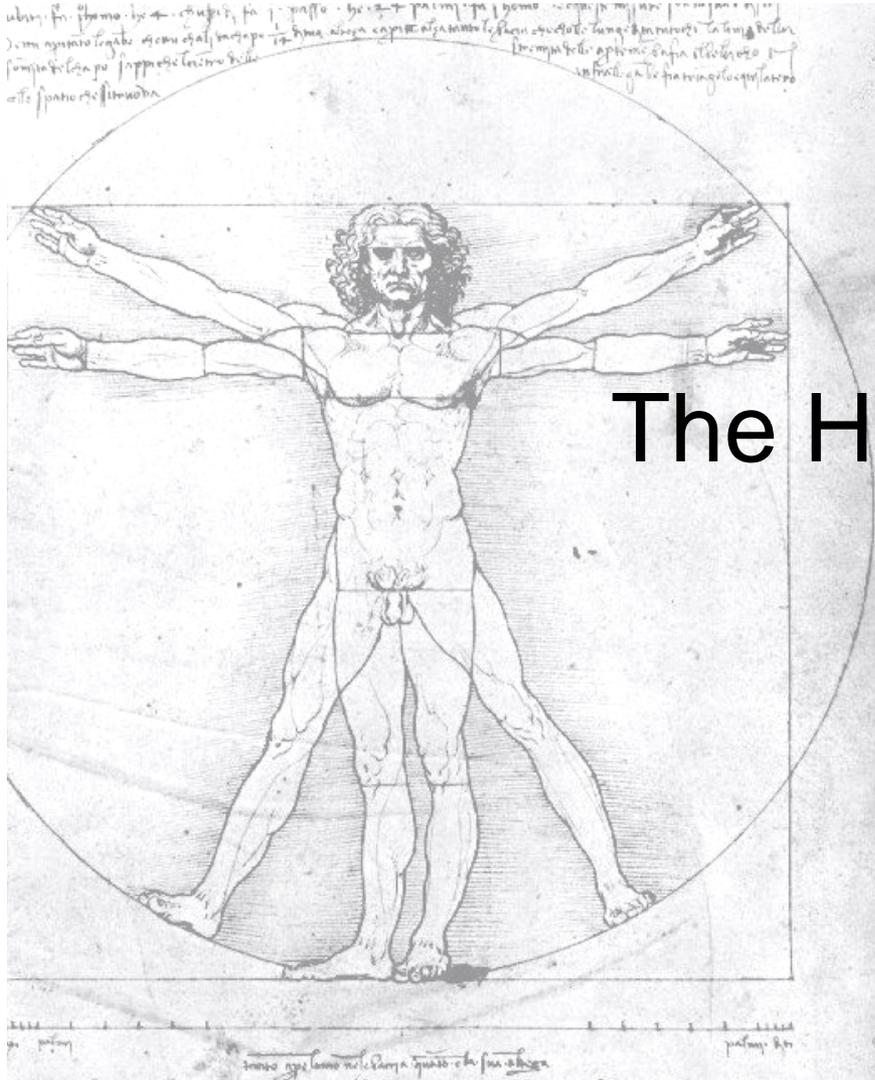


A C G T



The Human Genome and its upcoming Dynamics

Matthias Platzer

Genome Analysis
Leibniz Institute for Age Research
- Fritz-Lipmann Institute (FLI)

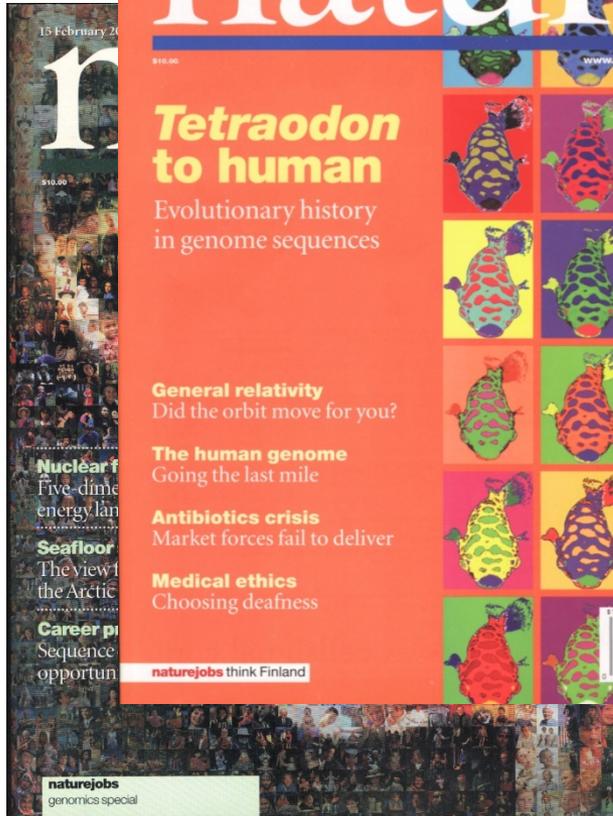
Sequencing of the Human Genome

Publications

2004



2001



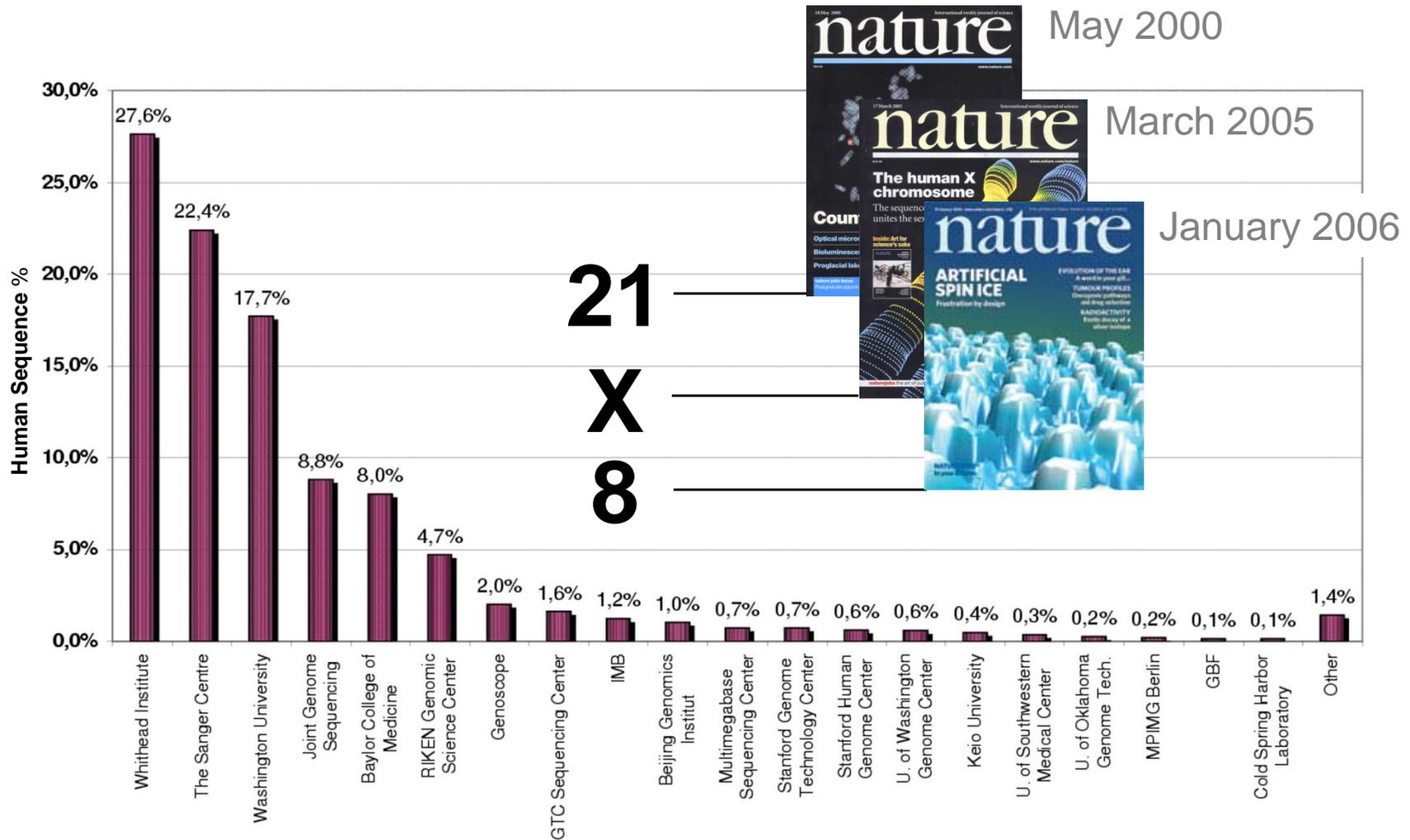
2001



International academic
consortium

International Human Genome Project

Contributions of the Members



Human Genome

Working Draft versions February 2001



Academic

Initial Sequencing & Analysis...

