

Curriculum Vitae

Frederick P. (Fritz) Roth, PhD

- Titles:** Professor, University of Toronto
Donnelly Centre for Cellular & Biomolecular Research
Department of Molecular Genetics
Department of Computer Science
- Senior Investigator, Lunenfeld-Tanenbaum Research Institute, Sinai Health
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- Citizenship:** Dual US and Canadian
- Education:** BA, Physics and Molecular & Cell Biology, Univ. of California, Berkeley, 1990
PhD, Biophysics, Harvard University, 1998
- Honors:** 2011 – 2017 Canada Excellence Research Chair
2008 – 2020 Senior Fellow, Canadian Institute for Advanced Research
1993 – 1996 National Science Foundation Graduate Fellowship
- Employment:**
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|----------------|--|
| 2011 – present | Professor, Donnelly Centre for Cellular and Biomolecular Research, University of Toronto, Toronto ON |
| 2011 – present | Senior Investigator, Lunenfeld-Tanenbaum Research Institute of Mt. Sinai Hospital, Toronto, ON |
| 2014 – 2020 | Co-Director, Genetic Networks Program, Canadian Institute for Advanced Research (CIFAR) |
| 2012 – 2014 | Associate Director, Genetic Networks Program, Canadian Institute for Advanced Research (CIFAR) |
| 2007 – 2010 | Associate Professor, Harvard Medical School, Department of Biological Chemistry and Molecular Pharmacology |
| 2000 – 2007 | Assistant Professor, Harvard Medical School, Department of Biological Chemistry and Molecular Pharmacology |
| 1998 – 2000 | Scientist, Millennium Pharmaceuticals Inc (w/ Dr. Chris Sander). |
| 1992 – 1998 | Graduate Student, Harvard University Program in Biophysics, (w/ Dr. George Church). |
| 1991 – 1992 | Staff Scientist, Operon Technologies, Inc. |
| Summer 1990 | Research Fellow, Scripps Inst. of Oceanography, (w/ Dr. Adrianus Kalmijn). |
| Summer 1989 | Research Assistant, UC Berkeley, Mol. Cell Biol. Dept. (w/ Dr. Jasper Rine). |
| 1987 – 1990 | Research Assistant, Space Sciences Lab, Berkeley, CA (w/ Dr. Kevin Hurley). |
| Summer 1987 | Research Assistant, Fly's Eye Cosmic Ray Facility, Physics Dept., U. of Utah. |

Academic Institutional Roles

2020 – present	Executive Committee, Atlas of Variant Effects Alliance
2020 – 2022	Director, Computational Biology in Molecular Genetics (CBMG) Track within the Molecular Genetics Graduate Program
2019 – present	Scientific Advisory Board, Mouse Genome Database, Jackson Laboratory, Bar Harbor, ME
2018 – 2019	Faculty of Medicine Decanal Promotions Committee
2017 – present	Department of Molecular Genetics Admissions Committee
2016	Co-founder, Computational Biology in Molecular Genetics (CBMG) Track within the Molecular Genetics Graduate Program
2014 – present	Department of Molecular Genetics Examination Committee
2013 – 2017	Donnelly Centre Promotions Committee
2011 – present	Cross-appointed to the Dept. of Computer Science, University of Toronto
2011 – present	Cross-appointed to the Dept. of Molecular Genetics, University of Toronto
2006 – present	Member, Center for Cancer Systems Biology, Dana-Farber Cancer Institute
2006 – 2011	Member, Joint Committee of the Harvard Countway & Boston Medical Libraries
2005 – 2011	Tutor, Harvard Undergraduate Concentration in the Biochemical Sciences
2003 – 2008	Admissions Committee, Harvard Program in Biophysics
2000 – 2011	Faculty, Biological & Biomedical Sciences Grad Program, Harvard Med School
2000 – 2011	Faculty, Harvard Graduate Program in Biophysics

Industry Roles

2021 – present	Strategic Advisory Board Member, BioSymetrics, Inc.
2021 – present	Scientific Advisory Board Member, Constantiam Biosciences, Inc.
2015 – present	Scientific Advisory Board Member, SeqWell, Inc.
2015 – 2017	Scientific Advisory Board Member, Ranomics, Inc.

