Date of birth: 15th February 1982

Nationality: French

ORCID: 0000-0001-9006-8436

NIH BioSketch

Github organization: https://github.com/IARCbioinfo

Website: http://rarecancersgenomics.com

Rare Cancers Genomics Team Genomic Epidemiology Branch International Agency for Research on Cancer (IARC-WHO)

Email: follm@iarc.who.int

Research Group: https://www.iarc.who.int/teams-rcg/

25 avenue Tony Garnier, 69007 Lyon, France

Professional Appointments

2021-on	Team Leader , International Agency for Research on Cancer (IARC-WHO), Lyon, France
2014-2020	Scientist, International Agency for Research on Cancer (IARC-WHO), Lyon, France
2012-2014	Staff Research Scientist, Swiss Federal Institute of Technology, Lausanne, Switzerland
2007-2012	Postdoctoral Researcher, Institute of Ecology and Evolution, University of Berne, Switzerland

Honors and awards

2024-on	Member of the WHO classification of turmours 6 th edition subcommittee on									
	computational pathology , International Agency for Research on Cancer (IARC-WHO), https://tumourclassification.iarc.who.int/									
2023	Young investigator award, International Mesothelioma Interest Group (iMig), https://imig.org									
2022-on	Chair of the of the Rare Cancers Working Group, European Prospective Investigation into Cancer and Nutrition (EPIC), http://epic.iarc.who.int									
2021-on	lung NET task force member, European NEuroendocrine Tumors Society (ENETS), https://www.enets.org									
2016-on	Scientific Committee member, French MESOBANK (virtual mesothelioma national biobank), https://www.netmeso.fr/netmeso/mesopath-et-mesobank									

Education

2004-2007	PhD	in	Population	Genetics	and	Evolution,	University	of	Grenoble,	France.
	https:/	/these	s hal science/te	1-00216192/						

2001–2004 **MSc Computer science and Applied Mathematics, specialization in Bioinformatics**, Grenoble Institute of Technology ENSIMAG, Grenoble, France.

Personal Statement

After studying applied mathematics and computer science, I completed a Ph.D. in population genetics and bioinformatics in 2007. During this time I gained a strong expertise in statistical genetics and bioinformatics methods, and I wrote several software now widely used in the field. After my PhD., I moved to the University of Berne as a postdoctoral researcher where I started to work more specifically on large human genomic data. I developed in particular several Bayesian models to identify genetic variants associated with phenotypes observed in different human populations or ethnic groups, being the consequence of local adaptation. In 2012, I was recruited as a Staff Research Scientist in the Swiss Federal Institute of Technology of Lausanne where I worked on new statistical and bioinformatics methods for the analysis Next Generation Sequencing data. In 2014 I been recruited as a staff scientist to develop computation cancer genomics projects and to coordinate the bioinformatics efforts in the International Agency for Research on Cancer (IARC-WHO). I am particularly involved in the molecular characterisation of rare thoracic tumors like lung neuroendocrine tumors and malignant mesothelioma, and since 2021 I am co-leading the IARC Rare Cancers Genomics team together with Dr. Lynnette Fernandez-Cuesta.

Publications

77 publications in international peer-reviewed scientific journals, including 9 publications as first/co-first author, and 14 publications as last/co-last/corresponding author.

13'415 total citations, h-index: 42 (source: Google Scholar).

Full list available at Google Scholar and My NCBI Bibliography.

Current Research Support

2024-2027 Understanding the spatiotemporal eco-evolutionary interactions in malignant pleural mesothelioma.

Worldwide Cancer Research, 252,246 GBP. Alcala N (PI), Fernandez-Cuesta L, Foll M (co-Investigator), Jaehee Kim.

Award: www.worldwidecancerresearch.org

2023-2024 Reconciling lung carcinoids histopathological and molecular classifications.

Investigator Award, Neuroendocrine Tumor Research Foundation, 270,000 USD. Foll M (PI).

Award: netrf.org/research/reconciling-lung-carcinoids-histopathological-and-molecular-classifications/

2022-2024 Intra-Tumour Heterogeneity of Pleural Mesothelioma at the Single-Cell Level.

Concept Award, Congressionally Directed Medical Research Programs, 100,000 USD. Foll M (PI), Fernandez-Cuesta L (co-PI).

Award: https://app.dimensions.ai/details/grant/grant.13055977

Selected Open Science Projects

I have been advocating for open science since the beginning of my career, with a focus on open-source software and open data sharing.

