

Mainak Bardhan

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· Gender: Male · Language: English, Bengali, Hindi · Nationality: Indian

RESEARCH INTEREST

Genetics, genomics, and pathology of human diseases with a focus on precision medicine and public health.

EDUCATION

AUGUST 2024-ONGOING

PhD (DOCTOR OF PHILOSOPHY In Human Genetics and Genomics), Howard Hughes Medical Institute, University of Miami, Miller School Of Medicine, Florida, USA

- Working at intersection genetics, genomics, epigenetics, proteomics & biochemistry under the joint mentorship of Prof. Danny Reinberg, PhD, and Prof. Ramin Shiekhattar, PhD at The John P. Hussman Institute for Human Genomics (HIHG)

SEPTEMBER 2014-MAY 2020

MBBS(Bachelor of Medicine & Bachelor of Surgery), Pt. Deendayal Upadhyay Memorial Health Sciences And Ayush University Of Chhattisgarh

PT. JAWAHARLAL NEHRU MEMORIAL MEDICAL COLLEGE, RAIPUR & DR. BHIM RAO AMBEDKAR MEMORIAL HOSPITAL, INDIA

- Percentage: 69.06; CGPA 3.84
- Gold Medal And Distinction in Anatomy
- USA equivalency First professional degree in medicine (Doctor of Medicine)
<https://badges.wes.org/Evidence?i=68569991-5315-45d4-bdfb-2ff63f28b830&type=us>

WORK EXPERIENCE

22ND JUNE 2023 – 25TH JULY 2024

RESEARCH ASSOCIATE 3, MIAMI CANCER INSTITUTE, FLORIDA, USA

- Clinical outcomes research in gliomas and brain metastases, with involvement in clinical trials.

13TH SEPTMEBER 2021 – 31ST MAY 2023

SCIENTIST B(MEDICAL), ICMR-NATIONAL INSTITUTE OF CHOLERA AND ENTERIC DISEASES

- Tenure Track Faculty recruited by Indian Council of Medical Research(ICMR) Government of India, apex body of India and one of the oldest research bodies in the world.
- Working in Molecular Microbiology with a focus on Bacterial Genetics of *Campylobacter* which is one of the most common causes of Guillain-Barré Syndrome.

- Official Page <https://niced.org.in/scientists/NICEDScientists/MainakBardhan.htm>

8TH JUNE 2020-12TH SEPTEMBER 2021

SENIOR RESEARCH FELLOW, NATIONAL INSTITUTE OF MENTAL HEALTH AND NEUROSCIENCES (NIMHANS), BENGALURU, KARNATAKA INDIA

- Worked to develop the ICMR National registry for Rare and other Inherited disorders (NRROID).
- Attended neuromuscular disorder clinics and worked in the neuromuscular lab, Department of Human genetics, to establish the Phenotype genotype correlation using NGS pathology, electrophysiological studies

1ST JULY 2019-23RD AUGUST 2019

AMGEN SCHOLAR, TSINGHUA UNIVERSITY, BEIJING, CHINA

- Transporter Pharmacology & Toxicology Lab In School Of Pharmaceutical Sciences and worked on Solute Carrier Transporter and potential drug target for neurodegenerative disorder <http://tsinghuaamgenscholars.com/AmgenScholars/2019ProgramBrochure.pdf>

10TH MARCH 2019-1ST APRIL 2019

MEDICAL STUDENT OBSERVER, NATIONAL INSTITUTE OF MENTAL HEALTH AND NEUROSCIENCES (NIMHANS), BENGALURU, KARNATAKA, INDIA

- Observership in Autonomic lab assessing autonomic status in various Neurological disorders
- Observership in Human Sleep Research laboratory assisting a project on various asanas on human cognition and memory
- Observership in cell culture and stem cell biology lab on a project of ALS

OCTOBER 2018, DECEMBER 2018

MEDICAL STUDENT RESEARCHER, CSIR-IGIB, NEW DELHI, INDIA

- Gained insights on Next Generation sequencing in Clinical diagnosis and rare diseases.
- Learnt about using zebra fish as model for studying various human diseases.