Private



The Sequence of ...

2.72 Gb	Sequenced Bases	2.65 Gb
1,000	Clone gaps	54,000
146,000	Sequence gaps	116,000
147,000	Gaps	170,000

overall coverage: **94%**
quality of unfinished data : **< 1 error/10kb in 91%**

human population heterogeneity: **1 SNP/kb**
variation between two individuals: **1 SNP/10kb**

Human Genome

Final version October 2004



Initial Seq...



Finishing the euchromatic sequence...

Academic

Private



The Sequence of ...

2.72 Gb	2.85 Gb	Sequenced Bases	2.65 Gb
1,000	283	Clone gaps	54,000
146,000	58	Sequence gaps	116,000
147,000	341	Gaps	170,000

near-complete sequence: **99%** of euchromatin
 extremely high quality: **< 1 error/100kb**

Segmental Duplications

Problems of the human reference sequence

~50%

of the

273 interior euchromatic gaps

located in

segmentally duplicated regions

Segmental Duplication

Definition

genomic regions **>1 kb**
with nt identity **>90%**

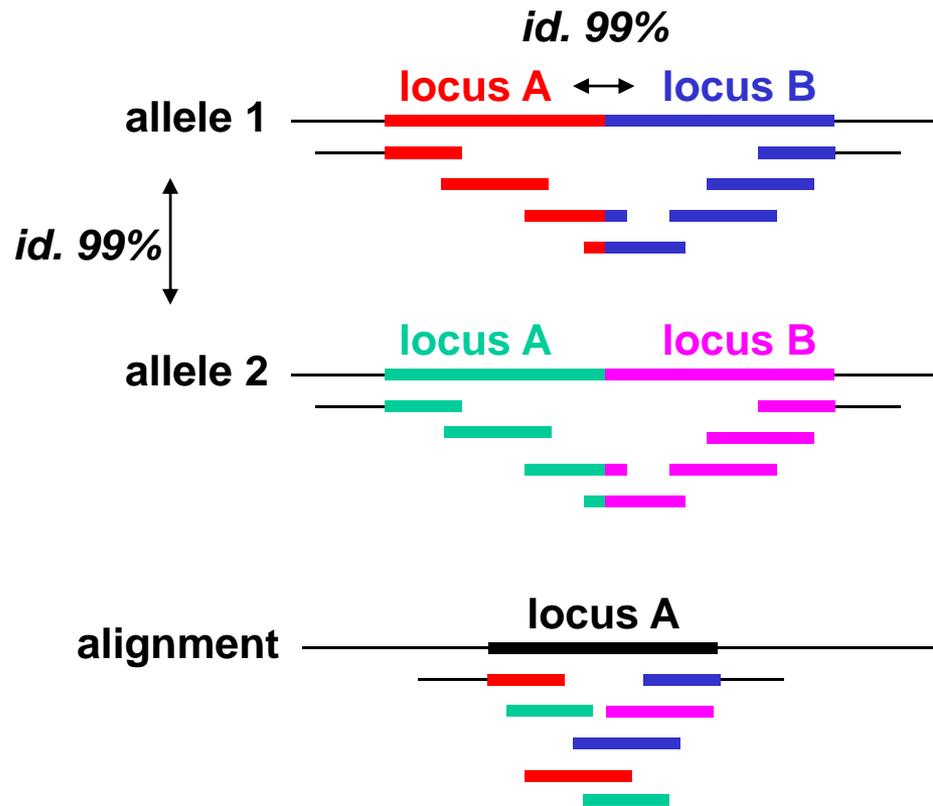
Human genome

5.3% segmentally duplicated

87% of all segmental duplications **>50 kb**

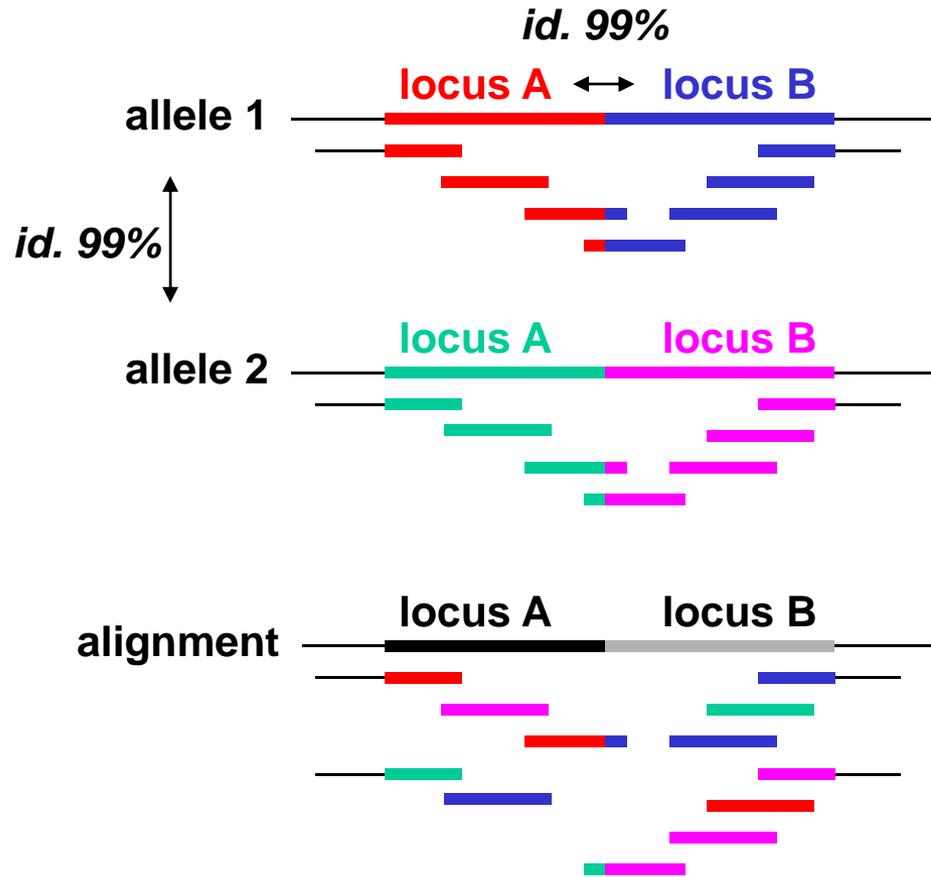
Segmental Tandem Duplications

Assembly problems



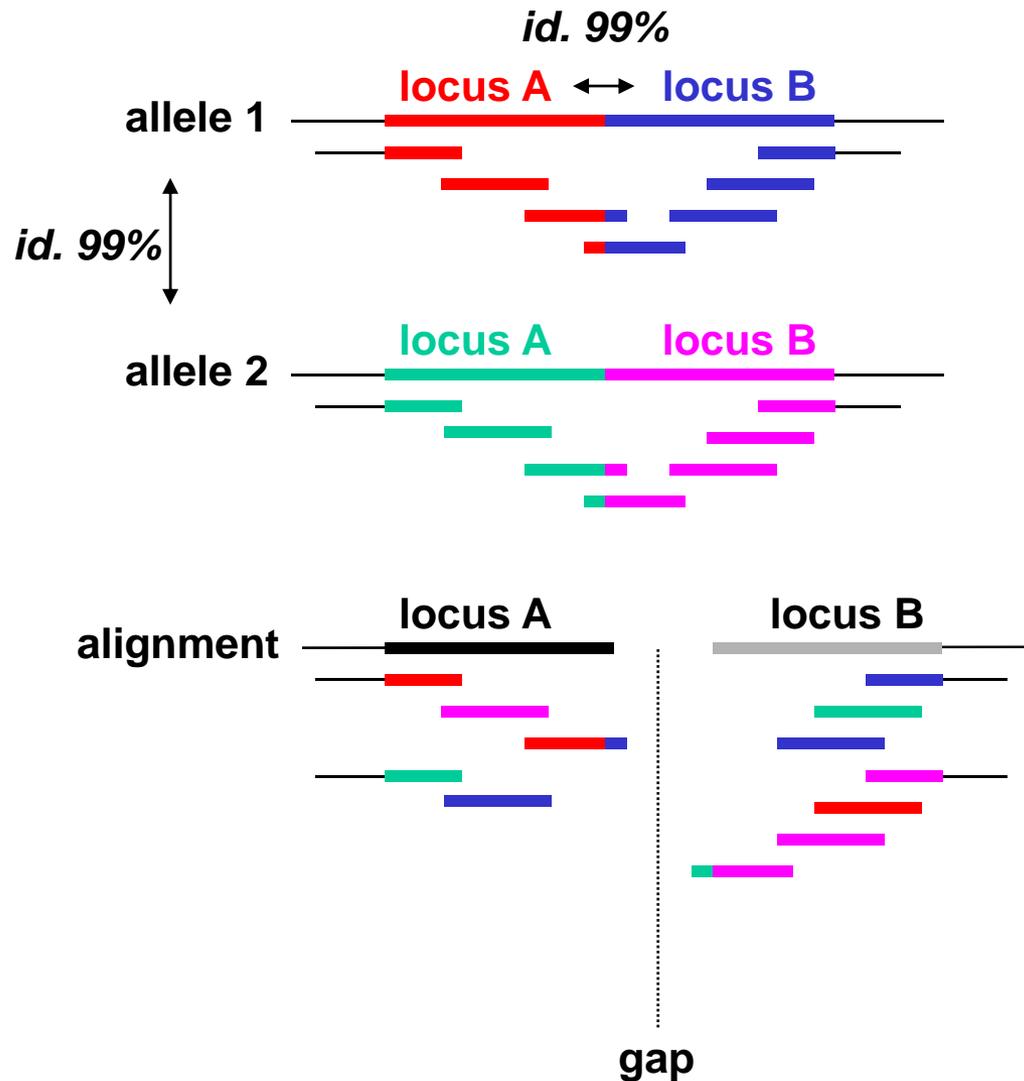
Segmental Tandem Duplications

Assembly problems



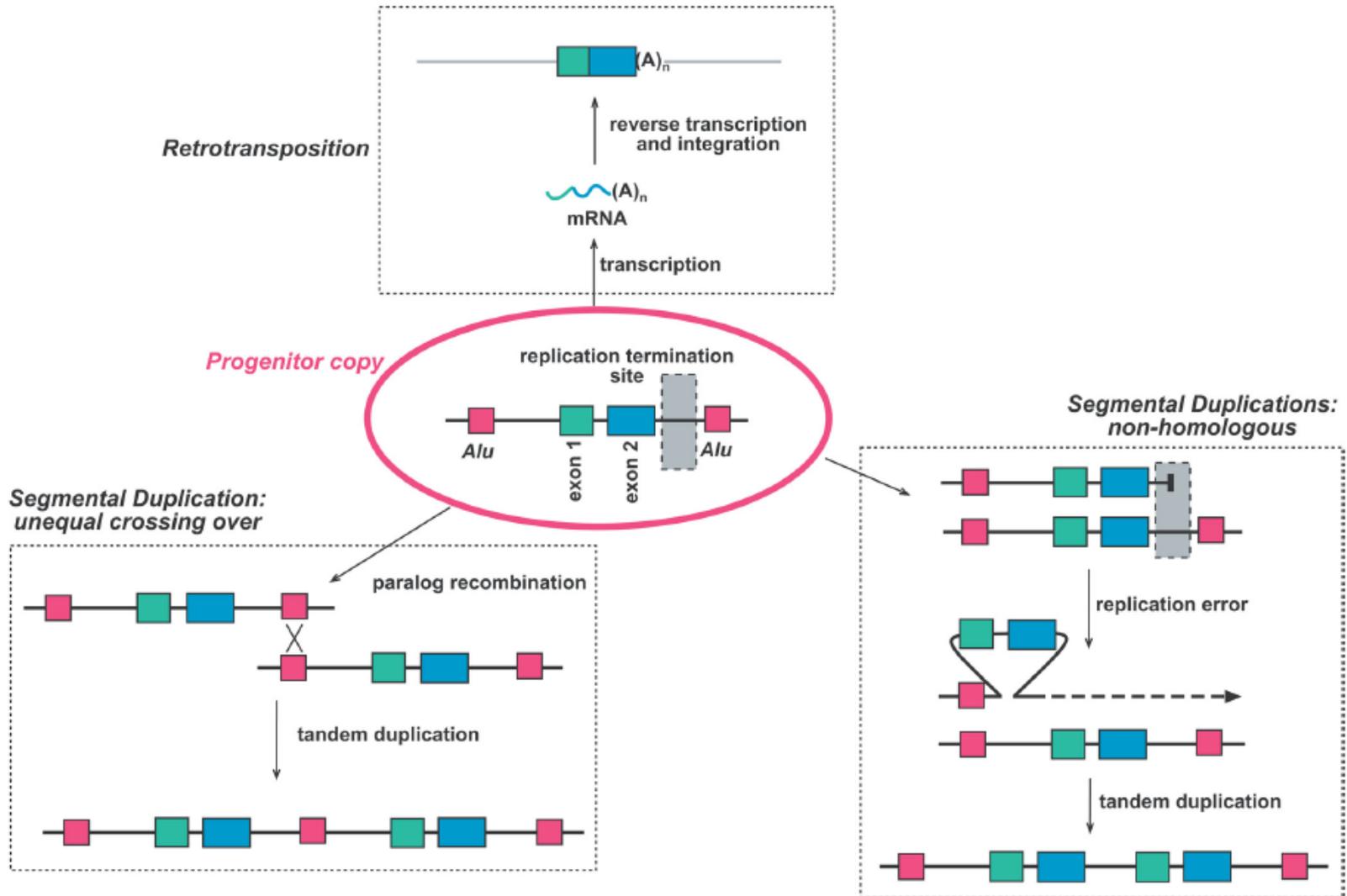
Segmental Tandem Duplications

Assembly problems



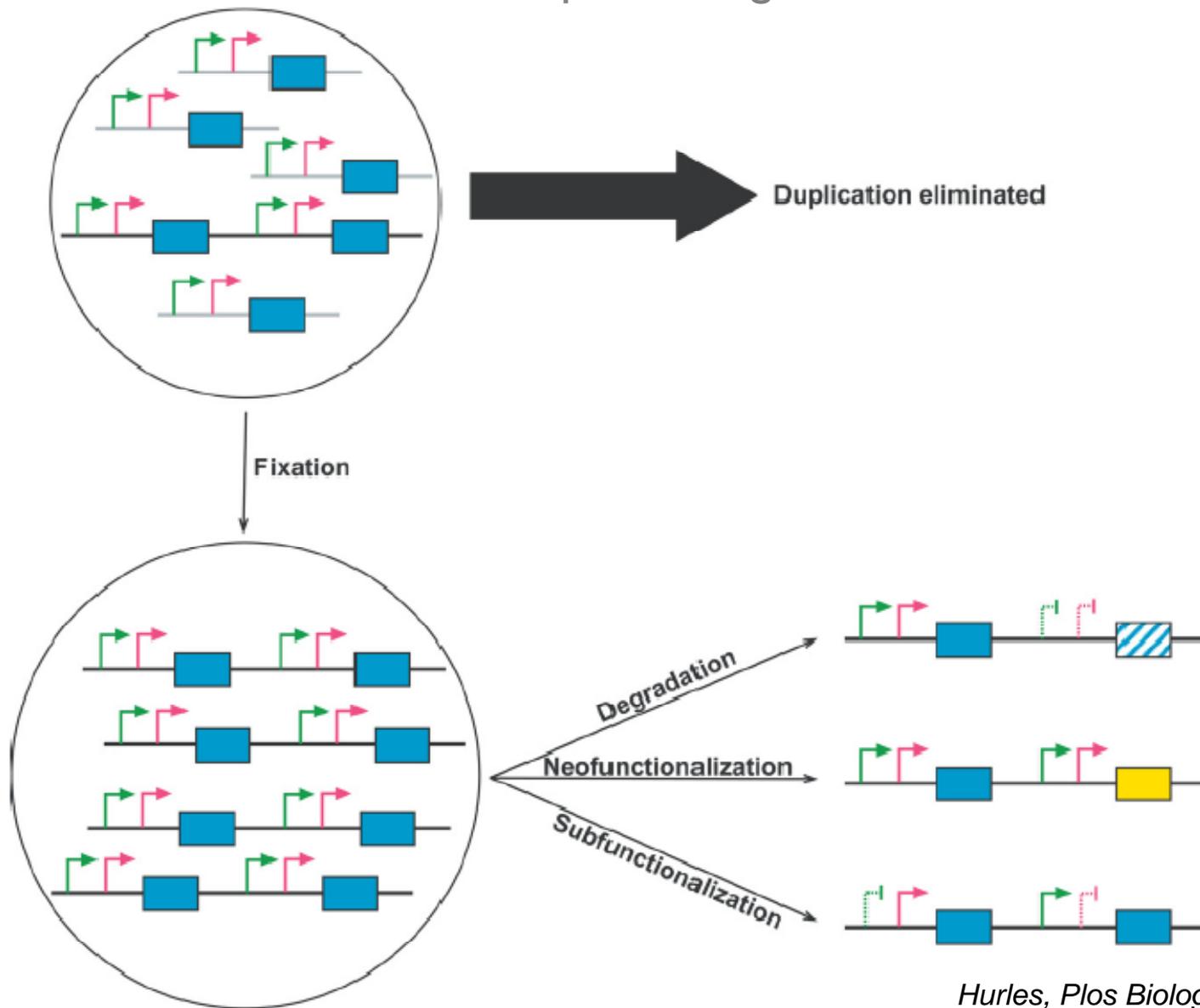
Segmental duplications

Mechanisms



Segmental duplications

Fate of duplicated genes



