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EFFECTS OF ATOMIC RADIATION

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NOTE

Throughout this report and its annexes cross-references are denoted by a letter followed by a number: the letter refers to the relevant technical annex (see Table of Contents) and the number is that of the relevant paragraph. Within each technical annex, references are made to its individual scientific bibliography by a number without any preceding letter.

Symbols of United Nations documents are composed of capital letters combined with figures. Mention of such a symbol indicates a reference to a United Nations document.
Annex H

THE GENETIC EFFECTS OF RADIATION

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I. MUTATION

1. The mechanics of mutation

The gene

1. The conventional concept of the gene has been that of a functional hereditary unit. In recent years this concept has required a more precise definition, since sensitive tests of allelism have indicated that a single functional gene may be separable into component elements by recombination and so shown to be capable of many pseudoallelic differences. Single mutational events which modify or prevent the action of the functional unit may affect different large or small parts of this unit. During the same period, it is notable that features of the genetics of natural populations have indicated the extent to which individual functional genes can be involved in larger complexes and lack complete autonomy. Possibly the most striking manifestation of this is at present in *Salmonella typhimurium*, in which it appears possible that there are integrated linear sequences of adjacent gene-structures responsible for whole sequences of biochemical operations, assembly-line fashion.

Gene mutations

2. Take in its widest sense, mutation means any change of the genetic constitution not due to recombination, ranging from whole genomes to alleles. Often mutation is used in a more restricted sense, viz. as change of the action of some specific gene. This is commonly referred to as point mutation, which, however, may be a misleading term, as it is known, especially from the work on *Drosophila* by Dubinin and others, that a change of the position of a gene may change its habit of action. Moreover, the idea of a point mutation, as distinct from a deletion or rearrangement, was formerly based upon the smallest unit of structure microscopically visible. Because recent structural analysis of the gene has seemed to penetrate almost as far as the much smaller ultimate units of its physico-chemical structure, believed to be the single nucleotides, it has already been suggested that the term "point mutation" be reserved for mutational events involving only one such unit. Such ideas do not by themselves affect the distinction between intragenic and intergenic mutations and may, indeed, clarify these: for example, it remains possible that further investigations of genes and chromosomes will lead to a distinction between a structural backbone and separate attached genes. There is no doubt that advances in this field will eventually add greatly to the refinement of current ideas concerning all aspects of mutation.

3. In man, the primary genetic concern is with all transmissible hereditary changes which simulate the change to a new allele. These are perhaps best grouped together under the term "apparent gene mutations", whatever their structural nature. However, other forms of genetic damage require consideration in connexion both with somatic effects and with the requirement that mutations must survive transmission through the germ cells if they are to be observed. These latter forms include both gene mutations and chromosome structural changes in somatic cells, which may well have a sensitivity to the radiation-induced process quite similar to that of germ line cells. Mutations and chromosome changes in these cells could bring about consequences, recognizable for the organism as serious somatic effects, ranging from death or incapacity of cells fulfilling vital specialized functions to unrestricted proliferation.

Chromosome breaks

4. It remains a major question to what extent chromosomal or other genetic effects may be responsible for cell death or damage in somatic or germinal tissues of man. Visible chromosomal alterations resulting from irradiation have been studied in cytologically favourable material, principally of plants and of insects. They commonly arise through one or more chromosome breaks in the cell rejoining in some new configuration. A frequent result is dominant lethality through loss of substantial chromosome parts or interference with cell division. In spite of the difficulties of objective numerical scoring of cytological phenomena, many quantitative investigations have been made upon them. It has been shown that the more densely ionizing radiations are relatively more effective in producing them and that the numbers observed or recovered can be considerably affected by various post-irradiation treatments if these are applied sufficiently early. In this way, recent work has suggested that there are some breaks at ionic bindings, which heal very rapidly, and others at co-valent bonds, which heal more slowly, as well as two separate effects of radiation, one in causing the breaks and the other in affecting the rejoining mechanism. The effects of oxygen, both at the time of irradiation and during the subsequent rejoining process, have played an important and controversial part in this advance. It would be of interest to learn to what extent investigation of post-irradiation modifiers of the rejoining process showed biochemical relationships parallel to those observed with modifiers of the cell lethality induced by irradiation.

5. Many investigations have connected ploidy with radiation resistance in unicellular organisms, especially the extensive work of Mortimer and his colleagues on yeast and this, together with the increased RBE of the more densely ionizing radiations, has led to the idea that much radiation-induced cell lethality has its origin in dominant genetic changes. Certain cases are, however, known in which this is not true; instead, lethality results from an imbalance or block in metabolism (as in very heavily irradiated Habrobacon eggs) or a generalized failure of the mitotic process hardly to be ascribed to individual processes of the break-rejoin type. On the basis of a two-hit killing curve for mammalian tissue culture cells of various ploidies, Puck has recently argued that radiation-induced death in these is chromosomal in origin, and that the number of fissions required before such a conclusion can be considered as finally established. However, the reduction in growth rate observed by Puck et al. in colonies derived from diploid mammalian tissue culture cells which had survived X-irradiation already provides prima facie evidence that even at doses of the order of 100 r, most surviving cells have suffered dominant deleterious changes. Moreover, Bender has recently demonstrated a rather high sensitivity of tissue culture cells derived from human kidney to chromatid breaks induced by X-rays.

The hereditary material

6. Recent years have remarkably advanced the knowledge of genetic material and of the role played in it by deoxyribonucleic acid (DNA). Indirect evidence from
different sources has long led cellular physiologists to believe that DNA, in close association with protein, forms part of genes and chromosomes; this has included the relative DNA content of haploid and diploid cells of various tissues of an organism,\(^{29}\) cytochemical evidence, including the almost complete restriction of the presence of DNA to the cell nucleus and the association of DNA synthesis with cell division.\(^{24}\) More recently, a very close association has been demonstrated between the assimilation of radioactive racers incorporated in DNA, and chromosome division.\(^{28}\) In addition, other evidence has inclined many geneticists to believe that DNA may be the actual material whose configuration constitutes genetic information; this evidence includes:

(a) The transformation of hereditary characters of cells of *Pneumococcus*\(^{22}\) and *Haemophilus*\(^{24}\) bacteria by application of solutions of pure DNA.

(b) The role played by DNA in the growth and heredity of the coliphages of the T series.\(^{25,6}\)

(c) Indications from current work that increased mutation in microbial systems occurs under conditions of deficiency for an essential constituent of DNA such as thymine, or in presence of a competitive analogue of a constituent, such as bromouracil.\(^{27}\)

All such phenomena carry the promise of new lines of investigation of the mechanisms of gene mutation.

7. Concurrent investigations have remarkably advanced understanding of the chemistry and structure of DNA, particularly the X-ray diffraction studies of Wilkins et al.\(^{18}\) and the complementary biochemical relationships uncovered by Chargaff and others,\(^{49}\) leading to the remarkable double helical structure proposed by Crick and Watson,\(^{49}\) so suggestive of the exact replicative process required for the transmission of hereditary characters, and already so productive of fresh ideas concerning the mechanics of mutation.

8. While none of these arguments is alone conclusive, and while it is recognized that the genetic material of cells of higher organisms is organized into very substantial stable structures, which must be more complex physically and chemically than the fine DNA fibrils visible only under the electron microscope,\(^{21}\) nonetheless very many geneticists believe that the ultimate carrier of genetic information is likely to be the arrangement of nucleotides in DNA.

9. In that event, total radiation-induced mutation rates, in the widest sense of change of the hereditary information, might be expected to be quantitatively correlated with the DNA content of the cells of the germ plasm together with the biochemical operations which construct and maintain DNA. It at least seems reasonable that when comparisons of mutation rates between different species or physiological conditions are made, parallel DNA comparisons should be kept in mind. The DNA contents of some relevant types of cell are listed in Table VII. Most kinds of cell nuclei contain enough DNA to form a structural molecule of great length which could only be packed inside the nucleus by much folding. This has given rise to the recent suggestion, now appearing on purely structural grounds, that the chromosome may consist of a multi-stranded structure.\(^{28}\) If the structure turned out to be, say, a proteinaceous backbone with attached DNA molecules as side arms forming the genes (a possibility which is not excluded), the distinction between inter- and intragenic mutations could eventually come to have a very real physical basis, and the two kinds of mutation could differ in mechanisms.

### Linearity of dose-mutation curve

10. The experimental justification for speaking of radiation-induced mutation rates at low doses rests upon *Drosophila* data, in which the linearity of the dose-mutation curve, when it is investigated under sufficiently rigorous conditions, has been confirmed down to X-ray doses of 25 rad for irradiation of spermatozoa by the painstaking work of Stern and his collaborators,\(^{31-32}\) following earlier experimenters.\(^{34-36}\) Muller\(^{21}\) has recently argued cogently that there is no point in pressing the test of linearity below 5 rad, and has indicated that this limit could be reached in *Drosophila* by techniques at present available. Many geneticists would agree with the implication of the cited passage, that linearity can already be safely accepted, without the enormous labour involved in its extension to still lower doses—at least in the absence of a definite proposed basis for expecting a non-linearity. However, it must be borne in mind that linearity has not been tested in this range of doses for spermatogonial irradiation. In the case of irradiation of these cells it is still difficult to conceive a priori of a non-linearity at low dose, followed by a linear portion of the curve at medium or high exposures. However, Oakberg\(^{28}\) has shown that some classes of spermatogonial cells of the mouse are very sensitive to the lethal effects of low doses (5 rad—100 rad) of gamma-radiation. If these same classes were to turn out also to be unusually sensitive to the induction of mutations by radiation, the curve of recovered mutations as a function of dose might turn out to be linear in the range of moderate doses, but to have considerably higher slope in the very low dose range where an appreciable proportion of the cells surviving irradiation belonged to the sensitive group.

11. The Committee has been informed of current experiments upon mice which will enable the linearity of the dose-mutation curve for irradiation of spermatogonia, oögonia and oöcytes to be checked down to 37.5 rad.\(^{49}\) Attention must, however, again be drawn to the dependence of the whole quantitative assessment of genetic effects of low doses upon an assumed linearity and for irradiation of a particular type of cell in a dose range not experimentally investigated.

### Mechanism of mutation

12. Many attempts have been made to affect the process of induced mutation after its initiation by exposure to ionizing radiation. Some of these have been successful to a greater or lesser degree,\(^{10-44}\) and this fact is of cardinal importance as demonstrating at least the possibility of interference between the irradiation and its principal genetic consequence. Unfortunately, in many of these cases the precise genetic nature of the mutational event is not known; association with chromosome breakage or rejoining may therefore be suspected. Moreover, many of the experiments refer to microbial material, in which it is possible that the genetic structures are far more exposed and more easily able to be reached and affected by external agents than are the mammalian chromosomes. Nevertheless, it is a hopeful sign that recent experiments reported to the Committee have extended the demonstration of post-irradiation interference to a well-known class of apparent gene mutations, the sex-linked recessive lethals of *Drosophila*.\(^{49}\) These experiments seem to show that a finite interval of at least some tens
of minutes exists in Drosophila before "fixation" of radiation-induced mutations.

13. In connexion with any possibility of ultimate practical use of chemical or other modifiers of induced mutations, it is well to remember that, in many populations, the largest man-made exposures of the gonads occur through comparatively large doses delivered relatively infrequently in the course of medical work at controlled times. The possibilities of modifying the mutational effects of radiation should be considered in the light of the more general discussion of modifiers of radiation effects in chapter IV and annex F of this report.

Other possibilities of interference between irradiation and its effects at the cellular level

14. Interference with and control of genetic consequences of irradiation does not end with the completion of the mutational process. However, to look further requires that the completed mutations be detectable. The number of conditions in which carriers of unexpressed deleterious genes can be detected has recently increased greatly;45,46 this trend is closely associated with advances in general biochemical and immunological genetics, and it is to be hoped that Governments will foster and encourage its progress. A second field closely related to this and other aspects of the present subject is that of human chromosomal cytology. We are indeed a long way removed from the beautiful situation which prevails in Diphera where giant salivary gland chromosomes can be studied in minute detail; nevertheless recent technical advances in the field have been considerable18,19 and can give us great hopes of progress. Such advances may bring about radical changes in human genetics and especially human radiation genetics.

15. Other radical possibilities for dealing with radiation-induced mutation, besides the cumbersome and often painful process of selection, beyond question exist. An example which must be considered, in the light of technological advance, is that of the natural or controlled transfer of genetic characters. This phenomenon is well-established in microbial materials,56 although usually but not always with very low frequency,57 and as an eventual aid in the elimination of harmful genes or their consequences it cannot be entirely dismissed as speculation.

Comparison between natural and radiation-induced mutations

16. There has been a widespread belief among geneticists, based largely upon the classical work of Stadler in corn58; that radiation produces in general a different type of mutant allele from those which occur naturally—more extreme, less likely to be reversible, more frequently a loss of function. However, Stadler's work may not be entirely typical even of plant material.185 Muller has recently reviewed the evidence against existence of such a distinction.6 Certainly, both the mechanism of production and the distribution among loci of radiation-induced and natural mutations differ;185 there is also some indication of small differences in the proportions of mutation to the different alleles at a single locus.44 Minute one-hit deletions do occur under the action of radiation.14 and some radiation-induced point mutations in Drosophila may be associated with breaks or structural changes near them.20 Moreover, evidence in Drosophila is against any appreciable correlation between natural mutation rate and radiation-induced mutability where either individual genes,17 strains,38 or physiological conditions59 result in altered natural rates. Very little correlation is also found between radiation-induced mutability and the natural rate in the sample of thirty biochemical back-mutations examined by Glover.19 However, the wide variations in the ratio of radiation-induced to natural mutability found both in the work of Glover on bacteria and in extensive work on plants60 do not seem to be correlated with the type or severity of the forward or back-mutation involved, and it is generally accepted that the ratio of visibles to lethals is much the same for natural and radiation-induced mutations in Drosophila, although no explicit study of this point has been made. Moreover, a very detailed investigation by Giles61 of purple-adenine and other mutants in Neurospora has shown no evidence for a qualitative or quantitative difference between radiation-induced and spontaneous mutations at the same locus. The evidence of Stadler primarily relates to the compound A locus; consequently, a possible explanation is that A has a very low sensitivity to radiation-induced point mutation. It is therefore reasonable to accept as a tentative assumption that spontaneous and radiation-induced mutations are qualitatively similar; wide differences in the two mutation processes exist but are functions of individual loci, and are not appreciably correlated with the type or severity of effect exerted by the mutant allele.

17. In connection with this problem, attention may be drawn to certain organisms such as Aspergilus,62 bacteria,43 and coliphage,64 in which very sensitive tests of allelism are possible; tests which may be calculated43 in some cases to be adequate for resolution of recombination distances corresponding to one nucleotide pair if genes are primarily constituted of DNA. Such investigations might eventually shed much light on the real magnitude of the structures disturbed by various types of mutational event of different origin, and indirectly on the "quality" of mutations caused by different agents. Unfortunately, all the above organisms are microbial and not necessarily representative of the larger chromosomes of higher organisms.

18. In man, little information yet exists concerning the relative sensitivities of genes to specific mutagens. However, a notable beginning has been made upon the problem by Penrose,67,189 who has analysed the mean parental age at birth of propositi showing various conditions, and correlated these shifts with hypotheses as to the principal kinetically different classes of mutagens, such as natural radiation (expected to raise both mean paternal and maternal ages by an equal small increment), copy-error (expected to raise mean paternal age somewhat), or chemical mutagens (which might under some circumstances raise the mean maternal age in such a way that incidence increased more than linearly with age). Thus the prospect already exists of the analysis of human genes in terms of sensitivities to different kinds of mutagen.

Detection of mutation

19. An apparent gene mutation can be detected if it results in a new allele which differs so much in its action from the original one that it can be scored by appropriate methods. There exist different alleles (isoalleles) whose phenotypic effects cannot at present be distinguished but which may differ in other respects as, for example, mutability.62 Studies of natural and induced mutations are restricted to those which can be distinguished pheno-
20. In Drosophila as well as in mice the rate of visible mutations at specific loci has been studied after matings of the stock to be tested with animals of the opposite sex containing the marker genes whose mutation frequency is to be examined. By this method the visibly scored include both those which are recessive lethals in homozygous condition and those which are homozygous viable, provided only that they are visible and viable as heterozygotes with the allele in the marker stock.\textsuperscript{189} As reported by Russell,\textsuperscript{29} six out of twenty-one tested mutants induced in spermatogonia of mice were lethal, seven were semi-lethals and eight were viable. The corresponding data from Alexander's\textsuperscript{29} test of mutations in spermatogonia in Drosophila yielded three lethals, one semi-lethal and four viable. Excluding rare heterozygotes combining a recessive viable visible with a recessive lethal visible, what could be scored in any corresponding study in man might be only those recessive visibles not rendered unscorable by their association with recessive lethals. Supposing the same relationship between viable and lethal visibles as in mice, one might easily understimate the total mutation rates of genes in man by a factor of two or three.

21. In estimating mutation rates it must also be borne in mind that the same phenotypic effect need not mean a genetically identical condition. In man, as in many other organisms, several different genotypes may exist which give rise to indistinguishable phenotypic expressions. In the case of man one must think of classes of genes each causing a similar effect, rather than of specific single genes. The number of genes in each such class may vary considerably, causing a strong variation between the observed rates of natural mutations in the various classes. Thus in man, because test breeding cannot be used to pin down an alteration to a specific locus, a mutation rate is always in fact measured for the whole class of genes giving rise to one altered condition, recognizable trait, or clinical entity.

22. In recent years many important studies of the mutational process have been made in unicellular organisms. There are, however, several major problems in the measurement of gene mutation rates in single cell material, including a lag between application of radiation or other mutagenic agent and the observable expression of mutations which enables them to be counted: this lag can be due to various factors, segregational or physiological.\textsuperscript{71} Furthermore, there is always a possible effect of non-mutant cells upon the survival of mutants during tests.\textsuperscript{29} A different problem, peculiar to back-mutations, is the difficulty of distinguishing apparent back-mutation at the same locus from suppressor or modifier effect. For this problem, which is related rather closely to the important question of the reversibility or otherwise of radiation-induced as compared to spontaneous mutations, there are great advantages to microbial material in which both kinds of forward and reverse mutations have been and are being explored. Both radiation-induced and spontaneous mutation rates have been measured with relatively high precision in unicellular organisms, especially bacteria, under a variety of conditions.\textsuperscript{73,74} It is to be hoped that the techniques and methods developed will yield equally valuable results when applied to the clones of mammalian tissue-culture cells now available.

