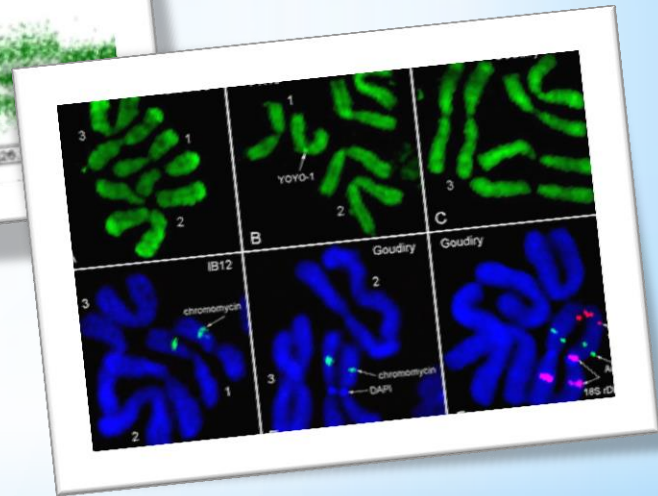
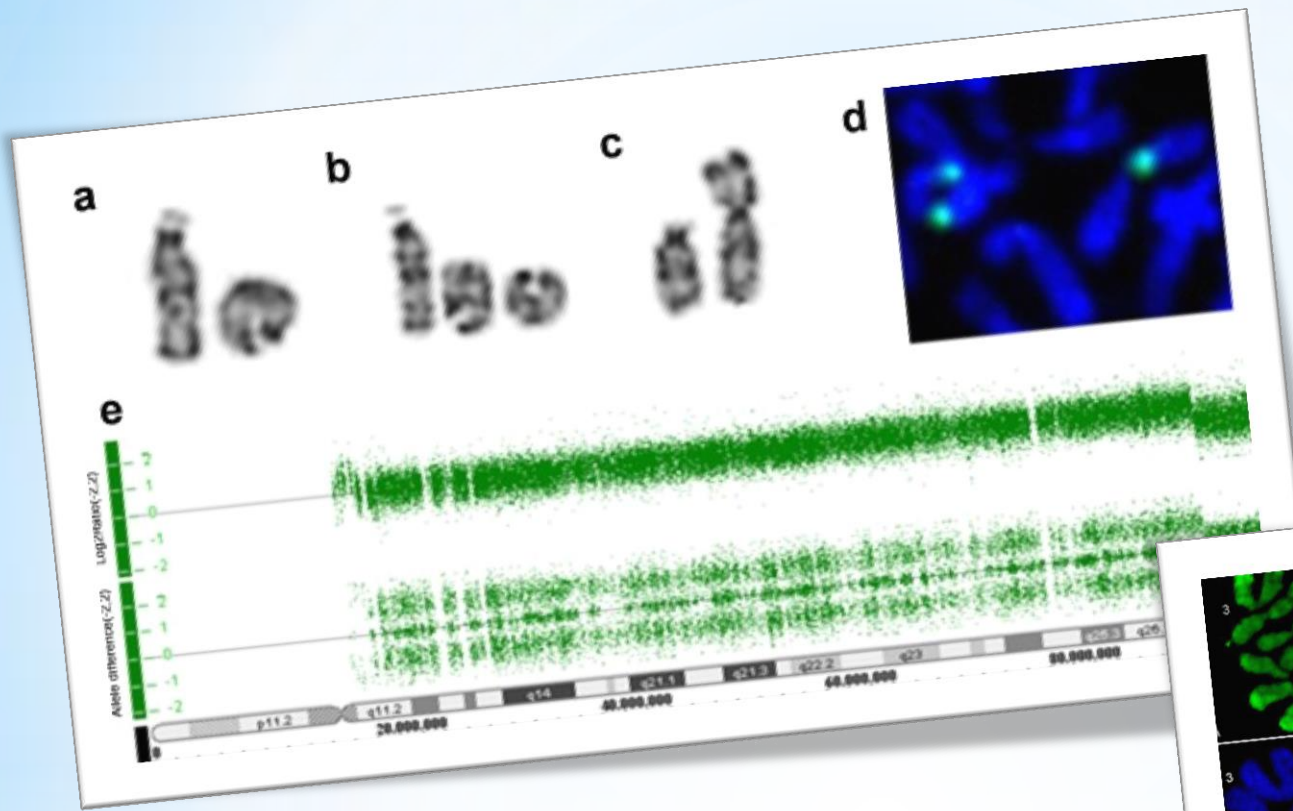