Structural Variations / Polymorphisms

Types

255 large-scale **copy number variations** (>100 kb).

Nat Genet 36:949 (2004)

Intermediate size variations >8 kb: 139 insertions, 102 deletions, 56 inversion break points

Nat Genet 37:727 (2005)

76 large-scale **copy number polymorphisms**; on average of 11 variations between two individuals with a median length of 222 kb.

Science 305:525 (2004)

586 **deletion polymorphisms** covering 276 genes; typical individual is hemizygous for 30-50 deletions >5 kb, totaling 550-750 kb.

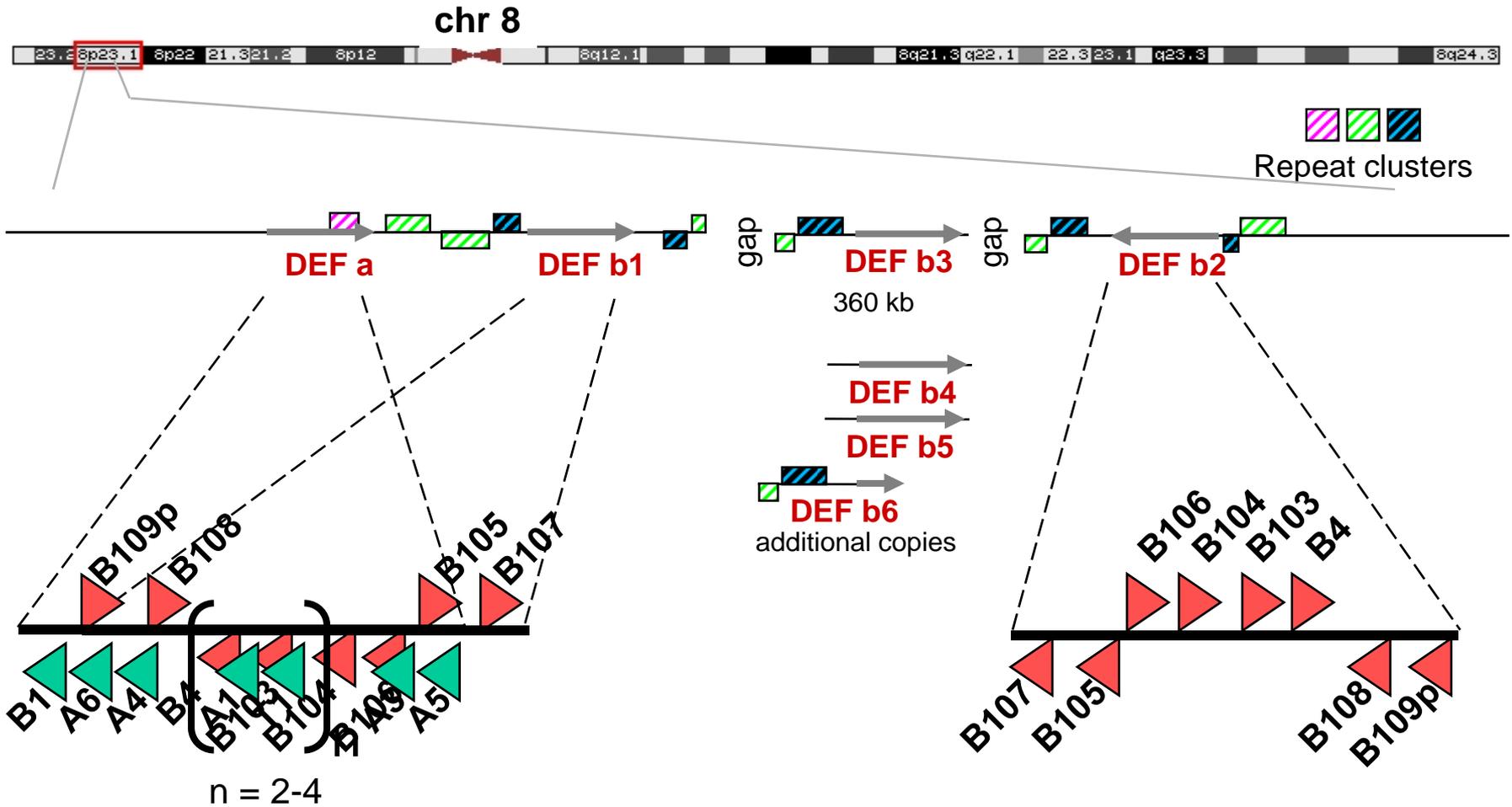
Nat Genet 38:75 (2006)

541 **deletion polymorphisms** covering 1-745 kb; 278 in multiple, unrelated individuals; 120 in homozygous state.