23. The basic difficulty in any quantitative study of natural mutation rates is to obtain large enough numbers, for these rates are low (tables I, II) and cannot of course, be raised artificially for purposes of study. Consequently, investigation has been confined to organisms which can be handled or are present in rather large numbers, such as bacteria, Drosophila, and humans. The limit to the information on natural mutation rates which can be derived from the very extensive and careful control observations in mice, in the work both of Carter, Lyon and Philips\textsuperscript{13} and of Russell\textsuperscript{89,74} illustrates the difficulty. Because chromosome structural changes occur naturally at much lower frequencies even than apparent gene mutations\textsuperscript{71,80,104} and the study of rates has been confined almost entirely to the latter events, only these will be considered here. In man, individual cases, once found, can be followed up with relative ease even in large populations, because the family and individual are indentifiable by name, etc. As a result, it is possible that more information about natural mutation rates for single phenotypic entities exists for man than for any other organism. In man, however, as in other organisms, the basic problem of small numbers governs consideration of the field.

The rate and variation of natural mutations in experimental organisms

24. In other organisms than man, it has been possible by experiment and test breeding to examine more closely the variations in natural mutation rates as well as the absolute magnitudes. The general ranges of the latter do not vary very widely (table II).

Physiological variations

25. As noted above in another context, physiological variables affecting natural mutation rates of individual loci have been examined in bacteria by Novick and Szilard\textsuperscript{13} who concluded that the number of mutations increased as a function of chronological time rather than cell division. This may, however, not be generally true.\textsuperscript{79} Moreover, the genetic material of bacteria may not be entirely representative of that of higher organisms. Moreover, the general lack of systematic variation of doubling dose among species of widely different generation times, militates against any assumed dependence of number of natural mutations upon chronological time.

26. Work on physiological variables in Drosophila has been carried out in relation to mutation at classes of loci, such as the recessive lethals, rather than at single loci. Differences between natural strains\textsuperscript{174} and between sexes\textsuperscript{88} and dependence upon age\textsuperscript{80} have been established for a number of organisms. These variations in natural mutability are not known to be correlated with variations in the radiation-induced rates.

*Strictly, the term mutation rate refers to the rate of occurrence of mutational events and not to the frequency of mutant gametes among tested gametes, although it is also commonly used to refer to this latter measure. The distinction must, however, be borne in mind in certain situations: for example, if it is desired to compare true natural mutation rates estimated for free living unicellular forms of life with the frequencies of appearance of mutant gametes in higher organisms, since the latter do not directly reflect the rates of occurrence of mutational events in the germ line cells (see table II).
27. The difficulty, even in Drosophila, of obtaining enough data to document significant variations in natural mutation rates between loci other than exceptional unstable genes further underlines the basic problem of numbers in the investigation of natural mutation rates. Variation between loci, and in certain cases between isoalleles at the same locus is, however, well-known in this organism. It has been far more extensively documented in the bacteria, at least for back-mutations; the rates of these vary from $10^{-8}$ to the lower limit of detection near $10^{-10}$; they are correlated with mutability by radiation to only a very small extent.

28. In extreme cases variations between loci may originate in genes which are themselves unstable or confer instability upon others. Where mutant genes affect all or a large part of the genome, they may in addition be partially responsible for variations in spontaneous mutability between strains. Again, such genetic modifications of spontaneous mutation rates is not known to be correlated with change in radiation-induced rates.

Natural mutation rates in man

29. Penrose, Neel and others have tabulated a number of calculated rates for single clinical entities in man (see table I). In examining these values, it is necessary to bear in mind the limitations of the data and of the methods of calculation by which they are obtained.

Direct methods: autosomal dominants and sex-linked recessives (table I)

30. In the case of clear-cut autosomal dominant visible entities, the mutation rate is in principle directly estimated by observation of propositi whose parents and other close relatives are normal. The various technical difficulties such as failures of ascertainment and occurrence of phenocopies, degree of penetrance, and the proportion of cases not due directly to fresh mutation have been discussed in the literature. The experimentally ideal dominant visible combining full penetrance, complete ascertainability and responsibility for total sterility would be of reduced value, since it could not be proved directly to be genetic in origin. Moreover, in practice studies are commonly made upon the natural mutation rates in those populations where they are known to be highest, simply in order to obtain enough documented cases to make the results statistically significant. It is therefore questionable whether the observed rates are representative. They cluster around $10^{-8}$ per gamete in a distribution which is rather skewed. If a population of $10^9$ is surveyed during five years for an ideal condition, observable during thirty years, it already constitutes a considerable labour, and yet significant results are unlikely to be obtained unless the mutation rate exceeds $10^{-4}$. In practice, no such ideal conditions exist. It is very probable that some of the well-documented human mutations have much lower frequencies. Perhaps the possibility should be faced that the sample of spontaneous mutation rates which have been measured in man isGrouped into 27368156945541874261003541891571537444319181928249700661077652060216018279218137786026817610678779194131216589794361631856371755928502805605357070367722797205971861824572895620059253322828629924979958971498472003229382529842301178

31. The mutation rate for autosomal visible recessives is calculated indirectly, by a process originally due to Haldane. The observed number of propositi, together with an estimated selective disadvantage in the homozygote, is used to calculate the rate of disappearance of the mutant alleles concerned from the population, and a balancing rate of forward mutation is inferred from an assumption of genetic equilibrium. The uncertainties concerning possible existence of small selective effects in the heterozygote and of large departures from equilibrium render extremely uncertain the values obtained in this way: indeed, perhaps the most notable use of such figures has been to deduce a priori expectation of heterosis from a few "unreasonably high" calculated mutation rates, although most of them lie in the same order of magnitude as those for dominant entities (see table I).

Lower limit to detection of recessives

32. An autosomal recessive with a selective disadvantage of only 1 per cent in the heterozygote, in a population whose coefficient of inbreeding was 0.01 per cent, would, if its mutation rate were $10^{-8}$, show up phenotypically in no more than about 1 in $10^8$ of the population. Even if the condition were fully penetrant, a mutation rate would be very difficult to estimate. Such genes, if their natural mutation frequencies were in the range of $10^{-6}$, could hardly be observed at all. There is therefore reason to believe that the best documented sample of recessives for which indirect estimates of mutation rate are available may be unrepresentative. If this is because they show very slight heterozygous advantage, the mutation rates calculated for them are also too high; but then there is a fallacy in the converse argument, that because many of these turn out upon investigation to be heterotic, most human mutant alleles are so.

Consanguineous marriages

33. The study of consanguineous marriages does not lead to estimates of natural mutation rates but to estimates of the numbers of recessive alleles present in populations. In principle, these marriages constitute a test-breeding for the presence of recessive alleles through the associated degree of homozygosity ($Y/4$ for first cousins) which they bring about. It may, however, be questioned whether a truly comparable control group can ever be obtained, although internal controls by comparison of different degrees of consanguinity are usually available. The limited number of studies made show as yet no very consistent picture. Of them, those by Sutter and Tabah and by Schull are the most extensive, and that by BóöK the most intensive. Morton, Crow and Muller, by an ingenious argument, have shown how to present the over-all reduction in viability, which is observed in three of the surveys, in the form of an equivalent number of alleles which would be lethal if homozygous, or lethal equivalents, carried per head of population. From the surveys analysed by them they conclude that 3-5 lethal equivalents acting before maturity were present per individual in the population, a figure with which the survey reported by Schull is in satisfactory agreement. Unfortunately, the intensive examination carried out by BóöK shows an entirely different picture of viability, although in a very small sample; the total deaths, including prenatal and up to age 30, in BóöK's sample, were almost identical in the cousin marriages and the controls.
34. The content of deleterious recessive genes of a population, whether expressed in lethal equivalents or otherwise, is an important parameter indicative of its genetic state. It is also a valuable standard of comparison for actual or postulated mutation rates. There is, however, another possible use for it. Comparison can be made of the total recessives in lethal equivalents, derived from vital statistics only, with intensive investigation of all the known recessive lethals present, such as that undertaken by Böök. (Ideally, the total reduction in viability and fertility up to the second generation beyond the cousin marriages should be employed, see paragraph 113 below) and the intensive examination should cover all known recessive conditions.) In this way it might be possible to obtain some idea of what proportion of recessive damage is covered by the known effects, and what proportion remains unknown; a factor of great importance to our confidence in any estimates or predictions, based as they must be upon current limited knowledge. This possibility is discussed in more detail in paragraph 113.

35. It is clear that improved recording of such consanguineous marriages, in maternity hospitals or centres of vital statistics, would be of great value and should be encouraged by Governments if they wish to be aware of the general state of genetic well-being of their peoples.

36. The Committee has been informed of large-scale current or planned surveys of consanguineous marriages both in Japan, where the frequency of these is high, and, as regards vital statistics, in Canada.95,96

37. It has, unfortunately, not been possible so far to establish total natural mutation rates in man for very large classes of genes, such as that formed by the sex-linked recessive lethals of Drosophila. Such large classes, if they could be investigated upon a firm genetic basis, might more easily provide adequate numbers for reliable statistical analysis than can be obtained from the laborious search for specific rare conditions. In this connexion, it is of interest that Lejeune and Turpin95 have recently attempted to interpret the decrease of sex-ratio at birth with age of the mother,186 and the combined data upon irradiated and aged fathers appears at present to involve contradictions. Since there does appear to be a decrease in sex-ratio with age of the father,186,38 it seems a reasonable possibility that mutations to sex-limited detrimental autosomal dominants are concerned and that they are due to natural irradiation or other non-cumulative, time-independent causative agents (Penrose's Class I; see paragraph 18 above). It would evidently be of great value if clear-cut interpretations could be established in some other mammal, such as the mouse, since secondary sex-ratio data are widely recorded in large populations, although not always in a form suitable for genetical analysis, and they are relatively free from the ambiguities of fine diagnostic distinctions. The possible interpretation of sex-ratio data is further discussed in paragraph 64 below.

*mutator and unstable genes*

38. In any consideration of variations in spontaneous mutation rates, the evidence of mutator genes and unstable genes, well-established in corn, in Drosophila and in bacteria,95 must be borne in mind, together with the fact that these commonly do not affect the rate of induc-

*Radiation-induced mutation rates*

39. Radiation-induced gene mutations have not yet been observed with certainty in man, and so no quantitative dose-mutation relation exists for the genes responsible for any specific clinical entity. In consequence, quantitative assessments of the mutational effects of the irradiation of human populations must rely upon extrapolations which are often of uncertain validity. In any event they depend upon the well-established results of the investigation of radiation-induced mutation in other organisms.

*magnitude and variation of radiation-induced mutation rates in organisms other than man*

40. Since the field of mutational radiation genetics was opened by Muller in 1927,95 it has been established in all the many organisms tested that ionizing radiations can induce apparent gene mutations: hence the same is believed true of man. X-ray induced mutation rates have been measured for a large number of single loci, especially in Drosophila. Both the range and average of such rates are known for a wide variety of individual visible markers through measurements made under very carefully controlled conditions, and so also is the total rate for certain large classes of markers such as the sex-linked recessives of Drosophila. A number of rates observed in experimental species are listed in tables III, IV and V.

41. In mammals, the most extensive investigation of the X-ray induction of mutations at single loci so far carried out is that for mice,60,75,10,96 in which the rates at seven autosomal recessive visible loci have been investigated in spermatogenesis; the average of these rates is found to be about fifteen times the average for a comparable group of loci in Drosophila.10

42. Extensive research has been conducted upon the variation in sensitivity to radiation-induced mutation with physiological condition. In the male it has now been established that the mutability is low in spermatogonia, rises to a peak during the time of formation of spermatozoids, falls to a second minimum in immature spermatozoa, and then rises up to the time of ejaculation, both in Drosophila57,95 and the mouse.99 In the female Drosophila, the oogonia show a mutability similar to that of spermatogonia while late oocytes are very mutable.37,100 The subject has recently been reviewed by Glass.160 Drosophila is also the only organism for which extensive determinations exist of the relative rates of mutations in different selective and other classes, either at single loci or summed over large parts of the genome.101,102

43. Muller93 has pointed out that evidence in Drosophila indicates that mutation rates in somatic and gonial
cells are about equal. Extension of this principle to other species and eventually to man might make possible very informative conclusions from investigations on somatic mutation rates in vivo in man.

44. Calculations have been made by Haldane and others concerning the practicability of observing not single locus rates but total rates over a large part of the genome in a mammal such as the mouse. Such an experiment upon the very large scale necessary might be of considerable value at this juncture in the process of extrapolation to man; it would, however, involve the expenditure of a great many scarce mouse-geneticist-years. The Committee has been informed of the existence of a pilot experiment on these lines.

45. The concept of genes as finite structures of different sizes which carry hereditary information largely in the form of different arrangements of nucleotides in DNA has recently made possible one particularly interesting interspecies comparison concerning induced mutations. There is evidence that in mice the total rate of induction of recessive lethal mutations in sperm is higher than the corresponding rate in Drosophila by a factor of about 20. The same is true for the rate of mutation per locus averaged over several different loci, and in addition there is a similar difference of about twenty-fold in the same direction in the DNA content per nucleus. This suggests that perhaps mouse genes are not more numerous but are larger than Drosophila genes—that the extra DNA has gone into building genes that are bigger and more complex rather than more numerous. The possible application of such an idea to man, an organism in which mutational events cannot in general even be assigned to definite loci by test crosses, but which has a DNA content per nucleus similar to that in the mouse, might lead one to expect rather high mutation rates, both spontaneous and induced, when measured "per clinical entity", as well as all the complexities and peculiarities of large multiple allelic series, of which a notable example has been uncovered by Dunn in the t-alleles of the mouse. Penrose has already drawn attention to the possibility of some unusually complex genes in the X chromosome of man, in connexion with very high observed natural mutation rates.

Radiation-induced mutation rates in man

46. Whatever approach is adopted to the problem of radiation-induced mutation rates, the gonad doses received both by control and by experimental groups will have to be known.

47. In principle, the simplest method to obtain a quantitative relation between dose and radiation-induced gene mutations in man is to make a comparative survey of the progeny of an irradiated ("experimental") and a comparable un-irradiated ("control") population. Those surveys published so far are concerned only with the first generation born of irradiated parents. However, it is easy to show that, as human matings cannot be controlled, examination of the first generation provides more information in itself than examination of subsequent generations.

48. In the last analysis, all the observed quantities come down to variations in frequency, and therefore:

(a) All studies must be accompanied by the examination of a control sample presumably issued from genetic stock identical to that of the irradiated sample. This condition greatly restricts the value of the results published so far.

(b) All the results obtained are subject to an inevitable sampling error which necessitates the collection of a very large amount of data.

A number of quantitative characters, such as birthweight, size and various anthropometric measurements, as well as statistical data, such as neo-natal mortality, have been suggested and examined. Unfortunately, the precise genetic component in these variables is not known; on the contrary, they are known to be dependent upon factors which are economic (standard of living), demographic (age of parents, order of birth, etc.) and sociological (medical care).

49. The characters that can be utilized may be grouped in two categories, according to whether they are connected with dominant (or sex-linked) visible mutations or with dominant (or sex-linked) lethal mutations. The detection of visible dominants is carried out in practice by the observation of malformations at birth. It is in fact reasonable to assume that an increase in the frequency of dominant mutations associated with visible effects would manifest itself to some unknown extent as an increase in frequency of malformations. The same would be true of visible sex-linked recessives in boys born to irradiated women. Lethal mutations may be revealed in four ways:

(a) Increase in frequency of miscarriages (virtually impossible to determine with certainty);

(b) Increase in frequency of still-births (much more feasible but subject to the demographic considerations mentioned in connexion with neo-natal mortality);

(c) Reduction in fertility, or even sterility (virtually impossible to measure in man);

(d) Disturbance in the ratio of the sexes at birth (deviation in the sex-ratio, an easily observable criterion).

50. The various studies which may be taken into account at the present time are listed, together with pertinent results, in table VI. Given the very uneven quality of the data presented by the various authors, and the particular way in which they were arranged by each of them, it is impossible to add together the figures from the separate surveys. In general, none of the investigations makes a definitive demonstration of a genetic phenomenon. Only the decrease in the sex-ratio, which is found in the three studies of irradiated mothers, seems to be acceptably established as a reality. Although no one of these studies concerning sex-ratio yields statistically significant results by itself, the fact that all three deviate in the same direction gives some confidence concerning the reality of the effect. Although several of the studies to date raise the possibility of an increase in congenital malformations among the offspring of irradiated persons, the findings in this regard are much less consistent than those concerning the sex-ratio. In this connexion, it must constantly be borne in mind that where many comparisons are being drawn between two groups, on the basis of chance alone in twenty of these comparisons will yield differences exceeding the 5 per cent level of significance. Further observations regarding the possibility of an increase in congenital defect or early death are highly desirable.

51. In summary, it seems possible, although only with great difficulty, to distinguish a detrimental effect of irradiation on the first generation issuing from irradiated parents. The possibility of firm demonstration and
measurement of this phenomenon suggest that all these studies be extended on the largest scale possible, wherever practicable surveys can be made with a reasonable probability of yielding positive significant results in a comparison with adequate controls.

52. In view of this possibility of future surveys of the progeny of irradiated persons, it seems worthwhile to indicate the criteria which determine the value or “resolving power” of any such study. In brief, five points must be considered:

(a) The dose to the parents of the individuals under study;

(b) The number of individuals whose parents have been so exposed;

(c) The number of characteristics of genetic significance to be recorded;

(d) The manner in which information on these characteristics is collected;

(e) The availability of a suitable control group.

