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1	M. Kavitha	Drug Discovery and Drug Development with Artificial Intelligence in Artificial Intelligence in Pharmaceutical Technology (volume 2)	-
2	M. Kavitha	Pharmacogenomics and Overview in Current Trends in multidisciplinary sciences -Vol 9	-
3	Dr. K. Bhavyasri	TEXTBOOK OF PHARMACEUTICAL QUALITY ASSURANCE	-
4	Dr. K. Bhavyasri	HYPHENATED TECHNIQUE - THERMOGRAVIMETRY-DIFFERENTIAL SCANNING COLORIMETRY COUPLED TO MASS SPECTROSCOPY	-
5	Dr. K. Bhavyasri	Recent Research Trends in Pharmaceutical Science (Volume - 3)	-
6	Dr. M. Vijaya Bhargavi	REVIEW: HYPERTROPHIC CARDIOMYOPATHY	-

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# ARTIFICIAL INTELLIGENCE IN PHARMACEUTICAL TECHNOLOGY

Volume  
2



  
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# Contents

Chapters	Page No.
1. A Comprehensive Review on Past, Present and Future Scenario of Medical Devices in Healthcare Department <i>(Bhimani Rushi K, Megha Gandhi and Dr. Shital Faldu)</i>	01-19
2. Design and Characterization of Clopidogrel Bisulfate Microspheres <i>(K. Muni Raja Lakshmi and S. Poojitha)</i>	21-47
3. Trending Role of Artificial Intelligence in Dental Radiology <i>(Dr. Kriti Garg, Dr. Vishal Mehrotra, Dr. Sunita Pathak and Dr. Ravi Seth)</i>	49-67
4. Artificial Intelligence in Psychiatry: The Way Forward <i>(Rajnish Raj and Bhavneesh Saini)</i>	69-84
5. Drug Discovery and Drug Development with Artificial Intelligence <i>(Dr. Narender Boggula, S. Raja Shekhar, Marati Kavitha, Aravinda Nalla and Dr. G. Umarani)</i>	85-111

# Chapter - 5

## Drug Discovery and Drug Development with Artificial Intelligence

Dr. Narender Boggula, S. Raja Shekhar, Marati Kavitha, Aravinda Nalla and  
Dr. G. Umarani

### Abstract

As expenditure on drug development increases exponentially, the overall drug discovery process requires a sustainable revolution. Since artificial intelligence (AI) is leading the fourth industrial revolution, AI can be considered as a viable solution for unstable drug research and development. Generally, AI is applied to fields with sufficient data such as computer vision and natural language processing, but there are many efforts to revolutionize the existing drug discovery process by applying AI. Artificial intelligence (AI) is becoming more widely adopted in the pharmaceutical industry, creating both excitement and questions about its potential and long-term success. In recent years, a plethora of companies-ranging from large pharmaceuticals to startups – have set expectations of AI as a panacea that will revolutionize the industry. While the thought of AI as such is an appealing one, it is not realistic. This review provides a comprehensive, organized summary of the recent research trends in AI-guided drug discovery process including target identification, Hit identification, ADMET prediction, lead optimization, and drug repositioning. The main data sources in each field are also summarized in this review. In addition, an in-depth analysis of the remaining challenges and limitations will be provided, and proposals for promising future directions in each of the aforementioned areas. The increasing relevance of AI in drug discovery and development is reflected by the growing number of start-up companies specialized in this field, the growing number of collaborations from pharma with AI platforms, and the high number of articles and reviews reporting current applications, their success and limitations. In this chapter, we focused on the recent data-driven based research trends of the fields that are effectively cost reducible with AI. This review will address the stages of drug discovery and development in which the application of AI and ML modelling has altered the traditional development of drugs.