Nat Genet 38:86 (2006)

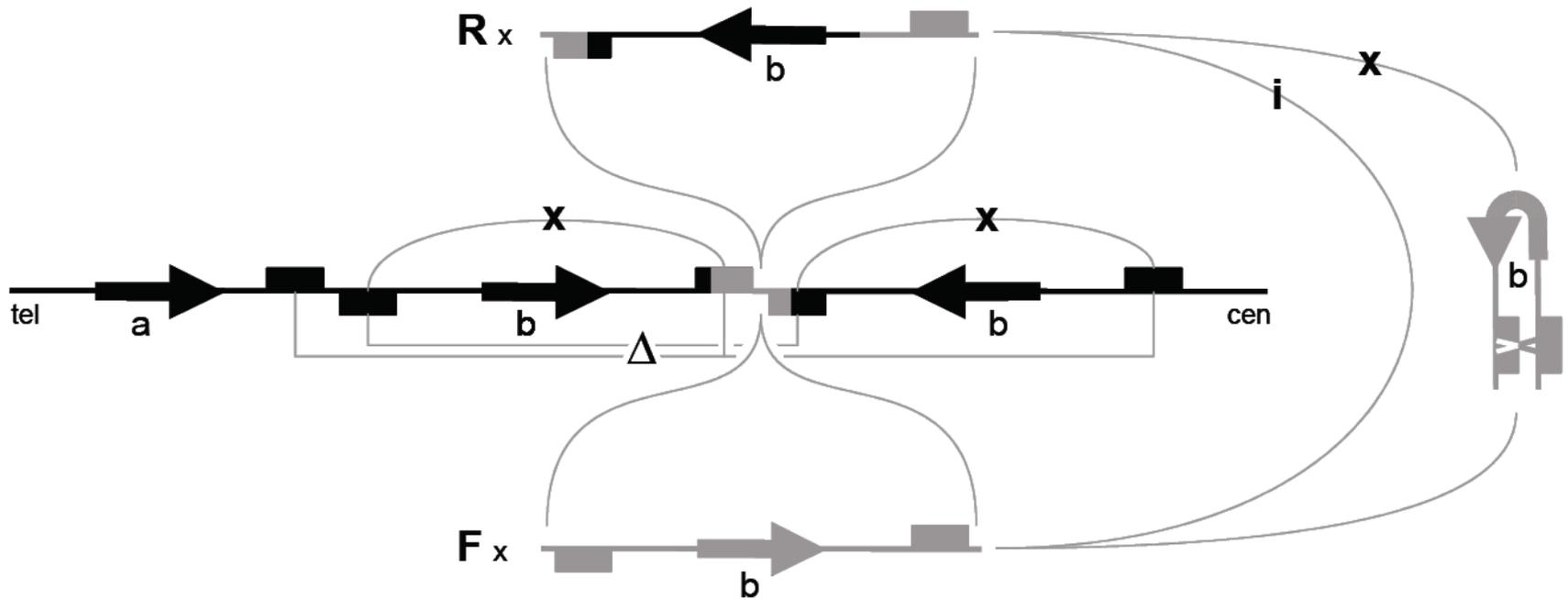
DEF cluster at 8p23.1

hg16: 6.3-8.3 Mb



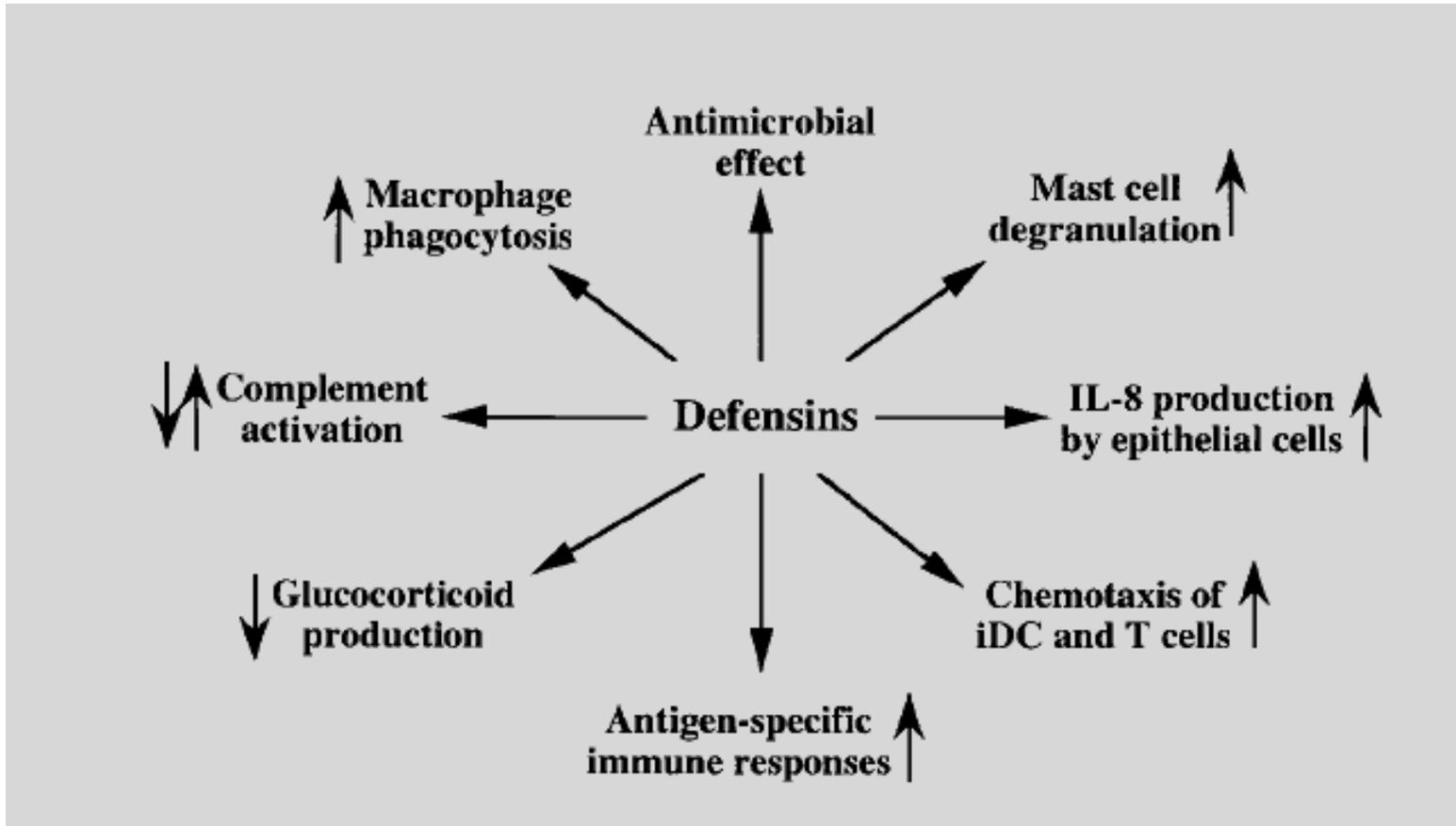
Genomic variability of 8p23.1 DEF locus

Hypothetical organisation



Defensins (DEF)

Multiple roles



Immunity & Cancer

Complex phenotypes / diseases

Structural variations

Gene	Type	Locus	Size (kb)	Phenotype	Copy number
<i>UGT2B17</i>	Deletion	4q13	150	Variable testosterone levels, risk of prostate cancer	0-2
<i>DEFB4</i>	VNTR	8p23.1	20	Colonic Crohn's disease	2-10
<i>FCGR3</i>	Deletion	1q23.3	>5	Glomerulonephritis, systemic lupus erythematosus	0-14
<i>OPN1LW/OPN1MW</i>	VNTR	Xq28	13-15	Red/green colour blindness	0-4/0-7
<i>LPA</i>	VNTR	6q25.3	5.5	Altered coronary heart disease risk	2-38
<i>CCL3L1/CCL4L1</i>	VNTR	17q12	Not known*	Reduced HIV infection; reduced AIDS susceptibility	0-14
<i>RHD</i>	Deletion	1p36.11	60	Rhesus blood group sensitivity	0-2
<i>CYP2A6</i>	Deletion	19q13.2	7	Altered nicotine metabolism	2-3

*Precise boundaries of the copy-number variant are not known.
VNTR, variable number tandem repeats.

