



R. J. COLLEGE of Arts, Science & Commerce (AUTONOMOUS)

(Hindi Vidya Prachar Samiti's RAMNIRANJAN JHUNJHUNWALA COLLEGE of Arts, Science & Commerce)

Opposite Ghatkopar Railway Station, Ghatkopar (West), Mumbai - 400 086, Maharashtra, INDIA.

Website : www.rjcollege.edu.in Email : rjcollege@rjcollege.edu.in Tel. No.: +91 22 25151763 Fax No: +91 22 25150957

College is recognized under Section 2(f) & 12(B) of the UGC Act, 1956

Affiliated to UNIVERSITY OF MUMBAI II NAAC Re-Accredited 'A' Grade (CGPA: 3.50)

First Year
Post Graduate

ADMISSION 2021-22

Click Here



PG (M.A/M.Sc/M.Com) and Post Graduate
Diploma Courses Admission 2021-22 (Started on
25th June 2021))

Post Graduate Diploma Courses Online Admission 2021-22

Admission will be on First cum First basis for Postgraduate Diploma Courses. Once the seats are full, the admission will close. No Merit List.

Important Instructions

For any Query regarding Postgraduate Diploma Course admission please come and meet Deviprasad Shetty, First Floor, Office between 11.00 to 4.00 PM

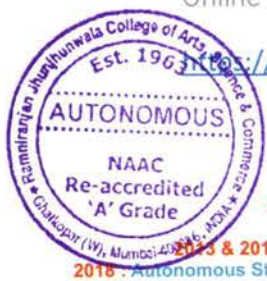
Admission Procedure Booklet

[admission21-ppt-PG-FY](#)

STEP 1: Filling of the Online admission Form

Online Registration on the portal

www.eazyinstitute.com//firstyear/#/login




PRINCIPAL
RAMNIRANJAN JHUNJHUNWALA COLLEGE
OF ARTS, SCIENCE & COMMERCE (AUTONOMOUS)
Ghatkopar (W), Mumbai-400 086, Maharashtra, INDIA

2019 : Star College Status by DBT

2008 : Best College by University of Mumbai 2010 : IMC RBNQ Award 'Performance Excellence' for the year 2009

2011 : 'Best Teacher Award' by Government of Maharashtra 2013 : DST-FIST 2014 : DBT STAR College

2013 & 2014 : 'Jagar Jaanivancha Award' by Govt. of Maharashtra 2016 : ISO 14001:2015 2016 : ISO 9001:2015 2017 : ISO 27001:2013

2016 : Autonomous Status by University Grants Commission (No. F 22-1/2018(AC) - 28.05.2018) & by University of Mumbai (No.Aff./ICD/18-19/440 - 08.06.2018)



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<https://eazyinstitute.com//firstyear/#/first-register>

Students who have passed TY in 2020-21 can use their login id and password of TY for Login <https://eazyinstitute.com//firstyear/#/login> and students of other college and previous years should register fresh on portal <https://eazyinstitute.com//firstyear/#/first-register>

1. If there are any changes in dates, the revised schedule will be notified on the website, and students are requested to abide by this and keep looking into this page for notification
2. Portal will open on 25th June 2021.
3. Admission to for PGD Programs will be based on Sem V Marks. Hence, Students applying are required to enter the Sem V marks obtained and total marks in the Educational Details Tab. If you don't have marks obtained and total marks in the marksheet get a conversion certificate from your parent college.
4. Students are required to enter the correct mobile no (WhatsApp no) and email id while filling the online form since all communication will be sent on the same email id and WhatsApp no.
5. Students have to fill all the details correctly.
6. Students of Hindi Linguistic Minority should tick on Hindi Minority Option as Yes and if you have marked Yes then only you will be considered under Hindi Linguistic Minority Quota.