53. To illustrate the manner in which (a) and (b) may be taken into consideration, a particularly simple hypothetical case has been selected, that of the detection of an ideal autosomal dominant visible allele causing complete sterility:

Suppose the gene concerned to mutate at a rate \( m \) per gamete in the control population and at an increased rate \( fm \) per gamete in the irradiated population. If the doubling dose for the mutational step concerned is \( D_r \) rad and the mean genetically significant exposure per parent of the irradiated group is \( D \) rad, then

\[
\frac{f - 1}{1} = \frac{D}{D_r}
\]

If \( P \) progeny of the irradiated group and \( Q \) of the unirradiated are examined with complete ascertainment for the visible allele, the numbers expected to be observed are respectively \( 2m(P) \) and \( 2m(Q) \). The observed difference in rate between the two groups is \( \Delta = 2m(f - 1) \) and has an approximate variance due to the limited sample size of

\[
\sigma^2 = 2m(1/P + 1/Q)
\]

In consequence, even if no other sources of error are considered,

\[
\chi^2 = \frac{\Delta^2}{\sigma^2} = \frac{2m(1-f)^2}{(1/P + 1/Q)}
\]

If we require \( \chi^2 \geq 4 \) for a significant increase of mutation rate in the irradiated group to be established, and denote \( \chi^2/4 \) by \( R \), then for a significant increase in mutation rate at a single locus,

\[
R = \frac{m}{2} \frac{(f - 1)^2}{(1/P + 1/Q)} \geq 1
\]

In terms of \( D \) and \( D_r \),

\[
R = \frac{m}{2} \left( \frac{D}{D_r} \right)^2 \left( \frac{1 + D/D_r}{P} + 1/Q \right)
\]

For example, in the study of Neel and Schull, the progeny of irradiated parents numbered \( 3.3 \times 10^4 \) and the progeny of control parents, \( 3.2 \times 10^4 \), while the average excess radiation exposure to the combined parents of the former group is about 17 rad. Because of the known heterogeneity of exposures, \( R \) for any single locus must be computed by adding together the calculated \( R \) values for the various exposure classes, which add up to \( 2.3 \times 10^2 \) on the assumption that the representative doubling dose is 30 rad. With respect to the possibility of significant findings based on mutation at any one locus, then this study (and any other study to date) is far below the level of significance.

54. Where multiple traits are involved in the inquiry, the power of the study is a function of the precise number of traits under consideration. For example, if one were to make the over-simplified assumption that mutation at any one of 100 loci resulted in completely penetrant, dominant mutations responsible for congenital defect, assuming independence in the expression of mutation at these loci, the calculated resolving power of the previously mentioned study becomes 2.3, and the failure to observe a significant effect of radiation on the frequency of congenital malformations in the aforementioned study might indicate that the assumed doubling dose was too low.

55. The sex-ratio is one of the more conveniently studied indicators of possible genetic damage. Information on this point is relatively easy to collect and has a high degree of objectivity. The calculations corresponding to those of paragraph 53 are relatively simple and proceed as follows:

Suppose a group of mothers receive gonad doses averaging \( D_m \) prior to conception of children, and suppose the irradiation causes a shift in the secondary sex-ratio \( s \) which is linear with the dose

\[
\Delta s_m = k_m D_m
\]

Suppose \( P_m \) progeny of these mothers are examined, the variance in the determination of the sex-ratio of the progeny of the group, due to limited sample size, will be

\[
\sigma^2 = \frac{s(1-s)}{P_m}
\]

Since \( s \) is always approximately \( 1/2 \), this may be written

\[
\sigma^2 = \frac{1}{4P_m}
\]

If such a group is compared with \( Q_m \) progeny of a control group the variance of the observed difference is

\[
\sigma^2 = \frac{1}{4P_m} + \frac{1}{4Q_m}
\]

and the significance of the observations is determined by

\[
\chi^2 = 4k_m^2 D_m^2 \left( \frac{Q_m}{P_m} + \frac{1}{Q_m} \right)
\]

If we require \( \chi^2 \geq 4 \) before the shift can be considered significant, then

\[
R_m = k_m^2 D_m^2 \left( \frac{Q_m}{P_m} + \frac{1}{Q_m} \right) \geq 1
\]

Similar formulae can be derived for comparison of the progeny of irradiated fathers with controls, where

\[
R_f = k_f^2 D_f^2 \left( \frac{Q_f}{P_f} + \frac{1}{Q_f} \right)
\]
A number of completed surveys, irrespective of the significance of their results, all show decreases in size when the mother is irradiated from which values of k of the order of $-1 \times 10^{-4}$/rad can be derived. If this figure is adopted for purposes of calculation, then

$$R_m = 10^{-6} D_m^2 \left( \frac{1}{P_m} + \frac{1}{Q_m} \right)$$

On the basis of the present limited information, values of $R_m$ have been calculated using a similar numerical value of k but of opposite sign

$$R_t = 10^{-6} D_t^2 \left( \frac{1}{P_t} + \frac{1}{Q_t} \right)$$

Clearly, if $k_t$ and $k_m$ do in fact differ in sign, then significant results may occasionally be obtained by the comparison of progeny of irradiated mothers with those of irradiated fathers, even where neither group differs significantly from the controls. On the basis of the numerical values adopted here, the same condition upon significance would then become

$$R_{t,m} = 10^{-6} (D_t + D_m)^2 \left( \frac{1}{P_t} + \frac{1}{P_m} \right)$$

where $P_t$ is the number of progeny of irradiated fathers examined and $P_m$ is the number of progeny of irradiated mothers examined. The resolving power of comparisons with controls of progeny both of whose parents have been exposed will, under these circumstances, involve $D_t - D_m$ and be relatively poor if the doses to the two parents are quite similar. If $k_t$ and $k_m$ were to have the same sign, the situation would be reversed. By way of example, the data of Turpin and Lejeune may be considered. In this study, $P_m$ is 136 and $Q_m$ is 236. For the purposes of this calculation, $D_m$ and $D_t$ will both be set at 450 rads. Then $R_m$ may be calculated to be 0.175. The calculated $R_t$ for the same data is 0.52. In passing, it might be noted that because of the many somatic factors thought to influence sex-ratio, one would as a matter of principle have more confidence in the genetic origin of a sex-ratio change among the offspring of irradiated fathers than among offspring of irradiated mothers.

56. That comparisons of the progeny of irradiated and non-irradiated groups must be carried out on rather a large scale, if there is to be any prospect that they will yield significant positive results, is emphasized by the high proportion of non-significant results obtained in the completed surveys of table VI. Moreover, they may require rigorous and complex analyses of controls, and therefore involve considerable effort of a very specialized kind. While negative results on a sufficient scale can be of great value in excluding the most alarming possibilities, only positive ones will suffice for a quantitative relation between dose and mutation frequency. In this connexion, a survey of the high radiation area of Kerala appears to have a potentially somewhat greater resolving power than any previously made, if an equally intensive investigation over a ten-year period is assumed.

57. At its first session, this Committee requested advice from the World Health Organization about the possibility of setting up a standard of recognition for one or more clearly recognizable medical conditions thought to be largely or solely genetic in origin. In their discussions of this, the geneticists of the study group which framed the reply of WHO made clear that they strongly questioned the feasibility of using a single condition as an indicator of the mutation level in large populations. Their feeling appeared to be based on the manifold uncertainties which exist concerning almost every single likely indicator condition, and in part upon the belief that reliability of results in this field depends upon intensive study of every case. The study group recommended that simultaneous investigations always be carried out on several conditions. Indeed, the sense of the document cited is such as to cast some doubt upon the practicability of such surveys, in view of the associated difficulties of obtaining sufficiently large numbers. It does not, however, rule out large-scale survey plans if the urgency of the situation warrants them. Moreover, if the objective were to survey one population serially in time so as to be able only to establish limits of possible relative increases in the mutation rate, without any interpretation as to cause, some of the difficulties might diminish.

One such difficulty seem to lie in combining the intensive examination of cases, which is the classical approach of human genetics, with the extensive survey of very large populations which is required if adequate numbers are to be obtained for studies of mutation rates at or near the spontaneous rate in man. This difficulty is emphasized by the sharp limit of about $3 \times 10^6$ set in discussion upon the size of human population which can be covered by an institute conducting epidemiological surveys of the classical type (see also ref. 11).

58. The difficulties of comparative surveys of high resolving power have led Penrose to propose a modified approach, by which a given class of mutant propositi would first be collected from a large population heterogeneous in radiation exposure as well as in other respects, and only then would personal histories, including radiation histories of the parents, be compiled for the propositi and a comparable control group. The method is a powerful one for the wider field of general human genetics, since it can serve as a basis for quantitative investigations of other mutagens than radiation. As applied to the radiation problem, this same possibility of alternative and perhaps unknown causes complicates the choice of a legitimate control group. Moreover, the burden of work is in part thrown into a sphere where rather considerable difficulty also prevails: the quantitative compilation of individual histories of irradiation. In order to obtain a quantitative dose-effect relation from a survey of this type, it is necessary to know not only the incidence of the condition under investigation in the general population, but also the general incidence in that population of individuals having similar radiation exposures to those of various classes of propositi. Many features of the approach are exemplified by the recent work of Steward et al. on a somatic radiation problem.

**Possible aids in extrapolation of radiation-induced mutation rates from other species to man**

59. In view of the difficulties of a formal human radiation genetics, it is necessary to consider possible ways in which radiation-induced mutation rates can be measured in systems closer to the *in vivo* germ cells of man. In this connexion a new field of work has been opened by the ability of Puck and his collaborators to grow colonies of tissue-culture cells, the majority of which are viable and able singly to give rise to fresh colo-
The well-developed methods of microbial genetics can in principle now be applied to such cultures both for natural and radiation-induced mutations, although certain features are believed by many workers still to limit the applicability of the material to this problem:

(a) Tissue-culture cells usually need a more complex medium than the whole organism from which they originate.

(b) Well-established lines tend to be poly- or aneuploid. They resemble both each other and the malignant HeLa strain, with which Puck first developed his techniques. In certain types of radiation experiment, this difficulty may be circumvented, as in the work of Bender, who used tissue-culture cells very recently derived from human kidney (within four transfers) in a cytological study of induced chromosome breaks. But the repeated propagation of lines of stable diploids from single cells appears to be a prerequisite for systematic studies of gene mutation in human tissue-culture.

(c) Some workers in the field doubt whether any line of normal (i.e. non-malignant) cells has really been successfully propagated as such (but see Puck). It is not yet known what is the exact relevance of studies on the mutational behaviour of somatic cells in vitro to that of mammalian germ cells in vivo.

Points (b) and (c) can perhaps be circumvented in part by applying the technique to cultures derived as freshly as possible from normal tissues. However, a difficulty of principle remains: the tissue-culture cell is a free living organism, whereas the ancestral tissue cell is part of an organism so that its growth, division and differentiation are subject to the developmental controls of that organism. In view of the close connexion of all, and especially the genetic, effects of radiation upon the cell with the process of cell division, some initial caution in interpretation is undoubtedly required. Nevertheless, the future role to be played by the tissue-culture methods in the making of comparisons between species so as to provide a basis for extrapolating from the known in vivo mutation rates or rates of occurrence of gross structural changes, does not seem open to doubt.

60. There is some evidence that the frequencies of radiation-induced mutations in somatic cells is similar to that in gonial cells. If this correlation could be extended to the variation between species, attempts to measure induced and/or natural mutation rates in human somatic cells in vivo might provide information of great value as a guide in estimating mutation rates in human genes.

Continued need of research in fundamental genetics

61. It cannot be too strongly emphasized that there is little basis either for planning or for interpreting ad hoc radiation genetic surveys in man, or for making calculations concerning radiation-genetic effects in man, except the great volume of fundamental research upon other organisms which has been carried out for its intrinsic interest alone. and directed wholly as a contribution to human understanding. This foundation must be extended and strengthened, and must not be weakened in the interests of the applied superstructure.

3. The representative doubling dose

62. Provided that the dose-mutation rate relation has a linear form

\[ m = m_0 + kD \]

the relative increase of mutation rate per unit dose is readily expressed by the ratio \( k/m_0 \). Another convenient parameter to use is the reciprocal of this ratio, \( m_0/k \), which is the radiation dose required to produce a number of mutations equal to those occurring naturally, or the "doubling dose" (\( D_2 \)). For a whole series of mutations \( m_i \) whose effects sum up or are collectively observed \( \Sigma m_i = \Sigma m_{o1} + D_2 k_i \), one can define a mean \( D_2 \) as

\[ \frac{\Sigma m_{o1}}{\Sigma k_i} = D_2. \]

This procedure can be used to estimate a \( D_2 \) for as representative a group of human genes as possible. It is not necessary to know how many genes are involved or of what kinds, provided that they can reasonably be assumed to be a representative sample and provided that there is assumed to be no correlation between \( D_2 \), \( k_i \), or \( m_{o1} \) and the degree or kind of manifestation. The representative \( D_2 \) should then express the dose-effect relation for any set of radiation-induced mutational events in so far as this itself depends upon a sufficiently representative sample of human genes: usually the sets will be of a kind in which the mutations at a very large number of loci are summed, both in calculating and in making use of \( D_2 \).

Estimates of the representative doubling dose for human genes

General levels in other species

63. It has been pointed out that a number of doubling doses calculated for different species cluster around the range 30-60 rad (table VIII). However, the significance of this fact for present purposes is limited by several considerations:

(1) The majority of the experimental radiation exposures concerned were of gametic cells. Where irradiation of gonial cells is concerned, it is true that the best estimate that can at present be made for a group of genes in the mouse (the only mammal so far investigated) is of the order of 30 rad, but this must be compared, for instance, with values for Drosophila ranging up to 400 rad (see table VIII).

(2) No satisfactory interpretation of the observed concurrence or range of values exists, and consequently any empirical extrapolation to man would have to rest upon an unsure basis.

(3) The lack of correlation of observed doubling doses with life-span can be interpreted as an indication that mutation at a constant rate in chronological time is not the dominant factor in determining the natural rates in the experimental species. But man is so much longer-lived than the experimental organisms that in his case an appreciable fraction of natural mutations is already quite likely to result from time-independent causes such as irradiation from natural sources. (See Penrose for a preliminary investigation of this point).

Sex ratio

64. Observations have been made of a shift of the sex-ratio in the progeny of irradiated mothers (see paragraphs 50, 55). In a first attempt to make use of the available data, Lejeune and Turpin have proposed a comparison between the effect of irradiation and the
effect of aging. These authors have calculated a significant decrease of the sex-ratio with the aging of the mother alone, the partial regression coefficient being \(-3.36 \times 10^{-4}\) for an aging of five years. Taking a value of \(-6 \times 10^{-4}\) for one rad as an estimate of the decrease of the sex-ratio following irradiation of the mother (table VI), and assuming that both decreases are related to the same extent to newly arising sex-limited detrimental mutations, they have proposed a doubling dose of

\[
-3.36 \times 10^{-4} \times 6 \div -6 \times 10^{-5} \text{ rad}\]

From age 0 to age 30 years.

Unfortunately, as these authors themselves recognize, such a calculation cannot be considered as legitimate before many problems have been solved. The needs include:

(1) A good estimate of the gonad dose effectively received by the mothers;

(2) A better estimate of the decrease of the sex-ratio with irradiation, including a test of linearity of the relationship between these quantities, which is implicit in all current calculations concerning sex-ratio;

(3) An explanation of the apparent contrast between the sex-ratio's decrease with the father's aging\(^{(114,128)}\) and the possible increase observed after acute irradiation of the father's gonads:\(^{(111,124)}\)

(4) The study of other variates such as birth rank\(^{(48)}\) which might interact with the real effect of the aging of the mother.

65. Only some preliminary data relevant to the problem of irradiation of the father are available, but these indicate that a sex-ratio decrease after chronic irradiation may perhaps have occurred in man\(^{(48)}\) and in the mouse.\(^{(48)}\) The latter body of data, although not significant at the 5 per cent level, yields at face value a representative doubling dose in satisfactory agreement with other data for this species.

66. In summary, while the possibility exists in principle of deriving a representative doubling dose by comparing the changes in secondary sex-ratio when parents either age or are irradiated, the relevant phenomena are, at present, not sufficiently well established either quantitatively or qualitatively for this procedure to be reliable. Yet relevant surveys of the secondary sex-ratio are more readily and widely carried out in human populations than are others which must depend upon finer diagnostic distinctions. Consequently, more extensive quantitative data concerning comparable irradiated and non-irradiated human populations should continue to be sought. In particular, it may be worth attempting to search for a decrease of sex-ratio among the progeny of not too heavily irradiated human males; the conclusion of such a test might go far to determine the utility of the parameter in considerations relevant to the human genetic radiation hazard.

67. It is not at present certain, even in Drosophila, whether the postulated genetic causes of shifts in the sex-ratio play the quantitative roles expected of them; and data of this kind are needed. It is also possible that further investigations upon experimental animals, especially among the progeny of male mice irradiated at low doses, together with similar observations upon irradiated female mice, may show that in both cases a doubling dose can be derived from sex-ratio shifts which is of the same magnitude as that calculated from purely mutational experiment. Establishment of such facts would greatly strengthen interpretation of corresponding observations upon man.

68. Although today it is not possible to assign any definite confidence to the use of the sex-ratio as an indication of mutation rates, it must be borne in mind that the parameter, even if not totally satisfactory, is the only one easily surveyed in entire populations, and that it represents the "cheapest" genetic trend available to research workers in terms of technical effort expended in surveys.

### Induction of leukemia

69. A reasonable probability now exists that, in an intermediate dose range, the radiation-induced incidence of leukemia is a linear function of the exposure of the bone marrow, whatever the manner of delivery of the dose. Upon this hypothesis, it has been calculated that 30-50 rad mean exposure of the red marrow might suffice to double the natural incidence of leukemia among an adult group.\(^{(128)}\)

70. Leukemia certainly involves a transmissible hereditary change in the tissue cells concerned, a "mutation" in the widest sense of the word. Whether the process of its induction in somatic cells corresponds qualitatively or quantitatively in any way to the process of apparent gene mutation as it is normally thought of in germ cells is extremely doubtful. Nevertheless, it is not entirely excluded from providing an indication of the relative sensitivity of human cells to natural and radiation-induced genetic changes. The indication must, however, be regarded with great reserve; even if the most helpful possibility eventually proved true, and leukemogenesis were primarily a process of somatic gene mutation, a single very atypical gene in a somatic cell might be responsible, and might be entirely unrepresentative of transmissible germ-line mutations.