Complex phenotypes / diseases

Structural variations

FCGR3 copy number & **glomerulonephritis** in humans
and rats

Nature 439:851 (2006)

Strong association of *de novo* **copy number** mutations
with **autism**

Science 316: 445 (2007)

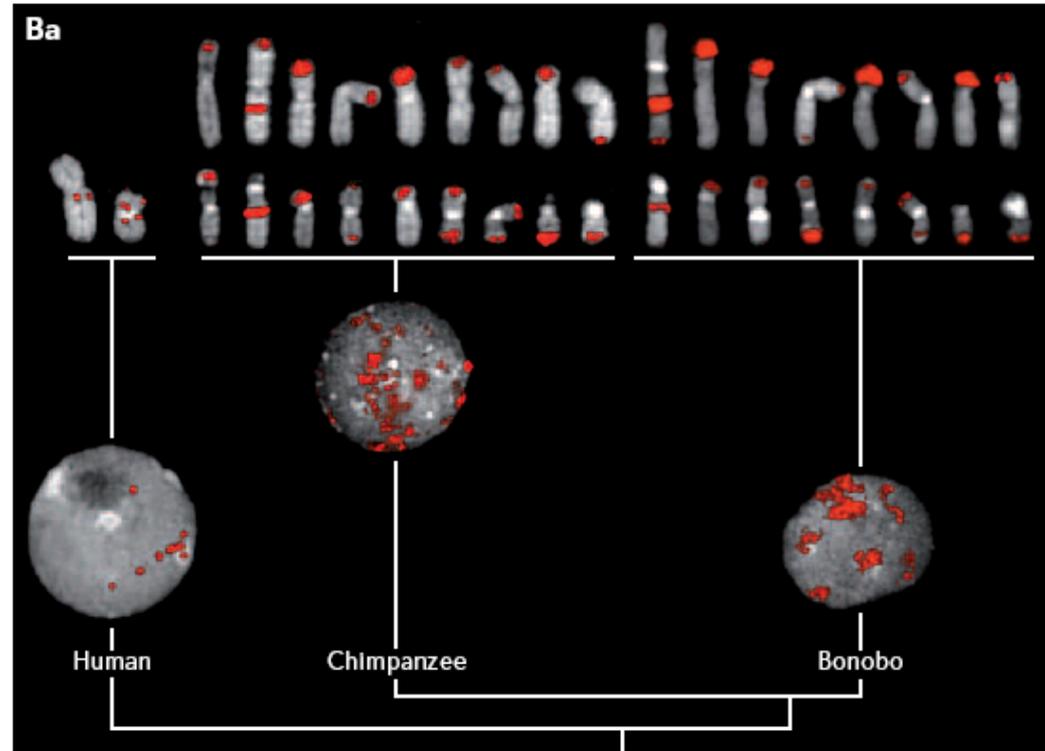
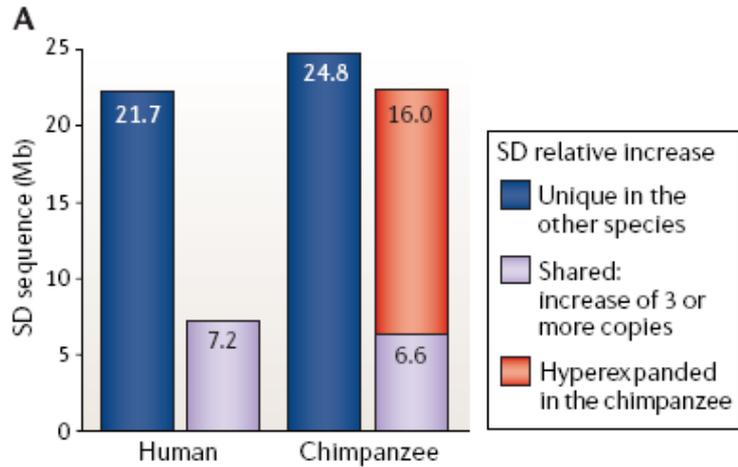
Segmental duplications

Content of sequenced animal genomes

	<i>Caenorhabditis elegans</i>	<i>Drosophila melanogaster</i>	Human	Mouse	Rat	Chicken	Chimpanzee*
SDs of >1 kb	4.3%	1.2%	5.2%	2.7%	1.6%	2.7%	N.D.
SDs of >10 kb	0.7%	0.1%	4.5%	2.2%	1.5%	0.3%	N.D.
SDs of >20 kb	N.D.	N.D.	4.0%	1.7%	0.9%	0.0%	~4.8%
Genome size	97	123	2,866	2,506	2,566	1,040	2,866

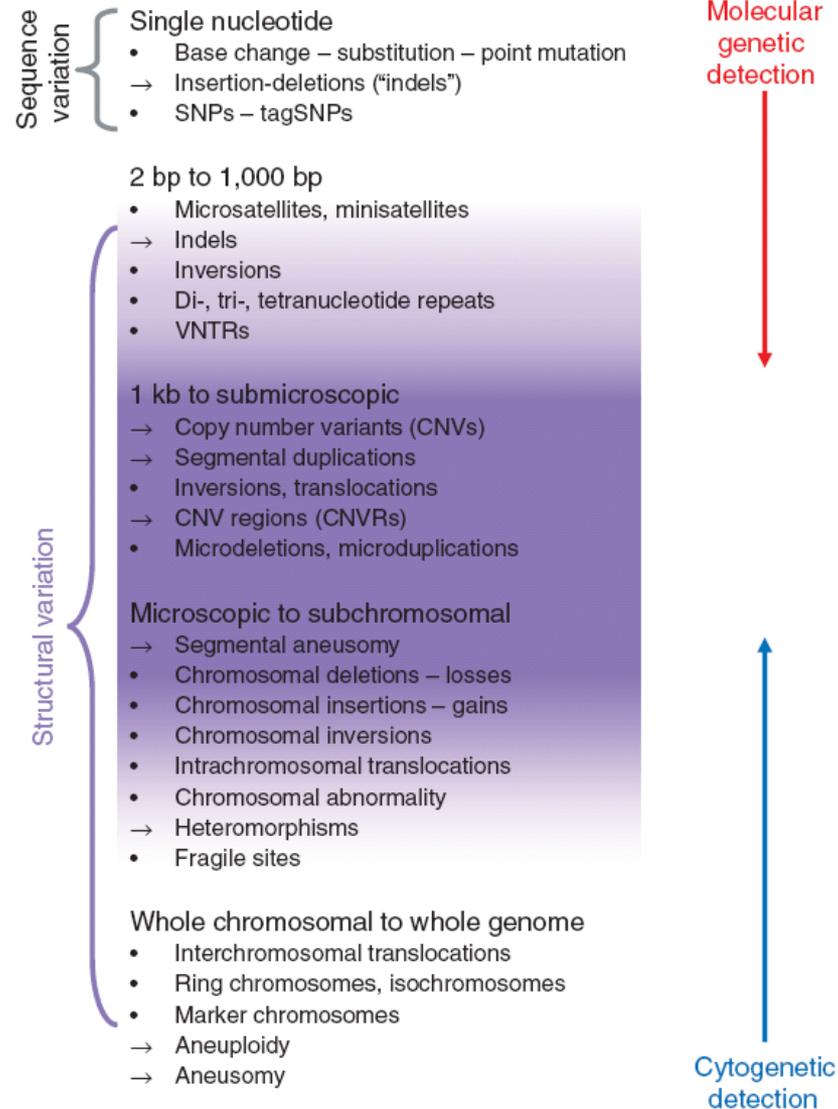
Segmental duplication content of hominoids

