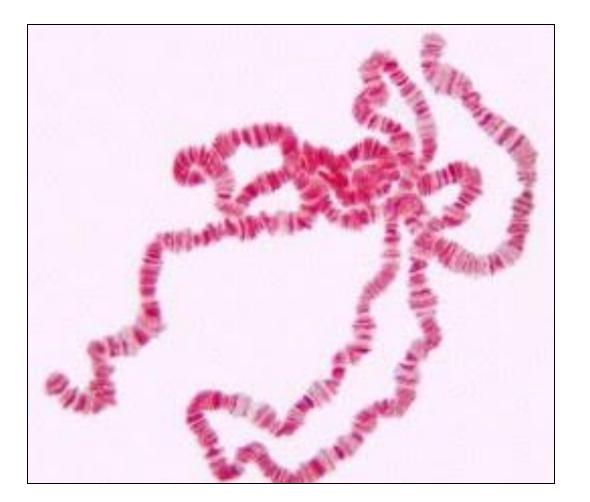
## POLYTENE AND LAMPBRUSH CHROMOSOMES

## What are these?

- ° Giant chromosomes (large sized)
- ° Can be both polytene (salivary gland chromosome) and lampbrush chromosomes



#### POLYTENE CHROMOSOMES

### INTRODUCTION

- Discovered in the salivary glands of *drosophila* (therefore called salivary gland chromosomes)
- Also present in insects like mosquitoes
- ° Kollar gave the term "polytene chromosome" because of high DNA content
- ° Also occur in rectum, gut, foot pads, fat bodies etc.

## HOW ARE THEY FORMED?

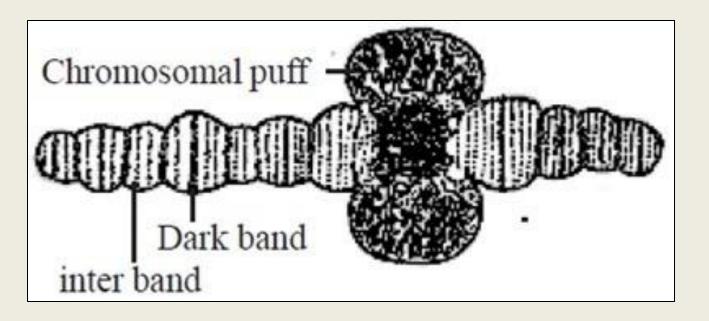
- ° Replication of chromosomal DNA several times
- ° Without nuclear division
- ° Resultant chromatids do not separate and remain joined side by side

POLYPLOIDY: excess DNA in nucleus but chromosomes separate after division

- ° Visible during prophase (of mitosis) and interphase
- Size in *drosophila*: 1000 DNA molecules long

#### HOW DO THEY LOOK LIKE?

- ° Series of dark bands (chromomere) alternate with interbands (where DNA is loosely coiled)
- Crossbanding pattern