MAY 2018

MEDICAL STUDENT RESEARCH TRAINEE, CSIR-CCMB, HYDERABAD, TELANGANA, INDIA

- Hands on training on various molecular techniques involved in wet lab experiments.
- Assisted a project to study role of karyotyping in the diagnosis of congenital disorders

2015-2017

MEDICAL STUDENT RESEARCHER, PT JNM MEDICAL COLLEGE, RAIPUR, CHHATTISGARH, INDIA

- Assess the relationship between various haematological factors resulting in vaso-occlusion and increased viscosity in sickle cell anaemia and osteonecrosis of femoral head(OFH).
- Sickle cell anemia and its role in growth and development in the pediatric age group.
- Serum proteome profiling in sickle cell crisis patients.

AWARDS

SCIENCE

- Certificate of Excellence, Department of Science and Technology(DST),Faculty Training program,Proteomics, Indian Institute of Technology(IIT), Bombay 2021
- ICMR Nurturing Clinical Scientists Scholarship 2020-21.
- Bursary winner XXV World Congress of Neurology (WCN 2021).
- Amgen scholars program scholarship awardee -2019.
- South Asian Medical Students' Association International conference poster presentation winner 19.
- Developing Indian Physician scientist workshop 2018 CSIR-IGIB, Newdelhi, INDIA
- Medical student research and training-2018 in CSIR-CCMB, Hyderabad
- Top 10 best oral paper presentations in the 25th International student congress of biomedical sciences 2018, UMCG, Groningen, Netherlands.
- Nobel Prize Series India, Department Of Biotechnology India,2017 Ideathon winner and delegate of the 1st Noble Prize series of India
- Indian Council of Medical Research, New Delhi, India short term studentship 2015-2016
- 17TH National Children's Science congress top 100 science project awardee.2009
- Bal Shree zonal level camp at creative scientific innovation category 2010

EDUCATION, ART, LITERATURE, LEADERSHIP

- Rhodes Scholarship Pre finalist India 2020
- Elsevier Student ambassador and mentor South Asia since 2016
- Plexus MD management internship 2017
- ISCOMS and UMCG, Groningen International student ambassador
- Leiden International Medial student conference ambassador 2017-18
- Gold medal winner in 6th International Child Art Exhibition-2008.

PUBLICATIONS

Full list of publications available through Google Scholar (**100+ publications, 7000+ citations, and h-index of 28**) at <https://scholar.google.com/citations?user=VCagZtIAAAJ&hl=en>

ORIGINAL ARTICLES

* - CORRESPONDING AUTHOR

1. **Bardhan M**, Polavarapu K, Baskar D, Preethish-Kumar V, Vengalil S, Nashi S, Ganaraja VH, Sharma D, Kulanthaivelu K, Nandeesh BN, Nalini A. **Identification Of A Novel Intronic Mutation In Vma21 Associated With A Classical Form Of X-Linked Myopathy With Autophagy.** *Global medical genetics.* 2024 May 10;11(2):167-174. doi: 10.1055/s-0044-1786815. PMID: 38736558; PMCID: PMC11087142.
2. **Bardhan M**, Anjanappa RM, Polavarapu K, Preethish-Kumar V, Vengalil S, Nashi S, Sanga S, Padmanabh H, Valasani RK, Nishadham V, Keerthipriya M, Geetha TS, Ramprasad V, Arunachal G, Thomas PT, Acharya M, Nalini A. **Clinical, genetic profile and disease progression of sarcoglycanopathies in a large cohort from india: high**

prevalence of SGCB c.544A > C. *Neurogenetics*. 2022 Apr 13. doi: 10.1007/s10048-022-00690-9. Epub ahead of print. PMID: 35416532. <https://doi.org/10.1007/s10048-022-00690-9>