### Survey of Japanese cities

71. Although the results were negative, the extensive observations of Neel and Schull\(^{(113)}\) in Nagasaki and Hiroshima provide some evidence of a lower limit for the representative doubling dose for human genes, at least for the dominant mutations which would have been observed by these authors. A difficulty of the type of survey conducted by Neel and Schull must be mentioned here: in order to obtain significant data, it is necessary to continue collection of it for some considerable time. Among a population who have been subjected to heterogeneous, heavy exposure, there may perhaps be some infertility of a progressive kind selectively induced among the most heavily exposed groups. In that event, incipient positive results may be masked by later data collected in the attempt to make the observations more significant. It is possible that the significance of the observations made in this kind of survey, because of its scale, complexity and uniqueness, can only be evaluated adequately by the authors. It therefore seems reasonable to accept the opinion of Neel and Schull that their negative results make it improbable that the representative doubling dose for human genes irradiated in gonial cells lies below 10 rad.

### The natural exposure

72. The representative doubling dose for human genes undergoing chronic irradiation cannot be less than the
genetically significant exposure of natural origin. In most areas this is about 3 rad per generation. In exceptional areas, the natural radiation may contribute so heavily to the natural mutation rate that the observed representative doubling dose would be increased.*

Current best estimates

73. Not one of the arguments in paragraphs 63-71 gives a reliable estimate of the representative doubling dose, yet each depends upon a different, independent set of unproven ideas. This Committee recognizes a need, in our existing state of knowledge, to make use of every available source of information, however tenuous. It considers that the separate arguments and repeated independent observations of small changes, in spite of the statistical limitations upon their significance, provide a reasonable indication when taken together; the representative doubling dose for human genes irradiated in pargametic cells is likely to lie between 10 and 100 rad. There is supplementary evidence that it cannot be less than 3 rad. The Committee notes that the value 30 rad is compatible with the whole of the probable range cited, within a factor of about 3; it therefore has a certain degree of utility for purposes of calculation wherever a 'most probable' value of the representative doubling dose is required.

4. Estimates of total rate of radiation-induced mutations in the genome of man

74. Because radiation-induced mutation has not yet been observed with certainty in man, it is not possible to give a satisfactory estimate of total induced mutation rate; indeed, this is hard enough even in Drosophila. Nevertheless, it could be hoped that the total rate might bear some relation to total genetic material: such a hope has recently been supported by the only available comparison, that between calculated total induced recessive lethal rates and DNA contents in the mouse and Drosophila.† The DNA content of human cells is about 6/5 that of mouse cells, according to Vendrelly.‡ Hence, upon the stated hypothesis, it might be expected that roughly one recessive lethal per 250 rad would be induced in human sperm by irradiation. Again by analogy with both the mouse and Drosophila, which behave alike, it might be expected that in spermatogonia only about one quarter as many gene mutations would occur. However, in Drosophila it has been estimated that the total rate of mutation to appreciably deleterious alleles is about four times the recessive lethal rate. In the assumptions made so far, it has been possible to rely upon common quantitative behaviour of two diverse species. But the induced mutation rates for single loci of mice, as well as the total recessive lethal rate, are greater than those of Drosophila by a factor of 20, corresponding approximately to the ratio of DNA contents per cell. This has suggested that perhaps the individual genes of the mouse are not more numerous but are larger and more complex than those of Drosophila. In turn, the ratio of total to recessive lethal mutations might be very greatly affected. That this is perhaps not so is suggested by comparison of the induced mutations at sets of visible recessive loci in the two organisms. In both cases, some two-thirds of the experimentally induced mutations have been found upon investigation to be lethal. The similarity could be a property peculiar to visible loci: but it at least suggests that the ratio of total to recessive lethal mutation rates may be the same for these two and possibly other species. If the Drosophila ratio is applied to man on these tenuous grounds, a total induced rate of appreciably deleterious mutations of about one for every 250 rad applied to the gonial cells is suggested. It will be clear to the reader that based as it is upon so many tenuous hypotheses, this figure, must be regarded with the very greatest reserve. In particular, it applies only to the sum of oligogenes with individually detectable effects and neglects the polygenes involved in quantitative inheritance, an especially serious omission for organisms which may have considerably larger and more complex genes than Drosophila, and may therefore be relatively much more liable to small changes giving rise to many isolales even at known loci.

11. The genetic consequences of irradiation

1. The connexion between mutation and genetic damage: Selection

75. The fate of a mutant allele newly introduced into a population is determined by selection. Hence the connexion between mutation and the genetic damage due to it depends primarily upon the selective properties of the mutant alleles concerned and, in particular, upon the degree of dominance or recessivity of these. Our ignorance of the relevant facts in man is very complete and urgently requires rectification.

76. It is useful to precede inquiry into the action of the selective process upon mutant alleles by an inquiry as to the origin of genetic variation in natural populations and its connexion with fitness. The question is an old one, especially in connexion with plant material, where the great extent of natural genetic variation was early observed, and where breeding experiments early gave rise to the controversial notion of 'hybrid vigour'. However, much of the agronomic literature is primarily concerned with the externally applied criterion of 'yield' rather than with fitness. Moreover, natural populations of plants differ decisively from those of animals in the aspects of genetic structure which are of immediate concern here.

77. What may be called the classical view of the adaptive norm of a natural population supposes the optimal allele to be homozygous at most loci: this situation, as maximum fitness is disturbed by mutation, continually restored by selection: rarely, due to chance, to change in external conditions in time or space, or to change in other parts of the genotype, a mutant allele will prove itself advantageous, displace the former predominant allele at the same locus, and become the new wild-type allele (see review in ref. 130). In recent years this view has been increasingly strongly challenged
by some, especially in connexion with the accumulation of extensive evidence concerning the prevalence and the superiority in many respects of structural heterozygotes in natural populations of *Drosophila*, a finding which is itself, however, compatible with the classical view of genic homozygosity as the adaptive norm. It has also been argued on more general grounds that heterozygosity is the adaptive norm at most loci and that heterozygotes are in fact intrinsically better able to adapt themselves and maintain their own stability in the face of changing environmental conditions. A recent experiment by Wallace seems to indicate that even random unselected radiation-induced heterozygosity in general confers an advantage, at least upon individuals otherwise homozygous for certain pairs of arbitrarily chosen chromosomes in laboratory populations of *Drosophila*.

78. These two views lead to different general expectations concerning the consequences of mutation. On the first, most mutants alleles will contribute to the limited degree of heterozygosity, will be harmful, and will require to be eliminated, diminishing the fitness of the population. On the second, mutational events, although the majority of them will still be harmful and will require to be eliminated, will scarcely affect the great degree of heterozygosity already existing, and will diminish the existing reproductive fitness to only a correspondingly small extent. However, this is a consequence of the fact that since the mating of diploid heterozygotes produces some homozygotes, on the second hypothesis the population must pay for its built-in adaptability and plasticity by a permanently reduced fitness due to these.

79. Unfortunately, while evidence now exists for the second view of natural populations of *Drosophila*, this particular organism has certain features (principally chromosomal inversions) which bestow upon it a special capacity for carrying structural heterozygotes, together with all the consequences which may flow from this capacity; these features include the absence of crossing-over in the male, coupled with a mechanism for eliminating undesirable products of cross-over between structurally different chromosomes from the egg in the female. There is no reason to suppose man to possess either this particular structural mechanism or an optimal degree of genetic heterozygosity, although the possibility is not excluded that equivalent mechanisms may be found. Hence the Committee is compelled to assume that the general genetic structure of human populations corresponds more closely to the classical model in so far as this relates to known genes having individually detectable effects. There is, however, no basis in our present limited state of knowledge for deciding whether the genes responsible for quantitative inheritance do or do not maintain themselves by overdominance in so far as they affect the over-all fitness. It must be emphasized that upon all the hypotheses discussed here, the great majority of radiation-induced mutations will be to alleles which are in the first instance harmful and unlikely to be retained in the population.

2. Approaches to Quantitative Assessment of the Genetic Consequences of Irradiation of Human Populations

80. On the classical basis, the irradiation of human populations is expected to result in mutations to alleles whose expressions are harmful and lead to their elimination: the expressions of these alleles also contribute to the genetic component of human ills.

81. As yet, nothing is known of the rate of induction by radiation of the mutations responsible for any specific condition in man. In consequence, the discussion which follows will be restricted to broad categories of effects. Only by such a grouping together of the consequences of mutation at a large group of loci can a representative rate of induction of mutations per gene, or a representative doubling dose, be applied: these are the only two parameters expressing a dose-effect relation so far available.

82. It is natural, in applying the results of an experimental science, to try to use a synthetic approach, assessing an effect from the accumulated knowledge of various causes. In the present instance, this means attempting to assess the magnitude of the social consequences of increased mutation by using mutation frequencies per rad at particular loci to build up a combined estimate from the effects of induced mutation at all loci. To use this method, let the total mutation rate to the set of alleles responsible for any specific condition denoted by $i$ be $k, D$, where $D$ is the genetically significant dose of radiation to the population. By a theorem originally due to Haldane there must on the average be $k, D$ subsequent eliminations of the mutant alleles through differential failure of reproduction. These are often referred to as genetic deaths, although they may take place through phenomena such as very early abortions, which are of no social significance, as well as through more or less severe disabilities or even premature death. Suppose a fraction $p_1$ are eliminated by socially serious expressions and think of $p_1$ as including some weighting factor whereby such qualitatively diverse end-results as death, physical disability, mental deficiency, etc. may somehow be quantitatively compared. Then the contribution to the social burden is $k, p_1, D$ and the whole contribution of the dose $D$ to the future social burden is $\sum k, p_1, D$ over all such specific conditions. The above argument continues to hold whether the mutation involved is to an allele which from the selective point of view is conditionally or unconditionally deleterious, although if the mutant allele is only conditionally deleterious then (a) it cannot be eliminated in those situations in which it is selectively favourable, and (b) the total elimination rate at any one time may greatly exceed the mutation rate, because the increased fertility of carriers under the selectively favourable conditions increases the gene frequency. If the natural mutation rate $m_1$ is known, then $k, m_1$ can be re-expressed in terms of the doubling dose $D_2$ by $k, D_2 = m_1$ and for all mutations or a large class of them a mean doubling dose $D_x$ can be defined by the equation $k, D_x = m$ where $k = \frac{7k}{m}$, $m = \frac{7m_1}$. It is unfortunate that in man we do not know any individual $k$, or $D_2$. Still more unfortunately, the fractions eliminated by socially serious expressions, $p_1$, are unknown and may depend upon rather small positive or negative fertility differentials in those who carry the mutant allele without expressing it, if they greatly out-number those in whom it is expressed. Nor can a mean $p_1$ be estimated for mutant human alleles. As a result, the synthetic approach leads to an estimate in such terms that it cannot as yet be satisfactorily related to the social consequences.

83. There is an alternative formulation of the problem by an analytic approach, based upon analysis of the
present social burden in terms of naturally occurring hereditary defects. In this, it is asked, (a) what is the social burden \( b \) due to a given condition denoted by \( i \), whose occurrence is related to the presence of adverse genes? (b) Of the genetic burden \( b_i \), what fraction \( f_i \) is due to recurrent mutation? (c) By what fraction \( g_i \) will this be increased immediately or in the future by a given fractional change \( c_i \) in the natural mutation rate \( m_i \)? If the change \( c_i \) is caused by a genetically significant dose \( D \) to the population,

\[
c_{i,m} = k_i D \text{ or } c_i = D/D_{21}
\]

For all conditions or a large class of them the total genetic burden may be written \( b = \frac{2}{F} b_i \), that due to recurrent mutation \( fb = \frac{2}{F} f_i b_i \), and that due to a given dose \( D \) as \( gb_i fb_i \). If it is assumed that \( g_i = c_i \), this may be written as

\[
D = \frac{f_i b_i}{D_{21}}
\]

It may be assumed that \( b_i \) and \( f_i \) are independent of \( D \). Then the increased burden may be written

\[
\frac{D}{D_{21}} = \frac{f_i b_i}{D_{21}} \text{ which may be written } (D/D_{21}) \frac{f}{b}
\]

where \( f = \frac{f_i b_i}{D_{21}} \). That is, the genetic burden due to a given dose equals

\[
\frac{\text{given dose}}{\text{(part of genetic burden doubled by recurrent dose mutation)}}
\]

The relation between induced mutation rate and exposure enters only through the representative doubling dose. In the present state of knowledge, the analytic approach is more certain than the synthetic approach, because the relation between induced mutation rate and exposure enters only through the representative doubling dose.

84. Even supposing the necessary quantitative relations between mutation rate and dose or radiation exposure to be known, calculation of the social consequences still requires knowledge of one of the sets of parameters, \( p_i \) or \( f_i \), dependent upon selective behaviour of the mutant alleles. The two approaches are compared from this point of view in table IX. It will be seen that, under conditions in which mutation contributes a large part of the social burden, \( f_i \) is relatively well known but \( p_i \) is not. Moreover, there is some reason to believe that most heterozygous carriers of individually detectable, socially deleterious recessive alleles are slightly less fertile than average. If this is true, most \( f_i \) are known but most \( p_i \) are not. It is concluded that, for most purposes, the analytic approach starting from the current social consequences of unfavourable alleles is to be preferred to the alternative method at the present stage of knowledge.

85. Certain assumptions are implicit but not stated in the analytic approach to the problem adopted here:

First, it has been assumed that the genetic component of the social burden is directly related to the expressed effects of unfavourable alleles. However, the actual social burden realized in a population will be modified by environmental factors such as the extent of care devoted to those affected. For this reason, the actual social burden resulting from a given genetic situation may be heaviest in those countries having the best medical care of the afflicted.

Second, the genetic component of today's social burden has been assumed to be related to the present natural rate of occurrence of mutations and to present selective conditions. Certainly this assumption is not true—the number and distribution of recessive alleles is determined by a long history of past mutation rates and past conditions of selection—yet with our present limited knowledge of the distant past and future no alternative assumption seems to present a possible basis of calculation. A number of considerations indicate that the errors involved may not be too serious:

(a) Because of recent improvements in medical care the present genetic burden may be below equilibrium with today's rates of elimination of undesirable alleles, so that the effects of a given increase in mutation rates are underestimated. On the other hand, further improvements in medical care are likely in the future to reduce the socially serious effects of mutations. This process cannot by itself affect the influence of mutation upon the Darwinian fitness of the population, but may affect the future social burden due to present mutations if it occurs without a corresponding effect upon the rates of elimination of the socially deleterious alleles. If this elimination takes place largely through rather trivial effects in heterozygous or other carriers of unexpressed alleles, alleviation of the expressions in grossly affected individuals might be accomplished with little influence upon the process of elimination. We would then have overestimated the future social burden from present mutations. Thus the two sources of error due to improving medical care act in opposite senses.

(b) In spite of changes in diet and living conditions of all kinds, there is no reason to suppose that natural mutation rates have changed very greatly; for example, chondrodystrophy, which, in man, is largely a dominant disease, has been prevalent at a low frequency since ancient times. Selection has, by contrast, certainly undergone great changes. This fact is relevant to the recommendation, contained in the report of a WHO study group submitted to this Committee, that research be initiated upon selection in primitive communities while the opportunity to do this still exists. But many of the specific detectable conditions with which we shall be concerned here either arise from dominant alleles, and hence do not in general persist for so many generations as recessives, or else they confer a reduction in selective fitness which has not yet been greatly modified by advances in medical practice. The working assumption may therefore be not too greatly in error for the broad categories of effects to be considered. In point of fact, the effect of improved living conditions and improved medical care is far from obvious. Penrose has pointed out that, besides preserving less fit individuals, this change may in recent years have removed the selective advantage of alleles which confer a degree of protection against an infectious disease in the heterozygote while being grossly deleterious in the homozygote: the classical example is sickle cell anaemia. How many such situations exist is debatable. However, the consequences of improved medical care could be called eugenic rather than dysgenic in such cases. It must also be borne in mind that the total potential intensity of selection in populations has, at least in recent years, not been changing anything like as rapidly as the qualitative basis of it. It may be observed here that the possible dys-
genic effect of future improvements in social and medical care is limited by the fact that no more deleterious mutant alleles can be saved for later generations than arise by mutation; moreover, a subsequent withdrawal of improved medical care by some social catastrophe will not cause more losses than would have occurred anyway had it never been present. Only the distribution in time will be altered. Thus, in a constant population, the dysgenic effect of a changing selection does not increase the total number of seriously affected individuals but by contrast, the dysgenic effect of increased mutation does increase the total number of seriously affected individuals. Finally, it has been assumed that radiation-induced mutations and spontaneous mutations are qualitatively similar: that there is no correlation between $D_m$ and the degree or kind of manifestation $(f_i, b, p)$ of a given mutation. This assumption has been discussed in another section and is acceptable to the Committee.

86. On the basis of the above arguments, the Committee considers:

(a) That the most satisfactory assessment of the genetic consequences of irradiation of human populations which can be attempted at the present time must be based on the present social burden due to hereditary conditions. Because it must employ the representative doubling dose, it must be restricted to rather broad categories of effects;

(b) That the sources of error in an assessment of this kind may not be too serious;

(c) That two principal sources of error are related to the extent to which selection changes in the transition from a technologically primitive to a technologically advanced environment and to the extent to which alleles responsible for socially serious conditions may confer small favourable differentials of fertility in the heterozygous, impenetrable or other "carrier" states. Both require to be investigated.

3. The current social burden of genetic origin in human populations, its connexion with mutation and radiation exposure

87. In order to make use of the representative doubling dose discussed earlier, there will be considered here only broad categories of damage, each of which may be caused by mutation at any one of many loci, such as the sums of specific clinical conditions or traits within various genetic categories, or biometrical characters such as intelligence, life-span or birthweight, each likely to be dependent upon many genes, or fertility.

Specific traits

88. For the present purpose, the available information concerning the incidence in man of specific diseases or disabilities of genetic origin is severely limited. Only very few sizeable populations have been surveyed, notably in Denmark,289 Michigan, U.S.A. and Northern Ireland. More, good quantitative data are only available for clear-cut traits or disorders, and, even here, the genetic interpretation of the facts is almost never straightforward. In the past, various estimates have been made of the frequencies of such specific traits, but the basis of the estimates has not always been clear. Sometimes it has been uncertain whether the trait frequencies referred to were those at birth or in the whole population. The latter estimate would always be expected to be lower, particularly if the trait was severe in its effects. Independent over-all estimates, both in the literature and in reports to this Committee, seem to be in reasonable superficial agreement with each other and are summarized in table X; each of these implies consideration of one or another category out of a total of some 500 clear-cut disorders or traits. However, it has seldom been specified which traits are included and which excluded in them.