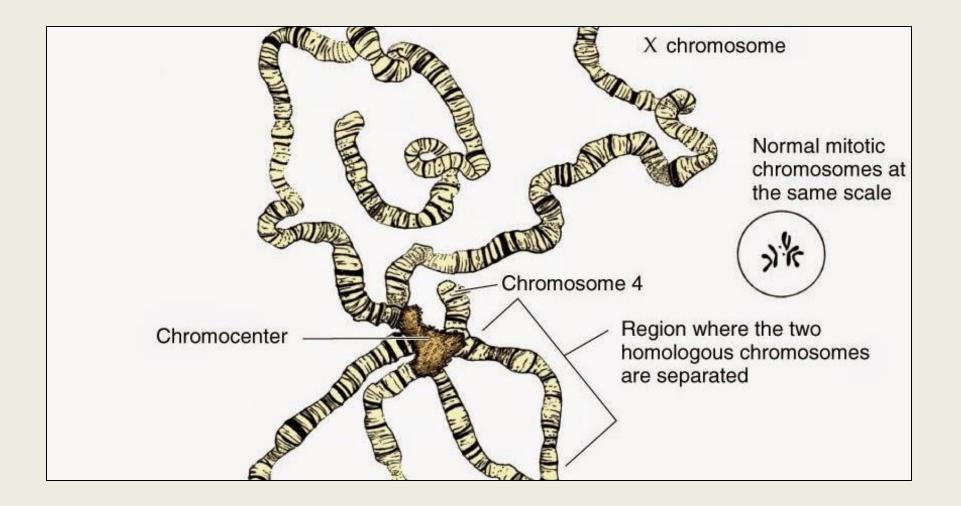


- Peculiar characteristic: maternal and paternal homologous chromosomes lie side by side →
  <u>Somatic pairing</u>
- ° In *drosophila*, heterochromatin of all polytene chromosomes--- coalesce--- <u>chromocentre</u>

#### CHROMOSOMAL PUFFS/ BALBIANI RINGS

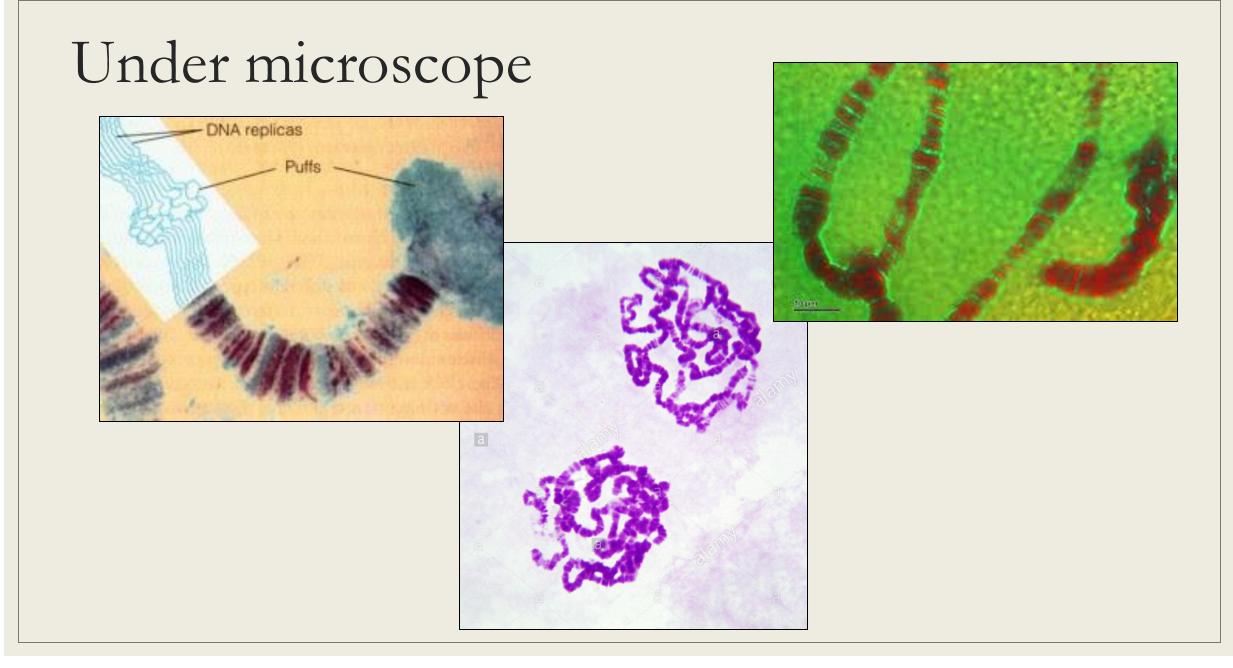
- ° Swellings of bands of polytene chromosomes
- ° Intense gene transcription area
- ° DNA unfolds --- open loops
- ° Distribution differ from one tissue to another
- ° Reversible phenomenon