**251 bot**  
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# \*Chromosome Banding

# Basic cytogenetic examinations

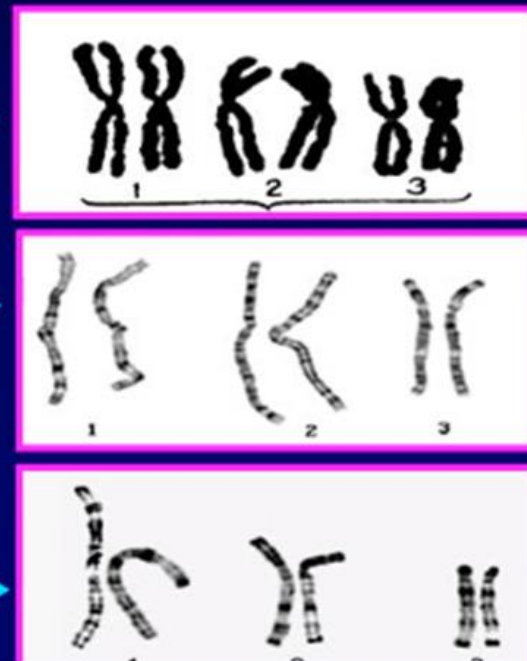
- **Interphase cells**

- Barr body (sex chromatin)



- **Metaphase cells** – staining of chromosomes

- Solid staining
- G-banding
- R-banding
- C-banding
- Q-banding
- Ag-NOR



# Classification of chromosomes banding

## Based on

- GC and AT rich regions
- Constitutive heterochromatin region
  
- Dark staining= +ve banding
- Light staining= -ve banding

## Dye used for stain

- **Giemsa dye**
- Quinacrine dye



# Number and Size of Bands

- The G-bands are black and the R-bands in white.
- Bands are numbered consecutively away from the centromere on both the short (p) and long (q) arms.
- The total number of bands or 'resolution' in the human karyotype depends on how condensed the chromosomes are, and at what stage of mitosis they are in..

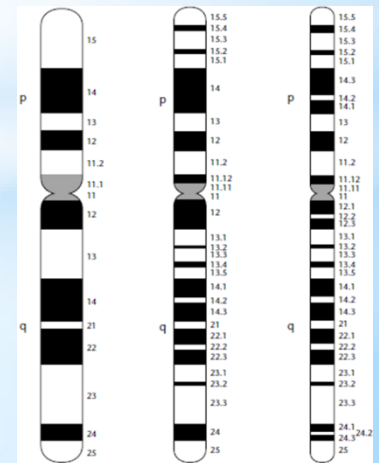


Figure 1 G-band ideograms of human chromosome 11 at (from left to right) 350, 550 and 850 band resolution.

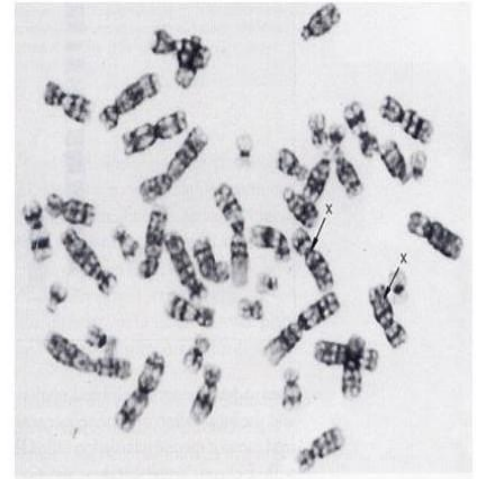
# \* Chromosome banding techniques

## G banding

- uses a stain Giemsa .
- For staining metaphase chromosomes.
- Chromosomes are pretreated with salt or proteolytic enzyme.
- Stains region of DNA rich in **Adenine** and **Thymine**
- gives light and dark stripes along the length of the chromosome

### G-Banding

G-banding of human female metaphase chromosomes



## G banding not used in Plants. Why?

- Human mitotic metaphase chromosome is 2.3 times shorter.
- Plant mitotic metaphase chromosome is 10 times more shorter than human chromosome.
- Hence difficult to demonstrate the arrangement of bands at this level with G banding technique

## R banding

- Reverse pattern of G bands.
- Pretreating cells with **hot salt solution** causing denaturation of DNA
- rich in **Adenine** and **Thymine**. Stained with Giesma stain.
- For analyzing the structure of chromosome ends.

