



Wild type female	Wild type male	Heterozygous male	Heterozygous female	Homozygous male	Homozygous female

# Biology If8765 Human Pedigrees

**Karl Pearson**



## **Biology If8765 Human Pedigrees:**

**Modern Analysis of Human Pedigrees** Laurence Hasbrouck Snyder,1934      **The Human Pedigree** Anthony Smith,1975      **The Human Pedigree** Anthony Smith,1975      **Some Extensions of the Gene Frequency Analysis of Human Pedigrees** Charles William Cotterman,1937      **The Use of Standard Nomenclature in Human Pedigrees** Heidi Ann Schoenhard,2004      *The Mendelian Collection of Human Pedigrees* George P. Mudge,1911\*      **Treasury of Human Inheritance: Pedigrees of physical, psychial and pathological characters in man** Karl Pearson,1911      Methods for Fine Mapping Complex Traits in Human Pedigrees Gonçalo Rocha Abecasis,William Cookson,2001      *The Pedigree of the Human Race* Harris Hawthorne Wilder,1926      **Treasury of Human Inheritance: Pedigrees of physical, psychial and pathological characters in man** Karl Pearson,1909      *The Treasury of Human Inheritance* ,1909      **The Treasury of Human Inheritance** Galton Laboratoty,University College, London,1912      **The Treasury of Human Inheritance** William Bulloch,2016-12-20 Excerpt from The Treasury of Human Inheritance With 11 Plates of Pedigrees and 1 Plate in Colours Plates 14 24 Plate F Pedigrees 73 192 Generation IV 1 6 children of III 2 and III 3 The three sons had each one foot enlarged the limbs of the three daughters being normal The order of birth of these children is not stated IV 7 and IV 8 the eight children of III 4 and III 5 One of the eight a boy has one enlarged foot the other seven being normal three of the seven are dead IV 9 IV 14 the five children of III 7 and III 8 the youngest sex l has an enlarged oedematous foot the others being normal IV 15 and 16 the four children of III 9 and III 10 The youngest sex not stated has an oedematous leg IV 17 IV 18 IV 19 three children of III 11 and III 12 Only the boy IV 19 was affected and suffered from enlarged foot When he reached maturity his testicles began to swell and progressed to such an extent that one of them had to be removed Strange to say as the testicle increased in size the oedematous foot decreased and after the operation became and remained normal IV 20 and IV 21 children of III 15 By her first husband the three children sex not stated were normal whereas by her second husband she had a son IV 21 with great oedema of both feet and both legs up to the level of the knees The left side was more oedematous however than the right IV 21 an American clergyman was the patient who consulted Milroy and whose condition led to Milroy s enquiry into the family history V 1 2 and 3 the 11 grandchildren of III 2 and III 3 Of them is stated that two viz A son and a daughter were affected with the family disease In both cases the foot alone on one side was affected The respective parents of these children are not given V 4 nine grandchildren of III 4 and III 5 sex and order of birth not stated none of them were oedematous V 5 V 6 V 7 nine grandchildren of III 7 and III 8 Their order of birth and in seven cases V 5 the sex not stated V 6 the son of the eldest daughter of III 7 has both legs enlarged while V 7 the son of the eldest son of III 8 has one foot enlarged The exact parentage of V 5 V 6 and V 7 is not clear from Milroy s account V 9 V 10 V 11 V 13 grandchildren of III 9 and III 10 One a son V 10 has an enlarged foot the other children being unaffected V 8 wife of V 9 and V 12 the wife of V 11 presumably normal V 14 and 15 two normal grandchildren sex of III 11 and III 12 V I6 13 normal grandchildren of III 15 The exact