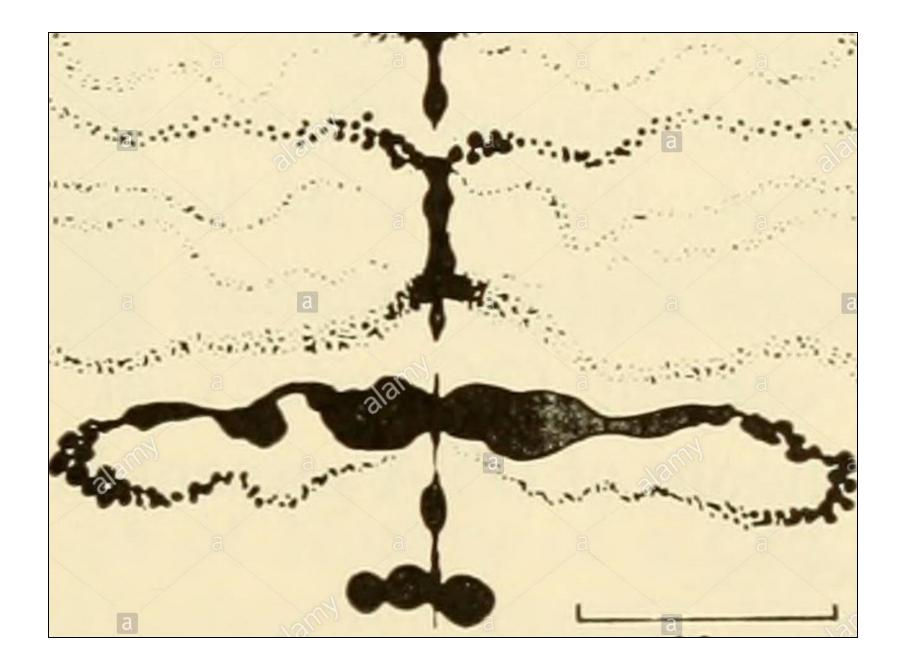




## Why are they formed? (functions)

- Possible explanation by Alberts *et all*
- Mainly three reasons:
- 1. To keep DNA organised
- 2. Isolate genes from their neighbours
- 3. Regulate gene transcription for cell differentiation
- ° Interbands: housekeeping genes
- Bands: cell-type specific genes





## LAMPBRUSH CHROMOSOMES

## Introduction

- $\circ~1^{st}\,observed$  in salamander oocytes
- ° Name--- looks like brushes used to clean old kerosene lamps
- ° Described in detail in shark oocytes
- ° Occurs at diplotene stage of prophase (meiosis I) (extended diplotene stage)
- ° Much larger/ longer than polytene chromosomes
- ° most conspicuous feature is widespread RNA transcription
- Not formed in mammals

## How are they formed?

- ° Formed during an extended diplotene stage of 1<sup>st</sup> meiotic division of female gametocyte
- The chromosomes go from a compact telophase form at the end of the last oogonial mitosis, become "lampbrushy"
- ° Then contract again to form normal first meiotic metaphase bivalents

