavitation in rubber (179/1995).

Geethamma and others described about the short coir fiber-reinforced rubber composites. They investigated the effects of fibre length, orientation and alkali treatment (180/1995).

In a study conducted by Gonzalez and others a new interpretation of the crosslinking mechanisms of natural rubber with dicumyl peroxide have described (181/1998). Diffusion of organic solvents in rubber membranes is measured (182/1995; 183/1995).

Thermal degradation of polythiophene-natural rubber is described by Hacaloglu (184/1997).

Hamed and Zhao described about the longitudinal cracking. During chemical stress relaxation of

carbon black filled natural rubber vulcanizate the longitudinal cracking takes place (185/1998).

An attempt is made to analyse the network structure of BR and SBR vulcanizates. Model vulcanization and sulfur vulcanization is described. NMR spectroscopy is used (186/1998).

Hardiman and Kartika conducted study on stabilizing mooney viscosity of narural rubber. This is conducted by using sodium phenolate, sodium metabisulphite and acetaldehyde (187/1995).

Proton NMR studies of molecular mobility in vulcanized and eposidised natural rubber is conducted (188/1996). Analytical electron microscopy study is conducted by Hofer and others. The characterisation of rubber-brass bonding layers have been studied (189/1996).

Swollenstate ¹³C spectroscopy has been used to study vulcanised blends of natural rubber/nitrile butadiene rubber and natural rubber/epoxidised natural rubber both qualitatively and quantitatively. The results obtained show that although the polymers present can be identified and the individual polymers characterised in terms of their copolymer composition, the use of this technique to determine the actual blend composition is unsatisfactory. The reasons

for the deviations of the results obtained from the true values are discussed (190/1994).

Hummel and others described about the rubber-brass bonding. Morphology of cross sections through the bonding layers as a possible basis for classification is described (191/1996).

A comparative study of the effect of thermooxidative aging on the physical properties of rice husk ash and commercial fillers in epoxidized natural rubber compounds have been explained (192/1997).

Job and Joseph described about the studies of the adhesives for rubber to rubber bonding (193/1995). FTIR analysis of silica-filled natural rubber is described by Kralevich and others (194/1998).

Linos and Stein Buchel describes about the microbial degradation of natural and synthetic rubber. They used a novel bacteria for this. This bacteria is belonging to the genus 'Gordona' (195/1998).

The density of chemical crosslinks and chains entanglements in unfilled EPDM vulcanizates studied with low resolution solid state IH NMR (196/1998).



The microstructure of carbon black is investigated by atomic force microscopy (206/1994). IHNMR analysis of thermal groups in the rubber chain is described by Ohye and Tanaka. The structural characterisation of rubber from tactarium mushrooms are identified (207/1998). The evaluation of the performance of polymeric roofing materials have studied by Paroli and Delgado. They used thermal analysis for this (208/1996). Quigley and others described about the viscoelastic constitutive models for rubber. They have determined the material constants (209/1995).Richards and Roberts described about the friction of various rubber articles lubricated with acid and alkaline water (210/1995).

Selden outlines the fracture mechanics analysis of fatigue of rubber (211/1995). The nonlinear elastic analysis of the adhesive strength between rubber like materials and regid surfaces are analysed by Sun and Chang (212/1995).

The gel content of rubber from high-ammona latex (HA-latex) decreased significantly after deproteinization with proteolytic enzyme. The addition of 1-2% ethanol in toluene solution reduced the gel content of rubber from HA-latex, deproteinized HA-latex and pale crepe. Transesterification of the rubber in toluene solution with

sodium methoxide dissolve the gel fraction. The gel fractions solubilized after transesterification showed molecular weight distribution rich in low molecular-weight fraction. The findings suggest that the branching and crosslinks are composed of two types of branch-points, one by association or aggregation of proteins or oligopeptides at the initiating end and the other by ester linkages including phospheric ester at the terminal end (213/1997).

Urabe and others conducted studies on rubber mixing. The action of zinc oxide during the mixing process is explained in this (214/1995). The rubber to metal bonding study is conducted by Van Ooji (215/1996). Veeralakshmanan and others described about the chlorination of liquid natural rubber (216/1995).

Nitrosamine formation in rubber is explained by Willoughby and others (217/1998).

Yeoh described about the rubber analysis. HE used hyperelastic material models for finite element analysis (218/1997).

4.26 LATEX

Aftab Alam conducted a study on the estimation of protein contamination in latex (219/1996). Akasawa and others outlines a new acidic allergen, Hev b 5, in latex. (220/1996). Amir-Hashim described about the effect of leaching and humidity as prevulcanized natural latex film. The most effective method is the use of a recentrifuged prevulcanized latex with a combination of wet gel leaching and dry-film leaching or direct water spraying (221/1997).

Azizah and others conducted study on latex protein allergy among factory workers. The finding of the study are described in this (222/1996). Beezhold and others described about the surface-bound latex protein antigens on surgical gloves (223/1995).

There is an urgent need to produce latex gloves with minimum extractable proteins. Stability of antigenic proteins in latex extracts is described by Bradley and others (224/1996).

The quality control of latex products used in the medial setting has to take into account the presence of allergens which induce a Type I hypersensitivity response. Assays to measure the total protein eluted from these products does not address the problem of allergenicity of the

proteins extracted. The study by cardosa and others introduces to rubber manufacturers the concept of enzyme immunoassay and describes various immunoassay formats by which antigens can be detected (225/1994).

The gloves produced by different manufacturers contain different amounts of antigens and antibodies. Monoclonal antibodies are used to show that different antigens are found in different latex gloves (226/1994).