3. **Bardhan M**, Polavarapu K, Bevinahalli NN, Veeramani PK, Anjanappa RM, Arunachal G, Shingavi L, Vengalil S, Nashi S, Chawla T, Nagabushana D, Mohan D, Horvath R, Nishino I, Atchayaram N. **Megaconial congenital muscular dystrophy secondary to novel CHKB mutations resemble atypical rett syndrome.** *J Hum Genet*. 2021 Mar 12. doi: 10.1038/s10038-021-00913-1. Epub ahead of print. Erratum in: J Hum Genet. 2021 Mar 26;; PMID: 33712684. <https://doi.org/10.1038/s10038-021-00913-1>
4. Polavarapu K*, **Bardhan M***, Anjanappa RM, Vengalil S, Preethish-Kumar V, Shingavi L, Chawla T, Nashi S, Mohan D, Arunachal G, Geetha TS, Ramprasad V, Nalini A. **Nemaline Rod/Cap Myopathy Due to Novel Homozygous MYPN Mutations: The First Report from South Asia and Comprehensive Literature Review.** *J Clin Neurol*. 2021 Jul;17(3):409-418. doi: 10.3988/jcn.2021.17.3.409. PMID: 34184449; PMCID: PMC8242322. <https://doi.org/10.3988/jcn.2021.17.3.409>
5. Nishadham V, **Bardhan M**, Polavarapu K, Vengalil S, Nashi S, Menon D, Ganaraja VH, Preethish-Kumar V, Valasani RK, Huddar A, Unnikrishnan GK, Thomas A, Saravanan A, Kulanthaivelu K, Nalini A, Nandeesh BN. **Thymic Lesions in Myasthenia Gravis: A Clinicopathological Study from India.** *J Neuromuscul Dis*. 2022 Apr 12. doi: 10.3233/JND-210785. Epub ahead of print. PMID: 35431258. <https://doi.org/10.3233/JND-210785>
6. Nashi S, Polavarapu K, **Bardhan M**, Anjanappa RM, Preethish-Kumar V, Vengalil S, Padmanabha H, Geetha TS, Prathyusha PV, Ramprasad V, Joshi A, Chawla T, Unnikrishnan G, Sharma P, Huddar A, Uppilli B, Thomas A, Baskar D, Mathew S, Menon D, Arunachal G, Faruq M, Thangaraj K, Nalini A. **Genotype-phenotype correlation and natural history study of dysferlinopathy: a single-centre experience from India.** *Neurogenetics*. 2023 Jan;24(1):43-53. doi: 10.1007/s10048-022-00707-3. Epub 2022 Dec 29. PMID: 36580222. <https://doi.org/10.1007/s10048-022-00707-3>
7. Nagabushana D, Polavarapu K, **Bardhan M**, Arunachal G, Gunasekaran S, Preethish-Kumar V, Anjanappa RM, Thomas P, Sadasivan A, Vengalil S, Nashi S, Chawla T, Warrier M, Keerthipriya M, Raju S, Mohan D, Nalini A. **Comparison of The Carrier Frequency of Pathogenic Variants of DMD Gene in an Indian Cohort.** *Journal of neuromuscular diseases*. 2021;8(4):525-535. doi: 10.3233/JND-210658. PMID: 33843695. <https://doi.org/10.3233/JND-210658>
8. Van Haute L, O'Connor E, Díaz-Maldonado H, Munro B, Polavarapu K, Hock DH, Arunachal G, Athanasiou-Fragkouli A, **Bardhan M**, Barth M, Bonneau D, Brunetti-Pierri N, Cappuccio G, Caruana NJ, Dominik N, Goel H, Helman G, Houlden H, Lenaers G, Mention K, Murphy D, Nandeesh B, Olimpio C, Powell CA, Preethish-Kumar V, Procaccio V, Rius R, Rebelo-Guiomar P, Simons C, Vengalil S, Zaki MS, Ziegler A, Thorburn DR, Stroud DA, Maroofian R, Christodoulou J, Gustafsson C, Nalini A, Lochmüller H, Minczuk M, Horvath R. **TEFM variants impair mitochondrial transcription causing childhood-onset neurological disease.** *Nature communications*. 2023 Feb 23;14(1):1009. doi: 10.1038/s41467-023-36277-7. PMID: 36823193. <https://doi.org/10.1038/s41467-023-36277-7>