Selected Editorial and Review Experience

2017 – present	Advisory Editorial Board Member, Molecular Systems Biology
2017 – 2019	Chair, NIH Genomics, Computational Biology and Technology Study Section
2014 – 2019	Member, NIH Genomics, Computational Biology and Technology Study Section
2011 – 2015	Associate Editor, G3: Genes Genomes Genetics
2010 – 2013	Associate Editor, PLoS Computational Biology
2001 – 2004	Associate Editor, Bioinformatics

Selected Teaching Contributions

2022 – present	Lecturer, MMG3204H: Practical Applications of Genome Interpretation (6-week graduate course)
2022 – present	Lecturer, MMG1001: Foundational Genetics Approaches (12-week graduate course)
2019 – present	Founding Co-Instructor/Co-Coordinator, Foundational Computational Biology I and II (2 6-week graduate topic courses).
2017- 2019	Founding Instructor, Computational Biology and Bioinformatics (12-week graduate course).
2016 – present	Lecturer, MGY360H1: Whole-Genome Sequencing and Analysis Laboratory
2013 – 2015	Co-founder & Course Coordinator, MGY360H1: Whole-Genome Sequencing and Analysis Laboratory
2011 – 2013	Lecturer, University of Toronto, Topic Course on Functional Genomics & Proteomics: Experimental Approaches

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- 2008 – 2010 Lecturer, Harvard Medical School, Microbiology 230: Analysis of the Biological Literature.
- 2007 – 2009 Lecturer, Harvard Medical School, BCMP 207: Molecular Approaches to Drug Action, Discovery and Design
- 2007 – 2009 Lecturer, Neurobiology 300. Tools for Statistical Inference in Experimental Science
- 2004 – 2010 Co-founder & Lecturer, Harvard Medical School, Biophysics 205: Computational and Experimental Functional Genomics
- 2001 – 2009 Lecturer, Harvard Medical School, BCMP 201: Proteins: Structure, Function and Catalysis

Selected Conference Organizing Roles (within past 5 years)

- 2023 Organizing Committee, Cold Spring Harbor Laboratories Meeting on Network Biology, New York, United States
- 2022 Lead Organizer, Mutational Scanning Symposium, Toronto, ON
- 2020 Program Committee, Yeast Genetics Meeting (YGM) and The Allied Genetics Conference (TAGC).
- 2020 Organizer, Atlas of Variant Effects Planning Workshop, Seattle, WA

Selected Speaking Roles (international events within past 5 years)

- 2023 Invited Speaker, Society for Inherited Metabolic Disorders (SIMD), Salt Lake City, UT
- 2023 Speaker, Quantitative Biology Meeting, Puerto Rico, US
- 2023 Speaker, Invitae, Inc., San Francisco, CA
- 2023 Speaker, Genentech, San Francisco, CA
- 2023 Speaker, Helix, Inc.
- 2022 Speaker, Ambry Genetics, Inc.
- 2022 Speaker, NIH/NHGRI Center for Multiplexed Analysis of Phenotypes Retreat
- 2022 Speaker, BridgeBio Inc.
- 2022 Invited Speaker, Workshop on Functional Analysis of Sequence Variants in Hereditary Breast and Ovarian Cancer Genes, Leiden, NL
- 2021 Speaker, Alexion Pharmaceuticals, Inc.
- 2021 Speaker, Amicus Therapeutics, Inc.
- 2021 Speaker, Beam Therapeutics, Inc.
- 2021 Speaker, Dynacure, Inc.,
- 2021 Seminar Speaker, Boston Children's Hospital, Boston, MA
- 2021 Invited Speaker, Belgrade Bioinformatics Conference, Belgrade, Croatia
- 2021 Speaker, NIH/NHGRI Center for Multiplexed Analysis of Phenotypes Retreat
- 2021 Speaker, Deep Genomics, Inc.
- 2021 Invited Speaker, NIH/NHGRI&NCATS Meeting on Acidemias and Homocysteinuria
- 2021 Invited Speaker, Cold Spring Harbor Asia Meeting on Yeast and Life Sciences
- 2021 Speaker, HTG Therapeutics, Inc.
- 2021 Speaker, Mutational Scanning Symposium
- 2021 Seminar Speaker, Weizmann Institute (Rehovot, Israel) [canceled due to pandemic]
- 2021 Seminar Speaker, Alnylam Pharmaceuticals, Inc. Boston, MA
- 2020 Speaker, Atlas of Variant Effects Alliance Kickoff Workshop
- 2020 Speaker, NIH/NHGRI Center for Multiplexed Analysis of Phenotype Retreat
- 2020 Speaker, U Washington, Seattle Genome Sciences Summer Program Seminar
- 2020 Keynote speaker, 28th Intelligent Systems in Molecular Biology Variant Interpretation Community of Special Interest (VarI-COSI)