Sulfur prevulcanization of natural rubber latex was conducted at 60,70,80 and 90 o C for different periods. At lower temperatures the rate of reaction was slow (227/1997; 228/1997).

Cook and other describes about the microstructure of natural rubber latex films (229/1997; 230/1997).

The properties of natural rubber latex concentrated by creaming method for gamma ray irradiation is described by Dafader and others (231/1996). The permeability and material characteristics of vulcanized latex film is described by Dillon and Schroedar (232/1997). The mechanisms of latex film formation is explained by Dobler and Holl (233/1996).

Elizabeth and others described about the treatment of examination of gloves for reducing extractable protein content (234/1996). Geurts and others described about the molecular mass control in methacrylic copolymer latexes (235/1997).

Low protein latex-has been found to be suitable for factory scale production of examination gloves. A post vulcanized formulation was used in the factory run. Unaged tensite strength of these gloves was in the region of 25 MPa-27MPa and a retention of more than 908 was obtained when they were aged at 70°C for 7 days and 100°C for 1 day (236/1994).

Gidrol and others described about the lectin-like protein called havein from the rubber trees. The use of havein in the coagulation of latex is explained (237/1994).

Hamsa and Amir-Hashim described about the latex processing. The changes of latex proteins during processing is explained (238/1997; 239/1998). They described about the elastomeric films from structured latexes (240/1997). They also conducted another study about the grafting behaviour of n-Butyl acrylate (241/1997). In another study they described about the hydrogenation of styrene-butadiene rubber latexes (242/1997).

Jacob and others described about the concentrated natural rubber latex. They points out the glove manufacture (243/1995). Karnika de Silva outlines about the natural rubber latex based adhesives (244/1996). Karnika de Silva and Hamielec conducted a Kinetic study of the graft polymexization of unsaturated monomers with natural rubber latex (245/1994). Khalid and others described about the dielectric properties of have a latex at various moisutre contents (246/1994).

The isomerization of isopentenyl diphosphate to dimethyl diphospate and the formation of oligoprenyl diphosphate as well as polyprenyl diphosphates, catalysed by IDP-Isomerase and polyprenyl transferases, respectively, were observed following bottom fraction which had been preincubated at 4°C overnight (247/1996).

Laliamma Jose and Rani Joseph suggest an accelerator system for transparent latex products (248/1996). Legge describes about the TPE condoms and described about the ozone level in it (249/1994). Llosent and others described about the particle size distribution of SBR and NBR latexes by UV-VIS turbidimentry (250/1996).

The allergenic responses in latex sensitive persons are evaluated by means of the skin-prick test.

Studies are in progress to identify the allergens. Allergenic repossess to radiation cured natural rubber latex is explained (251/1996).

Martin and Urban described about the spectroscopic studies of latexes (252/1996). Matthan explained about the underexploited export potential for the Indian latex gloves industry (253/1994). Mehta described about the cause of cessation of viral passage through artificially-induced holes in latex condoms (254/1998). Miller described about the latex sales (255/1996).

Considerable criticism has been made against the use of natural rubber latex for the maxifacture of elastic medical goods due to allergic reactions that can be used. A survey reveals that the proportion of the population affected by latex proteins is small and the effects are not usually serious. The inferior performance of PVC, Polyurethane, nitrile and neoprene, and styrene copolymers can lead to increased health and safety risks, and that they have also been known to cause toxi and allergic effects (256/1994).

Nagarajan and others described about the latex interpenetrating polymer networks. Their synthetic variations have explained (257/1996; 258/1996; 259/1996; 260/1996).

Prick test studies of raw materials and products of commercial natural rubber latex showed different threshold levels of residual extractable proteins. The threshold level of MRL products is currently 50-100 g/g, a level impossible to attain by leaching out the protein content. The threshold recommended is 30-40 g/g. (261/1998; 262/1997).

Naon and others describes about the drying of granules from natural rubber latex (263/1995).

With the reported incidence of Type 1 hypersensitivity associated with the presence of residual water-soluble proteins in latex products, there is an urgent need to produce latex gloves with minimum extractable proteins. The study shows that an additional leaching operation after drying is necessary the most effective means is the use of a recentrifuged prevulcanized latex with a combination of wet gel leaching and dry-film leaching or direct water spraying. Under the most favourable conditions, an extractable protein content of approximately 0.03 mg/g can be achieved (264/1994).

Niu and Urban outlines about the surface and interfacial FTIR spectroscopic studies of latexes (265/1995; 266/1996). Nordin Abdul Kadir Bakti and Isa described

about the efficient treatment system for rubber glove manufacturing plant (267/1997). Odeberg and others described about the influence of spacer groups on grafting ability, curing ability and film properties of water-based radiation curable latexes (268/1998; 269/1996).

Fluorescence method of studying void closure kinetics during film formation from high T latex particles are described by Pekcan and Canpolat (270/1997).

The problems currently facing by latex products manufacturers are protein allergy, and presence of nitrosamines and nitrosatable amines. Pendle in his paper discusses about the significance and scale of these problems in relation to the formulation of compounds and of production technique (271/1997).

Schloman and others described about the performance testing of guayule latex (272/1996). Shamsui Bahri describes about the immunocytochemical localisation of rubber membrance proteins in have a latex (273/1996).

Singha and others proposed a new method to hydrogenate nitrile rubber in the latex form (274/1995). Soedjanatmadia and others described about the processed products of the hevein precursor in the latex of the rubber tree (275/1995). Solichin described the preparation of

constant viscosity stabilized rubber from thin slab derived from latex in the presence of hydroxylamine neutral sulphate (276/1995).