# Current Trends in Multidisciplinary Sciences Volume - 9

Dr. S. Neelufar Shama

CURRENT TRENDS IN MULTIDISCIPLINARY SCIENCES VOLUME - 9

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# INDEX

S. No.	Chapter Name	Page No.
1	<b>Pharmacogenomics – An Overview</b> <i>Dr. Narender Boggula, Marati Kavitha, Santhoshipriya Dandamudi &amp; Kaveti Vamshi Sharathnath</i>	1
2	<b>Minimum Spanning Network Connectivity Problem</b> <i>Dr. P Madhu Mohan Reddy</i>	35
3	<b>Bioactive Natural Products &amp; Their Applications</b> <i>Sumit Kuma, Toshibaa and Kusumlata Goswami</i>	70
4	<b>Finite Strain Theory - Equation based on the Lagrangian &amp; the Eulerian Strain</b> <i>Dr. Sushil Kumar Pathak</i>	83
5	<b>A Logarithmic Equation of State - An Equation based on the Hencky Logarithmic Strain</b> <i>Dr. Sushil Kumar Pathak</i>	97

# PHARMACOGENOMICS – AN OVERVIEW

**Dr. Narender Boggula**

*Associate Professor, CMR College of Pharmacy,  
Kandlakoya, Medchal, Hyderabad, Telangana, India*

**Marati Kavitha**

*Assistant Professor, RBVRR Women's College of Pharmacy,  
Barkatpura, Hyderabad, Telangana, India*

**Santhoshpriya Dandamudi**

*Assistant Professor, School of Pharmacy,  
Anurag University, Venkatapur, Ghatkesar, Hyderabad,  
Telangana, India*

**Kaveti Vamshi Sharathnath**

*Associate Professor,  
Nalla Narasimha Reddy Education Society's Group of Institutions,  
School of Pharmacy, Chowdariguda, Ghatkesar,  
Telangana, India*

## **Abstract**

*The use of pharmacogenomics in clinical practice is becoming standard of care. However, due to the complex genetic makeup of pharmacogenes, not all genetic variation is currently accounted for. Pharmacogenomics is crucial for individualizing drug dosages and thereby improving drug therapy outcomes. Pharmacogenomics relies on inferred phenotypes based on known variants in pharmacogenes. Pharmacogenomics (constitute both branches i.e., pharmacology and genetic) is the study of the role of genetics in drug response. It deals with the effect of genetic variation on drug response in patient by correlating the gene expression with the pharmacokinetic parameter i.e., absorption, distribution metabolism and excretion. The Pharmacogenomics is used in the research to increase the safety and efficacy of the drug by targeting the drug at the particular site of*

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(Syllabus Prescribed by the Pharmacy Council of India)

**Dr. Khagga Bhavyasri** | Dr. Uttam Prasad Panigrahy  
Dr. M. Akiful Haque | Dr. Heena Farooqui





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### DESCRIPTION

Unit Title Page No. I Quality Assurance (QA) 1 II Organization Structure 51 III Introduction 113 IV Product Complaint 151 V Calibration 211

### CHAPTERS

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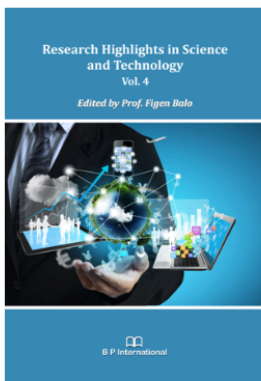


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Home / Books / Research Highlights in Science and Technology Vol. 4 / Chapters





## Hyphenated Technique - Thermogravimetry-Differential Scanning Colorimetry Coupled to Mass Spectroscopy

K. Bhavyasri

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### Abstract

A thermo-analytical approach which is known as thermogravimetric analysis uses weight changes that occur when a sample is heated to a consistent pace to ascertain a material thermal balance & percentage of the unstable additives present. DSC analysis is used for measuring a variety of quantities, including melting temperature, latent melting warmth, fusion warmth, reaction power, and many more. Mass spectrophotometry is a potent analytical tool containing numerous uses in the biomedical & pharmaceutical industries. A device's increased sensitivity and resolution have opened up new avenues for the examination of prescribed medications and complex organic structure metabolites. The chemical properties and thermal stability of plant-based polymers isolated from chemically unique soils were revealed using thermogravimetry in conjunction with DSC, quadrupole mass spectrometry, and other techniques. TGA/DSC, when used in conjunction along with MS, isolates several ions from the pattern under study in accordance with a specific mass-to-charge ratio. The linked instrument is used to evaluate the powder coating's physical characteristics and degradation balance while also identifying organic components.