STEP 2: Admission

1. Admission will be on First cum First basis for Postgraduate Diploma Courses. Once you submit your form wait for 24 hours. The fee payment tab will get activated. Once the seats are full the admission will close. No Merit List.
2. All Admission are provisional, subject to verification of documents and fulfilling the eligibility criteria for the course applied. If the documents and eligibility criteria is not fulfilled, the admission will be cancelled.
3. Your admission will be cancelled if any false information is provided for eg. Wrong entry of Sem V marks, not falling under Hindi Minority Quota, Failed or have pending ATKT in any of the semester.
4. The admission is based on merit and the College does not accept any donation for any admission.
5. The Principal decision will be final and principal reserves the right to refuse admission for non-fulfilment of eligibility criteria of UGC and University of Mumbai
6. First preference will be given to students of 2020-21 batch
7. Refund rules are available on the website for reference.

PGD in Clinical Studies Data Management and Medical Writing

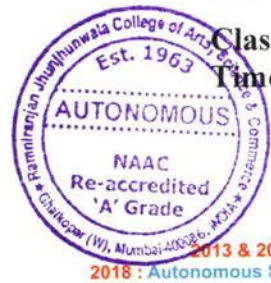
Graduates in Botany,Zoology,life Science, Microbiology,Biotechnology,

Eligibility : Biochemistry, Forensic Science, B Pharm, Medical and all allied medical graduates.

Duration : 1 Year

Class Time : Saturday & Sunday (10.00 am to 1.00.00 pm)

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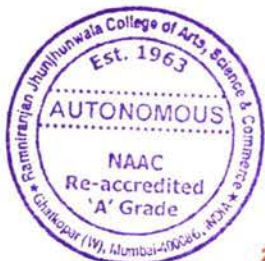
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Syllabus : [Link](#)



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Contact

diliptripathi1@yahoo.co.in

www.linkedin.com/in/dilip-tripathi-56991523 (LinkedIn)
www.vintagepoint.in (Other)

Top Skills

Analytical Chemistry
Pharmaceutics
Microbiology

Dilip Tripathi

Senior Manager R&D at Johnson & Johnson
Mumbai, Maharashtra, India

Experience

Johnson & Johnson

15 years 2 months

Senior Research Development Manager

August 2018 - Present (4 years 5 months)

India

Manager R&D

November 2007 - Present (15 years 2 months)

Mulund

FDC Limited

Asst Manager Analytical Development

May 2005 - November 2007 (2 years 7 months)

Ivax Pharmaceuticals

Executive Analytical Services

January 2003 - May 2005 (2 years 5 months)

Sandoz India Pvt Ltd

Executive

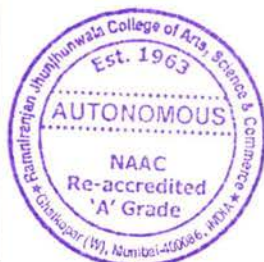
2003 - 2004 (1 year)

Education

St. Joseph High School, Colaba

SSC

St. Joseph's High School, Colaba, Mumbai




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Contact

Sunnyvale California USA
naad_archana@yahoo.co.in

www.linkedin.com/in/archana-shirpurkar-5aa1a740 (LinkedIn)
www.hvpsinternationalschool.com (Company)

Top Skills

Clinical Research
Research
Teaching

Languages

English (Full Professional)
Hindi (Full Professional)
Marathi (Full Professional)
Kannada (Professional Working)

Certifications

Project Management Program
University Teaching
Foundation of Teaching for Learning: Curriculum, Assessments

Publications

'Selective Oxidation of alcohols to carbonyl compounds with homogenous reagent system'

'A new reagent for oxidation of alcohols to carbonyl compounds (a homogenous system)'

'Phase transfer catalyzed oxidation of alcohols to carbonyl compounds'

Archana Shirpurkar

Course Coordinator, Clinical Research at Ramniranjan Jhunjhunwala College

Mumbai, Maharashtra, India

Experience

Ramniranjan Jhunjhunwala College
Course Coordinator, Clinical Research
June 2010 - Present (12 years 7 months)

Mumbai

HVPS International School

Campus Director

July 2010 - June 2012 (2 years)

Andheri, Mumbai India

Ramniranjan Jhunjhunwala College

Faculty in Chemistry

March 1993 - June 2007 (14 years 4 months)

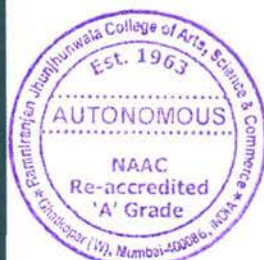
Education

Institute Of Science Mumbai

B.SC, M.SC, Ph.D., Chemistry · (1988 - 1994)