Natural Hev b 7, a 43-k Da patatin-like allergen was purified from the latex of Hevea brasiliensis and several internal peptide sequences were determined (277/1998).

Subramaniam and others described about the use of mechanistic principles to make modified natural rubber latex products using emulsion polymerization (278/1997).

A spectrum of proteins derived from Havea brasiliensis reacted with IgE in the plasma from persons allergic to natural rubber latex. Latex B-serum, C-serum and rubber particle membrane proteins were all found to contain allergenic proteins. Variations were observed in the reactivity of the latex IgE positive plasma to the different proteins. Monoclonal antibodies have been developed against some of these allergens (279/1994; 280/1995).

Suwardin describes about the simultaneous latex coagulation in the bamboo tube (281/1996). Tangboiboonrat and others explains the characterization of non-crosslinked natural rubber latex by phase transfer technique (282/1994; 283/1996). Tanrattanakul describes

about the toughening of polycarbonate with core-shell structured latex particles (284/1996). Tebelius and Urban conducted surface and interfacial FTIR spectroscopic studies of latexes (285/1995; 286/1996).

Thiandchanya and others described about the degradability testing of radiation – vulcanized natural rubber latex films (287/1994). Triwiyoso and Handoko described about the preservation of field latex. The study is aimed to increase the production of latex shares in NES project (288/1995). Tucci and others described about the structural features of latex gloves in dental practice (289/1996).

Varghese and others described about the use of sulphuric acid as a coagulant for natural rubber latex (290/1996).

Allergenic activity must take into account of the nature of allergenic proteins. White describes about managing the latex protein allergy (291/1996; 292/1995).

Yapa describes about the manufacture of low protein natural rubber pineapple juice or bromelain treatment of hevea latex (293/1995; 294/1995). Yeang describes about the new techniques in the quantitation of total proteins from latex gloves (295/1994). Yeng and others

conducted a latex allergy study by extraction of natural rubber latex proteins (296/1995).

With the awareness of Type' 1 allergy, improvements of latex-dipped products, particularly the medical gloves, invariably involve the reduction or removal of their residual extractable protein fraction containing the allergens. Studies have been carried out with 16 dry natural rubbers of various commercial grades and five rubber products including cut threads manufactured via processes quite different from those of latex-dipped articles. Findings reveal that they not only have extremely low extractable protein contents, but also show negligible or no allergic responses when skin-prick tested on a total of 28 latex hypersensitive persons. So dry natural rubber products are free from the protein allergy problem reported for some latex products (297/1997; 298/1997).

Yuen describes about the dithiocarbamyl compounds in natural latex concentrate (299/1995). Zehr and others described about the extraction of antigenic protein from latex films and gloves (300/1994).

4.3 RUBBER TECHNOLOGY

4.31 COMPOUNDING

Abo-Hashem and Elshokrofy explains about the thermal effects of carbon black-rubber composites (301/1994). Busfield and others described about the crack growth and strain induced anisotropy in carbon black filled natural rubber (302/1997; 303/1996). Dyrda and Meshchaniov describes about the dielectric permiability of rubber (304/1995).

The reinforcement mechanisms in carbon black and silica loaded rubber have been studied by Eggers and Schummer (305/1996). Ertong and others outlines about the flow properties of carbon black filled natural rubber. They described about the steady state shear flow properties (306/1994).

By using the thermal filed-flow fractionation and multi-angle laser light scattering Fulton and Groves determined the molecular architecture of synthetic and natural rubber (307/1997). Gent and Lai conducted a study on the adhesion and autohesion of rubber compounds (308/1995). Gerlach proposed some additives for the modification of silicone rubber (309/1995).

Gerspacher and others described about the high frequency viscoelasticity of carbon black filled compounds (310/1996).

A study conducted by Ghosh and others suggests modified carbon black as reinforcing filler for SBR (311/1996).

Gibala and others outlines about the cure and mechanical behaviour of rubber compounds containing ground vulcanizates. They explains about the mooney viscosity (312/1996). Gori and Nordsiek described about the rubber/filler batches in powder form. These batches will help to simplify the production of rubber compounds (313/1998).

Guerbe and Freakley gives a comparison of some indirect methods for measuring the carbon black disperson in rubber compounds (314/1998). Ismail and others describes about the different modes of action of a multifunctional additive in a carbon black filled natural rubber compound (315/1996; 316/1996).

Jurkowski and others conducted study on the methods of sulfur disperson and sulfur blooming in rubber compounds. They have used radioisotopic determination methods for their study (317/1996). Klasek and others

described about the rubber coupling agents (318/1996). Kluppel and Heinrich described about the fractual structures in carbon black reinforced rubbers (319/1995).

Koczorowska and Jurkowska conducted studies on sulfur behaviour in rubber compounds (320/1998). Li and others have studied about the carbon black aggregate structure. They proposed some simple methodologies for the modeling of carbon black aggregate structure (321/1996).

The evaluation of the external surface area of carbon black by nitrogen adsorption is described by Magee (322/1995).

Mantia and others conducted studies on the chemical interaction between surface oxidized carbon black and epoxidised natural rubber (323/1997).

Marwede and Sumner points out the new developments in the field of general purpose synthetic rubber (324/1998). Mass and Gronski conducted a study by atomic force microscopy. They have studied about the deformation of filler morphology in strained carbon black loaded rubbers (325/1995).

In a study conducted by Matheson and others the effect of carbon black filling on the pyrolysis of natural rubber and synthetic rubber have been identified (326/1996).

Medalia explains about the effects of carbon black on abrasion and treadwear (327/1994). Pond explains about the cavitation in bonded rubber cylinders (328/1995). Sangtani gives an introduction of practical rubber compounding (329/1997).