89. In order to formulate, upon a precise basis, overall estimates to which a representative doubling dose can reasonably be applied, the Committee has made use in the present report of a single, definite list of traits and their estimated frequencies of appearance in a single population, namely that compiled by Stevenson288 for the population of Northern Ireland. In so doing, it is recognized that the frequencies of specific traits will be different in other populations, so that some listed here may not occur at all, and others not in the present list will be prevalent. Nevertheless, such comparisons as can be made of the population frequencies of traits in different parts of Europe, North America and Japan suggest that, while the contributions of individual traits to the total may differ considerably in different populations, the totals, and their division into principal categories, will not vary appreciably so long as present methods of detection are employed.

90. The list of traits compiled by Stevenson has been broken down into separate categories in the following manner, which differs somewhat from that used in the original compilation.

Category I (table XI (a)-(b)): Category I includes traits determined by single, harmful mutant alleles. The majority of these are dominant with a high degree of penetrance, but some are autosomal recessive and a few are sex-linked. Most are not recognizable in the affected person at birth. It seems reasonable to assume that in respect of these traits there is no significant selective pressure in either direction against apparently unaffected carriers of the mutant alleles, although this cannot be proved in our present state of knowledge. It would therefore be expected that the ultimate consequence of an increase in mutation rate at each or all of these loci would be a direct effect upon trait frequency. About 110 different mutations are required to explain these traits. No doubt some similar, but separately identifiable, traits are determined by alternative alleles. Of these mutant alleles about 72 are dominant, 30 autosomal recessive and 8 sex-linked recessive. The estimated total of live-born affected is 1.1 per cent.

Category II (table XII): Category II includes a considerable number of traits mostly detectable at birth. A proportion of them sometimes determines intrauterine death, but this fraction of these conditions is ignored in the present context. Maternal health and intra-uterine environment appear to play a considerable part in determining whether and to what degree they are expressed. Their familial patterns in a community seldom satisfy the criteria of a single mutant expression. In all there is a familial concentration of cases greater than would occur by chance. In some, the family pattern approaches some of the criteria of those included in category I, and it will be clear to the reader that arbitrary decisions have had to be made. The estimated total of live-born affected is 1.0 per cent.

Category III (table XIII (a)-(b)): Category III consists of two unequal classes of traits. The first and smaller proportion (category III (a); table XIII (a)) consists of traits which appear to follow closely the ex-
predicted family patterns of a single recessive mutant gene, but show a frequency too high to be explained on a basis of mutation pressure alone, unless it is assumed that mutation occurs many times more frequently at the relevant loci than at those loci giving rise to dominant mutations in man or than in the general range of all types of mutation in experimental animals. In the data for Northern Ireland and elsewhere in the United Kingdom, only fibrocystic disease of the pancreas and deafness clearly fall into this category, although other conditions well-known elsewhere such as sickle cell anaemia and thalassemia also belong to it. It is possible, although neither provable or disprovable at present, that the gene frequencies in these conditions are maintained mainly by relative selective advantage in the heterozygous carriers. In deaf mutism several independent mutants contribute to the trait frequency. The two conditions together determine about 37 per cent of the total frequency of recessive traits at birth in the population studied by Stevenson (loc. cer.) and have a combined frequency of 0.09 per cent of all live births. The second and larger proportion (category III (b); table XIII (b)) of category III is difficult to define and limit. Six examples of serious, "constitutional," diseases are listed in the table, but it is difficult to know where to draw the line thereafter. In different communities the frequencies will vary considerably. It is impossible to estimate frequencies without making some arbitrary decisions as to what will be included; as an example, admission to hospital might be made a criterion. Furthermore, the frequencies depend on the ages to which people live in populations, as in the cases of diabetes and the primary pre-senile psychoses. Finally, there are environmental factors of importance which vary in different populations. In sum, at least 1.5 per cent of those liveborn will suffer from one or another of this group of disorders.

91. It must be emphasized that the list of traits, trait frequencies and categories outlined above and in tables XI-XIII:

(a) Represents only tangible or detectable genetic damage, which in principle, although in practice with great difficulty only, can be assessed by "counting heads";

(b) Includes only defects of such severity as to be at least very inconvenient to their possessors;

(c) Is certainly an incomplete list, even of such conditions;

(d) Ignores maternal/foetal incompatibility and mongolism; in the latter the genetic component appears to be weak, and in the former the relative frequency of the alleles, which is the most important factor in determining proportions of affected infants, would probably not be affected appreciably by increased mutation rates;

(e) Excludes a group of individually rare or mild traits which mostly appear to be determined by simple, irregular, dominant genes and are listed in table XIV. Nevertheless, the list gives rise to the expectation that some 4 per cent of the liveborn suffer or will suffer from defects predominantly of genetic origin. Certain comments are pertinent to this estimate:

(1) Any present over-all estimate of total genetic damage must of necessity be minimal. However, even though more sophisticated methods of detection can be expected to increase the present estimates, it is unlikely that in the near future more than a very small number of new specific traits will be discovered, relative to the total so far known. (See also paragraph 104 below.)

(2) The present estimates refer to those born alive. In addition, approximately another 3/4 to 3/2 per cent of foetuses alive after the twenty-eighth week of pregnancy are born dead mainly by reason of detectable developmental defects which may be of genetic origin.

(3) In about half of the affected liveborn, the defect will be detectable at or soon after birth, but in the other half the expression of the genotype will only be apparent in later childhood or in adult life.

92. The division of the 4 per cent affected live-births into categories as outlined above may be summarized as follows:

Category I: About 1 per cent of defects due to single mutants of classical type (majority not recognizable at birth);

Category II: About 1 per cent showing no consistent familial pattern compatible with a simple genetic hypothesis and often having an environmental component in their aetiology (majority recognizable at birth);

Category III: About 1.6 per cent either (a) show trait frequencies too high to be maintained by mutation pressure, or (b) determine constitutional illnesses whose frequency is also unexpectedly high in relation to their severity.

This division into categories is of great importance for predicting the results of increased exposures of populations to ionizing radiations. The supply of recognizably disadvantageous mutant alleles in a population may be maintained either by recurrent mutations balanced by selection or by selective advantage among individuals in whom the disadvantage is not expressed; that is, by a balance between opposing selective forces. A reasonably small increase in mutation rate cannot be expected to affect greatly the pattern of gene elimination and so should cause at equilibrium an equal fractional increase in the genetic damage due to alleles maintained in the first manner (corresponding to traits in category I above, together with an unknown fraction of those in categories II and III), but a much smaller increase in the genetic damage due to alleles mentioned in the second manner (corresponding to an unknown fraction of the traits in categories II and III above). It follows that permanent exposure of a population to an extra genetically significant dose D per generation may be expected eventually to give rise to an increase in the incidence of live births who are or will be affected of between 4P/D per cent and 4D/D per cent where D is the representative dose. If the increased irradiation were to occur in only one generation to a population of fixed breeding size P, it follows, by a principle of detailed balancing, that the calculated total number of affected live births is expected to lie between

$$\frac{D}{D_2} \times \frac{P}{100} \text{ and } \frac{D}{D_2} \times \frac{4P}{100}$$

93. It must be borne in mind that the mutant alleles concerned in the above estimates range all the way from severe dominant to true recessive, and the time during which the genetic damage either climbs to equilibrium or completes its expression after exposure of a single generation varies in turn from one or two to many tens of generations. Thus, in the case of irradiation of the present population, the damage may well become expressed under social and technological conditions which
cannot even be imagined today, and which may grossly affect the relation between gene elimination and its social consequences. Some geneticists therefore question the utility of assessment of a hazard so far in the future.\textsuperscript{143}

94. In conclusion, it must be emphasized that even for this most tangible kind of genetic damage, far more work is needed on family studies, on sib-correlations, incidence in consanguineous marriages, twin studies etc., so as to establish more accurately the genetic nature of the traits listed here and other conditions. If Governments wish to know the genetic health of their peoples it will be necessary for them to support the necessary work. It has been argued that, at present, populations under review by single institutes of human genetics cannot conveniently exceed $3 \times 10^9$.\textsuperscript{11} However, the problems related to both the scale and scope of such work involve questions of general medical education and co-operation as well as legal and administrative aspects which merit the attention both of Governments and public health authorities. For example, a number of human geneticists feel that the present Manual of the International Statistical Classification of Diseases, Injuries and Causes of Death is inadequate in its present scope and form for scientific purposes in the classification of congenital conditions.

\textbf{Biometrical characters}

95. Many important characteristics of man, among which specific mention must be made of intelligence, life-span and birthweight, vary continuously in natural populations about some mean which is often close to a selectively optimal value. Where this is true, selective action may act on the phenotype quite largely by reducing the variance, rather than by shifting the mean; to that extent, it is normalizing or stabilizing selection.\textsuperscript{148} Such quantitative variation is often influenced by many genes in combination whose separate effects cannot be distinguished, in contrast to those exhibiting specific qualitative effects and discussed above. These genes can only be studied statistically, principally through that part of the variance of the character for which they are responsible. This variance may be of considerable importance as a social burden or loss of population fitness. Discussion of the consequences of possible shifts in the mean of such characters will be deferred to paragraph 99 below.

96. The genetic component of the variance has been tentatively estimated in the case of birthweight by Penrose and by Robson as some 40 per cent.\textsuperscript{149,152} Half of it associated with maternal genotype, and in the case of intelligence as $\frac{3}{4}$, or perhaps as high as $\frac{4}{5}$.\textsuperscript{150} In each of these cases, the more extreme phenotypes of the distribution are observed to be associated with a loss in viability or reproductive fitness and with social burden. Thus on the basis of Penrose's\textsuperscript{149} and Karn and Penrose's\textsuperscript{151} work it can be estimated (see appendix) that the genetic component of this variance was associated with the occurrence of some 1.6 per cent of stillbirths and neo-natal deaths among males. Mather\textsuperscript{153} has calculated that on an intelligence quotient scale normalized to mean 100 and standard deviation 15, 2.3 per cent of children will fall below intelligence quotient 70, and a doubling of the heritable component of variance, assuming no shift in the mean, would increase this number by a factor which may lie between 2.2 and 2.9; this calculation depends upon the assumption of a Gaussian distribution of the measured variable at the tails of the distribution, where the assumption is itself least sure and confers greatest uncertainty. Mather's calculation is a useful guide to the upper limit of the social burden expected to be conferred by radiation-induced genetic changes in the variance of intelligence (but see footnote to paragraph 102 below).

\textbf{Relationship of genetic component of variance to mutation}

97. The relationships of selection, genetic variability and mutation in a character of relatively low selective importance (bristle number) have been studied by several authors in Drosophila. In particular, Clayton and Robertson\textsuperscript{144} have been able to show that the natural additive genetic variance in an outbred population, from which their experimental flies were originally drawn, exceeded the spontaneous increase in genetic variance per generation by a factor of 1,000, and observations by Paxman (quoted by Mather\textsuperscript{157,158}) support this conclusion. By comparing irradiated and unirradiated populations, Clayton and Robertson further showed that some $10^8\text{r}$ would have been needed to produce an increment equal to the natural variance. With selective neutrality such a genetic variance in the natural population is perfectly compatible with an established equilibrium between mutation and a degree of inbreeding due to limited effective population size. For a character of greater selective importance the genetic variance displayed by the population would exceed the increment per generation by a correspondingly smaller factor. Haldane\textsuperscript{146} has pointed out that, in the cited case of birthweight, selection removes 10 per cent of the observable variation per generation. If this selection makes no distinction between variation of genetic and of environmental origin, it poses the question: How is the genetic component of variance maintained?

98. Robertson\textsuperscript{155} has recently discussed the theoretical consequences of selection for optimal central phenotype. It appears that this process cannot of itself maintain genetic variability, even though heterozygotes have intermediate values of the character (see Fisher\textsuperscript{159}). The genetic variation must therefore be maintained either by the selective advantage, in some circumstances, of the heterozygotes as such (i.e., the majority of the genes are individually heterotic) or by mutation. Lerner\textsuperscript{163} has argued for expecting heterosis among genes of this kind, his argument being based partly upon an anticipation of improved buffering or canalization in the developmental processes of heterozygotes and partly upon experimental evidence (of which, however, a considerable fraction is drawn from Drosophila) and general experience of inbreeding. Paxman\textsuperscript{157,158} has, on the other hand, failed to find evidence of such heterosis, despite a search for it. Thus evidence of the necessary heterosis is by no means conclusive and further data are much needed. At the same time, the high rate of selective elimination does not seem compatible with replacement by mutation at the low rates observed in experiment. This difficulty may well, however, be less than it seems, because where many genes of similar effect contribute to the variation of a character, only a portion of the total genetic variability present in the population is manifest as variation actually observable by difference among the phenotypes of the individuals. In a polygenic system, alleles at different loci can exert their actions in opposite directions and thus balance out one another's effects, so that some of the variation lies hidden as balanced differences within the genotypes of the individuals.\textsuperscript{167} The proportion of the total variability so hidden increases directly with the
number of genes in the system, and it may go up even higher if the genes are linked. The hidden variability is released by recombination of the genes which balance one another, to become exposed as phenotypic differences, and this rather than mutation is the immediate source of replenishment of the observable variation eliminated by natural selection. Ultimately replenishment must depend on mutation; but, by virtue of the reservoir of hidden variability, the accumulation of new variation from mutation need balance loss through selection only in the long term. Thus the rate of selective elimination observed at any given time need not provide a reliable indication of the rate at which new variation is arising by mutation. Furthermore, the selective elimination of any fraction of the observable variation represents the loss of a much smaller fraction of the total genetic variability. Thus with birthweight, 10 per cent of the observable variation is eliminated in each generation, but this loss could represent as low a fraction as 1 per cent of the total genetic variability for this character in the population if it depended on the simultaneous action of no more than 10 polygenes. Mutational increments quite low in relation to the total variability might thus suffice to maintain the plyphic variation of a character against the erosion of selection. This is a matter on which more data are needed; but pending their appearance it would seem conservative to suppose that \( \frac{1}{10} \) of the genetic variability of most quantitative characters is the greatest fraction which it is necessary to envisage as replenished by mutation in each generation, and the fraction may indeed generally be very much smaller than this. The Committee emphasizes, however, that there is at present no satisfactory experimental basis for determining whether this fraction is large or small even in experimental species, much less in man. Clearly, further data are much needed in this whole area.

**Shifts in mean values of metrical characters**

99. Besides contributing to the variance of a metrical character, genetic factors may impose a social burden by affecting the position of its mean. Three quantities must be considered: the population mean, the selective optimum and the social optimum. The three may all differ, as is illustrated in table XV for the characters mentioned in paragraph 95.

100. The great majority of well-studied single-locus mutants in experimental organisms are hypomorphic.\textsuperscript{166,169} That is, they appear to lead to a reduction in the function or character most immediately affected. There is good a priori reason to expect this, as random interference with a complex machine will more often be destructive than constructive. In consequence it might be expected that most mutations and mutant alleles would act so as to diminish the population mean relative to the selective optimum. However, it must be questioned whether there are sufficient grounds for extrapolating this view to polygenes affecting quantitative characters. Provided that the changes are not so large that they excessively disrupt the organism's general control of the developmental channels concerned, is it not just as reasonable to suppose that a particular organ of social import—for example, the brain—may in fact benefit from hypomorphic changes in most other organs, due to a compensating diversion of resources, so that many such changes would be hypermorphemic for it? Among the characters of table XV, it is of interest that the facts concerning birthweight\textsuperscript{161} fit the classical expectation, but that those concerning intelligence\textsuperscript{161} possibly do not.

101. In the case of birthweight, it can be calculated (see appendix) that the difference by which Karn and Penrose observed the selective optimum to exceed the population mean in males is associated with 0.4 times as many deaths at or near birth as the total variance and about 0.7 times as many as the estimated genetic component of variance. What proportion of this deviation is genetic in origin is not known, but it is clear from the arguments outlined above that recurrent mutation could easily be the principal cause; if so, continued application of a doubling dose to every generation might eventually bring about an increased incidence of some 1.2 per cent in the deaths at or near birth. This selective optimum serves as to diminish the difference between the mean and the selective optimum by about 7 per cent per generation.\textsuperscript{16} If it does not distinguish between genetic and environmental components of the difference, the genetic effects of an altered mutation rate upon the mean must be expected to be spread over some ten generations, and any shift to a new equilibrium value will take a comparable period of time.

102. The case of intelligence is somewhat different. Here the social optimum lies far away from the selective optimum and it is not at all easy to decide what must be computed to assess the social implications of a given change. Moreover, the genetic picture is complicated by a high degree of phenotypic assortative mating.\textsuperscript{168} For the purposes of this report, Mather's calculations\textsuperscript{162} based on United Kingdom figures have been available. Mather based his calculations concerning the effect of increased variance upon an unchanged mean but he also considered a situation in which increased mutation was associated also with a falling mean, such that the effects mediated through mean and variance were roughly comparable in magnitude. However, there is no indication in the figures at present available for the United Kingdom that the population mean lies below the selective optimum; this gives rise to a presumption that increased mutation might not depress the mean appreciably. It seems important to try to find out if this situation is true and, if so, whether it is peculiar to the somewhat special demographic situation in the United Kingdom or is more general, since it raises a question as to how such a position might arise and be maintained.\textsuperscript{169} In the meantime, it seems premature to attempt here any assessment of the expected effects of increased mutation rate upon mean intelligence. The social consequence of hereditary shifts in intelligence probably occur mainly as a result of shift in the numbers at the extremes of the I.Q. distribution (of which only changes at the lower end are numerically cited in para. 96 above);\textsuperscript{*} a change in variance will in any event affect these more markedly than an equal change in the mean. Part of the difficulty in discussing shifts of the mean intelligence as measured by intelligence quotient may lie in the need to consider small intelligence quotient differences; it is possible that some of the effects of intelligence are not sufficiently well-developed and free from bias associated with other variables to serve as suitable material for close quantitative analysis. The problem of further progress in this field may thus depend upon developments in pure human biology. In any

\* An increase in variance without change in mean also causes an increase in the class frequency with highest I.Q. on which the social burden represented by a calculated increase in the numbers of individuals with I.Q. < 70. It must be borne in mind that there is some reason to believe that the distribution of variance due to new mutations would not be symmetrical, and that most of the increase would be in the direction of lowered intelligence.
over-all discussion of intelligence, it is necessary to bear in mind that it is affected not only as a biometrical character by many genes with small interacting effects, but by known specific loci, radiation-induced mutations at which will almost always cause serious harm to any individual in whom the mutant alleles are expressed.

