

Multilaminar chromatin model explains the structure of chromosomal aberrations in cancer cells

October 8 2015

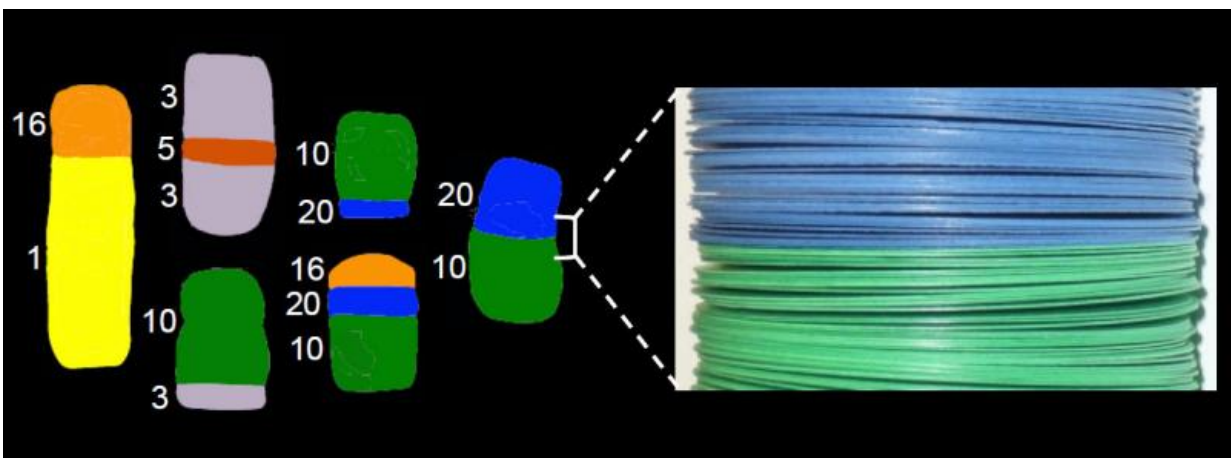


Figure: Direct relationship between the shape of translocations in cancer cells and the multilayer structure of chromosomes. The SKY images of different translocations involving chromosomes 1, 3, 5, 10, 16, and 20 in breast cancer cell line T47D were obtained from NCI-60 Drug Discovery Panel.

The organization of DNA packaged in chromosomes during cell division has been very difficult to study experimentally. Researchers from the Universitat Autònoma de Barcelona have found that the structure of translocations in cancer cells shows that chromosomes are formed by thin plates of chromatin.

During [cell division](#), each metaphase chromosome contains a single enormously long DNA molecule that is associated with histone proteins and forms a long [chromatin](#) filament with many nucleosomes. Current chromosome models consider that chromatin is folded forming loops or irregular networks. However, previous microscopy studies performed by researchers at the Chromatin Laboratory directed by professor Joan-Ramon Daban (Biosciences Unit of the Department of Biochemistry and Molecular Biology, UAB) showed unexpectedly that chromatin from metaphase [chromosomes](#) is folded into multilayered plates. This discovery led to the proposal of the thin-plate model, in which it is considered that chromosomes are formed by many stacked layers of chromatin oriented perpendicular to the chromosome axis. It was shown recently that this organization can justify the elongated cylindrical shape and mechanical properties of chromosomes.

Images obtained in many laboratories with different cytogenetic techniques have been used by Dr. Daban to investigate the internal structure of chromosomes. The orientation angle of G- and R-bands with respect to the chromosome axis has been measured; the mean values obtained for this angle range from 88° to 91°. This orthogonal orientation is observed even for the thinnest bands. The sub-bands produced by mechanical splitting of the original bands and the thin replication bands are also perpendicular to the chromosome axis. These results indicate that in three dimensions bands are disk-like structures that can be very thin and that short stretches of DNA can occupy completely the chromosome cross-section. Furthermore, in sister chromatid exchanges, the resulting connection surfaces are planar, and the observed orientation angle of these surfaces is approximately 90°. The connection surfaces observed in chromosome translocations occurring in different carcinomas and hematological malignancies are also planar; the mean values of the orientation angle of these surfaces in translocations analyzed using multicolor techniques (SKY and M-FISH) range from 90° to 92°.

All these observations impose geometric constraints that must be considered for the validation of models for chromatin organization in metaphase chromosomes. Models based on chromatin loops and irregular networks are not compatible with these constraints. The orthogonal orientation and planar geometry of the connection surfaces in chromosome translocations can be easily explained if chromosomes are formed by stacked thin layers of chromatin. This multilaminar model is also compatible with the orthogonal orientation of bands, with the existence of thin bands, and with band splitting produced by chromosome stretching.

This work has built a bridge between structural biology and cytogenetics. The results of this research are useful for the understanding of the three-dimensional organization of DNA in the chromosome and of the structural basis of the cytogenetic methods used for the diagnosis of hereditary diseases and cancers. The work has been published in the multidisciplinary journal *Scientific Reports* from Nature Publishing Group.

More information: "Joan-Ramon Daban. Stacked thin layers of metaphase chromatin explain the geometry of chromosome rearrangements and banding," *Scientific Reports* (2015). [DOI: 10.1038/srep14891](https://doi.org/10.1038/srep14891)

Provided by Universitat Autònoma de Barcelona

Citation: Multilaminar chromatin model explains the structure of chromosomal aberrations in cancer cells (2015, October 8) retrieved 24 April 2024 from <https://phys.org/news/2015-10-multilaminar-chromatin-chromosomal-aberrations-cancer.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private

study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.