The effect of adhesion on the equilibrium swelling of short sisal fiber reinforced natural rubber composites have been described by Varghese and others (330/1995). Srilathakutty and others described about the use of amine terminated liquid natural rubber as a plasticizer in filled natural rubber and NBR compounds (331/1996). Tokita et al studied about the carbon black elastomer interaction modelling (332/1994).

Waddell and others studied about the mechanism of precipitated silica. This silica improves the brass-coated wire-to-natural rubber adhesion (333/1996).

4.32 RUBBER PRODUCTS

Anon described about the natural rubber mountings for Pakistan (334/1995). Holownika describes about the rubber cylinders (335/1995). Huyberechts described about the truck tyre retreading (336/1995).

The analysis of extractable proteins in rubber products have been described by Karnika de Silva (337/1996).

Kuhn-Stoffers and others described the determination of carbonyl compounds in emissions from manufacture and storage of rubber products (338/1997). Lambillotte and Eiber described about the truck tyre technology (339/1994).

Leca and Runcan studied about the reclaiming of rubber from waste tyres by physical methods (340/1996). Mitra described about tyres and the properties of tyres (341/1995).

Mohn. Nor. Zairossani and others conducted studies on biosorption of zinc from the effluent of rubber product (342/1998). Mukhopadhyay and others points out the future developments of tyre technology (343/1998).

Nakauchi describes about the service life of rubber products (344/1996).

Reader gives an account of medical and anatomical models (345/1996). Veith described about the pneumatic tyres (346/1998). In a study conducted by Will and Giese points out the influence of plasticizer aerosols on sampling and analysis of air pollutants in the rubber industry (347/1996).

Yip and others conducted study on the non-allergenicity of NR dry rubber products. The dry natural rubber products are free from the protein allergy problem (348/1995).

4.33 VULCANIZATION

Abdul-Aziz and others described about the radiation vulcanization of styrene-butadiene rubber. They studied about the effect of polyfunctional monomers on radiation vulcanization (349/1996).

The efficiency of modified phthalocyanine compounds as antioxidants and UV stablilizers for SBR vulcanizates have been studied by Botros and Abd El-Ghaffar (350/1997).

Chandran and Kuriakose described about the utilization of coconut pith and coir waste as filler in natural rubber vulcanizates (351/1995).

Sulfur prevulcanization of natural rubber latex was conducted at 60, 70, 80, and 90°C for different periods. The extend of crosslinking was assessed. At lower temperatures, the rate of reaction was slow. At each temperature, tensile strength and elongation at break decreased when the prevulcanization time increased, whereas the modulus increased up to a maximum cross linking and thereafter decreased. The rate of stress relaxation slightly increased as the extent of prevulcanization increased (352/1997).

Colvin and Bull described about sulfur or olefin copolymers as vulcanizing agents (353/1995).

Conveney and others suggests a triboelastic model for the cyclic mechanical behaviour of filled vulcanizates (354/1995).

Deiber and Bortolozzi give a comparison of two oscillating bicone cells for vulcanization (355/1995)

Ding and Leonov suggests a kinetic model for sulfur accelerated vulcanization of natural rubber compound (356/1996; 357/1996).

Gibbesch and others outlines the product constancy in the vulcanization of rubber lining for corrosion protection (358/1995). Guriya and Tripathy described about the dynamic viscoelastic properties of silica-filled closed-cell microcellualr ethylene propylene diene rubber vulcanizates. The effect of blowing agent, the effect of temperature and strain are described in this study (359/1995; 360/1995).

Hill and others described about the mechanism of radiation vulcanization of natural rubber latex sensitized by monocylates (361/1995). In a study conducted by Isayev and others suggests a new ultrasonic technology for devulcanization of rubber waste (362/1995; 363/1996). Flammability of cis-1, 4-polyisoprene vulcanizates are described by Janowska (364/1997). Jurkowsaka and Jurkowski outlines about the computer simulation of changing the rubber network structure during vulcanization (365/1998).

Kawabata and others described about the mechanism of carbon-black reinforcement of rubber vulcanizate (366/1995).

The effect of crosslink structures as the fatigue crack growth behaviour of natural rubber vulcanizates with various aging conditions were described by Kim and Lee (367/1994).

Kresja and others conduct an analysis of the mechanism of N-t-butyl-2-Benzothiazole sulfenamide accelerated sulfur vulcanization of cis-polyisoprene (368/1994).

Kretzschmar and Weigel described about the isothiocyantes which is formed during accelerated vulcanization (369/1997). Levin and others described about effect of crosslink type on the the ultrasound devulcanization of SBR vulcanizates (370/1998; 371/1997). Mathur and others described about the quarternary ammonium salts, the new curatives which are used for the vulcanization of natural rubber (372/1996).

Menon outlines about the stress-relaxation characteristics of natural rubber modified with phosphorylated cashew nut shell liquid polymer (373/1997). Nieuwenhuizen and others described about the

homogeneous catalysis by zinc complexes during accelerated vulcanization (374/1998).

In a study conducted by Poh and others described about the cure characteristics of unaccelerated sulfur vulcanization of epoxidised natural rubber (375/1996). The accelerator activity of TBAB is described by Purohit and Mukhopadhyay (376/1997).

Shaker and others outlines about the UHMW-PE fibre as reinforcing materials in EPDS rubber vulcanized E-Beam radiation (377/1995).

Ramsingwani describes about sulfur as a vulcanising agent (378/1995).

Temple and Esyev have studied about the continuous ultrasonic devulcanization of unfilled natural rubber vulcanizates (379/1998). Tukachinsky and others described about the devulcanization of waste tire rubber by powerful ultrasound (380/1996).

