

Appel d'offres “ Plateforme Mutations ”

Résultats Réunion du Conseil Scientifique du 9 Juillet 2009

Liste des projets sélectionnés (ordre alphabétique)

AMIEL Jeanne	Identification of the disease causing gene in a familial sudden death syndrome
BELLANNE-CHANTELOT Christine	Analysis of a 1.9 Mb candidate region in two families with myeloproliferative neoplasms
FREBOURG Thierry	Identification of new types of alterations within the MMR genes in Lynch syndrome by high throughput sequencing
LAPORTE Jocelyn	High throughput mutations screening for the identification of novel genes in centronuclear myopathies
PETIT Christine	Identification of the gene responsible for non-syndromic hearing impairment, DFNB15
ROZET Jean-Michel	Identification of a gene responsible for non-syndromic autosomal recessive optic neuropathies
SMAHI Asma	Identification of the gene responsible for Bazex Syndrome