

Personal information

Name and surname: CINZIA CICCACCI

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Position

Assistant Professor (RTDb) in Medical Genetics at Saint Camillus International University of Health and Medical Sciences (UniCamillus)

Education and training

- 2001 Degree in Biological Science
- 2005 PhD in Evolutionist Biology and Ecology
- 2015 Specialty degree in Medical Genetics

National academic qualifications

- Associate Professor in Medical Genetics 06/A1 (31/03/2017)
- Associate Professor in Applied Medical Technologies 06/N1 (05/12/2014)

Research activity

The scientific activity of Dr. Cinzia Ciccacci is documented by 54 publications in peer-reviewed international journals, of which 51 indexed on pubmed, for a total IF of about 185, total citations = 760 (source: Scopus) and with h-index = 17 (Source: Scopus) and 44 communications to National and International Congresses.

This activity focused at the beginning of her academic career on the Genetics of human populations and evolutionary Genetics with particular regard to A) characterization of the variability pattern for SNS (Single Nucleotide Substitutions) MisSense and SameSense in coding sequences; B) Haplotypic variability of the CFTR (Cystic Fibrosis conductance Transmembrane Regulator) gene in various human populations.

In the last 12 years her activity has been focused on: A) the research of genetic factors contributing to the susceptibility to several multifactorial diseases, (in particular inflammatory bowel diseases, systemic lupus Erythematosus, Rheumatoid arthritis, Diabetes and its complications) by investigating the role of genetic variability in relation to the phenotypic variability of these pathologies and their different clinical manifestations and complications; B) Genetic variability in the response to drugs, both in terms of efficacy and toxicity (Pharmacogenetic/Pharmacogenomics), in particular by studying the genetic predictors of response in treatments with Nevirapine (anti-retroviral drug used in the treatment of HIV) and Warfarin (oral anticoagulant).

She has participated in several research projects (concerning the Genetics of multifactorial pathologies and Pharmacogenetics) financed by MIUR and other National and International Institutions.

She is “ad hoc” Reviewer for several Journals and Academic Editor for “Plos One” and “Journal of Immunology Research”.

Publications:

Modiano G, Bombieri C, Ciminelli BM, Belpinati F, Giorgi S, Georges MD, Scotet V, Pompei F, **Ciccacci C**, Guittard C, Audrezet MP, Begnini A, Toepfer M, Macek M, Ferec C, Claustres M, Pignatti PF. *A large-scale study of the random variability of a coding sequence: a study on the CFTR gene.* Eur J Hum Genet. 2005 Feb;13(2):184-92. IF=4.3

Pompei F, Ciminelli BM, Bombieri C, **Ciccacci C**, Koudova M, Giorgi S, Belpinati F, Begnini A, Cerny M, Des Georges M, Claustres M, Ferec C, Macek M, Modiano G, Pignatti PF. *Haplotype block structure study of the CFTR gene. Most variants are associated with the M470 allele in several European populations.* Eur J Hum Genet. 2006 Jan;14(1):85-93. IF=4.3

Ciminelli BM, Bonizzato A, Bombieri C, Pompei F, Gabaldo M, **Ciccacci C**, Begnini A, Holubova A, Zorzi P, Piskackova T, Macek M Jr, Castellani C, Modiano G, Pignatti PF. *Highly preferential association of NonF508del CF mutations with the M470 allele.* J Cyst Fibros. 2007 Jan;6(1):15-22. IF=3.85

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Borgiani P, **Ciccacci C**, Forte V, Romano S, Federici G, Novelli G. *Allelic variants in the CYP2C9 and VKORC1 loci and interindividual variability in the anticoagulant dose effect of warfarin in Italians.* Pharmacogenomics. 2007 Nov;8(11):1545-1550. IF=2.7

Perricone C, Borgiani P, Romano S, **Ciccacci C**, Fusco G, Novelli G, Biancone L, Calabrese E, Pallone F. *ATG16L1 Ala197Thr is not associated with susceptibility to Crohn's disease or with phenotype in an Italian population.* Gastroenterology. 2008 Jan;134(1):368-70. IF=18.2

Novelli G, **Ciccacci C**, Borgiani P, Amati M P, Abadie E. Genetic tests and genomic biomarkers: regulation, qualification and validation. *Clinical Cases in Mineral and Bone Metabolism.* 2008; 5 (2): 149-154.

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G. Novelli, P. Borgiani, C. **Ciccacci**, N. Di Daniele, G. Sirugo, M. Papaluca Amati. *Pharmacogenomics: role in medicines approval and clinical use.* Public Health Genomics 2010; 13(5):284-91 IF=1.5

Ciccacci C, Borgiani P, Ceffa S, Sirianni E, Marazzi MC, Doro Altan AM, Paturzo G, Bramanti P, Novelli G, Palombi L. *Nevirapine-induced hepatotoxicity and Pharmacogenetics: a retrospective study in a population of Mozambique.* Pharmacogenomics. 2010 Jan;11(1):23-31 IF=2.7

Predazzi IM, Martínez-Labarga C, Vecchione, L, Mango R, **Ciccacci C**, Amati F, Ottoni C, Crawford M, Rickards O, Romeo F, Novelli G. *Population differences in allele frequencies at the OLR1 locus may*

suggest geographic disparities in cardiovascular risk events. Ann Hum Biol. 2010 Apr;37(2):136-48. IF=1.57

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C Ciccacci, N Paolillo, D Di Fusco, G Novelli and P Borgiani. *EPHX1 Polymorphisms Are Not Associated with Warfarin Response in an Italian Population.* Clin Pharmacol Ther. 2011 Jun;89(6):791 IF=7.27

Ciccacci C, Biancone L, Di Fusco D, Ranieri M, Condino G, Giardina E, Onali S, Lepre T, Pallone F, Novelli G, Borgiani P. *TRAF3IP2 gene is associated with cutaneous extraintestinal manifestations in Inflammatory Bowel Disease.* J Crohns Colitis. 2013; 7 (1):44-52. IF=6.59

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