Kruger Research Center, Toronto, Ontario

CRA Certification, Clinical Research Professional Development Program




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Contact

www.linkedin.com/in/dr-amey-mane-7b11566 (LinkedIn)

Top Skills

Pharmacovigilance
Clinical Trials
Neurology

Publications

Safety and immunogenicity of single dose live attenuated Varicella vaccine (VR 795 Oka strain) in healthy Indian children: A randomized controlled study

Long-term Immunogenicity of Single Dose of Live Attenuated Hepatitis A Vaccine in Indian Children

Conducting clinical trials in emerging markets of sub-Saharan Africa: review of guidelines and resources for foreign sponsors

Changing epidemiology of hepatitis A virus in Indian children

Cross-Sectional Study for Prevalence of Non-Steroidal Anti-Inflammatory Drug-Induced Gastrointestinal, Cardiac and Renal Complications in India: Interim Report

Dr.Amey Mane

Cluster Head-Medical Affairs at Sun Pharma India
Maharashtra, India

Summary

I am a clinical pharmacologist with 16+ years of experience in multinational and Indian pharma. I have handled diverse responsibilities in medical affairs, clinical research, medical writing, ideation and pharmacovigilance. I have managed therapy shaping in different therapeutic areas like Neurology, Oncology, Cardiovascular, Diabetes, Dermatology, Vaccines, Rheumatology, Primary care and generic drugs. Stakeholder management including KOLs, regulatory authorities, internal & external customers and building up high performing teams have been the key activities of my profile.

Experience

SUN PHARMA

Cluster Head- Medical Affairs (India)
October 2022 - Present (3 months)
India

Lead and drive therapy shaping in cardiovascular, metabolics (diabetes), neurology, psychiatry, nephrology and dermatology; Lead Real World Evidence (RWE) for diverse therapy areas; Lead clinicoregulatory strategy for new drug approvals; Spearhead publication planning; Build and manage high performing teams; Collaboration with internal and external stakeholders including Thought Leaders, Key Opinion Leaders, physician associations and academic institutions

Dr. Reddy's Laboratories

Director Medical Affairs
January 2019 - October 2022 (3 years 10 months)

Head-Clinical Research, New product Ideation, Clinicoregulatory strategy & Publications for all therapeutic areas in India vertical

Accountable for the new product ideation process for NCEs/FTL, incremental innovation and differentiated products



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- Clinical trial (phase I to Phase IV) documents (Protocol, IB, ICF, CSRs, Clinical summary, CTD etc.) preparation in coordination with scientist, regulatory, DM and investigators.
- Medical Monitoring, medical advisor during conduct of trial
- SAE processing, Narratives, MSMP and Risk Management plan Preparation

UCB India Ltd.

Medical Advisor-Neuropsychiatry

June 2009 - September 2011 (2 years 4 months)

- To be a leading scientific / medical / technical resource within UCB India by acquiring scientific knowledge of CNS & Psychiatry therapeutic area.
- To collaborate with all internal and external stake holders by proactive and need based provision of medical-scientific information.
- Be an active member of the brand planning team to provide medical steer to the strategies
- Scientific and Academic liaisons with Key Opinion leaders in key therapeutic areas
- To provide medical and scientific support into the conduct of regional and national medical advisory board meetings, symposia and congresses by active participation.
- Conceptualization (wherever applicable), review and approval of promotional materials as per the local SOPs, UCB standards and local regulations.
- Identification and evaluation of new products in the area of UCB interest
- Medical training of field force, training and marketing group
- Regulatory affairs-Preparation of rationale for DCGI submissions for new products/indications, Prescribing and abbreviated prescribing information, liaison with regulatory manager for successful submissions
- Clinical research-To provide initial evaluation and continued support for clinical trials, investigator initiated studies(IIS),Non-interventional studies(NIS),Post-marketing surveillance (PMS)
- Adverse event reporting(Safety reporting) from the clinical trials & post-marketing experiences to concerned regulatory authorities in liaison with local safety officer

UCB India Ltd,

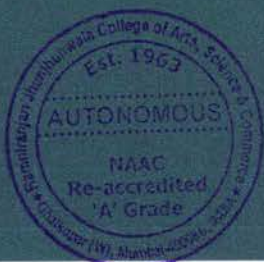
Clinical Pharmacologist

December 2008 - May 2009 (6 months)

Mumbai

- Overall responsibility for the Clinical Pharmacology aspects of any clinical program assigned to UCB India.