9. Pellerin D, Danzi MC, Wilke C, Renaud M, Fazal S, Dicaire MJ, Scriba CK, Ashton C, Yanick C, Beijer D, Rebelo A, Rocca C, Jaunmuktane Z, Sonnen JA, Larivière R, Genís D, Molina Porcel L, Choquet K, Sakalla R, Provost S, Robertson R, Allard-Chamard X, Tétreault M, Reiling SJ, Nagy S, Nishadham V, Purushottam M, Vengalil S, **Bardhan M**, Nalini A, Chen Z, Mathieu J, Massie R, Chalk CH, Lafontaine AL, Evoy F, Rioux MF, Ragoussis J, Boycott KM, Dubé MP, Duquette A, Houlden H, Ravenscroft G, Laing NG, Lamont PJ, Saporta MA, Schüle R, Schöls L, La Piana R, Synofzik M, Zuchner S, Brais B. **Deep Intronic FGF14 GAA Repeat Expansion in Late-Onset Cerebellar Ataxia.** *The New England journal of medicine.* 2023 Jan 12;388(2):128-141. doi: 10.1056/NEJMoa2207406. Epub 2022 Dec 14. PMID: 36516086. <https://doi.org/10.1056/NEJMoa2207406>
10. Ganaraja VH, Polavarapu K, **Bardhan M**, Preethish-Kumar V, Leena S, Anjanappa RM, Vengalil S, Nashi S, Arunachal G, Gunasekaran S, Mohan D, Raju S, Unnikrishnan G, Huddar A, Ravikiran V, Thomas PT, Nalini A. **Disease Progression and Mutation Pattern in a Large Cohort of LGMD R1/LGMD 2A Patients from India.** *Global medical genetics.* 2021 Nov 9;9(1):34-41. doi: 10.1055/s-0041-1736567. PMID: 35169782; PMCID: PMC8837411. <https://doi.org/10.1055/s-0041-1736567>
11. Siddiqui S, Polavarapu K, **Bardhan M**, Preethish-Kumar V, Joshi A, Nashi S, Vengalil S, Raju S, Chawla T, Leena S, Mathur A, Nayak S, Mohan D, Shamim U, Prasad C, Lochmüller H, Faruq M, Nalini A. **Distinct and Recognisable Muscle MRI Pattern in a Series of Adults harbouring an Identical GMPPB Gene Mutation.** *Journal of neuromuscular diseases.* 2022;9(1):95-109. doi: 10.3233/JND-200628. PMID: 34633329. <https://doi.org/10.3233/JND-200628>
12. Huddar A, Polavarapu K, Preethish-Kumar V, **Bardhan M**, Unnikrishnan G, Nashi S, Vengalil S, Priyadarshini P, Kulanthaivelu K, Arunachal G, Lochmüller H, Nalini A. **Expanding the Phenotypic Spectrum of ECEL1-Associated Distal Arthrogryposis.** *Children (Basel).* 2021 Oct 13;8(10):909. doi: 10.3390/children8100909. PMID: 34682174; PMCID: PMC8534696. <https://doi.org/10.3390/children8100909>
13. Beijer D, Polavarapu K, Preethish-Kumar V, **Bardhan M**, Dohrn MF, Rebelo A, Züchner S, Nalini A. **Homozygous N-terminal missense variant in PLEKHG5 associated with intermediate CMT: A case report.** *Journal of neuromuscular diseases.* 2022;9(2):347-351. doi: 10.3233/JND-210716. PMID: 34897098. <https://doi.org/10.3233/JND-210716>
14. Chawla T, Preethish-Kumar V, Polavarapu K, Vengalil S, **Bardhan M**, Puri R, Verma J, Christopher R, Supriya M, Nashi S, Prasad C, Nadeesh B, Nalini A. **Late Onset Pompe Disease with Novel Mutations and Atypical Phenotypes.** *Journal of neuromuscular diseases.* 2022;9(2):261-273. doi: 10.3233/JND-210728. PMID: 34864681. <https://doi.org/10.3233/JND-210728>
15. Vengalil S, Polavarapu K, Preethish-Kumar V, Nashi S, Arunachal G, Chawla T, **Bardhan M**, Mohan D, Christopher R, Bevinahalli N, Kulanthaivelu K, Nishino I, Faruq M, Nalini A. **Mutation Spectrum of Primary Lipid Storage Myopathies.** *Annals of Indian Academy of Neurology.* 2022 Jan-Feb;25(1):106-113. doi: 10.4103/aian.aian_333_21. Epub 2022 Feb 1. PMID: 35342266; PMCID: PMC8954319. https://doi.org/10.4103/aian.aian_333_21