**Keywords:** Mass spectroscopy; differential scanning calorimetry; thermogravimetry; quadrupole; additives; resolution

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# Hyphenated Technique - Thermogravimetry-Differential Scanning Colorimetry Coupled to Mass Spectroscopy

**K. Bhavyasri**<sup>a++</sup>

DOI:

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## ABSTRACT

A thermo-analytical approach which is known as thermogravimetric analysis uses weight changes that occur when a sample is heated to a consistent pace to ascertain a material thermal balance & percentage of the unstable additives present. DSC analysis is used for measuring a variety of quantities, including melting temperature, latent melting warmth, fusion warmth, reaction power, and many more. Mass spectrophotometry is a potent analytical tool containing numerous uses in the biomedical & pharmaceutical industries. A device's increased sensitivity and resolution have opened up new avenues for the examination of prescribed medications and complex organic structure metabolites. The chemical properties and thermal stability of plant-based polymers isolated from chemically unique soils were revealed using thermogravimetry in conjunction with DSC, quadrupole mass spectrometry, and other techniques. TGA/DSC, when used in conjunction along with MS, isolates several ions from the pattern under study in accordance with a specific mass-to-charge ratio. The linked instrument is used to evaluate the powder coating's physical characteristics and degradation balance while also identifying organic components.

*Keywords: Mass spectroscopy; differential scanning calorimetry; thermogravimetry; quadrupole; additives; resolution.*

## 1. INTRODUCTION

### 1.1 Thermogravimetric Analysis

TG analysis is a method where a mass of a substance is tracked as a key of time/temperature because the sample is put through regulated temperature software in a controlled environment.

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<sup>a</sup> Department of Pharmaceutical Analysis, RBVRR Women's College of Pharmacy, Barkatpura, Hyderabad-500027, India.

<sup>++</sup> Associate Professor and Head;

\*Corresponding author: E-mail: bhavya.khagga@gmail.com;

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# RECENT RESEARCH TRENDS IN PHARMACEUTICAL SCIENCE

Volume - 3



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## Contents

Chapters	Page No.
1. Isothermal Microcalorimetry <i>(Dr. K. Bhavya Sri, M. Nandhini and Mogili Sumakanth)</i>	01-16
2. Advances in Transungual Drug Delivery Systems: Exploring Novel Approaches for Effective Treatment of Nail Disorders <i>(Dr. Shaik Firoz and Dr. S. Prasanthi)</i>	17-39
3. Nanosponge: A Potential Nanocarrier for Novel Drug Delivery System <i>(Niyati Shah, Mamta Kumari, Piyushkumar Sadhu and Chitrali Talele)</i>	41-49
4. The Power of QSAR: Revolutionizing Drug Discovery <i>(Chitrali R Talele, Dipali R Talele, Niyati Shah, Mamta Kumari, Piyushkumar Sadhu and Chintan Aundhia)</i>	51-61
5. Nanoscale Cubosomes: Versatile Self-Assembled Carriers for Drug Delivery and Beyond <i>(Mamta Kumari, Piyushkumar Sadhu, Niyati Shah, Chitrali Talele and Chintan Aundhia)</i>	63-74
6. Edible Vaccines for Global Health: A New Frontier in Immunization <i>(Piyushkumar Sadhu, Mamta Kumari, Niyati Shah and Chitrali Talele)</i>	75-87
7. Zebra Fish: A Futuristic Model for Toxicity Studies & Novel Target for Drug Discovery <i>(Jeevan Jyoti Kaushik, Atul Kaushik, A. Sanjeeva Kumar, Filmon Debesay,</i>	89-98

# Chapter - 1

## Isothermal Microcalorimetry

Dr. K. Bhavya Sri, M. Nandhini and Mogili Sumakanth

### Abstract

Isothermal microcalorimetry (IMC) is an analytical technique for studying and assessing synthetic, physical and microorganisms responsible in real-time basis. IMC analyses the commencement, rate, extent and energetics of these phenomena for samples in tiny ampoules (e.g., 3-20 ml) at a steady specified temperature (ca. 15 °C-150 °C) over hours or days. IMC performs this dynamic simulation by recording and measuring the net rate of heat flow ( $J/sec = W$ ) to or from the specimen ampoule, in addition to the cumulative quantity of energy ( $J$ ) absorbed or created, vs. time elapsed. For four connected reasons, IMC is a powerful as well as adaptable analytical technique: All physical and chemical processes either are exothermic or endothermic, indicating they can generate or consume heat.