Van Duin and Soupahanthong conducted study on resin vulcanization. They described the chemistry of phenol-formaldehyde resin vulcanization of EPDM (381/1995).

Versloot and others outlines about the sulfur vulcanization of simple model olefins (382/1997; 383/1994; 384/1995).

Yushanov and others described about the ultrasonic devulcanization of SBR rubber (385/1998).

4.34 RUBBER MECHINERY

Othman and others described about the stress relaxation behavior of natural rubber vulcanisates (386/1996). Pingyue described the effect of stearic acid level on tear and heat build-up of natural rubber-based tread compounds (387/1994). Popovic and others outlines the stress-strain modelling (388/1996).

4.35 APPLICATION OF COMPOUNDING INGREDIENTS

Dias and Galuska described about the curative migration in rubber compounds (389/1996). Gorl and others investigate about the influence of white fillers on the formation of N-Nitrosamines (390/1994). Grtaft and others described about the application of promoters in rubber compounds (391/1996). Graf and Knorr give an account of the accelerator systems for the injection moulding

compounds (392/1996). The effect of Bis 3-Triethoxysilypropyl) tetrasulfide on silica reinforcement of styrene-butadiene rubber have been studied by Hashim and others (393/1998).

The influence of clearance variation between the rotors of an internal mixture on the mixing process have been studied by Michaeli and Meiertoberens (394/1994).

Nandavanam and others suggests linseed oil as a multipurpose ingredient in their study (395/1996). Terakawa and others described the application of epoxidized NR/diamine vulcanizates for tyre tread (396/1998). Rouse explains the application of crumb rubber modifier in asphaltic materials (397/1995).

The sorption and diffusion of aromatic hydrocarbons. Through filled natural rubber are described by Unnikrishnan and others (398/1996). Vande walle and others described about the carbon black reinforcement in rubber compounds (399/1996).

Wolf described about special carbon blacks for applications in mechanical rubber blacks for applications in mechanical rubber goods (400/1996).

Wolf and others described the effect of fillers on rubber reinforcement (401/1994; 402/1994).

4.36 RECLAIMED RUBBER

The processing, control and growth of rubber reclaiming is described by Biswas (403/1994; 404/1994).

Osborn explains about the reclaimed tire rubber in TPE compounds (405/1995).

4.37 RUBBER BLENDS AND COMPOSITES

Abbate and others described about the curing kinetics and mechanical behaviour of rubber blends. They described about the thermosetting bismaleimide rubber blends (406/1997). Abdullah and others described about the blending of natural rubber with linear low-density polyethylene (407/1995).

The partition coefficients of ester plasticisers between natural rubber and epxidised natural rubber were explained by Ahmad and others in their study (408/1995). A qualitative and quantitative analysis of SBR/BR blends by the thermogravimetric method is described in this study (409/1996). Asaletha and others outlines about the

technological compatibilization of natural rubber blends. This is done by the addition of natural rubber-graft-polystyrene (410/1995). The viscoelastic behaviour of NBR/EVA polymer blends were studied by Bandyopadhyay and others by the application of models (411/1998). Bensason and others described about the damage zone in PVC and PVC/MBS blends (412/1992; 413/1997).

Bhwomick outlines the influence of interaction on the properties of rubber-rubber and rubber-plastics blends (414/1996).

Biswas and others suggests new coupling agents for rubber blends in their study (415/1995). Blom and others described about the iPP/HDPE blends (416/1995).

Brown and Tinker explains the effect of modifying EPDM on the crosslink distribution in NR/EPDM blends (417/1996). In another study they described about the high damping epoxidised natural rubber blends (418/1997).

Bussi and Ishida explains the structural characterisation of the epoxidised rubber and the mechanical properties of the partially miscible blends (419/1994).

The effects of short-chain branching and comonomer type on the interfacial tension of polypropylene-polyolefin elastomer blends were described in a study conducted by Carriere and Silvis (420/1997). Chakraborty and Seth conducted studies on the blends of high styrene resin master batch and natural rubber (421/1998).

Cook and others described about the thermal behaviour of natural rubber and chlorinated rubber blends (422/1997).

Cook and others conducted another study about the morphology and property relationships in ABS/PET blends (423/1996; 424/1996).

The phase morphology of rubber and plastic blends were described by Coran and other (425/1998).

Elastomers are considered by many to be one of the most difficult groups of materials to prepare for electron microscopy. The study by Cudby and others describes the techniques used in scanning transmission imaging of elastomer blends using an unmodified conventional scanning electron microscope (426/1995).

Das and Banerjee explains the role of methods of blending on polymer-polymer compatibility (427/1996).

Dutta and others outlines the ionic elastomer blends of zinc salts of maleated EPDM rubber and carboxylated nitrile rubber (428/1998). Fujita and others describes the phase diagrams of the blends of natural rubber and petroleum resins (429/1998).

Natural rubber has a shorter scorch time than butadiene rubber, but results indicate that in blend materials the BR phase begins curing before and at a much faster initial rate than natural rubber. The cure rates are reversed as the cure progresses and the NR phase eventually attains the crosslinking levels expected from single polymer data (430/1996).

Huang and others conducted studies on the blends of carbondioxide copolymer using NBR/PPC system (431/1996). Ibar explains about the control of performance of polymers and their blends through melt vibration technology (432/1997). Thermo-oxidative ageing of poly (vinyl chloride) based thermoplastic elastomers were studied by Ishiaku and others (433/1997). Jansen and others studied about the effect of curing system on mechanical and thermal properties of the blends (434/1996; 435/1996). Kannan and others conducted studies on the oil distribution in polymer blends (436/1996). Kazno and others conducted studies on the distribution in Polymer blends (437/1996).