16. Polavarapu K, Mathur A, Joshi A, Nashi S, Preethish-Kumar V, **Bardhan M**, Sharma P, Parveen S, Seth M, Vengalil S, Chawla T, Shingavi L, Shamim U, Nayak S, Vivekanand A, Töpf A, Roos A, Horvath R, Lochmüller H, Nandeesh B, Arunachal G, Nalini A, Faruq M. **A founder mutation in the GMPPB gene [c.1000G > A (p.Asp334Asn)] causes a mild form of limb-girdle muscular dystrophy/congenital myasthenic syndrome (LGMD/CMS) in South Indian patients.** *Neurogenetics*. 2021 Oct;22(4):271-285. doi: 10.1007/s10048-021-00658-1. Epub 2021 Aug 1. PMID: 34333724. <https://doi.org/10.1007/s10048-021-00658-1>
17. Sanga S, Ghosh A, Kumar K, Polavarapu K, Preethish-Kumar V, Vengalil S, Nashi S, **Bardhan M**, Arunachal G, Raju S, Gayathri N, Biswas NK, Chakrabarti S, Nalini A, Roy S, Acharya M. **Whole-exome analyses of congenital muscular dystrophy and congenital myopathy patients from India reveal a wide spectrum of known and novel mutations.** *European journal of neurology*. 2021 Mar;28(3):992-1003. doi: 10.1111/ene.14616. Epub 2020 Nov 26. PMID: 33124102. <https://doi.org/10.1111/ene.14616>
18. Polavarapu K, Vengalil S, Preethish-Kumar V, Arunachal G, Nashi S, Mohan D, Chawla T, **Bardhan M**, Nandeesh B, Gupta P, Gowda VK, Lochmüller H, Nalini A. **Recessive VAMP1 mutations associated with severe congenital myasthenic syndromes - A recognizable clinical phenotype.** *European journal of paediatric neurology : EJPN : official journal of the European Paediatric Neurology Society*. 2021 Mar;31:54-60. doi: 10.1016/j.ejpn.2021.02.005. Epub 2021 Feb 16. PMID: 33631708. <https://doi.org/10.1016/j.ejpn.2021.02.005>
19. Rajula RR, Saini J, Unnikrishnan G, Vengalil S, Nashi S, **Bardhan M**, Huddar A, Chawla T, Sindhu DM, Ganaraja VH, Polavarapu K, Preethish-Kumar V, Kandavel T, Sathyaprabha TN, Nalini A. **Diaphragmatic ultrasound: Prospects as a tool to assess respiratory muscle involvement in amyotrophic lateral sclerosis.** *Journal of clinical ultrasound : JCU*. 2022 Jan;50(1):131-135. doi: 10.1002/jcu.23069. Epub 2021 Oct 5. PMID: 34609007. <https://doi.org/10.1002/jcu.23069>
20. Rajula RR, Saini J, Unnikrishnan G, Vengalil S, Nashi S, **Bardhan M**, Huddar A, Chawla T, Sindhu DM, Ganaraja VH, Polavarapu K, Preethish-Kumar V, Kandavel T, Nalini A. **Muscle ultrasonography in detecting fasciculations: A noninvasive diagnostic tool for amyotrophic lateral sclerosis.** *Journal of clinical ultrasound : JCU*. 2022 Feb;50(2):286-291. doi: 10.1002/jcu.23084. Epub 2021 Oct 15. PMID: 34653263. <https://doi.org/10.1002/jcu.23084>
21. Girija MS, Tiwari R, Vengalil S, Nashi S, Preethish-Kumar V, Polavarapu K, Kulanthaivelu K, Arbind A, **Bardhan M**, Huddar A, Unnikrishnan G, Kiran VR, Chawla T, Nandeesh B, Nagaraj C, Nalini A. **PET-MRI in idiopathic inflammatory myositis: a comparative study of clinical and immunological markers with imaging findings.** *Neurol Res Pract*. 2022 Oct 10;4(1):49. doi: 10.1186/s42466-022-00213-9. PMID: 36210472; PMCID: PMC9549636.
22. Sindhu DM, Huddar A, Saini J, Vengalil S, Nashi S, **Bardhan M**, Unnikrishnan G, Rajula RR, Kandavel T, Bathala L, Visser LH, Nalini A. **Cross-Sectional Area Reference Values of**

