

Dr. Rama Krishna Kancha

Education, Teaching & Research Experience

M.Sc.(2001) in Biotechnology, University of Calicut, India

Ph.D.(2010) in Cancer Biology, Technical University of Munich, Germany

Postdoctoral fellow at

(1) Technical University of Munich, Munich,

(2) University Medical Center Freiburg, Freiburg, and

(3) BioMedX Innovation Center, Heidelberg, Germany.

UGC-Assistant Professor (July 2014 till date) at the Molecular Medicine and Therapeutics Laboratory, CPMB, Osmania University

I/C. Director (November 2022 till date)

Research Funding & Publications

Total Funding as PI: approx. 1.25 crores(UGC-Start-Up, DST-SERB-EMR, ICMR-AdHoc)

Publications: 36

Books: 07

PhDs awarded: 01

Ph.D students/Project Assistants ongoing: 03

M.Sc/M.Tech/B.Tech dissertations completed/ongoing: 47

Training Programs conducted:

1. DBT-TSCOST Sponsored Skill Vigyan initiative Program in Telangana State "Training Program on – Training of Trainers (ToT)" (November 23-26, 2022), CPMB, Osmania University.
2. "Refresher Course in Molecular Medicine" (August 16-29, 2023) at the UGC-HRDC, Osmania University.
3. DBT-TSCOST Sponsored "Hands-on skill development training on advanced areas of life science and biotechnology for undergraduate and postgraduate faculty" (December 11-23, 2023), CPMB, Osmania University.
4. DBT-BUILDER Sponsored "Applications of computational tools to study cancer" (March 01-02, 2024), CPMB, Osmania University.

Scholarships, Fellowships & Memberships

Two-year M.Sc studentship (1999 to 2001) from the DBT, Govt. of India

Two-year Junior Research Fellowship(2002 to 2004)from the CSIR, Govt. of India

Three-month Visiting Fellow (2018-2019) at the JNCASR, Bengaluru

One-month Visiting Fellow (2021) at the Technical University of Munich, Germany

Member of American Association for Cancer Research (AACR)

Member and Ambassador of European Association for Cancer Research (EACR)

Associate Fellow of the Telangana Academy of Sciences (TAS)

Member of Indian Science Congress Association

Ten Select Publications

1. vanNoesel J, van der Ven WH, van Os TA, Kunst PW, Weegenaar J, Reinten RJ, [KanchaRK](#), Duyster J, van Noesel CJ (2013). Activating germline R776H mutation in the epidermal growth factor receptor associated with lung cancer with squamous differentiation. *Journal of Clinical Oncology*, 31(10): e161-4.
2. Akula S, Kamasani S, Sivan SK, Manga V, Vudem DR, [KanchaRK](#)(2018). Computational Analysis of Epidermal Growth Factor Receptor Mutations Predicts Differential Drug Sensitivity Profiles toward Kinase Inhibitors. *Journal of Thoracic Oncology*, 13(5): 721-726.
3. [KanchaRK](#), Peschel C, Duyster J (2011). The epidermal growth factor receptor-L861Q mutation increases kinase activity without leading to enhanced sensitivity toward epidermal growth factor receptor kinase inhibitors. *Journal of Thoracic Oncology*, 6(2): 387-92.
4. [KanchaRK](#), von Bubnoff N, Peschel C, Duyster J (2009). Functional analysis of epidermal growth factor receptor (EGFR) mutations and potential implications for EGFR targeted therapy. *Clinical Cancer Research*, 15(2): 460-7.
5. von Bubnoff N, Gorantla SH, [KanchaRK](#), Lordick F, Peschel C, Duyster J (2005). The systemic mastocytosis-specific activating cKit mutation D816V can be inhibited by the tyrosine kinase inhibitor AMN107. *Leukemia*, 19(9): 1670-1.
6. [KanchaRK](#), von Bubnoff N, Miething C, Peschel C, Götze KS, Duyster J (2008). Imatinib and leptomyacin B are effective in overcoming imatinib-resistance due to Bcr-Abl amplification and clonal evolution but not due to Bcr-Abl kinase domain mutation. *Haematologica*, 93(11): 1718-22.
7. Heidel F, Lipka DB, Mirea FK, Mahboobi S, Grundler R, [KanchaRK](#), Duyster J, Naumann M, Huber C, Böhmer FD, Fischer T (2009). Bis(1H-indol-2-yl)methanones are effective inhibitors of FLT3-ITD tyrosine kinase and partially overcome resistance to PKC412A in vitro. *British Journal of Haematology*, 144(6): 865-74.
8. [KanchaRK](#), von Bubnoff N, Duyster J (2013). Asymmetric kinase dimer formation is crucial for the activation of oncogenic EGFRvIII but not for ERBB3 phosphorylation. *Cell Communication and Signaling*, 11:39.
9. Subramanian J, Katta A, Masood A, Vudem DR, [KanchaRK](#) (2019). Emergence of ERBB2 Mutation as a Biomarker and an Actionable Target in Solid Cancers. *Oncologist*, 24(12): e1303-e1314.
10. Masood A, [Kancha RK](#), Subramanian J (2019). Epidermal growth factor receptor (EGFR) tyrosine kinase inhibitors in non-small cell lung cancer harboring uncommon EGFR mutations: Focus on afatinib. *Seminars in Oncology*, 46(3): 271-283.