103. In the case of the life-span, the data of Russell on the progeny of male mice irradiated by fast neutrons suggest the existence of the kind of effect which would be expected from classical hypomorphic mutations: that is, the occurrence of radiation-induced mutations to a series of weakly dominant alleles which collectively cause a shortening of the life-span. However, the magnitude of any corresponding effect which might be expected in man is entirely unknown. It might seem at first sight as if increased variance in the physiological processes, the reproductive period might be adversely affected, and the selective optimum for life-span might then be very long. It is essential that the work of Russell be confirmed and extended in order to have an adequate experimental basis in other organisms for consideration of the possible implications for man. Russell's experiments are in line with effects observed in irradiated mammalian tissue culture cells and other organisms, among which the survivors frequently carry slightly deleterious dominant alleles, as well as with observed correlations between the life-spans of related individuals suggestive of genetic influences.

Fertility

104. The most direct expression of the effect of undesirable mutations is through the net reproduction per generation or fertility differentials. Penrose has suggested that, in man, some 50 per cent of the zygotes of each generation fail to contribute to the next one by reproduction, and has suggested, by analogy with other metrical characters, that some half of this might be of genetic origin. Penrose also points out that, on the same analogy, much of the infertility might well be due to the presence of conditionally deleterious alleles which are not primarily maintained by mutation and are essentially unaffected by changes in mutation rate. However, one may compare such a rate of elimination with an estimate of the total rate of mutation to unconditionally deleterious alleles such as was derived in paragraph 74 above.

105. Applying a representative doubling dose of 30 rad to the estimate of paragraph 74, the natural rate of mutation to deleterious alleles would amount to some 14 per cent (i.e., approximately 30/250) per haploid gamete or 1/2 per diploid zygote. At equilibrium, these could be eliminated by 1/4 of zygotes failing to reproduce. These estimates of mutation are therefore consistent with that of Penrose concerning fertility, and with the assumption of genetic equilibrium, which suggests the possibility that at present 1/4 of all zygotes fail to contribute to the next generation because of the presence of deleterious alleles maintained by recurrent mutation. Taking this to be an upper estimate, indefinite application of a doubling dose to each generation might eventually extend the fraction of non-contributing zygotes from 1/2 to 1/4 and require a doubling of average family size for a previously constant population to maintain itself. This appears to be within human capacity. If it be further supposed that the mixture of dominants and recessives concerned has an average persistence in the population of 10-100 generations, then exposure of one generation to 10 or 100 times the doubling dose would impose the equivalent of the same load for a period of 10 or 100 generations. Such doses are of the magnitude 300-3000 rad, and in a range which is such as to render further considerations of genetic problems redundant. It therefore seems probable that the human race has ample breeding capacity to survive the genetic consequences of any foreseeable radiation exposure.

Pool of recessive mutants

106. Examination of the offspring of consanguineous marriages can give information concerning the total of deleterious recessive mutant alleles in a population, and Morton, Crow and Muller have recently shown how the results of statistical surveys of this kind can be expressed in the form of a number of lethal equivalents per member of the population. In the absence of a figure operationally equivalent to the total number of genes per individual, this information does not relate directly to the social burden upon the population nor, assuming an average dominance, can it be related to the natural mutation rate. The number of lethal equivalents per head is, however, in its own right, a most important parameter describing the genetic state of a population, derivable from a purely demographic type of information. Governments would do well to investigate it in their populations.

107. It is also possible in principle to compare the number of lethal equivalents, derived from vital statistical information, with the number of recessive deleterious genes found in the direct intensive surveys of smaller numbers of consanguineous marriages. Ideally, such studies should cover the whole period during which identical alleles are together and so liable to give rise to an effect through homozygosity; thus not only the number and viability but also the fertility of the progeny of consanguineous marriages should be investigated: a preliminary study of this kind has been initiated by Fraser. Such a comparison could be of great importance as indicating what fraction of the total recessive deleterious pool we know about, through recognizable specific effects. At the present time the evidence of both kinds is very scanty. By direct examination of a north Swedish population, Böök has estimated that about three recessive deleterious genes are carried per individual. However, using the criteria of Stevenson, this figure would be only 0.8-1.7, determined himself, in a somewhat smaller sample, has found 0.5-0.9. It is not possible to estimate accurately what is the reproductive fitness of the afflicted individuals, relative to the general population, but it is reasonable to suppose that the average would lie between 20 per cent and 80 per cent. Probably, therefore, the best direct intensive investigations today show up about 0.2-0.8 post-natal lethal equivalents per individual in the general population. Following Morton, Crow and Muller's analysis of the work of Sutter and Tabah but excluding stillbirths and neo-natal deaths, it is likely that a total of some 2-2.8 post-natal lethal equivalents per individual are present in their population. This suggests that present recognition encompasses somewhere between 7 per cent and 40 per cent of the total deleterious recessive damage which arises. These numerical figures reflect the strictness of the particular criterion employed by Stevenson in his use of the term "recessive".

108. The specific genetic conditions whose incidence is reported by Stevenson are divided into dominant and
recessive conditions; a most striking feature of the data is that the total incidence of rare dominant conditions exceeds that of recessives by a factor of 10. If a correction is applied for those recessive conditions not at present recognized, using the figures derived in the preceding paragraph, the ratio of total incidence of recessive conditions to total incidence of dominant conditions shifts from 0.1 to between 0.25 and 1.4 and the total incidence is increased by a factor of between 1.2 and 2.3. The calculation is now relatively insensitive to the exact criterion of recessivity employed, provided that it is the same throughout. It perhaps serves to give some idea of the limits of confidence which can be placed upon current estimates of the genetic social burden due to specific recognizable conditions.

109. It is of some interest to compare the ratio of the observed rates of elimination of deleterious recessive and dominant alleles, corrected as in the previous paragraph, with that to be expected from equilibrium with forward mutations. In mice, the ratio of recessive to dominant lethals occurring naturally appears to be about 2.5:1 or 3:1.\textsuperscript{167,168} (It is very different for \textit{Drosophila}, perhaps as low as 0.1:1, but if \textit{Drosophila} has much less complex genes than mouse or man, it may be a poor guide.) If the ratio of natural rates is similar in man and mouse, the corrected ratio of elimination rates, between 0.5:1 and 2.8:1, is in reasonable agreement with it, but suggests that the recessive alleles might, if anything, tend to be on the average slightly deleterious rather than advantageous in the heterozygous state.
TABLE I. MEASURED OR CALCULATED VALUES OF NATURAL MUTATION RATES IN MAN

<table>
<thead>
<tr>
<th>Class of mutants studied</th>
<th>Mutants per tested gamete</th>
</tr>
</thead>
<tbody>
<tr>
<td>Autosomal dominants (direct observation)</td>
<td></td>
</tr>
<tr>
<td>Epilalia..........................</td>
<td>8 x 10^{-4}</td>
</tr>
<tr>
<td>Achondroplasia....................</td>
<td>45 x 10^{-4}</td>
</tr>
<tr>
<td>Amniria..........................</td>
<td>5 x 10^{-4}</td>
</tr>
<tr>
<td>Retinoblastoma...................</td>
<td>4-23 x 10^{-4}</td>
</tr>
<tr>
<td>Partial albinism with deafness.....</td>
<td>4 x 10^{-4}</td>
</tr>
<tr>
<td>Microphthalmos....................</td>
<td>5 x 10^{-4}</td>
</tr>
<tr>
<td>Neurofibromatosis..................</td>
<td>1.3-2.5 x 10^{-4}</td>
</tr>
<tr>
<td>Average of 7 loci..................</td>
<td>4 x 10^{-4}</td>
</tr>
<tr>
<td>Rare dominants</td>
<td></td>
</tr>
<tr>
<td>Porcupine..................................</td>
<td>&lt;10^{-9}</td>
</tr>
<tr>
<td>Sex-linked recessives (direct)</td>
<td></td>
</tr>
<tr>
<td>Hemophilia..........................</td>
<td>3 x 10^{-5}</td>
</tr>
<tr>
<td>Duchenne's type muscular dystrophy..</td>
<td>4-10 x 10^{-5}</td>
</tr>
<tr>
<td>Autosomal recessives (indirect)</td>
<td></td>
</tr>
<tr>
<td>Albinism............................</td>
<td>2.8 x 10^{-5}</td>
</tr>
<tr>
<td>Ichthyosis congenita................</td>
<td>1.1 x 10^{-5}</td>
</tr>
<tr>
<td>Total colour blindness.............</td>
<td>2.8 x 10^{-5}</td>
</tr>
<tr>
<td>Infantile amaurotic idiocy.........</td>
<td>1.1 x 10^{-5}</td>
</tr>
<tr>
<td>Amyotonia congenita................</td>
<td>2.0 x 10^{-5}</td>
</tr>
<tr>
<td>True microcephaly...................</td>
<td>4.9 x 10^{-5}</td>
</tr>
<tr>
<td>Phenylketonuria....................</td>
<td>2.5 x 10^{-5}</td>
</tr>
<tr>
<td>Average of 7 loci...................</td>
<td>2.4 x 10^{-5}</td>
</tr>
</tbody>
</table>

* Very rough estimate; see ref. 83.

TABLE II. MEASURED OR CALCULATED VALUES OF NATURAL MUTATION RATES AT SINGLE LOCI OF ORGANISMS OTHER THAN MAN

<table>
<thead>
<tr>
<th>Class of mutants studied</th>
<th>Mutants per tested gamete</th>
</tr>
</thead>
<tbody>
<tr>
<td>D. melanogaster</td>
<td></td>
</tr>
<tr>
<td>Average for 9 sex-linked recessive visibles in XXY female..........</td>
<td>3 x 10^{-6}</td>
</tr>
<tr>
<td>Average for 4 autosomal recessive visibles in Oregon-R females..</td>
<td>2.5 x 10^{-6}</td>
</tr>
<tr>
<td>Average for 4 autosomal recessive visibles in Oregon-R males...</td>
<td>4.3 x 10^{-6}</td>
</tr>
<tr>
<td>Average for about 12 sex-linked recessive visibles in Oregon-R females........</td>
<td>2.4 x 10^{-6}</td>
</tr>
<tr>
<td>White eye.........................</td>
<td>0.7-3.7 x 10^{-5}</td>
</tr>
<tr>
<td>8 sex-linked recessive visibles in mutable Florida stock..........</td>
<td>3 x 10^{-5}</td>
</tr>
<tr>
<td>Mice</td>
<td></td>
</tr>
<tr>
<td>Average of 7 autosomal recessive visibles in male ca. 7 x 10^{-6}</td>
<td>Russell, Carter et al.</td>
</tr>
<tr>
<td>Bacteria</td>
<td></td>
</tr>
<tr>
<td>Average of about 30 biochemical back-mutations.</td>
<td>4.5 x 10^{-9}</td>
</tr>
<tr>
<td>Range of above... 10^{-14} to 4 x 10^{-4}</td>
<td>4.5 x 10^{-9}</td>
</tr>
</tbody>
</table>

* But approximately 3 x 10^{-4} if allowance is made for the fact that the rate of sex-linked recessive lethals was abnormally high in this experiment. Drosophila rates vary very widely with stage of life, cell-development, etc.

TABLE III. MEASURED OR CALCULATED VALUES OF TOTAL NATURAL MUTATION RATES FOR CLASSES OF LOCI IN ORGANISMS OTHER THAN MAN

<table>
<thead>
<tr>
<th>Class of mutants studied</th>
<th>Mutants per tested gamete</th>
</tr>
</thead>
<tbody>
<tr>
<td>D. melanogaster</td>
<td></td>
</tr>
<tr>
<td>Sex-linked recessive lethals:</td>
<td></td>
</tr>
<tr>
<td>young sperm..............</td>
<td>1.0 x 10^{-3}</td>
</tr>
<tr>
<td>aged sperm...............</td>
<td>2.0 x 10^{-3}</td>
</tr>
<tr>
<td>range for various wild type stocks........</td>
<td>0.7-11 x 10^{-3}</td>
</tr>
<tr>
<td>mutable Florida stock.....</td>
<td>1.1 x 10^{-3}</td>
</tr>
<tr>
<td>XXY females...............</td>
<td>7.0 x 10^{-3}</td>
</tr>
<tr>
<td>1.8 x 10^{-3}</td>
<td>Muller</td>
</tr>
</tbody>
</table>

TABLE IV. RATES OF RADIATION-INDUCED MUTATIONS AT SINGLE LOCUS IN ORGANISMS OTHER THAN MAN

<table>
<thead>
<tr>
<th>Loci studied</th>
<th>Mutations/locus/r</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>D. melanogaster</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average of 9 recessive visible autosomals in oocytes, oogonia</td>
<td>1.4 x 10^{-8}</td>
<td>Muller, Valencia and Valencia</td>
</tr>
<tr>
<td>Average of 9 recessive visible autosomals:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>spermatogonia...</td>
<td>1.5 x 10^{-5}</td>
<td>Alexander</td>
</tr>
<tr>
<td>mature sperm...</td>
<td>6 x 10^{-5}</td>
<td>Alexander</td>
</tr>
<tr>
<td>mature sperm...</td>
<td>4.4 x 10^{-6}</td>
<td>Patterson</td>
</tr>
<tr>
<td>mature sperm...</td>
<td>5.2 x 10^{-8}</td>
<td>Demerec</td>
</tr>
<tr>
<td>White eye mature sperm..................</td>
<td>0.8-1.2 x 10^{-7}</td>
<td>Bonnier and Lünung</td>
</tr>
<tr>
<td>D. virilis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average of 7 sex-linked recessive visibles:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>mature sperm...</td>
<td>7.6 x 10^{-6}</td>
<td>Girvin</td>
</tr>
<tr>
<td>E. coli</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average of about 30 biochemical back-mutations........</td>
<td>2.7 x 10^{-9}</td>
<td>Glover in Demerec et al.</td>
</tr>
<tr>
<td>Mice</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average of 7 recessive visible autosomals:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>spermatogonia...</td>
<td>2.5 x 10^{-7}</td>
<td>Russell</td>
</tr>
</tbody>
</table>

TABLE V. TOTAL RATES OF RADIATION-INDUCED MUTATIONS IN CLASSES OF LOCI IN ORGANISMS OTHER THAN MAN

<table>
<thead>
<tr>
<th>Mutations/r</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>D. melanogaster</td>
<td></td>
</tr>
<tr>
<td>Sex-linked recessive lethals in</td>
<td></td>
</tr>
<tr>
<td>aged sperm..............</td>
<td>2.3 x 10^{-5}</td>
</tr>
<tr>
<td>young sperm...............</td>
<td>2.8 x 10^{-5}</td>
</tr>
</tbody>
</table>
### Table VI. Surveys of human populations for purposes of radiation genetics

<table>
<thead>
<tr>
<th></th>
<th></th>
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<th></th>
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<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Survey of pregnancies in Hiroshima and Nagasaki</td>
<td>Neel and Schull111</td>
<td>approx. 27,000 ♀</td>
<td>8-200</td>
<td>546</td>
<td>408</td>
<td>300</td>
<td>234</td>
<td>regression lines based upon several doses: for raw data see Ref. 111, Chapter VII.</td>
<td>Observations include sex-ratio, frequency of stillbirths and neonatal deaths, birthweight, occurrence of congenital malformations, and, for a random 30% sample reexamined at age 9 months, certain bodily measurements and the occurrence of additional malformations not apparent at birth.</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>approx. 14,000 ♂</td>
<td>8-200</td>
<td>33,181</td>
<td>31,559</td>
<td>33,527</td>
<td>31,904</td>
<td></td>
<td>Non-significant decrease in sex-ratio k = (-5.5 \times 10^{-6}) rad for irradiated mothers: among earlier births1012 k was (-8 \times 10^{-6}) and significant.</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Survey of offspring of French patients</td>
<td>Turpin, Lejeune and Rethoré17, 155</td>
<td>289 ♂</td>
<td>4-450(^a)</td>
<td>47</td>
<td>46</td>
<td>8</td>
<td>7</td>
<td>9</td>
<td>8</td>
<td>225</td>
<td>358</td>
<td>Diminution of sex-ratio for irradiated mothers significant compared to irradiated fathers (x^2 = 4.2), not significant compared to controls. k between (6 \times 10^{-4}) rad and (12 \times 10^{-4}) rad.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>97 ♀</td>
<td>40-450(^b)</td>
<td>26</td>
<td>18</td>
<td>7</td>
<td>5</td>
<td>5</td>
<td>1</td>
<td>63</td>
<td>130</td>
<td>No account taken of age and parity. Inquiry by questionnaire.</td>
</tr>
<tr>
<td>Follow-up on progeny of women treated for sterility</td>
<td>Kaplan159</td>
<td>311 ♀</td>
<td>ca. 60</td>
<td>91</td>
<td>2</td>
<td>3</td>
<td></td>
<td></td>
<td></td>
<td>191</td>
<td></td>
<td>No control as yet. Very low sex-ratio, significantly different from 0.515, k (\sim -8 \times 10^{-4}) rad.</td>
</tr>
<tr>
<td>Surveys of children of American radiologists</td>
<td>Crow160</td>
<td>654 ♂</td>
<td>Unknown accumulation of many small doses</td>
<td>766</td>
<td>548</td>
<td>328</td>
<td>216</td>
<td>2,090</td>
<td>1,766</td>
<td>Non-significant increase in still-births and abortions. Significant increase in malformations (x^2 = 6.7) includes many very slight malformations or other diseases, but remains significant if restricted to cardiac malformations only. Inquiry by questionnaire.</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>5,461 ♀</td>
<td>Unknown accumulation of many small doses</td>
<td>5,461</td>
<td>4,484</td>
<td>4,484</td>
<td>4,127</td>
<td>3,390</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

\(^a\) Earlier literature surveys by Maurer108 and by Murphy and Goldstein107 respectively revealed 7, 229 and 7, 417 malformations among mothers who had received heavy doses, but without controls.