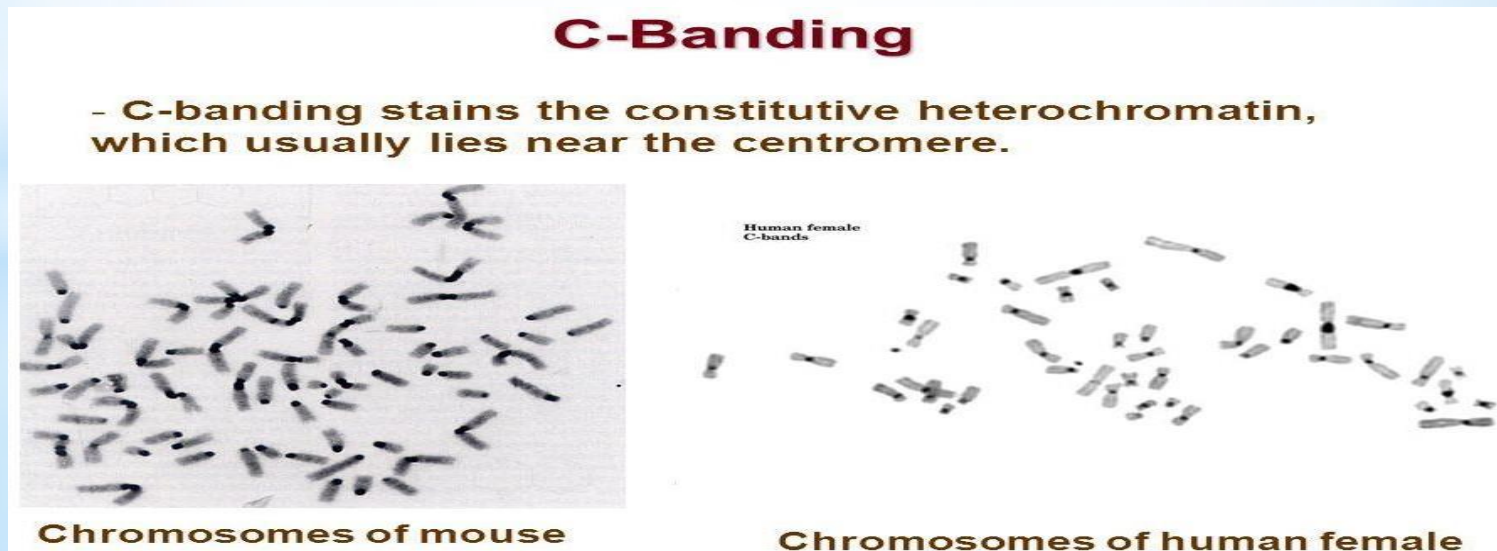
It is the opposite of C-banding.,, R-banding stains non-centromeric regions.





# C banding

- Staining only **heterochromatic regions** close to the **centromeres** and rich in satellite DNA stain
- Useful in humans for staining centromeric chromosome regions.
- A technique is presented for C-banding plant chromosomes with a modified **Wright stain**
  
- Dark banding = constitutive heterochromatic regions
- Light banding = remaining regions