Contact

www.linkedin.com/in/sagar-vaidya-76b8a9b (LinkedIn)

Top Skills

PCR
GCP
CTMS

Certifications

Planning an Effective Presentation
CEO Excellence Award 2020
Ensuring Successful Presentation Delivery
Building Your Presentation

Sagar VAIDYA

Project Manager at PPD

Mumbai, Maharashtra, India

Experience

PPD

12 years 6 months

Project Manager

September 2021 - Present (1 year 4 months)

Mumbai, Maharashtra, India

Sr. Clinical Trial Manager

April 2019 - September 2021 (2 years 6 months)

Mumbai Area, India

Clinical Trial Manager

April 2016 - March 2019 (3 years)

Mumbai

Principal CRA

April 2015 - March 2016 (1 year)

Mumbai

Senior Clinical Research Associate II

April 2012 - March 2015 (3 years)

Dynasty, Kanakia Spaces, Andheri-Kurla Road, Andheri (East), Mumbai 400 059

- Lead CRA role for APAC region for Major Oncology Project of the client since Dec 2011.
- Successfully accomplished EMA inspection for Site recently in Sep 2013
- Perform and coordinate assigned aspects of the clinical monitoring process in accordance with GCPs and global SOPs to assess the safety and efficacy of investigational products and/or medical devices.
- Conduct site visits to determine protocol and regulatory compliance, and prepare required documentation.
- Develop collaborative relationships with investigative sites and client company personnel.
- Perform all duties of a CRA, serving as the primary contact with individual investigative sites that conduct clinical research for PPD



- Perform study-specific training with project team
- Develop study-specific monitoring tools and forms for use by monitoring team to assist in efficient review of study data
- Perform Serious Adverse Event reconciliation and worked with sponsor, study sites and CRAs to resolve discrepancies
- Review outstanding data reports and worked with CRAs to ensure data collection was met per contractual guidelines
- Assist sponsor, study sites, and CRAs with audit preparation/responses and quality issues
- Monitors investigator sites, with particular ability to manage complex studies and/or challenging sites, to ensure the accuracy and validity of CRF entries
- Provide mentoring and support to less experienced members of the project team and advice on training and quality issues.
- Perform and coordinate assigned aspects of the clinical monitoring process in accordance with GCPs and global SOPs to assess the safety and efficacy of investigational products.
- Conduct site visits to determine protocol and regulatory compliance, and prepared required documentation

Piramal LifeSciences Limited

Senior Executive

April 2008 - July 2010 (2 years 4 months)

Nirion Complex, Piramal Life Sciences, Western Express Highway, Goregaon

- Was responsible for Initiations, Monitoring, Closures of Phase I, II trial in Ovarian Cancer, Pancreatic Cancer & Head-Neck Cancer.
- Was Responsible for Initiations, Monitoring of Phase IV trial in obesity.
- Handled 8 clinical sites of oncology across the country.
- Was involved in Investigational Product Management at site as well as clinical trial warehouse.
- Accomplished the recruitment target as per project plan at study start up level.
- Also worked as study start up champion for different projects. Conducted Site Feasibility and Site Assessments visits in various therapeutic areas like skin diseases, oncology, diabetes etc.
- Was responsible to maintain the Study files at Sponsor as well as Investigator's Site Level
- Achieved all requirements for Monitoring visit report, follow up letters finalization.
- Completed the co-monitoring/training visits to train new Clinical Research Associates/ Clinical Trial Assistants.
- Has participated in arranging the investigator's meet for various trials.