Kida and others conducted studies on the structure and formation mechanism of carbon gel in the carbon black filled polyisoprene rubber composites (438/1996).

Kim describes the effect of blending thermoplastic elastomer with natural rubber compound (439/1996). Kim and Do outlines the effect of viscosity ratio, rubber composition, and peroxide/coagent treatment in PP/EPR blends (440/1996). Kole describes the influence of the nature of diene in EPDM rubbers on the properties of silicone-EPDM blends (441/1994).

Compounding the ground rubber tire (GRT) with commercial polyolefins is a possible and effective way of used tirte recycling which can provide new rubber/plastic blends at a much tower cost. In the study GRT and ultra sonically deveulcanized GRT are incorporated into a polypropylene. GRT is compounded and dynamically relvulcanized in polypropylene. Mechanical properties, rheology and morphology as well as the thermostability and melting behaviour of the resulting rubber/plastic blends are studied (442/1998).

Mallick and others explains the effect of HAF carbon black filler on chemorheological behaviour of expoxidised natural rubber blend (443/1997; 444/1997).

Oshinski and others describes the effect of polyamide end-group configuration on morphology and toughness of blends with maleated elastomers (445/1996).

Petrovic and others describes the effect of addition of polyethylene on properties of polypropylene/ethylene – propylene rubber blends (446/1996).

The effect of blend ratio on mooney scorch time of rubber blends were described by Poh and Wong (447/1998). The mechanical properties, crosslink density and surface morphology of SBR/silicone rubber blends were described by Popovic and others (448/1997).

Radusch and Merseburg outlines the morphology formation in dynamic vulcanized PP/EDM blends (449/1996). Rahman and others outlines the influence of phase morphology on the properties of natural rubber and epoxidised natural rubber blends (450/1994). Roy and Das describes the speciality polyblend stem (451/1995).

Rozman and others describes the rubber wood-polymer composites. The study is based on glycidyl methacrylate and diallyl phthalate blends (452/1998).

Sankarapandian and Kishore gives a comparison of the miscibility stratetgies in poly (benzyl methacrylate) – polystyrene blends (453/1996).

Sorption and diffusion of n-allcanes into different blends of natural rubber such as NR/BIIR, NR/CIIR, NR/ neoprene, NR/EPDM, NR/polybutadiene and NR/SBR have been studied from 25 to 60°C. Arrhenius parameters for the diffusion process and the thermodynamic quantities for the process of equilibrium sorption have been estimated (454/1998; 455/1998).

Thomas and Gopinathan Nair describes the viscosity behaviour of PVC-modified liquid natural rubber blends (456/1996).

Tripathy and Das describes the effect of blending on the dispersion and flow behaviour of carbon black filled NBR and chlorosulphonated polyethylene rubber (457/1995; 458/1994).

Tsai and Chang outlines the polymer blends of PBT and PP compatibilized by ethylene-co-glycidyl methacrylate copolymers (459/1996).

Vainio et al describes about the functionalised polypropylene (460/1996). Varkey and Rao explains the

flow behaviour of natural rubber and expoxidised natural rubber latex blends (461/1996). Vilgis and Heinrich explains about the crosslinked polymer blends (462/1995).

Vinod and others conducted study on the effect of adhesion on the equilibrium swelling of natural rubber-aluminium powder composites (463/1998).

Wang and others conducted study on the effect on the mechanical properties of blends (464/1997). Wefringhaus and Gruber described about the water based rubber to metal bonding agents (465/1995). Youn and others describes about the blends of brominated EPDM and natural rubber (466/1996).

The mechanical properties and fracture behaviour of natural rubber/poly (methyl methacrylate) blends were investigated as a function of composition, graft copolymer concentration, and mixing conditions. The mechanical properties and failure behaviour vary with the blend ratio, graft copolymer concentration, and mixing conditions (467/1997).

4.38 PHYSICAL TESTING

Aboutorabi et al explains the fracture energy calculations (468/1998). Azemi Bin Samsuri describes the tearing of unvulcanized black-filled natural rubber (469/1995). Charlton and Teh describes the athermomechanical FEA for rubber components (470/1995).

A double-networked natural rubber (DNNR) was prepared by a "two-step cross linking method, in which the cross linking was achieved while the natural rubber was in stretched condition. The swelling behaviour, tensile properties, crepe, recovery and permanent set were investigated (471/1997).

Kasner and Meinecke describes the effect of porosity on mechanical properties of rubber (472/1995). Kim and others conducted study on the preparation and physical properties of rubber-modified epoxy resin (473/1997). Konar conducted a kinetic and rheometric investigation of hexamethylol melamine derivative on the cure of natural rubber compounds (474/1997).

Lee explains the effect of heating and pressuring on PBA/PS linear composite polymer (475/1998).

Mariani et al conducted study about the thermal properties of elastomeric compounds (476/1997).

The flow properties of thermally depolymerized liquid natural rubber were explained by Nair and others (477/1995; 478/1994).

Okel explains the effect of precipitated silica and physical properties on silicone rubber performance (479/1995). Okieimen and Alighodion describes the estimation of dilute solution viscosity parameters of rubber seed oil alkyds (480/1998). Santangelo and Roland explains the mechanical behaviour of double network elastomers (481/1994). Wang and others describes about the carbon-silica dual phase filler (482/1998). Youn and others explains the effect of bromination on the properties of ethylene-propylene-diene rubber (483/1990).

