PUBLICATIONS

Scientific publications


Weinreich SS, Rigter T, van El CG, Dondorp WJ, Kostense P, van der Ploeg AT, Reuser AJJ, Cornel MC, Hagemans MLC. Public support for neonatal screening for Pompe disease, a broad-phenotype condition. Orphanet Journal of Rare Diseases 2012, 7:15

Published abstracts and presentations


Rigter T. Reflecting on earlier experiences with unsolicited findings: Points to consider for next generation sequencing and informed consent in diagnostics. Video highlight website Human Mutation: www.youtube.com/watch?v=gh4qc0Ucy3Y.

Rigter T. European initiatives for unified informed consent. Oral presentation. January 13-14 2013 Tel Aviv, Israel. E-Rare Workshop: Ethical aspects of exome and whole genome sequencing studies (WES/WGS) in rare diseases.