- Nerves in the Upper and Lower Extremities using Ultrasonography in the Indian Population.** *Ann Indian Acad Neurol.* 2022 May-Jun;25(3):449-456. doi: 10.4103/aian.aian_727_21. Epub 2022 Apr 1. PMID: 35936619; PMCID: PMC9350782.
23. Nishadham V, Rao S, Saravanan A, Kulanthaivelu K, Vengalil S, Venkatappa HA, Valasani RK, **Bardhan M**, Pruti N, Nalini A, Mahadevan A. **Inflammatory myofibroblastic tumors: A short series with an emphasis on the diagnostic and therapeutic challenges.** *Clin Neuropathol.* 2023 Apr 13. doi: 10.5414/NP301540. Epub ahead of print. PMID: 37051870. <https://doi.org/10.5414/NP301540>
24. Inbaraj, G., Arjun, K., Meghana, A., Preethish-Kumar, V., John, A. P., Polavarapu, K., Nashi, S., Sekar, D., Udupa, K., Prathyusha, P. V., Prasad, K., **Bardhan, M.**, Raju, T. R., Kramer, B. W., Nalini, A., & Sathyaprabha, T. N. (2023). **Neuro-Cardio-Autonomic Modulations in Children with Duchenne Muscular Dystrophy.** *Journal of neuromuscular diseases*, 10(2), 227–238. <https://doi.org/10.3233/JND-221621>
25. Ghosh D, Chowdhury G, Samanta P, Shaw S, Deb AK, **Bardhan M**, Manna A, Miyoshi SI, Ramamurthy T, Dutta S, Mukhopadhyay AK. **Characterization of diarrhoeagenic Escherichia coli with special reference to antimicrobial resistance isolated from hospitalized diarrhoeal patients in Kolkata (2012-2019), India.** *Journal of applied microbiology*. 2022 Jun;132(6):4544-4554. doi: 10.1111/jam.15548. Epub 2022 Apr 6. PMID: 35338762. <https://doi.org/10.1111/jam.15548>
26. Thakrar V, **Bardhan M**, Chakraborty N. **Early intervention in psychosis: An analysis of the characteristics and service needs of patients over the age of 35.** *Early intervention in psychiatry*. 2022 Jun 23. doi: 10.1111/eip.13318. Epub ahead of print. PMID: 35739609. <https://doi.org/10.1111/eip.13318>
27. Hesarur N, **Bardhan M**, Taallapalli A, Nashi S, Udupi GA, Kulkarni GB. **Lichtenstein-Knorr syndrome: A rare case of ataxia with sensorineural hearing loss.** *Annals of Indian Academy of Neurology*. Sep-Oct;25(5):970-973. doi: 10.4103/aian.aian_288_22. Epub 2022 Oct 31. PMID: 36561016; PMCID: PMC9764880. https://doi.org/10.4103/aian.aian_288_22
28. Juneja K, Chauhan A, Shree T, Roy P, **Bardhan M**, Ahmad A, Pawaiya AS, Anand A. **Self-medication prevalence and associated factors among adult population in Northern India: A community-based cross-sectional study.** *SAGE open medicine*. 2024 Mar 25;12:20503121241240507. doi: 10.1177/20503121241240507. PMID: 38533200; PMCID: PMC10964435. <https://doi.org/10.1177/20503121241240507>
29. Girija MS, Menon D, Polavarapu K, Preethish-Kumar V, Vengalil S, Nashi S, Keertipriya M, **Bardhan M**, Thomas PT, Kiran VR, Nishadham V, Sadasivan A, Huddar A, Unnikrishnan GK, Inbaraj G, Krishnamurthy A, Kramer BW, Sathyaprabha TN, Nalini A. **Qualitative and Quantitative Electrocardiogram Parameters in a Large Cohort of Children with Duchenne Muscle Dystrophy in Comparison with Age-Matched Healthy Subjects: A Study from South India.** *Annals of Indian Academy of Neurology*. 2024 Jan-