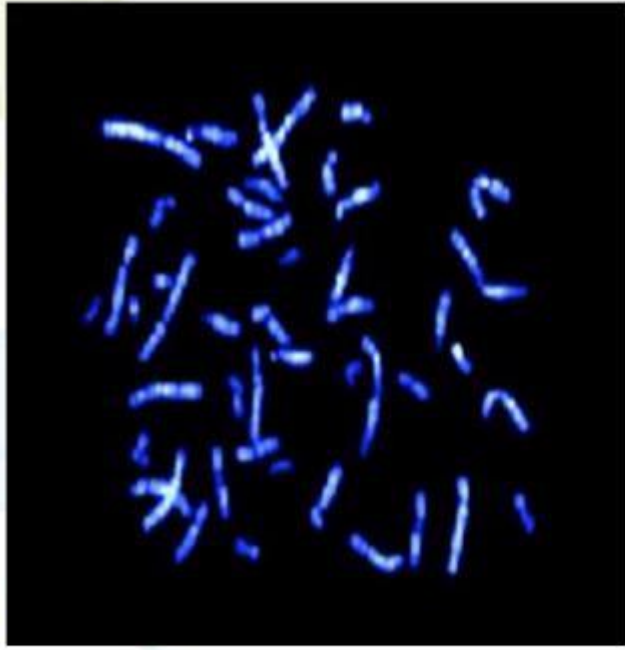
## Q-banding

- uses a stain Quinacrine
- Q-banding yields a fluorescent pattern.

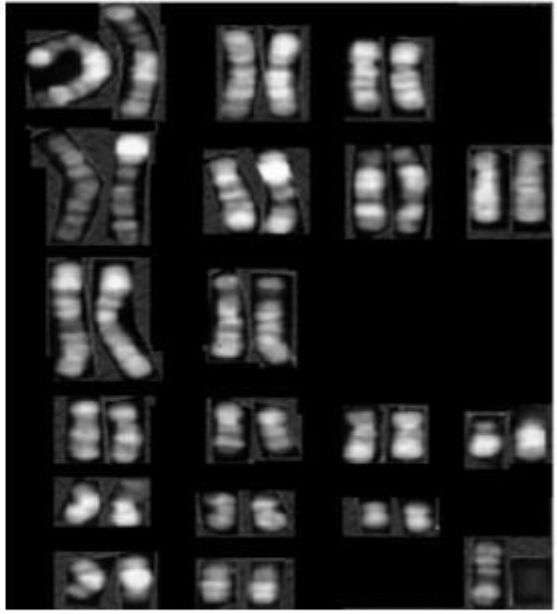
It is similar in pattern to G-banding, but glows yellow.

- Bright bands, regions of DNA rich in **Adenine** and **Thymine**.
- Dull bands, regions with **Guanine** and **Cytosine**.
- For identifying human chromosome Y and different polymorphisms.

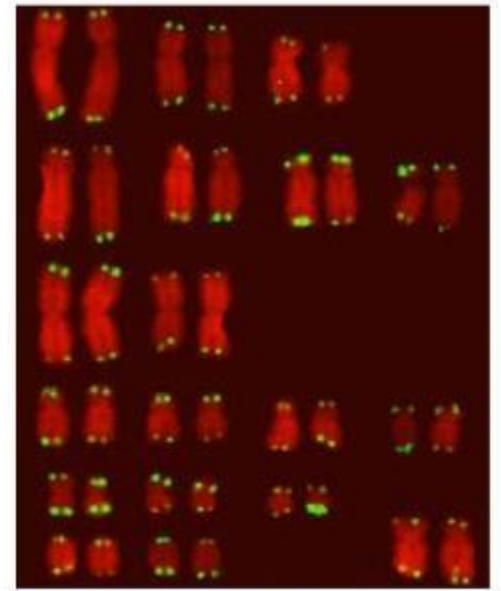




Q banding



Cat Q band



Cat telo

## O bands

- Used in plants. since in plants G, C and Q bandings have relatively limited application
- quick method using trypsin-orcein for banding plant chromosomes
- The technique is directly applicable to meristematic tissues (e.g. root tips)

## T bands

- used to stain the telomeric regions of chromosomes for cytogenetic analysis.
- By **Giemsa stain** or **Acridine orange**, after controlled thermal treatment.

