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

Computational Cancers Genomics Team Genomic Epidemiology Branch International Agency for Research on Cancer (IARC-WHO) 25 avenue Tony Garnier, 69007 Lyon, France

Email: follm@iarc.who.int

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

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									
2022-2024	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 P	PhD	in	Population	Genetics	and	Evolution,	University	of	Grenoble,	France.
h	nttns://t	heses	hal science/tel	-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 thoracic tumors like lung neuroendocrine neoplams and malignant mesothelioma. In 2016, Dr. Lynnette Fernandez-Cuesta and I launched the Rare Cancers Genomics initiative, and and since 2021 we have been co-leading the IARC Computational Cancers Genomics team.

Publications

80 publications in international peer-reviewed scientific journals, including 9 publications as first/co-first author, and 15 publications as last/co-last/corresponding author.

14'113 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 IARC and part of the INSA Lyon engineering school Master curriculum (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 Assessment of the current and emerging criteria for the histopathological classification of lung neuroendocrine tumours in the lungNENomics project. 3

ESMO Open. DOI: https://doi.org/10.1016/j.esmoop.2024.103591; PMID: 38878324

Mathian E, Drouet Y, Sexton-Oates A, Papotti MG, Pelosi G, ..., Foll M.

GitHub: https://github.com/IARCbioinfo/LNENBarlowTwins.

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. 8

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. 3

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. T147 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. 3

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. ♥ 141 citations ∂

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. * 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. 3

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, $\underline{\text{Foll M}}$, 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. T 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^{\star} 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. Table 218 citations

9

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

Foll M, Gaggiotti O.

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

- *: first/co-first author.
- #: last/co-last/corresponding author.
- 🝸: >20 citations/year.
- T: >50 citations/year (on average since publication, source: Google Scholar).
- 3: Open Access publication.