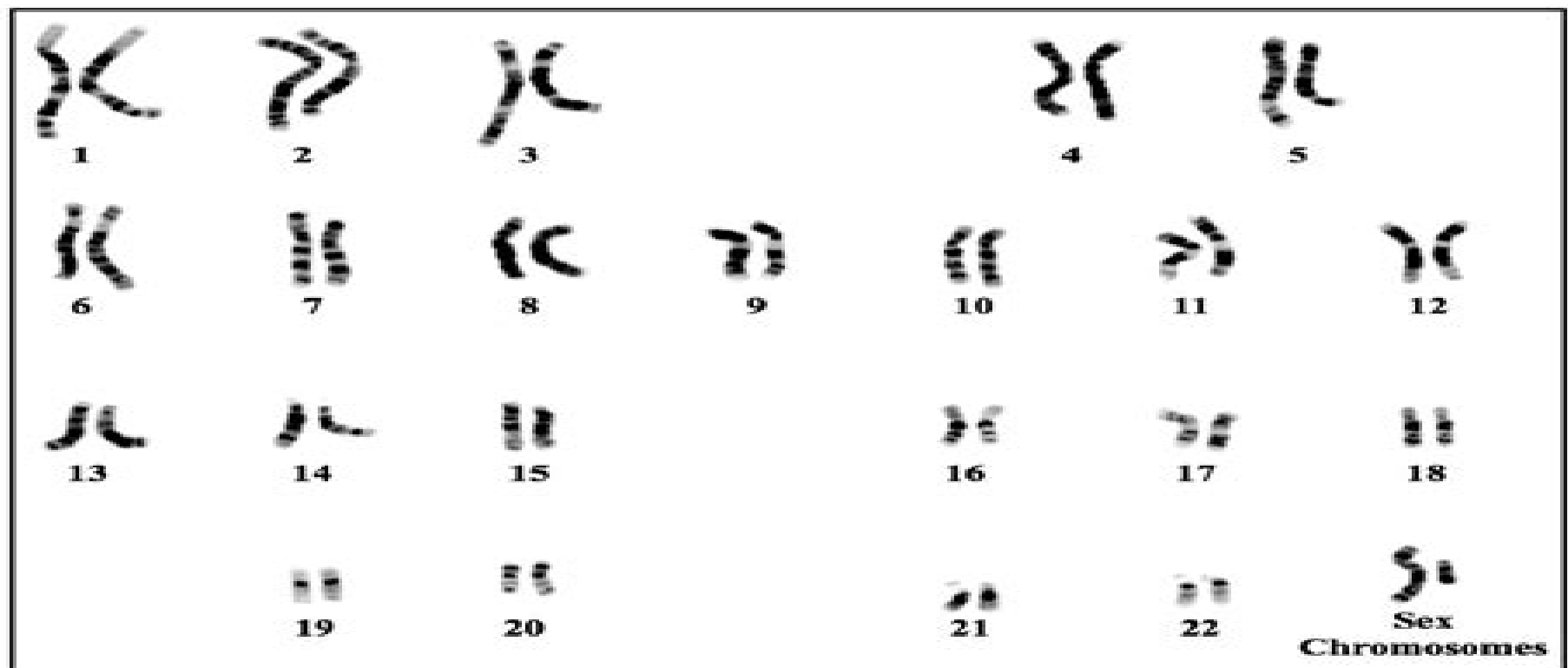


## Karyotype Activity Worksheet

**Background:** Karyotyping is routinely used to diagnose disorders caused by abnormal chromosomes. A **karyotype** is the arrangement and classification of the complete set of chromosomes in an individual. Cells are suspended while in mitosis, when the chromosomes have condensed and are easily visualized using a light microscope. A dye is introduced to the cell, which stains certain regions of the chromosomes, producing a banding pattern (also called "G-bands"). An image is taken of the chromosomes, and they are digitally rearranged so that members of each pair are next to each other and chromosomes are arranged according to size. The analysis involves comparing chromosome number, length, placement of the centromere, and the banding pattern. Below is an example of a karyotype taken from a normal individual.



Modified with permission from the National Human Genome Research Institute

1. How are the chromosomes arranged?
2. Is this karyotype of a male or female individual?

# Karyotype Activity Cut And Paste Matching

**Anna Morgan**



## **Karyotype Activity Cut And Paste Matching:**

*Chromosome Abnormalities and Genetic Counseling* R. J. M. Gardner, Grant R. Sutherland, 1996 There have been many advances in clinical cytogenetics since the first edition of this book appeared in 1989 The authors have written more expansively on segregation and risks in reciprocal translocations X autosome translocations inversions insertions and prenatal diagnosis The deletion syndromes have become more numerous and more precisely defined and this required treatment Fluorescence in situ hybridization has become a routinely applied methodology and its use has extended the power and increased the sophistication of the discipline of clinical cytogenetics The Human Genome Project is unravelling the complexity of our genetic inheritance and readers will find evidence of its impact on the practicalities of human cytogenetics throughout the book Two phrases not used at all in the first edition are dynamic mutation and genomic imprinting The fragile X syndrome now has the status of the prototypical dynamic mutation The authors completely rewrote this chapter giving it a much more molecular character The Prader Willi and Angelman syndromes likewise have become the classic examples of genomic imprinting and uniparental disomy Compared with the tentative comments on these syndromes in the first edition they now receive more extensive treatment Yet while the subject has become more complicated the book's aim remains the same to furnish a straightforward scientific description that will help readers understand the various chromosome abnormalities encountered in clinical practice and to provide practical advice that can be passed on to the people who have or whose families have these abnormalities

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