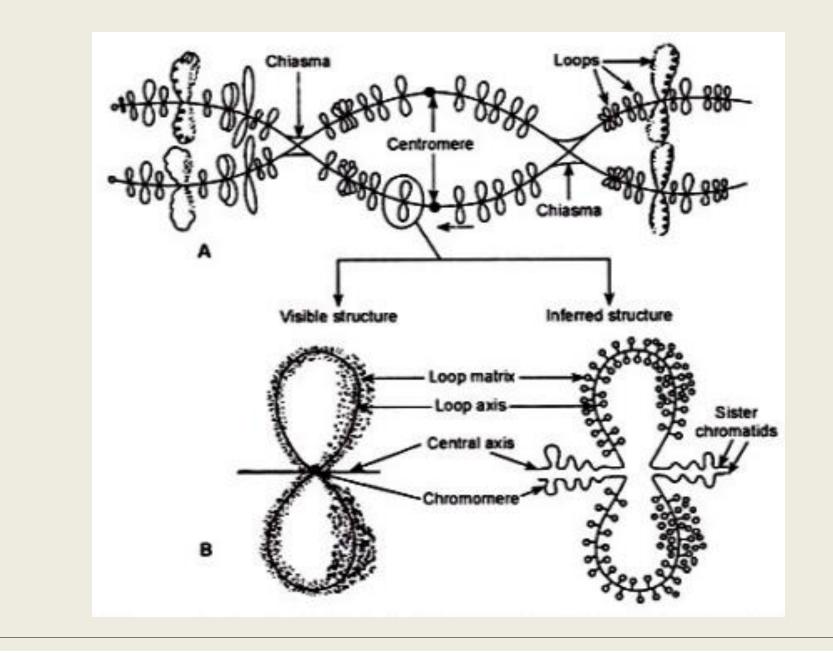
Leptotene  $\rightarrow$  zygotene  $\rightarrow$  pachytene  $\rightarrow$  **Diplotene** 

## How do they look?

- Brush like (RNA polymerase + nascent RNA + proteins)
- Bivalent (2 homologous chromosomes) held together by chiasmata
- ° Length can be up to 1mm

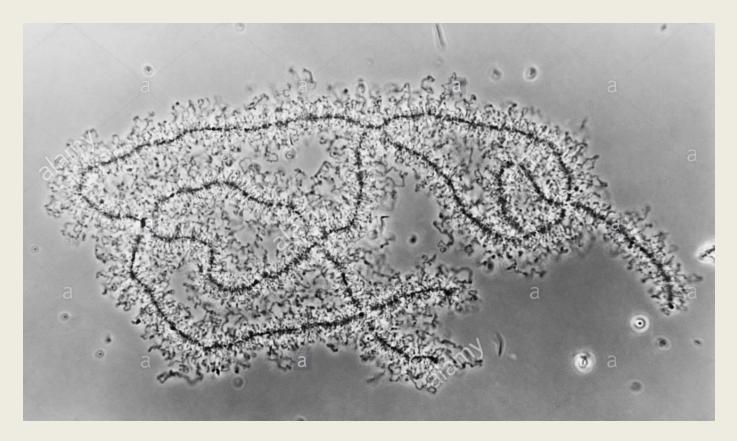
LOOPS:

- Always symmetrical
- $\circ\,$  Not formed at centromere
- About 10,000 loops in each chromosome set
- Each loop has an axis (made of DNA  $\rightarrow$  RNA polymerase present here)



#### FUNCTION OF THE LOOPS

- Each loop = intense transcription or hn RNA (heterogenous nuclear RNA)
- $\circ$  Proteins attached to RNA  $\rightarrow$  ultimately released as ribonucleoproteins
- $\circ$  Loops may be static or dynamic (spinning out and retraction hypothesis)  $\rightarrow$  rejected



# Hypothesis regarding loops of lampbrush chromosomes

#### 1. <u>Spinning out and retraction hypothesis (Gall and Callan)</u>

In this, new material "spin out" of one side of the chromosome forming a loop and retract on the other side.

Meaning all genome can be expressed

Rejected

#### 2. Master and slave hypothesis (Callan and Lloyd)

One loop contain a number of copies of one gene out of which one is the master copy Information is transferred from it to the 'slave' copies Slave copies  $\rightarrow$  transcribes to RNA

Many copies means higher rate of RNA synthesis is possible

## Functions of lampbrush chromosome

 $\circ\,$  Involved in the production of mRNAs for early development

- ° Giant granular loops could either be the sites where such mRNAs are packaged
- ° Could be sites where specific alterations of the deoxyribonucleoprotein fiber take place.
- Gene amplification  $\rightarrow$  required during growth phase of oocytes

# THANK YOU