



ICMR - Centre for Research, Management and Control of Haemoglobinopathies, Chandrapur

Activities conducted at Haemoglobinopathies satellite centre

Mandate

To serve as a leader in research on haemoglobinopathies in the country and become a Centre of Excellence for Teaching, Innovative Research, Technology Development and translating this for the benefit of the patients

Inauguration of centre

Inauguration of Haemoglobinopathies
Satellite Centre at T.B, hospital campus,
Chandrapur on 25th October' 2015.

Establishment of diagnostic facility

- Complete blood count
- High performance liquid chromatography
- Liver function test, Kidney function test
- Molecular confirmation of sickle cell, α and β-thalassemia

Community screening and awareness

- 1448 villagers were screened and counselled in 10 tribal villages for haemoglobinopathies.
- Stalls were set up in 3 community and health fairs where more than 300 individuals visited the stalls.
- More than 500 students in tribal hostels and colleges were screened for haemoglobinopathies.















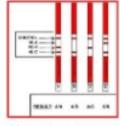




Validation of Point of care (POC) sickle cell diagnostic kits

- Sickle SCAN (POC) kit andHemotype SC (POC) kit were validated for sickle cell diagnosis by testing of 300 adults and 500 newborns.
- Results of both kits were confirmed against paper electrophoresis, HPLC and molecular diagnosis.





Sickle SCAN kit

Hemotype SC kit

Training programmes

- Training programmes are annually conducted for DMLT and MSW students.
- Training programs of POC kits and HPLC test were conducted for NHM staffs of Chandrapur, Gadchiroli and Palghar districts.





Research publications

- M Mukherjee, R Colah, et al. Multicenter Evaluation of HemoTypeSC as a Point-of-Care Sickle Cell Disease Rapid Diagnostic Test for Newborns and Adults Across India, American Journal of Clinical Pathology, Vol 153, Issue 1, Jan' 20, Pg 82–87
- P Warang, T Homma, et al. Potential involvement of ubiquitin-proteasome system dysfunction associated with oxidative stress in the pathogenesis of sickle cell disease, British journal of haematology, Vol 182, Issue 4, Aug' 18, Pg 559-566

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