## Hy banding

- For plant cells.
- Applied to somatic chromosomes of members of *Liliflorae*

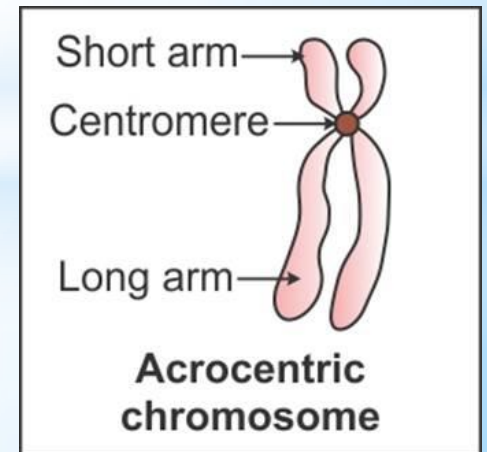


## N or NOR staining

- For nuclear organizing regions.
- Silver nitrate solution binds (secondary constrictions of **Acrocentric** chromosomes)

## CT bands

Banding of centromeric and telomeric segments

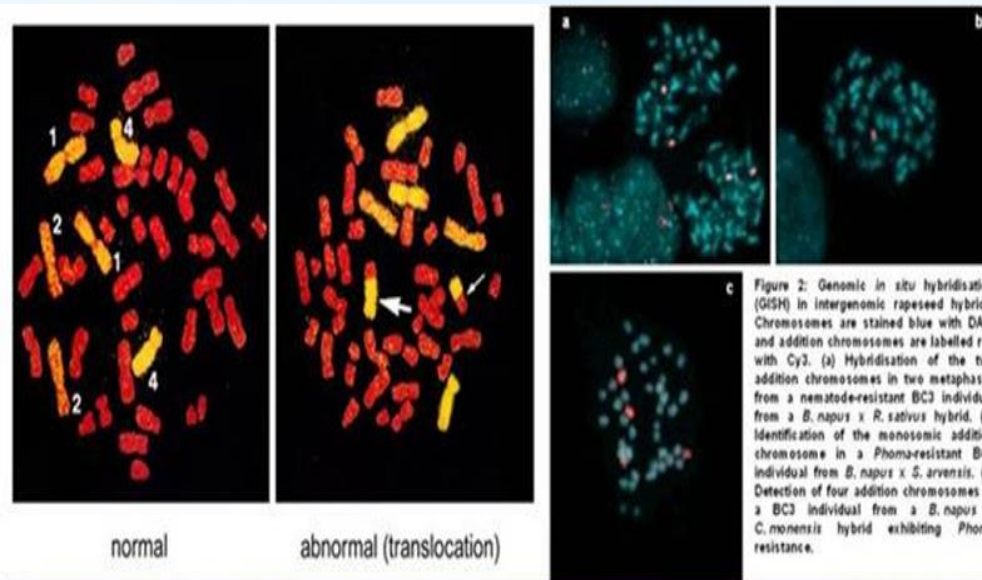


Other techniques

**In situ hybridization- ISH**

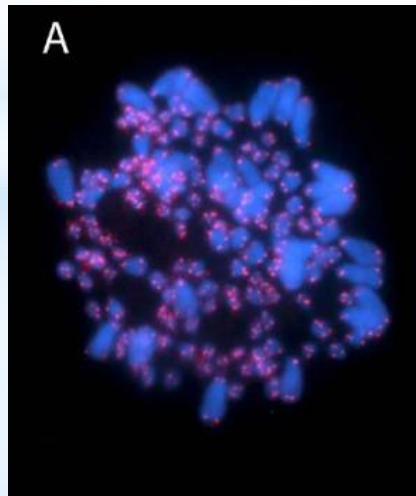
**(FISH) and (GISH).**

Use probe sequence tagged with **radioisotopes** or **fluorescent compounds**..

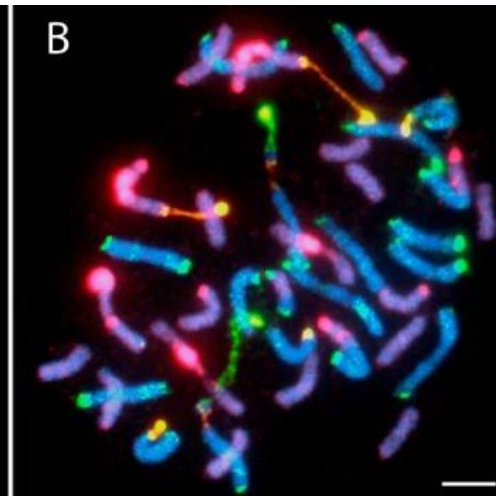


## Difference between FISH and GISH

FISH



GISH



The 9 lecture ended

# Homework

Write 3 questions for the lecture

The next lecture