- Feb;27(1):53-57. doi: 10.4103/aian.aian_989_23. Epub 2024 Feb 6. PMID: 38495238; PMCID: PMC10941898
30. Chowdhury G, Debnath F, **Bardhan M**, Deb AK, Bhuina R, Bhattacharjee S, Mondal K, Kitahara K, Miyoshi SI, Dutta S, Mukhopadhyay AK. **Foodborne Outbreak by *Salmonella enterica* Serovar Weltevreden in West Bengal, India.** *Foodborne pathogens and disease.* 2024 Apr;21(4):220-227. doi: 10.1089/fpd.2023.0064. Epub 2024 Jan 8. PMID: 38190304. <https://doi.org/10.1089/fpd.2023.0064>
31. Baskar D, Veeramani-Kumar P, Polavarapu K, Nashi S, Vengalil S, Menon D, Thomas A, Bhargava Sanka S, Muddasu Suhasini K, Huddar A, Unnikrishnan G, **Bardhan M**, Thomas PT, Manjunath N, Atchayaram N. **Clinical spectrum, biochemical profile and disease progression of Kennedy disease in an Indian cohort.** *Internal medicine journal.* 2024 Mar;54(3):455-460. doi: 10.1111/imj.16205. Epub 2023 Aug 14. PMID: 37578398. <https://doi.org/10.1111/imj.16205>
32. Vengalil S, Pruthi N, Bhat D, Uppar AM, Polavarapu K, Preethish-Kumar V, Nashi S, Rajesh S, Aswini NS, Behera BP, Vandhiyadevan GD, Prasad C, Baskar D, Kulanthaivelu K, Saravanan A, Kandavel T, Nishadham V, Huddar A, Unnikrishnan G, Thomas A, Keerthipriya MS, Sanka SB, Manjunath N, Valasani RK, **Bardhan M**, Nalini A. **Monomelic Amyotrophy/Hirayama Disease: Surgical Outcome in a Large Cohort of Indian Patients.** *World neurosurgery.* 2024 Mar;183:e88-e97. doi: 10.1016/j.wneu.2023.11.087. Epub 2023 Nov 23. PMID: 38006932. <https://doi.org/10.1016/j.wneu.2023.11.087>
33. Chawla T, Reddy N, Jankar R, Vengalil S, Polavarapu K, Arunachal G, Preethish-Kumar V, Nashi S, **Bardhan M**, Rajeshwaran J, Afsar M, Warrier M, Thomas PT, Thennarasu K, Nalini A. **Myotonic Dystrophy Type 1 (DM1): Clinical Characteristics and Disease Progression in a Large Cohort.** *Neurology India.* 2024 Jan 1;72(1):83-89. doi: 10.4103/neuroindia.NI_1432_20. Epub 2024 Feb 29. PMID: 38443007. https://doi.org/10.4103/neuroindia.NI_1432_20
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ABSTRACT PUBLICATION(selected ones)

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CONFERENCES & PRESENTATIONS

- American Academy of Neurology 2024: Oral Presentation.
- American Academy of Clinical Oncology 2024: Poster Presentation.
- 16th Asian Conference on Diarrhoeal Disease and Nutrition 16th ASCODD. November 11-13, 2022, Kolkata, India
- 19th Asian-Oceanian Myology Center Meeting, Busan Korea: Poster presentation
- Tsinghua University, Beijing-Amgen scholars Program 2019: Oral and poster presentation.
- National University Singapore Amgen Scholars symposium 2019: Poster presentation.
- South Asian Medical Students' Association International conference Kolkata 2019 poster presentation
- 25th International student congress of biomedical sciences 2018, The University Medical Center Groningen, Netherlands-Oral Presentation.
- Armed Forces Medical College UG research conference 2018, India-Oral presentation

MEMBERSHIPS

- American Academy of Neurology
- European Academy of Neurology
- American Society of Clinical Oncology
- Sigma Xi, The Scientific Research Honor Society

CERTIFICATIONS & MOOCS

- Basics of Extracellular Vesicles by the University of California, Irvine on Coursera.
- Understanding Clinical Research: Behind Statistics by the University of Capetown on Coursera.
- Writing in the Sciences: Stanford University on Coursera.

- Whole-genome sequencing of bacterial genomes-tools and application: Technical University of Denmark(DTU) on Coursera.
- Probability and Statistics: To p or not to p By the University of London on Coursera.
- LaTeX for Everyone and Everything on Udemy.
- Python for everybody-from the University of Michigan on Coursera.
- Data Science -from The Johns Hopkins University on Coursera
- System Biology and Biotechnology: Ichsan School of Medicine at Mount Sinai on Coursera
- The Unix work Bench: The Johns Hopkins University on Coursera
- Genomic Data Science with Galaxy: The Johns Hopkins University on Coursera
- Introduction to R : Data Camp
- Intermediate R : Data Camp
- Artificial intelligence in Healthcare: Stanford University
- Fundamentals of Clinical Communication Skills: Glocal Academy UK

VOLUNTEERING

- Lions Club Raipur old age home geriatric health camps under the banner of MSAI.
- Health awareness and social camps in leprosy colony in Raipur and in The Leprosy Mission Hospital, Chandkhuri, The Leprosy Mission Trust, India
- Regular health awareness camps in nearby urban slums of Raipur, India
- Illustrator and blogger in Lexicon-The Medical Magazine and blog <http://www.lexiconin.com>
- Illustrator, blogger, marketing head of Scintilla online magazine for medical students.
- Content reviewer and student contributor in Elsevier publications

PEER-REVIEW & EDITORIAL ACTIVITIES

>200 reviews done. Invited peer-reviewer for:

