

Clinical trials using genome editing based therapeutics and gene delivery therapies

Successfully audited by the FDA, EMA and Health Canada



LPL-gene replacement therapy (Glybera)

The first in-human gene therapy to reach the market

Co-development

Execution of pivotal trials

Publications

15 years Follow-up



Next generation LPL-gene replacement therapy

With the National Research Council Canada (NRC)

Co-development

Clinical Sponsor



Human gene editing clinical trials

Verve therapeutics, Undisclosed

Scientific Input

Clinical Input

Regulatory Input

Execution



Multiple Phase IIa and Investigator-initiated studies

Conception

Conduct

SCIENTIFIC
OUTPUTS

REGULATORY-RELATED
EXPERIENCE

REMOTE
MONITORING CAPABILITIES

PoC-PHASE III

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