Hyperexpansions in chimpanzee



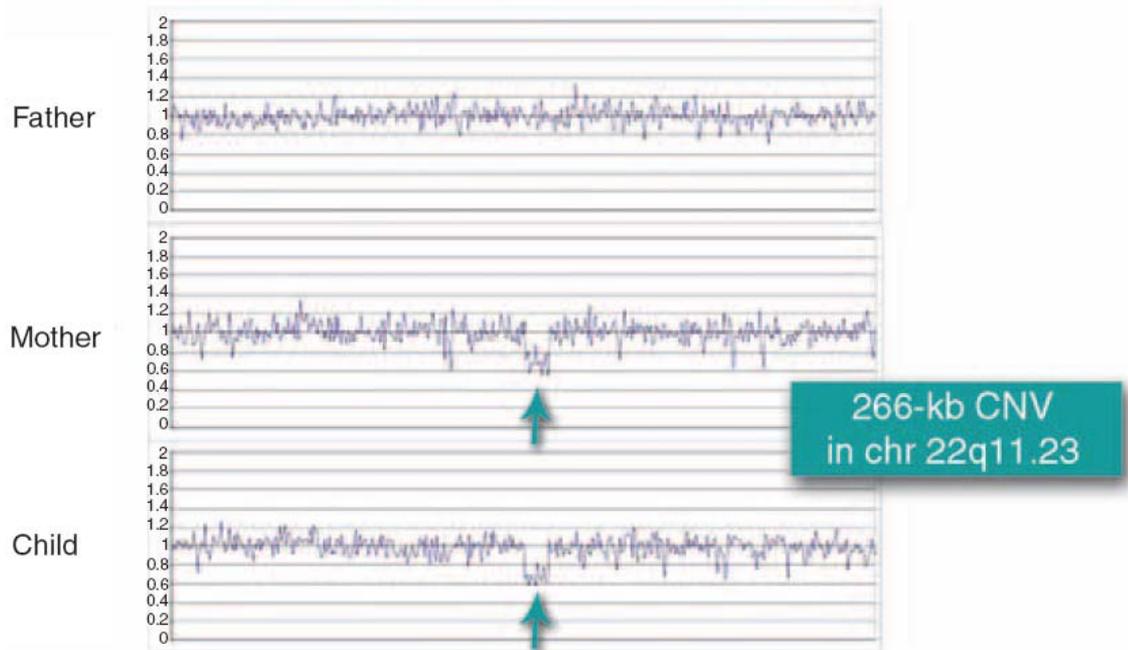
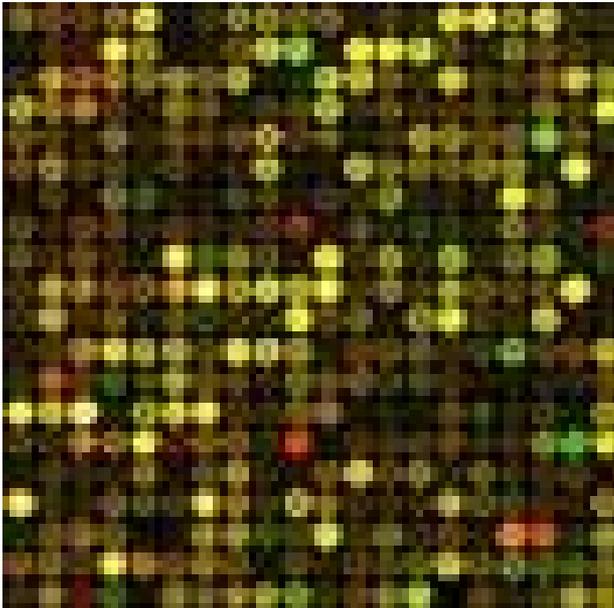
Genomic variations

Lexicon



Copy number variation (CNV)

Detection by DNA microarrays

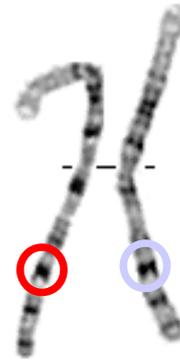
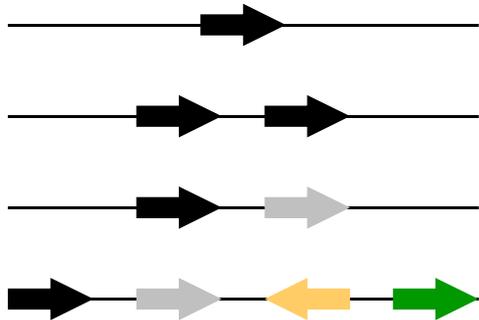


- 0.5-2 Mio data points
- comparative hybridization vs. a reference

Genetic Variability

Structural variations

Chromosome **A**



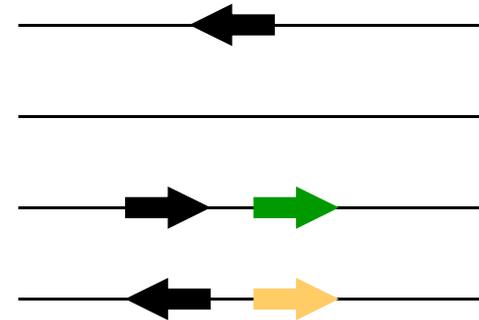
Inversion

InDel

Allele variation

Combination

Chromosome **B**



Conclusions

Genomes of any two individuals in the human population **differ more at the structural level** than at the nucleotide sequence level.

Differences between individuals

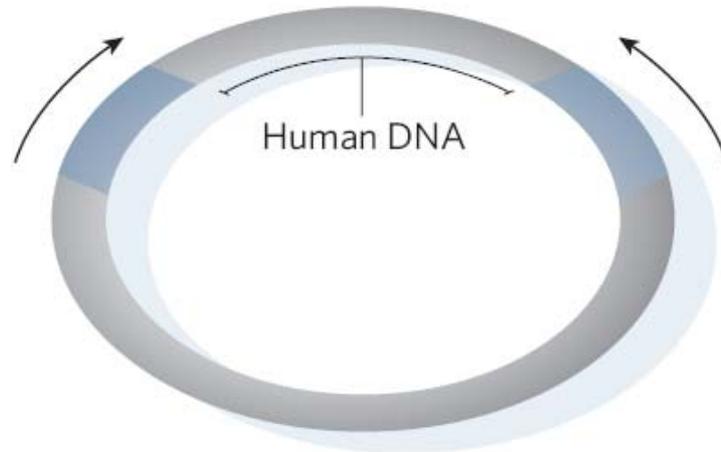
- CNV: >4 Mb >1/800 bp **> 0.12 %**
- SNP: 2.5 Mb 1/1,200 bp **0.08 %**

Conclusions

Copy number polymorphism at orthologous regions of diverse genomes ... suggests that genome plasticity... is a more common **cause of genetically complex phenotypes** than has hitherto been observed.

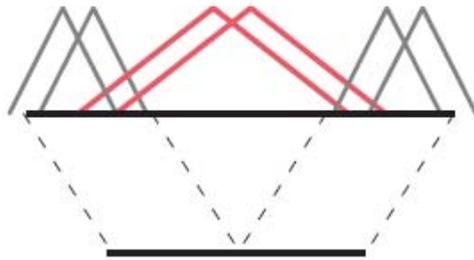
Completing the map of human genetic variation

Mapping structural variations



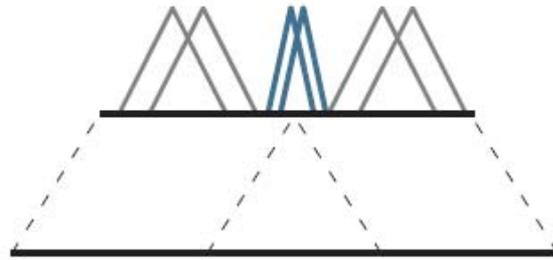
BAC/Fosmid vector

Span > mean +3 s.d.



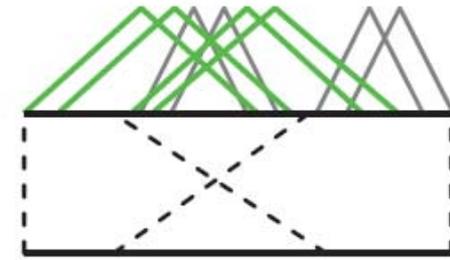
Deletion

Span > mean -3 s.d.



