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Molecular diagnostics

is a collection of

techniques used to

analyse biological

markers in the

genome and

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individual's genetic code and how their cells express their genes as proteins—by applying molecular biology to medical testing. The technique is used to diagnose and monitor disease, detect risk, and decide which therapies will work best for individual patients.

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changes. As discussed in the section on [Basics](#), chromosomes are composed of DNA.

Genetic Testing

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Molecular diagnostics

is the outcome of the

fruitful interplay

among laboratory

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sample of the tumour that has already been removed as part of your treatment. The sample of blood, saliva or body tissue is sent to a genetic testing laboratory to be analysed.

Genetic and genomic testing - NHS

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biology level. It is mainly concerned with the interrelationships between DNA, RNA and protein synthesis. A molecular biologist studies the processes of replication, translation and transcription of genetic material on a wide scale. Given below are molecular biology mcqs for the

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tests may be useful or

even necessary to

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study single genes or short lengths of DNA

to identify variations or mutations that lead to a genetic disorder.

Chromosomal genetic tests analyze whole chromosomes or long lengths of DNA to see if there are large genetic changes,

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such as an extra copy of a chromosome, that cause a genetic condition.

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