4.39 DYNAMIC MECHANICAL PROPERTIES

Abd-El-Messieh and Younan explains the dielectric relation and mechanical properties of natural and chloroprene rubber with some nitroaniline additives. A systematic dielectric study over a frequency range from 400 to 10MHz was carried out on natural rubber and

chloroprene rubber with a small amount of nitroaniline additives (484/1996).

The non-linear stress-strain characteristics of fille rubber, and attempts at producting a non-linear model have been reviewed. Alternative methods of calculating the equivalent linear parameters: secant, harmonic, or skeleton methods, give different answers, as demonstrated for a particular rubber (485/1997).

Bandyopadhyay describes the low strain dynamic properties of carboxylated nitrite rubber-carbon black system (486/1996).

Bhowmic and Jha explains the dynamic mechanical properties of thermoplastic elastomers (487/1997).

The dynamic properties of batch polyomerized SSBR based compounds were described by Bortolotti and others (488/1998).

Dick and Pawlowski explains the application of new dynamic mechanical rheological tester in measuring carbon black (489/1995).

Good child and Pond explains the dynamic mechanical properties of certain natural rubber vulcanisates (490/1997). Hanhi and Stenberg explains the friction and

dynamic mechanical and thermal properties of polyurethane elastomers (491/1994). Leblanc explains the dynamic strain methods to probe the morphology of uncured rubber compounds (492/1996).

Luo conducted study on the dynamic properties of polymeric isolators (493/1996). Mallic and Tripathey explains the effect of carbon black filler on strain dependent dynamic mechanical properties (494/1994). Mandal explains the dynamic mechanical properties of an ionic elastomer based on carboxylated nitrile rubber. The effect of calcuim carbonated and clay were explained (495/1997; 496/1996).

Metteo and others studied about the dynamic mechanical properties of NR/BR blends at different cure levels (497/1995). Schaefer explains the dynamic properties of rubber (498/1995).

The dynamic mechanical properties and morphology of silica-reinforced butadiene rubber by the solgel process were described by Sugia and others (499/1997). The electrical and mechanical properties of ethylene propylene diene monomer-chloroprene rubber blend loaded with white and black fillers were explained by Younan and others (500/1998).

CHAPTER V

FINDINGS AND CONCLUSION

5.0 INTRODUCTION

This chapter is a summing up of the major findings and inferences drawn from 'Bibliometric Analysis' and 'Trend Report'. The findings are fruitful in substantiating the objectives formulated in chapter one. The findings are grouped under two major headings:

- (1) Findings based on bibliometric analysis
- (2) Research trends in Rubber Chemistry and Technology.

5.1 FINDINGS BASED ON BIBLIOMETRIC ANALYSIS

The findings of this major heads are grouped under the following subheadings

5.11 YEAR-WISE DISTRIBUTION

From year-wise analysis of publications during the period under study, the following conclusions were drawn.

- 1. The year 1996 is the highest productive year, produced 154 articles (30.8%) out of the total 500 publications.
- 2. Year-wise distribution of publication on rubber chemistry and technology is relatively low in 1994, 69(13.8%).
- 3. The number of publications has decreased during 1997 and 1998.

5.12 CORE PERIODICALS

The periodical, "Journal of Applied Polymer Science" is the most productive journal in the field of study that published 109 articles (21.8%) during the period of study, followed by "Rubber Chemistry and Technology" which published 88 articles (17.6%) and "Kautschuk Gummi Kunststoffe" that published 75 articles (15%). These periodicals are the core journals in the field of Rubber chemistry and Technology.

5.13 SUBJECT-WISE DISTRIBUTION

In the case of subject-wise distribution, the literature on latex comes first with a percentage of 14 and the second position goes to Rubber Blends and composites with 12.2%.

5.14 AUTHORSHIP PATTERN

Authorship pattern shows that majority of publications are brought out by multiple authors (55.4%). The contribution by single author constitutes 15.2%.

5.2 RESEARCH TRENDS IN RUBBER CHEMISTRY AND TECHNOLOGY.

The findings based on research trends in Rubber Chemistry and Technology are grouped under the following sub-heads.

RUBBER RHEOLOGY.

Recent trends in the field of study show that the rheological behaviour of selfvulcanizable blends on chlorosulphonated polyethylene and carboxylated nitrile

rubber has been investigated in terms of effect of blend ratio, carboxyl content of carboxylated nitrile rubber, carbon black filler loading, rate of shear and temperature on viscosity, die swell and power law constants. The blend system follows the power law model in the experimental range of rate of shear and shows pseudoplastic flow behaviour.

PROPERTIES OF RUBBER

The degree of compatibility of two rubbers is important in determining the morphology and mechanical properties of blends made from them. The mechanical properties of the blends are largely determined by phase morphology and this is particularly marked with regard to air permeability and tear strength.

RUBBER IN ENGINEERING APPLICATION

The engineering applications of rubber is in the field of engine mounts, bridge bearings, piers and fenders, as flexible composites, noise control and structural dampling and in vibration isolation.

A study of the behaviour of laminated rubber bearings is described with particular emphasis on important aspects concerned with seismic isolation. The effect of compressive load on damping and the height reduction behaviour of a bearing under combined compression and shear were studied. Shear stiffness and the tilting of individual rubber layers in a bearing indicates that a departure from the normal linear shear stiffness behaviour occurs at large deformation.

ELASTOMERS

Thermoplastic elastomers have been prepared by blending poly (vinyl chloride) and epoxidised natural rubber in the presence of a plasticiser. Low temperature resistant thermoplastic elastomers were identified.

A novel preparation method of functionally graded interpenerating network polymer (IPN)- type elastomers were prepared by electrophoresis of ionic manomers. Abrasion resistance of the surface of IPN-type PUEs evaluated by DIN abrasion test was better than that of nontread polyurethane elastomers.

