

**Name:** Aathmaja D

**Age/Sex:** 8 Month/Female

**Cytogenetics lab No:** C547/2024

**Clinical History:** Microcephaly, development delay, streaky pigmentation.

**Referred by:** Dr. Sheela Nampoothiri

**Collected on:** 06/06/2024

**Received on:** 06/06/2024

**Reported on:** 13/06/2024

**Time:** 4.14pm

**Time:** 4.30pm

**Time:** 1:30pm

**Method:** Peripheral venous blood Culture

**Number of cells analyzed:** 100

**MRD No:** 2717163

**DOB:** 05/10/2023

**Nature of sample:** Peripheral venous blood

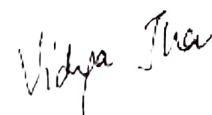


**Result**

**ISCN 2020 Karyotype:** 46,XX

**Interpretation:**

Chromosome analysis (GTG-banding) revealed an apparently normal female chromosome complement in all the cells examined.



**Dr. Vidya Jha**

Consultant (I/C), Cytogenetics

**\*END OF REPORT\***

**Limitations:**

All laboratory investigations have their limitations of sensitivity and specificity, and therefore the final diagnosis of the disease should be done in correlation with other clinical and laboratory findings.

Subtle chromosome alterations cannot be ruled out with the conventional band length of 400-450bp.

All medical reports must be viewed and reproduced as a whole.

**WISH YOU A GOOD HEALTH**