\(^b\) Taking the gonad dose to be 1/3 the skin dose.
TABLE VII. CONTENT OF DNA IN VARIOUS TYPES OF CELLS

<table>
<thead>
<tr>
<th>Organism and cell types</th>
<th>gm DNA-phosphorus per cell</th>
<th>gm DNA per cell</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria.............. B. lact. aerog.</td>
<td>2 x 10^{-15}</td>
<td></td>
</tr>
<tr>
<td>E. coli</td>
<td>2.3 x 10^{-15}</td>
<td></td>
</tr>
<tr>
<td>(compare T2 bacteriophage)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Microbes.............. Penicillium</td>
<td>1.5 x 10^{-13}</td>
<td></td>
</tr>
<tr>
<td>Aspergillus</td>
<td>1.9 x 10^{-12}</td>
<td></td>
</tr>
<tr>
<td>Yeast</td>
<td>6.2 x 10^{-15}</td>
<td></td>
</tr>
<tr>
<td>Drosophila.............. Salivary glands</td>
<td>2.6 x 10^{-11}</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2.8 x 10^{-11}</td>
<td></td>
</tr>
<tr>
<td>Rat...................... Diploid cells</td>
<td>0.6-1.0 x 10^{-12}</td>
<td>1.7 x 10^{-13}</td>
</tr>
<tr>
<td>Mouse..................... Submaxillary glands (diploid)</td>
<td>0.7-1.4 x 10^{-12}</td>
<td></td>
</tr>
<tr>
<td>Man....................... B.M.</td>
<td>8.7 x 10^{-12}</td>
<td></td>
</tr>
<tr>
<td></td>
<td>8.6 x 10^{-12}</td>
<td></td>
</tr>
<tr>
<td></td>
<td>7.0 x 10^{-12}</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.0 x 10^{-12}</td>
<td></td>
</tr>
<tr>
<td></td>
<td>8.7 x 10^{-13}</td>
<td></td>
</tr>
</tbody>
</table>

*For a further extensive table, see ref. 21.

TABLE VIII. CALCULATED DOUBLING DOSES IN ORGANISMS OTHER THAN MAN

<table>
<thead>
<tr>
<th>Organism</th>
<th>Loci</th>
<th>Conditions of irradiated cell</th>
<th>Doubling dose (rad)</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zea mays</td>
<td>4 recessive visibles</td>
<td>Pollen</td>
<td>28</td>
<td>179</td>
</tr>
<tr>
<td>Oenothera, Prunus</td>
<td>Self-incompatibility</td>
<td>Pollen</td>
<td>60</td>
<td>180, 181</td>
</tr>
<tr>
<td>Drosophila</td>
<td>Sex-linked lethals</td>
<td>spermatozoa</td>
<td>50</td>
<td>31-33</td>
</tr>
<tr>
<td></td>
<td></td>
<td>aged spermatozoa</td>
<td>140</td>
<td>31-33</td>
</tr>
<tr>
<td></td>
<td></td>
<td>oocytes and oogonia</td>
<td>390</td>
<td>171</td>
</tr>
<tr>
<td>Mouse</td>
<td>7 recessive autosomal visibles</td>
<td>spermatogonia</td>
<td>30</td>
<td>76, 69, 75</td>
</tr>
<tr>
<td></td>
<td>Dominant lethals</td>
<td>through spermogenesis except time of peak sensitivity</td>
<td>&lt;50</td>
<td>164</td>
</tr>
<tr>
<td></td>
<td>Sex-ratio*</td>
<td>spermatogonia</td>
<td>50</td>
<td>125</td>
</tr>
</tbody>
</table>

*Approximate calculation for natural rate corresponding to age of mice used in 76, 69, 75.

TABLE IX. COMPARISON OF APPROACHES TO QUANTITATIVE ASSESSMENT OF MUTATIONAL DAMAGE

<table>
<thead>
<tr>
<th>Fertility of carriers of the sex-repressed mutant allele</th>
<th>Knowledge of fn</th>
<th>Knowledge of fn</th>
<th>Relative effect of mutation upon frequency of condition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Higher than average........................................</td>
<td>p = 1</td>
<td>Small but unknown</td>
<td>Small</td>
</tr>
<tr>
<td>Lower than average.........................................</td>
<td>Small but unknown</td>
<td>f = 1</td>
<td>Large</td>
</tr>
</tbody>
</table>

TABLE X. SOME OVER-ALL ESTIMATES OF SOCIAL BURDEN

<table>
<thead>
<tr>
<th>Author</th>
<th>Class of traits</th>
<th>Incidence</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>In population</td>
</tr>
<tr>
<td>Slemston1144</td>
<td>Rare heterozygotes..................................................</td>
<td>1.36 x 10^{-2}</td>
</tr>
<tr>
<td></td>
<td>Rare homozygotes....................................................</td>
<td>1.0 x 10^{-3}</td>
</tr>
<tr>
<td></td>
<td>Rare sex-linked.....................................................</td>
<td>1.6 x 10^{-4}</td>
</tr>
<tr>
<td></td>
<td>Common traits of hard interpretation................................</td>
<td>1.0 x 10^{-2}</td>
</tr>
<tr>
<td></td>
<td>TOTAL...................................................................</td>
<td>2.7 x 10^{-2}</td>
</tr>
<tr>
<td>U.S.A. Panel1145</td>
<td>Tangible defects of genetic origin (1/2 total)....................</td>
<td></td>
</tr>
<tr>
<td>Kemp125, 191</td>
<td>Physical malformations and defects..................................</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Severe hereditary afflictions........................................</td>
<td></td>
</tr>
</tbody>
</table>
### Table XI. List of Specific Traits, with Estimated Incidences: Category I

#### (A) Autosomal Dominant Traits

<table>
<thead>
<tr>
<th>Trait</th>
<th>Remarks</th>
<th>Phenotype frequency per million</th>
<th>Births</th>
<th>Living</th>
</tr>
</thead>
<tbody>
<tr>
<td>Achondroplasia</td>
<td>Chondrodystrophy ‘Foetalis’</td>
<td></td>
<td>28</td>
<td>28</td>
</tr>
<tr>
<td>Arachnodactyly</td>
<td>Marfan’s syndrome</td>
<td></td>
<td>60</td>
<td>26</td>
</tr>
<tr>
<td>Brachydactyly (major)</td>
<td>Hands and feet affected—mean stature reduced</td>
<td></td>
<td>6</td>
<td>6</td>
</tr>
<tr>
<td>Ectrodactyly</td>
<td>Including all types of ‘split hand’</td>
<td></td>
<td>30</td>
<td>20</td>
</tr>
<tr>
<td>Multiple exostoses</td>
<td>Only a minority are troublesome</td>
<td></td>
<td>400</td>
<td>400</td>
</tr>
<tr>
<td>Osteitis deformans</td>
<td></td>
<td></td>
<td>30</td>
<td>23</td>
</tr>
<tr>
<td>Osteogenesis imperfecta</td>
<td>Frailities ossium. Several types, all irregular dominant—genetical relationship not known</td>
<td></td>
<td>60</td>
<td>25</td>
</tr>
<tr>
<td>Cranio-facial, cranio-bleidal, mandibulo-facial dysostoses</td>
<td>A series of separate disorders individually uncommon</td>
<td></td>
<td>30</td>
<td>20</td>
</tr>
<tr>
<td>Hypertelorism</td>
<td>Dominant hereditary ataxias—a group of which Friedrich’s is the best defined</td>
<td></td>
<td>200</td>
<td>110</td>
</tr>
<tr>
<td>Ataxia</td>
<td></td>
<td></td>
<td>20</td>
<td>8</td>
</tr>
<tr>
<td>Epilopa</td>
<td>Tuberose sclerosis (9 living sporadic cases in N.I.)</td>
<td></td>
<td>30</td>
<td>7</td>
</tr>
<tr>
<td>Huntington’s chorea</td>
<td>(Three families in N. Ireland known)</td>
<td></td>
<td>10</td>
<td>8</td>
</tr>
<tr>
<td>Hydrocephaly internal obstructive</td>
<td>Includes stenosis of and forking of aqueduct of Sylvius—probably each due to irregular dominant gene</td>
<td></td>
<td>1,230</td>
<td>25</td>
</tr>
<tr>
<td>Peroneal muscular atrophy</td>
<td>Charcot-Marie-Tooth disease</td>
<td></td>
<td>40</td>
<td>24</td>
</tr>
<tr>
<td>Spastic diplegia</td>
<td></td>
<td></td>
<td>400</td>
<td>20</td>
</tr>
<tr>
<td>Dystrophia myotonica</td>
<td></td>
<td></td>
<td>40</td>
<td>24</td>
</tr>
<tr>
<td>Muscular dystrophy, limb girdle</td>
<td>Faces affected</td>
<td></td>
<td>25</td>
<td>14</td>
</tr>
<tr>
<td>Myositis ossificans</td>
<td></td>
<td></td>
<td>20</td>
<td>10</td>
</tr>
<tr>
<td>Deaf mutism (Deafness total hereditary)</td>
<td>Estimated 3 per cent of all hereditary deaf mutism due to dominant genes</td>
<td></td>
<td>46</td>
<td>46</td>
</tr>
<tr>
<td>Deafness perception</td>
<td>Early onset dominant type</td>
<td></td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>Deafness and cataract</td>
<td>Severe early onset deafness and cataract</td>
<td></td>
<td>6</td>
<td>6</td>
</tr>
<tr>
<td>Deafness</td>
<td>Absence of or atresia of external auditory meatus</td>
<td></td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>Neurofibromatosis</td>
<td>Von Recklinghausen’s disease</td>
<td></td>
<td>300</td>
<td>200</td>
</tr>
<tr>
<td>Polyposis of colon, multiple</td>
<td></td>
<td></td>
<td>100</td>
<td>55</td>
</tr>
<tr>
<td>Alopecia areata</td>
<td></td>
<td></td>
<td>700</td>
<td>700</td>
</tr>
<tr>
<td>Anhidrotic syndrome</td>
<td>Anhidrotic “ectodermal” Dysplasia</td>
<td></td>
<td>34</td>
<td>5</td>
</tr>
<tr>
<td>Cephalo-facial haemangiomatosis</td>
<td>Naevoid Amentia</td>
<td></td>
<td>30</td>
<td>7</td>
</tr>
<tr>
<td>Epidermolysis bullosa</td>
<td></td>
<td></td>
<td>100</td>
<td>40</td>
</tr>
<tr>
<td>Pityriasis rubra pilaris</td>
<td></td>
<td></td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>Telangiectasia haemorrhagica</td>
<td></td>
<td></td>
<td>100</td>
<td>12</td>
</tr>
<tr>
<td>Tylosis palmaris et plantaris</td>
<td></td>
<td></td>
<td>35</td>
<td>35</td>
</tr>
<tr>
<td>Urticaria pigmentosa</td>
<td></td>
<td></td>
<td>90</td>
<td>90</td>
</tr>
<tr>
<td>Xanthoma tuberosum multiplex</td>
<td>Cutaneous xanthomatosis and essential hypercholesteraeimia</td>
<td></td>
<td>40</td>
<td>25</td>
</tr>
<tr>
<td>Willebrand’s disease</td>
<td>Haemophilia—like syndrome</td>
<td></td>
<td>25</td>
<td>8</td>
</tr>
<tr>
<td>Polycthcaemia vera</td>
<td></td>
<td></td>
<td>45</td>
<td>20</td>
</tr>
<tr>
<td>Spherocytosis</td>
<td></td>
<td></td>
<td>60</td>
<td>25</td>
</tr>
<tr>
<td>Thrombocytopenia chronic recurrent</td>
<td></td>
<td></td>
<td>60</td>
<td>45</td>
</tr>
<tr>
<td>Porphyria</td>
<td>Dominant type genotype detectable but seldom causes illness</td>
<td></td>
<td>200</td>
<td>130</td>
</tr>
<tr>
<td>Diabetes (insipidus)</td>
<td></td>
<td></td>
<td>40</td>
<td>20</td>
</tr>
<tr>
<td>Cystic disease of lungs</td>
<td>(Included here “congenital” bronchiectasis)</td>
<td></td>
<td>500</td>
<td>400</td>
</tr>
<tr>
<td>Megacolon</td>
<td>Hirschsprung’s disease</td>
<td></td>
<td>100</td>
<td>10</td>
</tr>
<tr>
<td>Aniridia</td>
<td>Dominant very irregular degree of manifestation and probably several dominant mutants can cause</td>
<td></td>
<td>60</td>
<td>60</td>
</tr>
<tr>
<td>Cataracts “congenital”</td>
<td>Types detected at birth or early—probably several different types</td>
<td></td>
<td>160</td>
<td>150</td>
</tr>
<tr>
<td>Cataracts, senile and pre-senile</td>
<td>Several types varying in severity and depending largely on location for disability caused</td>
<td></td>
<td>1,500</td>
<td>1,500</td>
</tr>
<tr>
<td>Choroidal sclerosis</td>
<td>Common—vary from slight iris defect to big defects of iris chorioid and retina involving macula</td>
<td></td>
<td>500</td>
<td>500</td>
</tr>
<tr>
<td>Colobomata</td>
<td></td>
<td></td>
<td>250</td>
<td>200</td>
</tr>
<tr>
<td>Corneal dystrophies</td>
<td>Several types of very variable severity</td>
<td></td>
<td>140</td>
<td>140</td>
</tr>
<tr>
<td>Fundal dystrophies</td>
<td></td>
<td></td>
<td>150</td>
<td>150</td>
</tr>
<tr>
<td>Glaucomas, infantile and juvenile</td>
<td></td>
<td></td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Hypermetropia</td>
<td>Can only be arbitrarily accepted as a segregating trait at about 10</td>
<td></td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Keratoconus</td>
<td>At least two dominant types occur</td>
<td></td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Macular dystrophies</td>
<td>Familial idiopathic non-albinotic usually lateral</td>
<td></td>
<td>700</td>
<td>700</td>
</tr>
<tr>
<td>Nystagmus</td>
<td>Relatively mild regular dominant type</td>
<td></td>
<td>150</td>
<td>150</td>
</tr>
<tr>
<td>Retinitis pigmentosa</td>
<td></td>
<td></td>
<td>58</td>
<td>14</td>
</tr>
<tr>
<td>Retinoblastoma</td>
<td></td>
<td></td>
<td>6</td>
<td>6</td>
</tr>
<tr>
<td>Subluxation of the lens</td>
<td>Primary and not part of Marfan’s syndrome</td>
<td></td>
<td>7</td>
<td>7</td>
</tr>
</tbody>
</table>

**Total** | | | 9,555 | 7,100 |
### TABLE XI. LIST OF SPECIFIC TRAITS, WITH ESTIMATED INCIDENCES: CATEGORY I (continued)

#### (B) AUTOSOMAL RECESSIVE TRAITS

<table>
<thead>
<tr>
<th>Trait</th>
<th>Remarks</th>
<th>Phenotype frequency per million</th>
<th>Births</th>
<th>Living</th>
</tr>
</thead>
<tbody>
<tr>
<td>Albinism</td>
<td>Usual type with ocular signs. More than one mutant (allele), can cause</td>
<td></td>
<td>130</td>
<td>130</td>
</tr>
<tr>
<td>Alkaptonuria</td>
<td></td>
<td></td>
<td>5</td>
<td>3</td>
</tr>
<tr>
<td>Methaemoglobinemia</td>
<td></td>
<td></td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>Phenylpyruvic acid amonia</td>
<td></td>
<td></td>
<td>100</td>
<td>30</td>
</tr>
<tr>
<td>Porphyria congenital</td>
<td>Recessive light sensitive type</td>
<td></td>
<td>50</td>
<td>5</td>
</tr>
<tr>
<td>Galactosuria</td>
<td></td>
<td></td>
<td>50</td>
<td>2</td>
</tr>
<tr>
<td>Gargoylism</td>
<td></td>
<td></td>
<td>20</td>
<td>4</td>
</tr>
<tr>
<td>Hepato-lenticular degeneration</td>
<td>Wilson's disease</td>
<td></td>
<td>10</td>
<td>3</td>
</tr>
<tr>
<td>Lawrence-Moon-Biedl syndrome</td>
<td></td>
<td></td>
<td>40</td>
<td>6</td>
</tr>
<tr>
<td>Microcephaly, true</td>
<td>Microcephalic imbecility</td>
<td></td>
<td>40</td>
<td>21</td>
</tr>
<tr>
<td>Ataxia</td>
<td></td>
<td></td>
<td>40</td>
<td>20</td>
</tr>
<tr>
<td>Choreo-athetosis</td>
<td></td>
<td></td>
<td>70</td>
<td>15</td>
</tr>
<tr>
<td>Myoclonic epilepsy</td>
<td></td>
<td></td>
<td>50</td>
<td>6</td>
</tr>
<tr>
<td>Spastic diplegia</td>
<td>Spastic diplegia familial often with oligophrenia</td>
<td></td>
<td>50</td>
<td>18</td>
</tr>
<tr>
<td>Muscular dystrophy limb girdle type</td>
<td>Face not affected</td>
<td></td>
<td>30</td>
<td>16</td>
</tr>
<tr>
<td>Polioderma</td>
<td></td>
<td></td>
<td>10</td>
<td>3</td>
</tr>
<tr>
<td>Epidermolysis bullosa dystrophica</td>
<td></td>
<td></td>
<td>20</td>
<td>6</td>
</tr>
<tr>
<td>Ichthyosis congenita</td>
<td>May be more than one type</td>
<td></td>
<td>10</td>
<td>—</td>
</tr>
<tr>
<td>Anophthalmos</td>
<td></td>
<td></td>
<td>100</td>
<td>50</td>
</tr>
<tr>
<td>Corneal dystrophies</td>
<td>Severe recessive type</td>
<td></td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>Glaucomas</td>
<td>More than one recessive type with buphthalmos</td>
<td></td>
<td>15</td>
<td>15</td>
</tr>
<tr>
<td>Macular dystrophies</td>
<td>Juvenile and adult types</td>
<td></td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Microphthahmas</td>
<td>Pure type as distinct from those associated with other eye defects. Mental deficiency often associated.</td>
<td></td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Myopia, high</td>
<td>Segregating traits overlapping with ordinary refraction variations</td>
<td></td>
<td>150</td>
<td>150</td>
</tr>
<tr>
<td>Optic atrophy</td>
<td>Very early onset type</td>
<td></td>
<td>50</td>
<td>50</td>
</tr>
<tr>
<td>Retinitis pigmentiosis</td>
<td>Probably several independent mutants contribute</td>
<td></td>
<td>60</td>
<td>60</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td></td>
<td>1,260</td>
<td>738</td>
</tr>
</tbody>
</table>