parentage of these 13 children not stated VI 1 three normal children sex 1 of V 8 and V 9 VI 2 has one enlarged foot VI 3 sex not stated but alleged not to be oedematous VI 4 has both feet enlarged VI 5 two great grandchildren of III 15 age and sex not given but stated not to suffer from oedema In Milroy s family of 97 individuals there were 22 cases of oedema or about 23 o of the whole number With three exceptions the disease was congenital and progressive the exceptions being i III 9 whose normal leg began to enlarge after a carriage accident III 15 who was normal till the age of 12 3 IV 19 whose oedema disappeared after the removal of a testicle With reference to the sex incidence there were twelve affected males eight females and two the sex of which was not determined see Bibl No About the Publisher Forgotten Books publishes hundreds of thousands of rare and classic books Find more at [www.forgottenbooks.com](http://www.forgottenbooks.com) [Pedigree Studies of the Inheritance of Certain Human Characteristics](#) Paul Daniel Hickson,1953 [Dealing with Relatives](#) Dominic Nelson,2021 All of humanity and indeed all living things are connected by a complex web of ancestral relationships How much of this shared pedigree is known how much can be inferred and how completely it can be described are important factors in interpreting present day genetic diversity and discovering genetic contributions to disease The challenges of using pedigree information are twofold First in human populations genealogical data is often unavailable or incomplete Second when such datasets are available there are significant computational challenges to using them In both cases the response has often been to use approximate models of pedigree structure which removes the need for complete historical data and simplifies analysis But genetic cohorts are growing and increased sample sizes necessitate more detailed models of relatedness The work presented here represents a first step in explicitly modeling the relatedness of millions of individuals and using it to understand the fine scale diversity of whole populations as well as the genetic architecture of rare diseases First we developed a software package ISGen Importance Sampling in Genealogies for inferring the ancestral origin of rare alleles in population scale pedigrees as well as their distribution among present day individuals in a fully Bayesian framework Similar analysis had never been possible before at that scale and as a proof of concept we analyzed the distribution of the allele causing Chronic Atrial and Intestinal Dysrhythmia CAID within the province of Quebec Using observations of only 11 patients and 4 heterozygous carriers of the allele we estimated the allele frequency in 26 geographic regions of the province finding the highest expected frequency in Charlevoix where no carriers had yet been observed Despite its value for supporting the design of screening programs for rare diseases ISGen is limited to rare alleles and a single locus Extending to whole genome inference is a daunting challenge and can be aided significantly by high quality data with a known transmission history Simulations are a natural source of such data yet the most efficient simulation frameworks were based on coalescent models with known biases for long genomic regions and large sample sizes We developed two tools to address this issue One is an extension to the state of the art msprime coalescent simulator which allows simulations under a Wright Fisher model This extension makes use of the highly efficient algorithms and data structures of msprime and we showed more realistic relatedness among simulated individuals

as well as improved performance over coalescent simulations at whole genome scale We then continued to develop this extension to allow simulations to be performed within a predefined pedigree and when used with the population scale genealogy of Quebec Canada we showed that simulated cohorts captured much of the structure of a real dataset from the same population and significantly more than comparable simulations of a randomly mating population To show the potential of large scale pedigree simulations more generally we further described how pedigree simulations can aid in the design of sequencing cohorts for imputation and be used to detect rare variant associations while explicitly accounting for relatedness among cohort members

**Treasury of Human Inheritance, Vol. 1 (Classic Reprint)** Karl Pearson, 2017-10-22 Excerpt from Treasury of Human Inheritance Vol 1 For a publication of this kind to be successful at the present time it should as I have indicated above be entirely free from controversial matter The Treasury of Human Inheritance therefore contains no reference to theoretical opinions It gives in a standardised form the pedigree of each stock This is accompanied by a few pages of text describing the individual members of the stock giving references to authorities and if the material has been published to the locus of original publication When necessary the characteristic is illustrated by photography or radiography In this way it is hoped in the course of a few years to place a large mass of material in the hands of the student of human heredity It will not cut him off from but directly guide him to original and fuller sources of information Further the Treasury will provide students of eugenics and of sociology medical men and others with an organ where their investigations can find ready publication and where as time goes on a higher and more complete standard of family history than has hitherto been usual can be maintained Each pedigree and its description appears under the name of the author responsible for its completeness and accuracy and by aid of a key number to a confidential manuscript register of names and localities it is hoped that it may be occasionally possible for future investigators to recover traces of individual stocks or to ascertain whether newly discovered cases can be linked on to previously recorded families No one who has attempted a collection of this kind drawn from many quarters and prepared by different writers will be over severe on discrepancies and omissions in the earlier issues The full work of standardisation can only be carried out as the diverse needs of different types of family characters are better appreciated It is not always possible to maintain a proper balance between the graphic and verbal descriptions but I wish most strongly to insist on the point that neither are to be interpreted alone they are component parts of one whole and the reader who draws conclusions from the engraved pedigrees without consulting the verbal accounts is certain to be led into error Presence or absence of a character cannot be settled by the simple blacking or omitting to black a circle The description is practically that of the original observer whereas the pedigree is the work of the author of the special section of the Treasury and he may under or over estimate the statement presented to him As general editor I feel sure that much care has been taken to reach an unbiassed judgment and I know that contributors and members of this laboratory have Spent many days of labour in following up both original pedigrees and pedigrees in dissertations books and journals very

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**Human Pedigree Studies** Robert A. Peel,1999 [Phenotypic Variation, Fingerprints, and Human Behavior: An Application of the Family-Pedigree Paradigm](#) ,1974 *Mapping Human History* Steve Olson,2002 Olson traveled through four continents to gather insights into the development of humans for this sweeping history of humanity based on a new understanding of genetics Maps *A Catalog for Family Pedigrees, Cell Lines and DNA for Research* ,1994

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