2023 HaloAE software | GitHub: IARCbioinfo/HaloAE

A local transformer Auto-Encoder for anomaly detection and localization.

Role: Project co-founder, developer supervision.

2020 Needlestack bioinformatics pipeline | GitHub: IARCbioinfo/needlestack

An ultra-sensitive variant caller for multi-sample next generation sequencing data.

Role: Project founder, lead developer and supervision.

2019 Medical Genomics Open Educational Resource | GitHub: IARCbioinfo/medica_genomics_course

Medical Genomics course held annually at the INSA Lyon engineering school (French National Institute of Applied Sciences of Lyon). Includes all lecture slides and practicals for self-paced learning.

Role: Project co-founder, lecturer and supervision.

2014 Cancer Genomics bioinformatics pipelines | rarecancersgenomics.com/tools & github.com/IARCbioinfo

Best-practices pipelines for WGS, RNA-seq, and methylation data analysis and integration, mostly based on Nextflow.

Role: Project founder, developers supervision.

2014 Rare Cancers Genomics data | rarecancersgenomics.com/datasets

WGS, RNA-seq, and methylation data for rare cancers.

Role: Project co-founder.

2008 BayeScan software | GitHub: mfoll/BayeScan & GitHub: mfoll/BayeScanHierachical

Software to identify candidate loci under natural selection from genetic data. Publication cited >2,000 times, one of the most widely used method in the field.

Role: Project founder, developer.

Selected Oral Presentations

2024 Understanding cancer biology through multi- omics genotype-phenotype tumour maps.

Keynote speaker, BioSyl Computational Biology of Cancer 2024, Grenoble (France).

2024 The Rare Cancers EPIC database: a gateway to rare cancer epidemiological research.

Proffered Paper, ESMO Sarcoma and Rare Cancers Congress 2024, Lugano (Switzerland).

DOI: 10.1016/j.esmoop.2024.102440

2023 Understanding cancer biology through multi-omics genotype-phenotype tumour maps: applications in rare cancers.

Invited Speaker, EMBL conference Cancer Genomics, Heidelberg (Germany).

2023 Malignant pleural mesothelioma: from inter- to intra-tumor heterogeneity.

Young Investigator Award, 16th International Conference of the International Mesothelioma interest group (iMig), Lille (France).

2023 Multi-omics characterization of rare heterogeneous tumors.

Invited Presentation, 31st Annual Intelligent Systems For Molecular Biology (ISMB) / 22nd Annual European Conference on Computational Biology (ECCB), Lyon (France).

Video: https://youtu.be/BoeBCkxckLw

2023 Biocomputational approaches for the study of rare endocrine cancers.

Invited Speaker, European Congress of Endocrinology (ECE), Istambul (Turkey).

- The lungNENomics Project: a Comprehensive Multidisciplinary Characterisation of Pulmonary Carcinoids.

 Abstract presentation, Annual European Neuroendocrine Tumor Society (ENETS) annual conference, Vienna (Austria).
- 2023 Molecular features of lung neuroendocrine tumors.

Invited Speaker, European Association of Nuclear Medicine (EANM) annual congress, Barcelona (Spain).

2022 Multi-omics characterization of heterogeneous tumors.

Invited Speaker, European Meeting on Molecular Diagnostics (EMMD), Noordwijk (Netherlands).

Selected Publications

2024 Multi-omic dataset of patient-derived tumor organoids of neuroendocrine neoplasms.

Gigascience. DOI: 10.1093/gigascience/giae008; PMID: 38451475

Alcala N, Voegele C, Mangiante L, Sexton-Oates A, Clevers H, Fernandez-Cuesta L, Dayton TL, Foll M.

GitHub: IARCbioinfo/MS_panNEN_organoids.

2023 Druggable growth dependencies and tumor evolution analysis in patient-derived organoids of neuroendocrine neoplasms from multiple body sites. 8

Cancer Cell. DOI: 10.1016/j.ccell.2023.11.007; PMID: 38086335

Dayton TL, Alcala N, Moonen L, den Hartigh L, Geurts V, ..., Foll M, Fernández-Cuesta L, Clevers H.

2023 Spotlight on Small-Cell Lung Cancer and Other Lung Neuroendocrine Neoplasms.

American Society of Clinical Oncology Educational Book. DOI: 10.1200/EDBK_390794; PMID: 37229617

Fernandez-Cuesta L, Sexton-Oates A, Bayat L, Foll M, Lau SCM, Leal T.

