£53m for research to probe genetic links to health and disease

Research into human genetics has received a £53 million boost from the Medical Research Council (MRC).

Funds will support work at the MRC Human Genetics Unit, University of Edinburgh for the next five years.

Scientists there are using the latest technologies to study how our genes work and to uncover the genetic basis of both rare and common diseases.

Experts are examining the entire genetic make-up of patients and their families to discover how variations in people’s DNA code can lead to diseases, including those that affect childhood development.

Researchers are also investigating how genes are switched on and off to reveal how these processes contribute to diseases such as cancer, brain and eye disorders.

Professor Wendy Bickmore, Director of the MRC Human Genetics Unit at the University of Edinburgh, said: “The scale and long-term nature of this funding from the Medical Research Council will enable our scientists to tackle the most challenging questions in human genetics.

“We are using the latest computational and experimental techniques to investigate how our genomes work to control the function of cells and tissues in people and populations.

“We will also be able to train the next generation of scientists to apply their expertise to improving health and the lives of patients and their families.”

The MRC Human Genetics Unit was first established more than 60 years ago as the Clinical Effects of Radiation Unit. It was renamed in 1988 and joined the University of Edinburgh in 2012.

Today, The MRC Human Genetics Unit is a major partner in the MRC Institute of Genetics and Molecule Medicine at the University of Edinburgh.

Dr Lindsay Wilson, MRC Programme Manager for Genetics and Genomics, said: “The Human Genetics Unit is the MRC’s biggest Unit investment and we’re pleased to continue to
support Director Wendy Bickmore’s vision for world-leading research across the next five years and look forward to many exciting discoveries.”

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