Web of Science (<https://www.webofscience.com/wos/author/record/aal-3008-2020>)

- PLOS ONE
- International journal of health promotion and education
- Medical Education
- Neuroscience Informatics
- Annals of Medicine and Surgery
- BMC Health Services Research
- Clinical Case Reports
- Clinical Epidemiology and Global Health
- Children
- Clinical and Translational Neurosciences
- Current Neuropharmacology
- Journal of Clinical Ultrasound: Sonography and other Imaging Techniques (JCU)
- Journal of Neurosciences in Rural Practice
- BMC Public Health
- Clinical Neurology and Neurosurgery
- Genes
- Gene Reports
- Nutrients
- F1000 Research
- Brain Sciences

- Frontiers in Neurology
- Gene
- Global Pediatrics
- Journal of Clinical Medicine
- Journal of Clinical Neurosciences
- Neuroscience Informatics
- Annals of Internal Medicine: Clinical Cases
- BMJ Open
- Biomarkers in Medicine
- Frontiers in Human Neuroscience
- Cureus
- Gut Pathogens
- Global Medical Genetics
- Global Health Journal
- Gastrointestinal Oncology: Management and Care
- Neurobiology of Disease
- Neuroradiology
- International Journal of Health promotion and education
- Journal of Epidemiology and Global Health
- Neurology International
- Preventive Medicine Reports
- International Medical Case Reports Journal
- Infection and Drug Resistance
- Journal of Pain Research

EDITORIAL BOARD MEMBER

1. Journal: **BMC Neurology**
 - a. About: BMC Neurology is an open access, peer-reviewed journal that considers articles on all aspects of the prevention, diagnosis and management of neurological disorders, as well as related molecular genetics, pathophysiology, and epidemiology.
 - b. My Role: **Editorial Board Member:** Handling peer review process of submitted manuscripts
 - c. Website: <https://bmcneurol.biomedcentral.com/about/editorial-board>
2. Journal: **BMC Neurosciences**
 - a. About: BMC Neuroscience is an open access, peer-reviewed journal that considers articles on all aspects of neuroscience, welcoming studies that provide insight into the molecular, cellular, developmental, genetic and genomic, systems, network, cognitive and behavioral aspects of nervous system function in both health and disease.
 - b. My Role: **Editorial Board Member:** Handling peer review process of submitted manuscripts
 - c. Website: <https://bmcneurosci.biomedcentral.com/about/editorial-board>
3. Journal: **PLOS ONE**
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- b. My Role: **Academic Editor**; overseeing the peer review process for the journal, including evaluating submissions, selecting reviewers and assessing their comments, and making editorial decisions.
 - c. Website: https://journals.plos.org/plosone/static/editorial-board?ae_name=Mainak+Bardhan
4. Journal: **European Journal of Medical Research**
- a. About: European Journal of Medical Research publishes translational, preclinical and clinical research of international interest across all medical disciplines, enabling clinicians and other researchers to learn about developments and innovations within these disciplines and across the boundaries between disciplines.
 - b. My Role: **Associate Editor**: Handling peer review process of submitted manuscripts
 - c. Website: <https://eurjmedres.biomedcentral.com/about/editorial-board>
5. Journal: **Therapeutic Advances in Chronic Disease**
- a. About: Therapeutic Advances in Chronic Disease (TACD) is a peer-reviewed open-access journal that focuses on the highest quality research, reviews, and scholarly comment on pioneering efforts and innovative studies across all chronic diseases.
 - b. My Role: **Editorial Board member**: Handling peer review process of submitted manuscripts
 - c. Website: <https://journals.sagepub.com/editorial-board/TAJ>
6. Journal: **Frontiers in Neurology**
- a. About: Frontiers in Neurology is a multidisciplinary journal that addresses all areas of neurology to better understand, treat, and prevent neurological disorders.
 - b. My Role: **Review Editor**: Handling peer review process of submitted manuscripts
 - c. Website: <https://loop.frontiersin.org/people/1810255/editorial>
7. Journal: **Frontiers in Human Neuroscience**
- a. About: Frontiers in Human Neuroscience publishes research that advances our understanding of the brain mechanisms supporting cognitive and social behavior in humans, in both healthy and diseased states.
 - b. My Role: **Review Editor**: Handling peer review process of submitted manuscripts
 - c. Website: <https://loop.frontiersin.org/people/1810255/editorial>

PROFESSIONAL REFERENCES

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