#### (C) SEX-LINKED RECESSIVE TRAITS

<table>
<thead>
<tr>
<th>Trait</th>
<th>Remarks</th>
<th>Phenotype frequency per million</th>
<th>Births</th>
<th>Living</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diabetes insipidus</td>
<td></td>
<td></td>
<td>50</td>
<td>5</td>
</tr>
<tr>
<td>Haemophilia</td>
<td></td>
<td></td>
<td>100</td>
<td>66</td>
</tr>
<tr>
<td>Christmas disease</td>
<td></td>
<td></td>
<td>10</td>
<td>4</td>
</tr>
<tr>
<td>Ichthyosis vulgaris</td>
<td></td>
<td></td>
<td>6</td>
<td>6</td>
</tr>
<tr>
<td>Muscular dystrophy</td>
<td>Duchenne's type</td>
<td></td>
<td>176</td>
<td>24</td>
</tr>
<tr>
<td>Megalocornea</td>
<td>Only sex limited</td>
<td></td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>Optic atrophy</td>
<td>Leber's type— really sex linked</td>
<td></td>
<td>15</td>
<td>10</td>
</tr>
<tr>
<td>Retinitis pigmentiosis</td>
<td></td>
<td></td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td></td>
<td>397</td>
<td>155</td>
</tr>
</tbody>
</table>

#### (D) SUMMARY OF TRAITS OF CATEGORY I

<table>
<thead>
<tr>
<th>Inheritance mechanism</th>
<th>Frequency per million</th>
<th>Births</th>
<th>Living</th>
</tr>
</thead>
<tbody>
<tr>
<td>Autosomal dominant</td>
<td>9,555</td>
<td>7,100</td>
<td></td>
</tr>
<tr>
<td>Autosomal recessive</td>
<td>1,260</td>
<td>738</td>
<td></td>
</tr>
<tr>
<td>Sex-linked recessive</td>
<td>397</td>
<td>155</td>
<td></td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td>11,212</td>
<td>7,993</td>
<td></td>
</tr>
</tbody>
</table>
### Table XII. List of specific traits, with estimated incidences: Category II

<table>
<thead>
<tr>
<th>Trait</th>
<th>Remarks</th>
<th>Births</th>
<th>Living</th>
</tr>
</thead>
<tbody>
<tr>
<td>Absence of limbs or parts of limbs</td>
<td>Congenital endogenous amputations</td>
<td>200</td>
<td>80</td>
</tr>
<tr>
<td>Cleft palate and hare lip, together or separately</td>
<td>Not including these anomalies occurring as parts of syndromes or associated with other gross defects</td>
<td>970</td>
<td>700</td>
</tr>
<tr>
<td>Congenital dislocation of hip</td>
<td>Mostly limited in effects to females</td>
<td>900</td>
<td>900</td>
</tr>
<tr>
<td>Osteonecrosis</td>
<td>Includes osteochondritis dissecans and local, e.g. diseases of, Kienbock, Kohler, Perthe and Schlatter</td>
<td>200</td>
<td>200</td>
</tr>
<tr>
<td>Radio-ulnar defects</td>
<td>Varying degrees of absence and deformity, radius usually primary and determining also hand defects</td>
<td>205</td>
<td>205</td>
</tr>
<tr>
<td>Talipes-equino-varus</td>
<td>Excluding, where recognized, those with neurological determining causes and when part of severe syndromes e.g. anencephalus</td>
<td>800</td>
<td>700</td>
</tr>
<tr>
<td>Vertebræ, defects and fusions</td>
<td>A large group including Klippel—Fiel syndrome, Sprengel's anomaly etc.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Psoriasis</td>
<td>Not known if all of one origin—varying age of onset, duration of attacks and severity</td>
<td>3,000</td>
<td>3,000</td>
</tr>
<tr>
<td>Ichthyosis vulgaris</td>
<td></td>
<td>1,100</td>
<td>1,100</td>
</tr>
<tr>
<td>Deafness, otosclerotic</td>
<td></td>
<td>200</td>
<td>200</td>
</tr>
<tr>
<td>Anencephalus</td>
<td>i.e. The live born with these defects usually dying shortly after birth, with and without spina bifida or rachischisis</td>
<td>360</td>
<td>—</td>
</tr>
<tr>
<td>Occipital meningocoele</td>
<td></td>
<td>80</td>
<td>—</td>
</tr>
<tr>
<td>Hydrocephalus (Arnold—Chiari)</td>
<td></td>
<td>300</td>
<td>—</td>
</tr>
<tr>
<td>Lumbo-sacral spina bifida</td>
<td></td>
<td>800</td>
<td>100</td>
</tr>
<tr>
<td>Other central nervous system malformations</td>
<td></td>
<td>320</td>
<td>—</td>
</tr>
<tr>
<td>Cardiac malformations</td>
<td></td>
<td>1,200</td>
<td>400</td>
</tr>
<tr>
<td>Digestive tract malformations</td>
<td></td>
<td>630</td>
<td>100</td>
</tr>
<tr>
<td>Urogenital tract malformations</td>
<td></td>
<td>200</td>
<td>40</td>
</tr>
</tbody>
</table>

**Total trait frequency**

9,825 | 8,725

### Table XIII. (A) List of specific traits, with estimated incidences: Category III

<table>
<thead>
<tr>
<th>Trait</th>
<th>Remarks</th>
<th>Phenotype frequency per million</th>
</tr>
</thead>
<tbody>
<tr>
<td>Deafness total from birth</td>
<td>97 per cent of all genetic deafness at birth. A number of independent mutants involved. Relative fertility of homozygote about 1/3</td>
<td>264</td>
</tr>
<tr>
<td>Fibrocystic disease of pancreas</td>
<td>A generalized disorder of external secretory glands. For practical purposes, relative fertility of homozygote is zero</td>
<td>600</td>
</tr>
</tbody>
</table>

**Total trait frequency**

864 | 279

### Table XIII. (B) List of specific traits, with estimated incidences: Category III (continued)

<table>
<thead>
<tr>
<th>Trait</th>
<th>Remarks</th>
<th>Phenotype frequency per million</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anaemia, pernicious</td>
<td>Addison's anaemia</td>
<td>1,300</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td></td>
<td>4,000</td>
</tr>
<tr>
<td>Exophthalmic goitre</td>
<td>Graves's or Basedow's disease</td>
<td>1,700</td>
</tr>
<tr>
<td>Manic depressive reactions</td>
<td>Based on severity requiring hospital admissions</td>
<td>4,000</td>
</tr>
<tr>
<td>Schizophrenia</td>
<td>Based on severity requiring hospital admissions</td>
<td>1,300</td>
</tr>
<tr>
<td>Epilepsy</td>
<td>Secondary to disease or injury</td>
<td>2,500</td>
</tr>
</tbody>
</table>

**Total phenotype frequency**

14,800 | 10,300

199
TABLE XIV. Dominant conditions identified in the Northern Ireland population but not included in Categories I and II for the reasons stated

A. Because their effects are slight (most are common)

**Hand defects:** Brachydactyly thumbs, brachydactyly 1st finger, brachydactyly 1st, 3rd, and 4th fingers; camptodactyly; clinodactyly; polydactyly (not part of syndrome) of radial side and of ulnar side (more common) of hands; syndactyly and symphalangism mostly 3rd and 4th finger (hundreds of cases of the above types are known but have not been sought out for special investigation); Dupuytren's contracture familial.

**Foot defects:** Garber's toe deformity; hallux valgus (familial cases may be associated with metatarsal anomalies); hammer toes (many are familial); syndactyly and symphalangism.

**Other skeletal:** Diaphyseal achalasia, epiphysitis punctata.

**Teeth anomalies** (Other than parts of syndromes): Defective or absent enamel (various types); opalescent dentine; additional teeth (many types); absence of permanent incisors and pre-molars; some such anomalies are present in about 1.3 per cent of the population.

**Skin and hair anomalies:** Adenoma, cystic multiple benign; cysts, epidermoid; dermatomyomat multiple; anonychia and hypoplasia of nails; leukonychia totalis; pachyonychia congenita; hair, white patches; hair kinky; hair woolly; hydros aestival; porokeratosis

**Eye anomalies:** Eyelids—spasm, absence of tarsal plates, uncomplicated ptosis; absence fistula of lacrimal ducts; retina, opaque fibres; strabismus convergent and divergent (primary).

**Miscellaneous:** Pelger's anomaly, elliptocytosis.

**Ear anomalies:** Cat's ears, microtia; pre-helice pits; lobule pits; accessory auricles.

B. Because even if the effects are severe, the anomalies are probably present in less than five persons per million

**Skeletal:** Osteoporosis (Albers-Schönberg); phocomelia; ankylosing spondylosis; polyostotic fibrous dysplasia (Albright's disease); multiple enchondroma; fibula absent or defective; oxycephaly; acrocephaly; syndactyly.

**Skin:** Ichthyosiform congenital erythroderma; keratosis follicularis spinulosa (Darier's disease); monilethrix; urticaria pigmentosa; tylosis palmaris et plantaris; pili torti; mal de Meleda; lipodystrophy progressiva without gargoilyism.

**Miscellaneous:** Milroy's disease: periodic paralysis; dominant microcytic anaemia; Waardenberg's syndrome: anotia.

TABLE XV. Classes of biometrical character

<table>
<thead>
<tr>
<th>Character</th>
<th>Presumed position of social optimum</th>
<th>Position of selective optimum</th>
<th>Position of population mean</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birthweight</td>
<td>At selective optimum</td>
<td>Intermediate finite value</td>
<td>Below selective optimum</td>
</tr>
<tr>
<td>Intelligence (measured as IQ)</td>
<td>+∞ *</td>
<td>Intermediate finite value</td>
<td>Near and possibly even above</td>
</tr>
<tr>
<td>Life-span</td>
<td>+∞ *</td>
<td>Unknown; perhaps +∞ *</td>
<td>Below social optimum; probably below selective optimum</td>
</tr>
</tbody>
</table>

* "+∞ *" implies positive and indefinitely large, always greater than the population mean.

References

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50. Zinder, N., and J. Lederberg, J. Bacteriol., 64, also op. cit. ref. 19 and 20.
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**APPENDIX**

*Calculations concerning survival at or near birth and the distribution of birth-weights*

110. Both Karn and Penrose\(^{150}\) and Fraccaro\(^{184}\) have found in samples of several thousand births that the distributions both of survivors S and non-survivors N through birth and the subsequent 30 days are Gaussian. Under these conditions the influence of the mean and variance of the over-all birth-weight distribution upon survival at or near birth can, at least approximately, be treated algebraically. Suppose birth-weight \(w\) to be measured from the birth-weight at which S is maximal.

Let \(S = S_0 \exp \left(-\frac{w^2}{2\sigma'^2}\right)\)

\(N = N_0 \exp \left(-\frac{1}{2} \left(\frac{w - m'}{\sigma}\right)^2\right)\)

Then the curve determining survival is

\[
S = \frac{S_0}{N_0} \exp \left(-\frac{1}{2\sigma'^2} \left(w + \frac{m'\sigma'^2}{\sigma^2 - \sigma'^2}\right)^2\right) \exp \left(\frac{1}{2} \left(\frac{m'^2}{\sigma^2 - \sigma'^2}\right)\right)
\]

where \(\frac{1}{\sigma'^2} = \frac{1}{\sigma^2} - \frac{1}{\sigma'^2}\)

and the over-all survival is \(1 - k\) where

\[
k = \frac{\sigma S_0}{\sigma S_0 + \sigma' N_0}
\]

Moreover, optimal survival is at \(\omega_{opt} = -\frac{m'\sigma'^2}{\sigma^2 - \sigma'^2}\) and at this point

\[
\left(\frac{S}{N}\right)_{opt} = \frac{S_0}{N_0} \exp \left(-\frac{1}{2} \left(\frac{m'^2}{\sigma^2 - \sigma'^2}\right)\right)
\]

Then the survival at \(\omega = \omega_{opt}\) is \(1 - k_{min}\)

\[
N_0/S_0 \exp \left(-\frac{1}{2} \left(\frac{m'^2}{\sigma^2 - \sigma'^2}\right)\right)
\]

where \(k_{min} = \frac{1 + N_0/S_0 \exp \left(-\frac{1}{2} \left(\frac{m'^2}{\sigma^2 - \sigma'^2}\right)\right)}{1}
\]
It is desirable to express the relation between \( k_m \) and \( \bar{k} \) in terms of (1) the variance \( \sigma^2 \) of the over-all distribution of birth-weights

\[
T(W) = S(W) + N(W)
\]

(2) the difference \( m \) between the mean of the over-all distribution of birth-weights and the birth-weight for optimal survival, and

(3) the variance \( \sigma_1^2 \) which determines the shape of the birth-weight-survival relation.

In terms of the parameters describing \( S(W) \) and \( N(W) \)

\[
m' = m' \left( \frac{\sigma^2}{(\sigma_1^2 - \sigma^2)} + \bar{k} \right)
\]

\[
\sigma_1^2 = \sigma^2 \left( 1 - \bar{k} \right) + \sigma^2 \bar{k} + m'^2 \bar{k} \left( 1 - \bar{k} \right)
\]

If it is assumed that \( \bar{k} \) is small by comparison with unity, it is possible to write

\[
r = \frac{k}{k_m} = \frac{\sigma_1}{\sigma_1} \exp \frac{1}{2} \left( \frac{m^2}{\sigma_1^2 - \sigma^2} \right) + 0 \left( \bar{k} \right)
\]

and so since \( \sigma_1^2 = \sigma^2 + 0(\bar{k}) \) and \( \sigma_1^2 \) can be eliminated in terms of \( \sigma_1^2 \) and \( \sigma^2 \)

\[
r = \frac{\sigma_1}{(\sigma_1^2 - \sigma^2)^{1/2}} \exp \frac{1}{2} \left( \frac{m^2}{\sigma_1^2 - \sigma^2} \right) \ldots \ldots (1)
\]

Comparison of \( \bar{k} \) as observed by Karn and Penrose with values calculated from the above formula and the parameters of their experiments gives

<table>
<thead>
<tr>
<th>Survey Sample</th>
<th>( r_{\text{obs}} )</th>
<th>( r_{\text{calc}} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Males</td>
<td>2.07</td>
<td>2.23</td>
</tr>
<tr>
<td>Females</td>
<td>2.21</td>
<td>2.36</td>
</tr>
</tbody>
</table>

111. On the basis of equ. (1) it is possible to estimate the consequences of small shifts in the mean or variance of the distribution of birth-weights, assuming that the survival curve \( (k_m, \sigma_1) \) remains constant, by the relations

\[
\frac{dr}{r} \frac{d\sigma^2}{\sigma^2} = \frac{1}{2} \left( \frac{\sigma^2}{\sigma_1^2 - \sigma^2} \right) + \frac{1}{2} \left( \frac{m^2 \sigma^2}{(\sigma_1^2 - \sigma^2)^2} \right)
\]

and

\[
\frac{dr}{m} = -\left( \frac{r^2}{\sigma_1^4 - \sigma^2} \right)
\]

Calculated numerical changes in \( r \) and \( \bar{k} \) for 1 per cent changes in variance and in departure of mean from optimal birth-weight are given in table XVI calculated from the data of Karn and Penrose and Fraccaro.

112. It has been estimated by Robson\textsuperscript{185} and by Penrose\textsuperscript{19} that some 40 per cent of the variance of birth-weight in a United Kingdom sample was due to genetic factors, either of the mother or of the foetus. Possibly only a small fraction of this is maintained by recurrent mutation. The other extreme possibility is that recurrent mutation maintains the whole of the genetic component of the variance \( \sigma^2 \). In that event a 10 per cent change in mutation rate might lead to a 4 per cent change in \( \sigma^2 \) and so to changes in survival at and near birth amounting to 0.2-0.7 per cent. If the representative doubling dose for the polygenes concerned were to be 30 rad, this would then correspond approximately to the genetical influence of natural sources of irradiation upon survival at or near birth.

113. The part played by genetic factors in maintaining the difference between the mean birth-weight and that for optimal survival is not known, but the most extreme possibility is again that recurrent mutation may be responsible for the whole of \( m \). In that event a similar change in mutation rate might lead to changes in survival at or near birth amounting to 0.2-0.8 per cent. These calculated upper limits apply to regions in which the total loss of infants at or near birth is in the range 4-7 per cent. They are illustrative of the need to resolve the underlying and more fundamental problem of the part played by mutation in maintaining the current distribution of birth-weights against the pressure of selection acting through this phenotype.

<table>
<thead>
<tr>
<th>Survey Sample</th>
<th>Changes due to 1 per cent change in ( \sigma^2 )</th>
<th>Changes due to 1 per cent change in ( m )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Fractional change in ( \sigma^2 ) (per cent)</td>
<td>Absolute change in ( \sigma^2 ) (per cent)</td>
</tr>
<tr>
<td>Karn and Penrose\textsuperscript{185}</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Males</td>
<td>1.5</td>
<td>0.072</td>
</tr>
<tr>
<td>Females</td>
<td>1.6</td>
<td>0.11</td>
</tr>
<tr>
<td>Fraccaro\textsuperscript{185}</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Males</td>
<td>1.6</td>
<td>0.11</td>
</tr>
<tr>
<td>Females</td>
<td>2.9</td>
<td>0.18</td>
</tr>
</tbody>
</table>

TABLE XVI. CALCULATED CONSEQUENCES OF CHANGES IN THE PARAMETERS GOVERNING BIRTH-WEIGHT DISTRIBUTION
Appendix

LIST OF SCIENTIFIC EXPERTS

The scientific experts who have taken part in the preparation of the report while attending Committee sessions as members of national delegations are listed below. The Committee must also express its appreciation to the many individual scientists not directly connected with national delegations whose voluntary co-operation and good will contributed in no small measure to the preparation of the report.

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