University of Mumbai

Master of Marketing Management - MMM · (2010 - 2013)

Welingkar Institute of Management

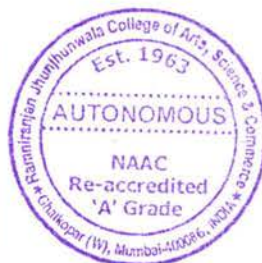
Diploma in Business Management, Basic Concepts of Management (Part Time) Weekend Course · (2006 - 2007)

Cranfield University

Master of Clinical Research, Clinical Research · (2005 - 2007)

Nagpur University

Bachelor of Pharmacy, Pharmaceutical Sciences · (2001 - 2005)




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Contact

drvishal.narang@gmail.com

www.linkedin.com/in/vishal-narang-ph-d-mba-0193245 (LinkedIn)

Top Skills

Microsoft Outlook

Clinical Research Organization functioning

Medical Writing

Certifications

Pharmacist

Publications

<https://pubmed.ncbi.nlm.nih.gov/?term=narang+vs>

Vishal Narang Ph.D., MBA

Sr.General Manager, Clinical Drug Development at Nivagen Pharmaceuticals, Inc.

Mumbai, Maharashtra, India

Summary

Ongoing:

Nivagen, a virtual pharmaceutical company, has given me this terrific opportunity to learn and contribute in holistic areas of drug development including pharmaceutical formulation, tech-transfer, manufacturing and marketing.

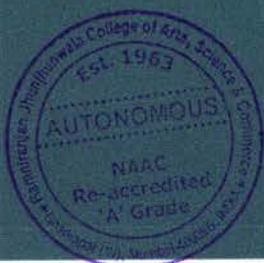
Freshly working on:

1. Developing small animal models for proof-of-concept studies
2. Validating IVIVC data that serve as gatekeepers for bioequivalence/bioavailability studies in healthy/patient volunteers
3. Authoring and reviewing multiple Pre-investigational new drug (PIND) and new drug applications (NDA), prior to submission to either critique or initiate preparations for anticipated deficiencies.
4. Leading due-diligence activities to raise funds, and
5. Strategic-, Project-, Operational- and Vendor-mangement of global clinical trials with severely complex protocols for ongoing 505b2 projects

Earlier:

Completed first ever clinical trial with narcotics in India. Consequently, developed deep understanding of workings of regulatory bodies and bureaucracy of multiple states to manage disposition of narcotic investigational products.

Played a significant role in successful completion of the largest global clinical trial in the history of Quintiles. Briefly, managing a team of~ 30 colleagues spanning a battery of activities including clinical operations, data cleaning and queries resolution management, safety reporting to DSMB.



3. Development and validation of small animal models, and other in vitro techniques to improve the predictability of success during early clinical phase studies

4. Similar responsibilities mentioned below (whilst at Cipla). In addition, seek strategic inputs from thereapeutic-KOLs, regulatory bodies whilst developing clinical trial protocols, ePRO tools, study manuals and statistical analysis plans. Provide critical inputs and facilitate technical discussions while investors conduct their due diligence.

RAMNIRANJAN JHUNJHUNWALA COLLEGE

Visiting Faculty

July 2007 - Present (15 years 6 months)

Mumbai

Design and develop clinical-staff specific courses for various study personnel
Routinely conduct work-shops and corporate training for industry and clinical trial sites.

Deliver lectures on a spectrum of clinical trial related areas.

Cipla

Director, Clinical Project Leadership and Operations

April 2018 - August 2021 (3 years 5 months)

Mumbai Area, India

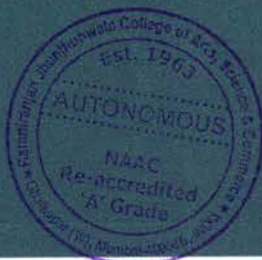
- Evaluate and recommend global and local strategic partner/s specifically in respiratory 'clinical' part of drug development
- Evaluate partners specializing in generating and analyzing data from varied sources including ePRO and eCOA.
- Manage and routinely report performance to CEO and Board of Directors for global clinical projects worth 60-75 mn USD
- Develop specific capabilities (human resources and technology) that promote better management of partners, collation of data and subsequent insights.
- Identify and establish specialized site-networks that can reduce the reliance on CRO and in-turn may provide a superior operating model.
- Inculcate a project management-driven culture, curate patient recruitment tools, and create robust processes while managing global resources that enhance sponsor's oversight during conduct of clinical trials.