2023* Multiomic analysis of malignant pleural mesothelioma identifies molecular axes and specialized tumor profiles driving intertumor heterogeneity. 7 24 citations 8

Nature Genetics. DOI: 10.1038/s41588-023-01321-1; PMID: 36928603

Mangiante L, Alcala N, Sexton-Oates A, Di Genova A, Gonzalez-Perez A, ..., Foll M#, Fernandez-Cuesta L#.

2023 HaloAE: A Local Transformer Auto-Encoder for Anomaly Detection and Localization Based on HaloNet. 8
Proceedings of the 18th International Joint Conference on Computer Vision, Imaging and Computer Graphics
Theory and Applications (VISIGRAPP 2023). DOI: 10.5220/0011865900003417

Mathian E, Liu H, Fernandez-Cuesta L, Samaras D, Foll M, Chen L.

GitHub: IARCbioinfo/HaloAE.

2022# A molecular phenotypic map of malignant pleural mesothelioma.

Gigascience. DOI: 10.1093/gigascience/giac128; PMID: 36705549

Di Genova A, Mangiante L, Sexton-Oates A, Voegele C, Fernandez-Cuesta L, Alcala N, Foll M.

GitHub: IARCbioinfo/MESOMICS_data.

2021[#] Challenges in lung and thoracic pathology: molecular advances in the classification of pleural mesotheliomas.

Virchows Archiv. DOI: 10.1007/s00428-020-02980-9; PMID: 33411030

Fernandez-Cuesta L, Mangiante L, Alcala N, Foll M.

2020# A molecular map of lung neuroendocrine neoplasms.

Gigascience. DOI: 10.1093/gigascience/giaa112; PMID: 33124659

Gabriel AAG, Mathian E, Mangiante L, Voegele C, Cahais V, ..., Foll M.

GitHub: IARCbioinfo/DRMetrics.

2020 Needlestack: an ultra-sensitive variant caller for multi-sample next generation sequencing data.

NAR Genomics and Bioinformatics. DOI: 10.1093/gigascience/giac128; PMID: 36705549

Delhomme TM, Avogbe PH, Gabriel AAG, Alcala N, Leblay N, ..., Foll M.

GitHub: IARCbioinfo/needlestack.

2020 EURACAN/IASLC Proposals for Updating the Histologic Classification of Pleural Mesothelioma: Towards a More Multidisciplinary Approach. T 147 citations 8

Journal of Thoracic Oncology. DOI: 10.1016/j.jtho.2019.08.2506; PMID: 31546041

Nicholson AG, Sauter JL, Nowak AK, Kindler HL, Gill RR, ..., Foll M, ..., Galateau-Salle F.

2019# Molecular studies of lung neuroendocrine neoplasms uncover new concepts and entities.

Translational Lung Cancer Research. DOI: 10.21037/tlcr.2019.11.08; PMID: 32038931

Fernandez-Cuesta L, Foll M.

2019# Redefining malignant pleural mesothelioma types as a continuum uncovers immune-vascular interactions.

EBioMedicine. DOI: 10.1016/j.ebiom.2019.09.003; PMID: 31648983

Alcala N, Mangiante L, Le-Stang N, Gustafson CE, Boyault S, ..., Foll M#, Galateau-Salle F#, Fernandez-Cuesta L#.

2019[#] Integrative and comparative genomic analyses identify clinically relevant pulmonary carcinoid groups and unveil the supra-carcinoids. T 141 citations 8

Nature communications. DOI: 10.1038/s41467-019-11276-9; PMID: 31431620

Alcala N, Leblay N, Gabriel AAG, Mangiante L, Hervas D, ..., Foll M#, Fernandez-Cuesta L#.

2019 Linking a mutation to survival in wild mice. T 172 citations 8

Science. DOI: 10.1126/science.aav3824; PMID: 30705186

Barrett RDH, Laurent S, Mallarino R, Pfeifer SP, Xu CCY, Foll M, ..., Hoekstra HE.

2018 Prediction of acute myeloid leukaemia risk in healthy individuals. * 778 citations 3

Nature. DOI: 10.1038/s41586-018-0317-6; PMID: 29988082

Abelson S, Collord G, Ng SWK, Weissbrod O, Mendelson Cohen N, ..., Foll M, ..., Shlush LI.

