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1.1.2 Mendel's law of
segregation 1.2

Distinguish between:

1.2.1 Complete,

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incomplete and co-
dominance 1.2.2

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The principle of
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Mendel's law of
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Distinguish between:

1.2.1 Complete,
incomplete and co-
dominance 1.2.2

Heterozygous and
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Phenotype and genotype

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1.4 Give ONE reason why the process mentioned in

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causative genes can be tracked through cancer-prone families via genetic linkage and positional cloning. Several of the genes discovered have subsequently been proved to play critical roles in normal growth and development. There are also implications for the families themselves in terms of genetic

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testing with its attendant dilemmas, if it is not clear that useful action will result. The chapters in *The Genetics of Cancer* illustrate what has already been achieved and take a critical look at the future directions of this research and its potential clinical applications.

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Experiments which in previous years were made with ornamental plants have already afforded evidence that the hybrids, as a rule, are not exactly intermediate between the parental species. With some of the more striking characters, those, for instance, which relate to the form and size of the leaves,

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the pubescence of the several parts, etc., the intermediate, indeed, is nearly always to be seen; in other cases, however, one of the two parental characters is so preponderant that it is difficult, or quite impossible, to detect the other in the hybrid.

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important scientific works ever written, the 1865 paper Experiments in Plant Hybridisation was all but ignored in its day, and its author, Austrian priest and scientist GREGOR JOHANN MENDEL (1822-1884), died before seeing the dramatic long-term impact of his work, which was rediscovered at the turn of the 20th

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century and is now considered foundational to modern genetics. A simple, eloquent description of his 1856-1863 study of the inheritance of traits in pea plants. Mendel analyzed 29,000 of them. This is essential reading for biology students and readers of science history. Cosimo presents this compact

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The purpose of this manual is to provide an educational genetics resource for individuals, families, and health professionals in the New York - Mid-Atlantic region and increase awareness of specialty care in genetics. The manual begins with a basic introduction to genetics concepts,

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followed by a description of the different types and applications of genetic tests. It also provides information about diagnosis of genetic disease, family history, newborn screening, and genetic counseling. Resources are included to assist in patient care, patient and professional education, and

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identification of specialty genetics services within the New York - Mid-Atlantic region. At the end of each section, a list of references is provided for additional information.

Appendices can be copied for reference and offered to patients.

These take-home resources are critical to

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helping both providers
and patients understand
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