IQVIA

Next Generation CLOPS Lead

March 2017 - April 2018 (1 year 2 months)

Mumbai Area, India



Ensure acceptable engagement, motivation, utilization and realization amongst team members

Identify, and represent Quintiles for new business opportunities within the Northern Region

Support project delivery at various levels

Resource allocation and management

Vendor management

Quintiles Research Private Limited

Sr.Clinical Project Manager

October 2009 - November 2012 (3 years 2 months)

All duties and responsibilities mentioned below as Clinical Team Lead.

Identify quality risks and issues associated with execution of global clinical trials (involving at least 7 countries in Asia-Pacific and Australia-New Zealand). Consequently create and execute corrective action plans to prevent or correct deficiencies in performance of the study.

Ensure utilization of the personnel are above acceptable limits of 85% and that staff is meeting defined workload through regular review of deliverables

Ensure realization of the project are above acceptable limits of 85% and that execution of deliverables is within the contracted scope of work

Diligently track Out-of-scope activities and ensure remittance for the same is received by the organization.

Evaluate the training needs of personnel, and equip them with various study tools, systems access to enable them to complete job responsibilities.

Responsible for development and execution of plethora of plans mandatory for the execution of global clinical trials as per National and International regulatory guidance.

Quintiles Research Pvt. Limited

Clinical Team Leader/Project Manager

June 2008 - October 2009 (1 year 5 months)

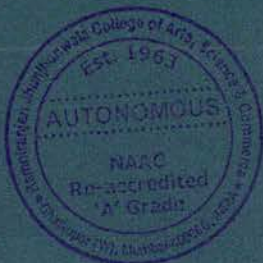
Communicate and establish customer service relationship with clinical client representative.

Routinely participate in bid-defenses with a successful track record and decent 'win: participation' ratio.

Conduct feasibility for potential studies in various indications for various clients

Work collaboratively with various functional groups and third-party vendors to support achievement of milestones.

Preparation of submission documents, resolution of queries and procurement of eventual approval from ethics committee and regulatory agencies



Cipla Ltd

Clinical Research Scientist

October 2003 - January 2005 (1 year 4 months)

Responsibilities

Design, initiate, conduct, monitor and supervise bioequivalent and clinical studies for routine submissions to US-FDA:

Instrumental in developing and medical writing of pharmacokinetic section of the ANDA.

Protocol development for submissions to various regulatory authorities such as EMEA.

Initiated 'department of clinical trials' by incorporating basic elements of SOPs and personnel-development/training and electronic data capture systems.

To design and review documents such as protocols, statistical analysis, diary cards, IB, ICFs, CRFs. Facilitate translations into local languages.

To assist in site-selection and site-development endeavors: Impart training to PIs, and development of site-SOPs.

Management of the overall conduct and operational activities of one or more clinical studies (phase III) within a project.

Univ of Washington, Seattle

Clinical Research Associate

January 2003 - January 2005 (2 years 1 month)

Routinely present clinical data and experiences at project team meetings.

Provide operational updates, plans and recommendations to the site-staff to facilitate smooth conduct of the trial.

To draft and review protocols, statistical analysis, informed consent forms, case record forms and other relevant documents pertaining to clinical research. Ensure compliance with ICH and HIPAA guidelines with adequate supervision.

Periodically assist the site in facing numerous audits by external and internal funding agencies such as NIH. Audits were found to be satisfactory by the management with no significant finding.

Keep the senior management informed of significant issues such as changes in budget/spending, and timelines for the completion of trials.

Education

University of Cincinnati




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REPORT

The PG Diploma in Clinical Studies, Data Management and Medical Writing commenced on Sunday, 5th September 2021 with an orientation about the course by Dr. Archana Shirpurkar the course coordinator. In view of the Covid pandemic the entire programme was conducted online using Zoom platform. 51 students successful completed the program. The program was conducted in two semesters semester I and Semester II. Eminent invited faculty trained the students in different aspects of Clinical studies. The eminent faculty from the industry were Dr. Vishal Narang, Dr. Amey Mane, Mr. Sagar Vaidya. From the month of April every week one session was conducted in college premises and one session was conducted online. Thus, the program later on was conducted in hybrid mode. All students had to attend all the continuous evaluation, and semester end examinations for qualifying for the PG Diploma Program.




