

SURESH KUMAR R, PhD

Scientist B

Jubilee Centre for Medical Research

Jubilee Mission Medical College &

Research Institute, Thrissur, Kerala, India. 680 005

Mobile: +919633590310

E-mail: sureshraveendran2015@gmail.com

Valiyaveedu, Attinkuzhy,

Kazhakuttom.P.O

Thiruvananthapuram, Kerala,

India. 695 582

ACADEMIC QUALIFICATIONS

2016: Ph.D in Biotechnology (Cancer Cytogenetics) (Regional Cancer Centre, Thiruvananthapuram/ University of Kerala)

2005: M.Sc Biotechnology (University Campus, Kariavattom/ University of Kerala)

2003: B.Sc Biotechnology (St.Xaviers College, Thumba/ University of Kerala)

FELLOWSHIPS/AWARDS

2005: MSc biotechnology 5th Position, Kariavattom University campus

2007: Qualified in National Educational Testing, University Grants Commission, New Delhi

2008: Qualified in Junior Research Fellowship Examination, Indian Council of Medical Research, New Delhi

AREAS OF INTERESTS

- Cancer Genetics and Epigenetics
- Spatial organization of genome in genetic diseases

RESEARCH PUBLICATIONS

1. Vijay S, Sarojam S, **Raveendran S**, Syamala V, Leelakumari S, Narayanan G, Hariharan S. (2012). Recurrent isochromosome 21 and multiple abnormalities in a patient suspected of having acute myeloid leukemia with eosinophilic differentiation - a rare case from South India. **Chin J Cancer**, 31(1):45-50.
2. Santhi Sarojam, **Sureshkumar Raveendran**, Geetha Narayanan, Hariharan Sreedharan. (2013). Novel t(7;10)(p22;p24) along with NPM1 mutation in patient with relapsed acute myeloid leukemia. **Ann Saudi Med**, 33(6): 619-622.
3. **R Sureshkumar**, S Santhi, V Sangeetha, N Geetha, S Hariharan. (2014). Significance of

- nucleophosmin1 (NPM1) gene mutation status on acute myeloid leukaemia patients with normal karyotype in South India. **Molecular Cytogenetics**, 7 (Suppl 1):P71.
4. **Sureshkumar Raveendran**, Santhi Sarojam, Geetha Narayanan, Hariharan Sreedharan.(2014). A novel chromosomal abnormality t(9;14)(p24;q13) in B-acute lymphoblastic leukemia. **Indian J Hum Genet**, 20(1):79-81.
 5. Santhi Sarojam, Sangeetha Vijay, **Sureshkumar Raveendran**, Jayadevan Sreedharan, Geetha Narayanan, Hariharan Sreedharan. (2014). FLT3 Mutation as a Significant Prognostic Marker in de novo Acute Myeloid Leukemia Patients: Incidence, Distribution and Association with Cytogenetic Findings in a Study from South India. **Middle East Journal of Cancer**, 5(4): 185-196.
 6. **Sureshkumar Raveendran**, Santhi Sarojam, Sangeetha Vijay, Aswathy Chandran Geetha, Jayadevan Sreedharan, Geetha Narayanan, Hariharan Sreedharan. (2015). Mutation Analysis of *IDH1/2* Genes in Unselected *De novo* Acute Myeloid Leukaemia Patients in India – Identification of a Novel IDH2 Mutation. **Asian Pac J Cancer Prev**,16(9):4095-101
 7. Santhi Sarojam, **Sureshkumar Raveendran**, Sangeetha Vijay, Jayadevan Sreedharan, Geetha Narayanan, Hariharan Sreedharan.(2015). Characterization of CEBPA Mutations and Polymorphisms and their Prognostic Relevance in De Novo Acute Myeloid Leukemia Patients. **Asian Pac J Cancer Prev**,16(9):3785-92.
 8. **Sureshkumar Raveendran**, Santhi Sarojam, Sangeetha Vijay, Shruti Prem, Hariharan Sreedharan.(2015). A case report of concurrent *IDH1* and *NPM1* mutations in a Novel t(X;2)(q28;p22) in Acute Myeloid Leukaemia without maturation (AML-M1). **Malays J Med Sci**,22(5): 93-97.
 9. Sangeetha Vijay, Geetha N, Santhi sarojam, **Sureshkumar Raveendran**, Hariharan sreedharan.(2016). Enigmatic Inv(9): A case report on rare findings in hematological malignancies. **Iran Red Crescent Med J**, 18(4): e25062.

SCIENTIFIC PRESENTATIONS

- 2011:** International Symposium on Translational Research, Rajasthan, India
- 2013:** Indian Association for Cancer Research (IACR), New Delhi, India
- 2014:** Indian Society of Human Genetics (ISHG), Gujarat, India
- 2017:** Kerala Science Congress, Thiruvalla, Kerala, India