Insertion

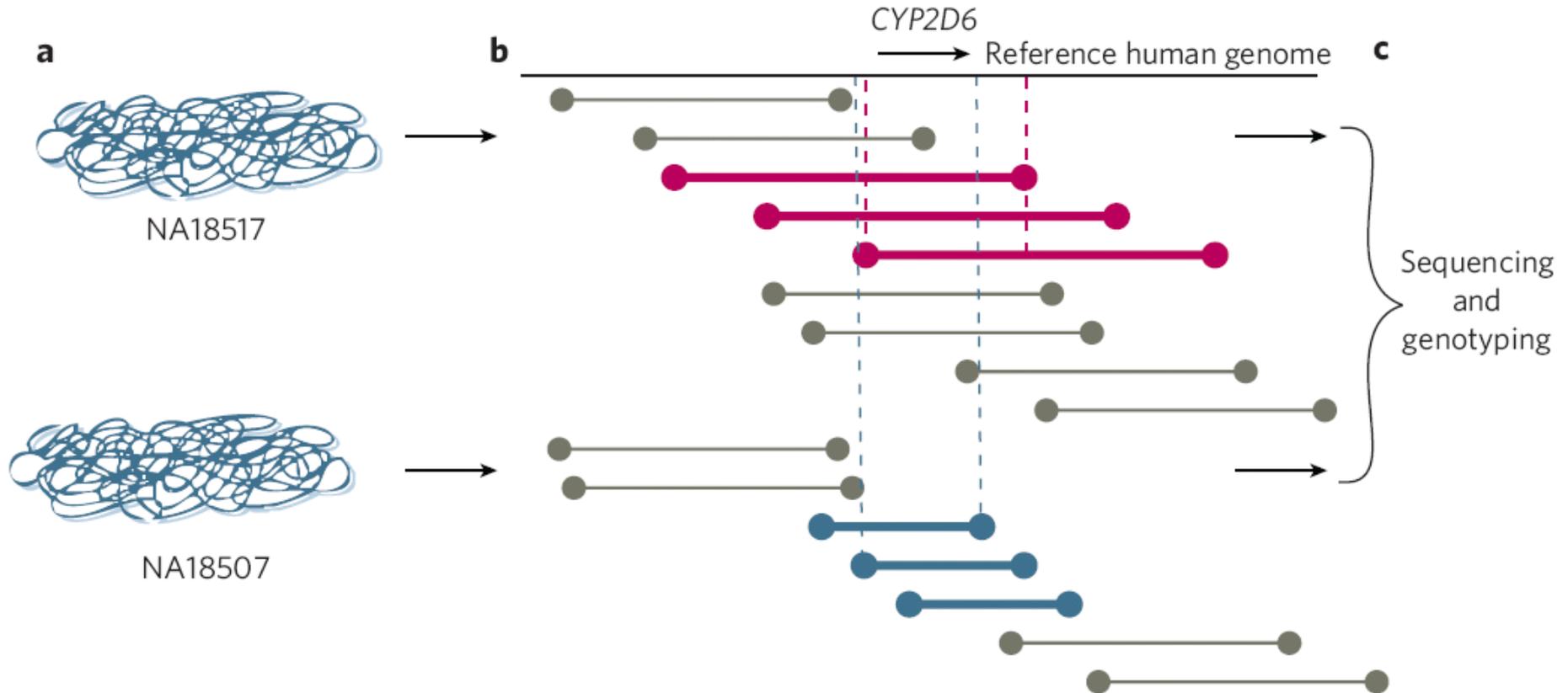
Inverted orientation



Inversion

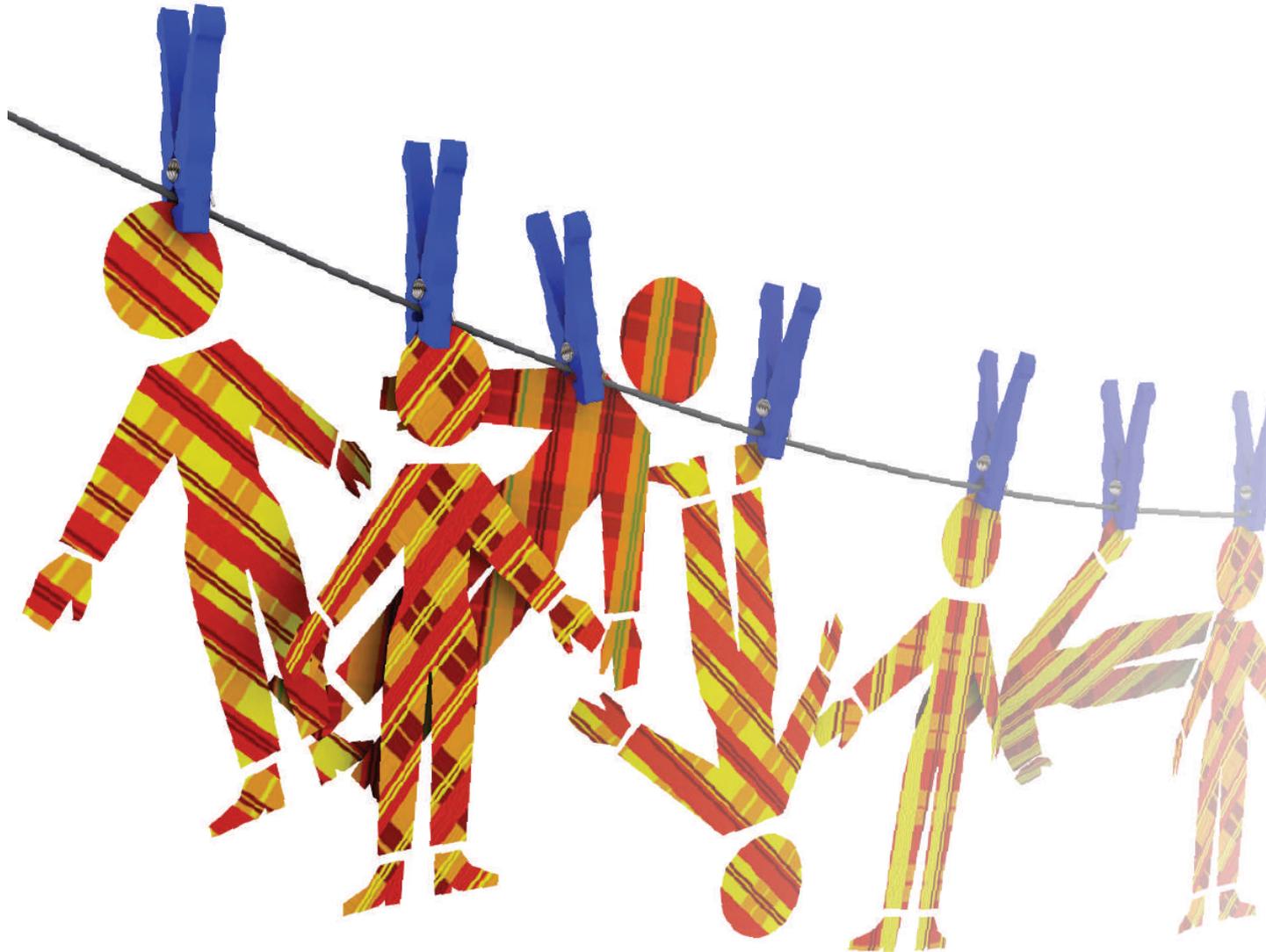
Completing the map of human genetic variation

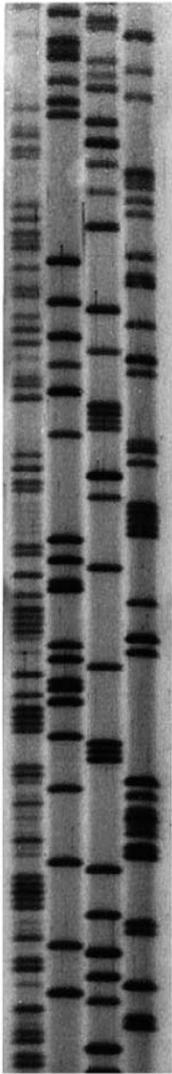
Sequencing structural variations



Genome Dynamics

Patchwork people ?





A C G T

genome.fli-leibniz.de Lectures