

## LISTA PUBBLICAZIONI

1. Functional loss of *Ccdc151* leads to hydrocephalus in a mouse model of primary ciliary dyskinesia.

Chiani F, Orsini T, **Gambadoro A**, Pasquini M, Putti S, Cirilli M, Ermakova O, Tocchini-Valentini GP.

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2. Identification of genes required for eye development by highthroughput screening of mouse knockouts.

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3. Identification of genetic elements in metabolism by highthroughput mouse phenotyping

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4. A large scale hearing loss screen reveals an extensive unexplored genetic landscape for auditory dysfunction

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## 5. Genome wide conditional mouse knockout resources

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6. Three-Dimensional microCT imaging of murine embryonic development from immediate post-implantation to organogenesis: application for phenotyping analysis of early embryonic lethality in mutant animals

Ermakova O, Orsini T, **Gambadoro A**, Chiani F, Tocchini- Valentini GP. Mamm Genome. 2017 Nov 23. doi: 10.1007/s00335-017-9723-6. [Epub ahead of print]

7. Disease model discovery from 3,328 gene knockouts by The International Mouse Phenotyping Consortium

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## 8. High-throughput discovery of novel developmental phenotypes

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12. Nucleo-mitochondrial interactions in *Saccharomyces cerevisiae*: characterization of a nuclear gene suppressing a defect in mitochondrial tRNAAsp processing.

T. Rinaldi, **A. Gambadoro**, S. Francisci, L. Frontali