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2018 : Autonomous Status by University Grants Commission (No. F 22-1/2018(AC) - 28.05.2018) & by University of Mumbai (No.Aff./ICD/18-19/440 - 08.06.2018)



R. J. COLLEGE of Arts, Science & Commerce (AUTONOMOUS)

(Hindi Vidya Prachar Samiti's RAMNIRANJAN JHUNJHUNWALA COLLEGE of Arts, Science & Commerce)

Opposite Ghatkopar Railway Station, Ghatkopar (West), Mumbai - 400 086, Maharashtra, INDIA.

Website : www.rjcollege.edu.in Email : rjcollege@rjcollege.edu.in Tel. No.: +91 22 25151763 Fax No: +91 22 25150957

College is recognized under Section 2(f) & 12(B) of the UGC Act. 1956

Affiliated to UNIVERSITY OF MUMBAI II NAAC Re-Accredited 'A' Grade (CGPA: 3.50)

List of Students who Completed the Program

| Sr No. | Name | Division | Roll No |
|--------|---|----------|---------|
| 1 | MAYUGHA MANILAL SINDHU | A | 1 |
| 2 | SHIVANI NARESH SUVARNA | A | 2 |
| 3 | AHIR JAYSHRI SHANKAR REKHA | A | 3 |
| 4 | ANSARI NAZIA MOHAMMED AMIN MUMTAZ | A | 4 |
| 5 | ANSARI AAYSHA MEHFUZ AKBARI | A | 5 |
| 6 | CHAUDHARI PRIYANKA PRITHVIPAL GUNINDER | A | 6 |
| 7 | DESAI HINDAVI RAVINDRA GEETA | A | 7 |
| 8 | DUBEY SONALI ASHOK KIRAN | A | 8 |
| 9 | GAVANDE JUHI SANJAY MANSI | A | 9 |
| 10 | GUPTA NIDHI SHANKARLAL SAVITADEVI | A | 10 |
| 11 | JAISWAL PANKAJ RAMRATHI NEELAM | A | 11 |
| 12 | KAMBLE CHARUSHILA MAHESH MANISHA | A | 12 |
| 13 | KESARKAR NISHA SHAMRAO KANCHANA | A | 13 |
| 14 | KHOND NEHA RAJENDRA SAVITA | A | 14 |
| 15 | MAHAJAN AKSHAYA DHANANJAY APARNA | A | 15 |
| 16 | MENON VAISHNAVI VIKRAM HEMA | A | 17 |
| 17 | MULLA JUVERIA FERAZ NUZHAT | A | 18 |
| 18 | NAIK AISHWARYA PRASAD SAMRUDHI | A | 19 |
| 19 | NARVEKAR JAGRUTI TUSHAR TEJASWI | A | 20 |
| 20 | PASI HITESH KAMLESH SAVITRI | A | 22 |
| 21 | PATIL GAURAV MANISH MITALI | A | 23 |
| 22 | PATIL MANASI KRISHNAKANT NEELAM | A | 24 |
| 23 | PILLAI DIVYA ALAGESAN SHANTI | A | 25 |
| 24 | PRAJAPATI KAJAL CHOTELAL SANGEETA | A | 26 |
| 25 | RAJANI IRAM MUNIR DILSHAD | A | 27 |
| 26 | REDEKAR SAURABH MARUTI SUNITA | A | 28 |
| 27 | SAGVEKAR ANIKET SHASHIKANT SHYAMAL | A | 29 |
| 28 | SAIYYAD SHAHZADI PARVEEN MOHD ASLAM SULTANA BEGUM | A | 30 |
| 29 | SANTOSH SHUBHAM SAWANT CHHAYA | A | 31 |
| 30 | SAWANT GOVIND YASHWANT YOGITA | A | 32 |
| 31 | SHAIKH SOPHIYA MUNAWAR RABIA | A | 33 |
| 32 | SHAIKH UZMA ATIQUE MUMTAZ | A | 34 |

PRINCIPAL

RAMNIRANJAN JHUNJHUNWALA COLLEGE OF ARTS, SCIENCE & COMMERCE (AUTONOMOUS) Ghatkopar (W), Mumbai-400 086, Maharashtra, INDIA