2017 BAP1 Is Altered by Copy Number Loss, Mutation, and/or Loss of Protein Expression in More Than 70% of Malignant Peritoneal Mesotheliomas. 8

Journal of Thoracic Oncology. DOI: 10.1016/j.jtho.2016.12.019; PMID: 28034829

Leblay N, Leprêtre F, Le Stang N, Gautier-Stein A, Villeneuve L, ..., Foll M, Fernandez-Cuesta L, Brevet M.

2016 The past, present and future of genomic scans for selection.

Molecular Ecology. DOI: 10.1111/mec.13493; PMID: 26745554

Jensen JD, Foll M, Bernatchez L.

2015* WFABC: a Wright-Fisher ABC-based approach for inferring effective population sizes and selection coefficients from time-sampled data.

Molecular Ecology Resources. DOI: 10.1111/1755-0998.12280; PMID: 24834845

Foll M, Shim H, Jensen JD.

GitHub: mfoll/WFABC.

2014* Widespread signals of convergent adaptation to high altitude in Asia and America.

The American Journal of Human Genetics. DOI: 10.1016/j.ajhg.2014.09.002; PMID: 25262650

Foll M, Gaggiotti OE, Daub JT, Vatsiou A, Excoffier L.

GitHub: mfoll/BayeScanHierachical.

2014* Adaptive, convergent origins of the pygmy phenotype in African rainforest hunter-gatherers.

Proceedings of the National Academy of Sciences. DOI: 10.1073/pnas.1402875111; PMID: 25136101

Perry GH, Foll M, Grenier JC, Patin E, Nédélec Y, ..., Barreiro LB.

2014* Influenza virus drug resistance: a time-sampled population genetics perspective.

PLOS Genetics. DOI: 10.1371/journal.pgen.1004185; PMID: 24586206

Foll M, Poh YP, Renzette N, Ferrer-Admetlla A, Bank C, ..., Jensen JD.

2013# Robust demographic inference from genomic and SNP data. 🏆 1350 citations 8

PLOS Genetics. DOI: 10.1371/journal.pgen.1003905; PMID: 24204310

Excoffier L, Dupanloup I, Huerta-Sánchez E, Sousa VC, Foll M.

2013 Approximate Bayesian computation.

695 citations

PLOS Computational Biology. DOI: 10.1371/journal.pcbi.1002803; PMID: 23341757

Sunnåker M, Busetto AG, Numminen E, Corander J, Foll M, Dessimoz C.

2012 Genomic data reveal a complex making of humans. 8

PLOS Genetics. DOI: 10.1371/journal.pgen.1002837; PMID: 22829785

Alves I, Srámková Hanulová A, Foll M, Excoffier L.

2011# fastsimcoal: a continuous-time coalescent simulator of genomic diversity under arbitrarily complex evolutionary scenarios. • 419 citations 8

Bioinformatics. DOI: 10.1093/bioinformatics/btr124; PMID: 21398675

Excoffier L, Foll M.

Code: cmpg.unibe.ch/software/fastsimcoal2.

2010# Quantifying population structure using the F-model.

Molecular Ecology Resources. DOI: 10.1111/j.1755-0998.2010.02873.x; PMID: 21565093 Gaggiotti OE, Foll M.

2009 Genetic consequences of range expansions. **?** 1330 citations

Annual Review of Ecology, Evolution, and Systematics. DOI: 10.1146/annurev.ecolsys.39.110707.173414 L Excoffier, Foll M, RJ Petit

2009[#] Detecting loci under selection in a hierarchically structured population. ₹ 921 citations 8

Heredity (Edinb). DOI: 10.1038/hdy.2009.74; PMID: 19623208

Excoffier L, Hofer T, Foll M.

Code: cmpg.unibe.ch/software/arlequin35/.

2008* A genome-scan method to identify selected loci appropriate for both dominant and codominant markers:

a Bayesian perspective. T 2867 citations 8

Genetics. DOI: 10.1534/genetics.108.092221; PMID: 18780740

<u>Foll M</u>, Gaggiotti O. GitHub: mfoll/BayeScan.

2006^{*} Identifying the environmental factors that determine the genetic structure of populations. ₹ 418 citations

8

Genetics. DOI: 10.1534/genetics.106.059451; PMID: 16951078

Foll M, Gaggiotti O.

Code: leca.osug.fr/-Genomique-des-populations-.

🝸: >20 citations/year.

T: >50 citations/year (on average since publication, source: Google Scholar).

a: Open Access publication.

^{*:} first/co-first author.

^{#:} last/co-last/corresponding author.