POLYMERIZATION

The non-rubber substances present in natural rubber latex seem to have influence on the rate of polymerization reaction, degree of grafting and grafting efficiency of graft copolymerization reaction of methyl methorylate on to natural rubber. The semicountinuous emulsion polymerization of vinyl acelate in the presence of potassium persulfate and octyl aldehyde sodium bisulfite as a newly developed redox initiation system was studied using polyvinyl alcohol as a unique surface active agent. It shows that very small amounts of polyvinyl alcohol in the presence of the newly developed octyl aldehyde sodium bisulfite led to the formation of highly stable emulsions, with 40% solid content.

COMPOUNDING INGREDIENTS

The incorporation of filler generally reduce the permeability. But certain fillers such as mica, graphite and clay produced substantial reduction which was attributed to the particular morphology of these fillers.

CHEMICAL MODIFICATION

Aerobic ageing is found to increase the dynamic modulus of the vulcanizates. Results at longer ageing times indicate the modulus may, depending up on the vulcanisates, ingredients, pass through a maximum, however the modulus remains greater than for unaged material.

The number of branch-points in natural rubber was analysed for the rubber from deproteinised fresh field and high-ammonia latices, using ¹³C-NMR and osmometric measurement before and after transesterification of the rubber.

CHEMICAL ANALYSIS

Epoxide groups in natural rubber, which are derived from oxidative degradation of polymer are found to be mainly in cis configuration. Solid state C¹³-NMR has been used to compare the chemistry of accelerated vulcanization of natural rubber using N-t-butyl-2-benzothiazole-sulfenamide in the presence and absence of carbon black. Swollen state ¹³C spectroscopy has been used to study vulcanised blends of natural rubber/nitrile

butadiene rubber and natural rubber/epoxidised natural rubber both qualitatively and quantitatively.

The application of gamma-ray attenuation techniques to characterize natural rubber, rubber compounds and latex has yet to be fully exploited. DRC of latex at various concentrations has been measured using the multi-component system and the results show a good correlation between the attenuation method and the standard laboratory method.

LATEX ALLERGY

By identifying the proteins of various latex compounds that are responsible for the allergenicity, it might be possible to eliminate, or at least greatly reduce these proteins. The strongest antigenic responses to the antibodies were from the B-serum proteins which arise from the lutoids. Two allergenic proteins have been isolated from the B-serum of natural rubber latex by dialysis- induced protein precipitation followed by gel filtration.

The Finnish Natural Public Health Institute and a team from the universities of Helsinki and Tampere reported a step forward in establishing which proteins in latex cause allergenic reactions.

A study revealed that dry natural rubber products are free from protein allergy problem reported for some latex products.

COMPOUNDING

Rubber compounding is a term covering a wide scope including the art and procedures that are necessary to produce useful products from natural or synthetic elastomers. A component rubber compounder must be able to use the proper elastomers and additives available to meet the conditions required for the finished product.

Different carbon black master batch blending procedure were used to achieve a wide range of black distribution. The black distribution and blending procedures influenced the physical properties significantly. Modified carbon black can be used as reinforcing filler for SBR.

RUBBER PRODUCTS

India is one of the leading countries which manufacture recycled rubber in the form of reclaim rubber and crumb rubber. Whole tire reclaim is the main product.

About 33% of reclaim rubber is used in bicycle tires, the result in battery, containers, footwear etc.

VULCANIZATION

Sulfur prevulcanization of natural rubber latex was conducted at 60, 70,80 and 90°C for different periods. At each temperature, tensile strength and elongation at break decreased when the prevulcanization time increased. The rate of stress relaxation slightly increased as the extent of prevulcanization increased.

Sulfur/olefin copolymers are known as vulcanizing agents for rubber, but synthetic procedures for these materials are poor. The relationship between the sulfur/olefin copolymer and vulcanizate properties are discussed in terms of the structure of the copolymer and other reaction products.

The radiation vulcanization of natural rubber latex has been investigated for many years. Recent trends shows that certain antioxidants are capable of improving the ageing properties of radiation vulcanization of natural rubber latex films.

MACHINERY

Stress-strain modeling is used to find out the resolution of natural rubber /poly dimethylsiloxane binary blend properties.

APPLICATION OF INGREDIENTS

Surface energy of fillers and its effect on rubber reinforcement have been determined.

The applications, performance and mode of action of promoters in rubber compounds have identified.

RECLAIMED RUBBER

A trend in the reclaimed rubber reveals the processing, control and growth of rubber reclaiming.

RUBBER BLENDS

A swollen-state NMR method of rubber blend analysis has been used to study cross linking in gum vulcanisates for cure times compatibility of natural rubber/polystyrene blend is poor and can be enhanced by the addition of a graft copolymer of natural rubber and

polystyrene. Light and optical microscopy and nuclear magnetic resonance have been used to study the phase contrast and crosslink distribution of the elastomer blends.

The mechanical properties and fracture behaviour of natural/poly (methyl/methacrylate) blends were investigated as a function of composition, graft copolymer concentration and mixing conditions.

DYNAMIC MECHANICAL PROPERTIES

Dynamic mechanical analysis of silica-filled closed-cell microcellular EPDM vulcanizates has been studied as a function of temperature and double strain amplitude. The viscoelastic behaviour of silica-filled microcellular EPDM is similar to that of solid rubber vulcanizates.

5.3 CONCLUSION

The above findings and discussions are fruitful in substantiating the objectives of the study. This study will help to examine the growth of research studies in Rubber chemistry and Technology during 1994-98.

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