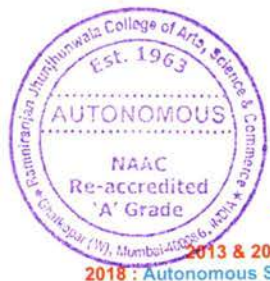
2019 : Star College Status by DBT

2008 : Best College by University of Mumbai 2010 : IMC RBNQ Award 'Performance Excellence' for the year 2009

2011 : 'Best Teacher Award' by Government of Maharashtra 2013 : DST-FIST 2014 : DBT STAR College

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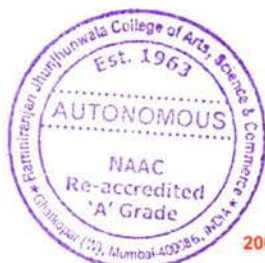
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| | | | |
|----|---|---|----|
| 33 | SHELAR ULKA PRAKASH JYOTI | A | 35 |
| 34 | SINGH HARSHITA PRAMOD VANDANA | A | 36 |
| 35 | SINGH MAHEE CHANDRABHADUR NEELAM | A | 37 |
| 36 | SINGH PRIYA HARISHCHAND SUMAN | A | 38 |
| 37 | THAKUR POOJA PRADEEP PRAJAKTA | A | 39 |
| 38 | TUNGE SHREYA JITENDRA SHAILA | A | 41 |
| 39 | TURBADKAR SANJANA MANOJ PAWAR SHOBHA PAWAR | A | 42 |
| 40 | VISHWAKARMA KIRAN SHIVKUMAR BINDU | A | 43 |
| 41 | PATEL TITHI HASMUKH BINDUBEN | A | 44 |
| 42 | DIXIT PRIYANKA RAMAKANT SAROJINI | A | 45 |
| 43 | YADAV ANKITA JAIGOVIND SEETADEVI | A | 46 |
| 44 | VISHWAKARMA RIDDHI BHARAT JAYANTI | A | 47 |
| 45 | KAMBLE KUSUM DAULU SAGUNA | A | 48 |
| 46 | KAMBLE PRANALI ARUN UJWALA | A | 49 |
| 47 | YADAV ABHINAV GHANSHYAM UMA | A | 50 |
| 48 | MORE SHRIYA MOHAN SAMIKSHA | A | 51 |
| 49 | PISAL UTKARSH NANDKUMAR MANGALA | A | 52 |
| 50 | MORAJKAR SHITAL MAHADEV SHUBBHANGI | A | 53 |
| 51 | JONDHALE MAYURI RAVI RAJARAM JONDHALE AARTI RAVI JONDHALE | A | 54 |





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Hindi Vidya Prachar Samiti's
RAMNIRANJAN JHUNJHUNWALA COLLEGE
of Arts, Science & Commerce
(Autonomous)

Affiliated to **UNIVERSITY OF MUMBAI**

CERTIFICATE OF MERIT



This is to certify that

MAYUGHA MANILAL SINDHU

has secured

'A+' Grade

in Postgraduate Diploma in Clinical Studies Data Management and Medical Writing

in the examination held in Aug 2022.

Controller of Examination

**Certified as
TRUE COPY**

Principal

**Ramniranjan Jhunjhunwala College,
Ghatkopar (W), Mumbai-400086.**

Principal

Certificate ID: PGDCSDMMW-2021-22-01

Date of Declaration: 30th Aug, 2022

Hindi Vidya Prachar Samiti's
RAMNIRANJAN JHUNJHUNWALA COLLEGE
of Arts, Science & Commerce
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Affiliated to **UNIVERSITY OF MUMBAI**

CERTIFICATE OF MERIT



This is to certify that

WALUNJ SHIVANI NARESH SUVARNA

has secured

'A' Grade

in Postgraduate Diploma in Clinical Studies Data Management and Medical Writing

in the examination held in Aug 2022.

Controller of Examination

**Certified as
TRUE COPY**

Principal

**Ramniranjan Jhunjhunwala College,
Ghatkopar (W), Mumbai-400086.**

Principal

Certificate ID: PGDCSDMMW-2021-22-02

Date of Declaration: 30th Aug, 2022