**Keywords:** Cumulative energy, real time basis, isothermal microcalorimetry, exothermic and endothermic

### Introduction

The technique of detecting heat generated by chemical processes as well as other physical effects is termed calorimetry. A device known as calorimeter is employed to perform calorimetry. A device known as calorimeter is employed to perform calorimetry. measure the accumulating quantity of heat ( $J$ ) that such a sample inserted inside an IMC apparatus consumes or produces under virtually fixed temperature in real-time. These alterations inside a specimen's chemical or physical component are what are causing heat. The thermal transfer varies with direct proportion to the total pace of changes occurring in any moment. The total number of aggregate changes that have taken place during a certain period of time are inversely related to the entire quantity of heat generated throughout that period. Current IMC equipment are generally semi-adiabatic, which means that heat exchange among both its specimen and also its environment is not zero (adiabatic), as IMC determination for heat transfer is dependent upon

# Recent Advances in Pharmaceutical Sciences (Volume 11)



Dr. ANJU GOYAL



<b>Chapter No.</b>	<b>Title</b>	<b>Author(s)</b>	<b>Page No.</b>
15.	<b>EFFECTS OF AYURVEDIC THERAPY FOR THE MANAGEMENT OF AUTISM IN CHILDREN: A REVIEW</b>	Nitu Sinha, Nisha Kumari Ojha	270-294
16	<b>REVIEW: HYPERTROPHIC CARDIOMYOPATHY</b>	P. Soni Dixitha, Dr. M. Vijaya bhargavi, Prof. M. Sumakanth	295-318
17	<b>ACUTE VERSUS CHRONIC HEMODYNAMIC RESPONSE OF CARVEDILOL IN CHRONIC LIVER DISEASE WITH PORTAL HYPERTENSION</b>	Shaheen Nazir, Zeeshan Ahmed Wani, Muzzafer Mir, Wasim Ahmed, Afaq Ahmad, Altaf Hussain	319-333

## **Chapter-16**

### **REVIEW: HYPERTROPHIC CARDIOMYOPATHY**

P. Soni Dixitha, **Dr. M. Vijaya bhargavi**, Prof. M.  
Sumakanth

Department of Pharmaceutical Chemistry, RBVRR women's  
college of pharmacy, Barkatpura, Hyderabad, India

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**ABSTRACT:** Hypertrophic cardiomyopathy (HCM) is a genetic disorder that is characterized by left ventricular hypertrophy unexplained by secondary causes and a nondilated left ventricle with preserved or increased ejection fraction [1]. It is commonly asymmetrical with the most severe hypertrophy involving the basal interventricular septum. The histological features of HCM include myocyte hypertrophy and disarray, as well as interstitial fibrosis. Hypertrophy is also frequently associated with left ventricular diastolic dysfunction. It is also an important cause of sudden cardiac death, particularly in adolescents and young adults. Non-sustained ventricular tachycardia, syncope, a family history of sudden cardiac death, and severe cardiac hypertrophy are major risk factors for sudden cardiac death. Atrial fibrillation is also a common complication and is not well tolerated. Mutations in over a dozen genes encoding sarcomere-associated proteins cause HCM. *MYH7* and *MYBPC3*, encoding  $\beta$ -myosin heavy chain and myosin-binding protein C, respectively, are the 2 most common genes involved.

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### **INTRODUCTION**

Hypertrophic cardiomyopathy (HCM) is a heterogeneous myocardial disease, most often caused by autosomal dominant sarcomeric gene mutations, representing the most common monogenic cardiomyopathy in humans. It is characterized by increase in left ventricular wall thickness (hypertrophy) which causes left ventricular outflow obstruction, diastolic dysfunction, myocardial ischemia, and mitral regurgitation, hyper myocardial contractility, and reduced compliance, myofibrillar disarray, and fibrosis [2,3]. These structural and function abnormalities can produce fatigue, dyspnoea, chest pain,