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- 2020 Invited Speaker, Hereditary Breast and Ovarian Cancer Variants of Uncertain Significance Workshop, Leiden, Netherlands [canceled due to pandemic]
- 2020 Invited Speaker, From Basic to Life Sciences X, Cavtat, Croatia [canceled due to pandemic]
- 2020 Invited Speaker/Panelist, NIH/ NCI SeqSPACE, "Sequencing Strategies for Population and Cancer Epidemiology studies" Webinar and Panel Discussion
- 2019 Participant, Illumina Workshop on Variants of Functional Significance, Seattle WA
- 2019 Speaker, One Brave Idea Foundation Meeting, Boston, MA, USA
- 2019 Seminar Speaker, Molecular Biosciences Department, U Texas, Austin, TX
- 2019 DFCI Center for Cancer Systems Biology/UMassMed Program in Systems Biology Retreat, Gloucester, MA
- 2019 Seminar Speaker, Univ. of Copenhagen, Copenhagen, Denmark
- 2019 Keynote speaker, 29th International Conference on Yeast Genetics and Molecular Biology (ICYGMB), Goteborg, Sweden
- 2019 Seminar Speaker, European Bioinformatics Institute, Hinxton, UK
- 2019 Invited Speaker, Myriad Genetics, Salt Lake City, UT
- 2019 Ambry Genetics Webinar
- 2019 American College of Medical Genetics and Genomics Annual Meeting, Seattle, WA
- 2019 Speaker, CIFAR Workshop on Personalized Genomics, Santa Cruz, CA
- 2019 Speaker, CIFAR Genetic Networks Program Meeting, Santa Cruz, CA
- 2019 Keynote Speaker, Cold Spring Harbor Laboratory Meeting on Network Biology, Cold Spring Harbor, NY
- 2019 Seminar Speaker, UCSF Dept of Bioengineering & Therapeutic Sciences, CA
- 2019 Speaker, Invitae, Inc., San Francisco, CA
- 2019 18th Annual Workshop at Bellairs - Single Cell and Massively Parallel Approaches, Holetown, Barbados
- 2018 NIH/NHGRI Centers of Excellence in Genomic Science Grantee Meeting
- 2018 CIFAR Genetic Networks Program Meeting, Toronto, ON
- 2018 Invited Speaker, Mt Sinai Icahn School of Medicine, New York City, NY
- 2018 CIFAR Molecular Architecture of Life workshop, Toronto, ON
- 2018 NIH Centers of Excellence in Genome Sciences Meeting, Chicago, IL
- 2018 Invited speaker, Dana Farber Cancer Institute Center for Cancer Systems Biology/UMassMedical Program in Systems Biology Retreat, Gloucester, MA
- 2018 Jasper Rine Symposium, Berkeley, CA
- 2018 One Brave Idea Webinar
- 2018 Speaker and Lecturer, CSHL Yeast Genetics and Genomics Course
- 2018 CIFAR-GETx Conference, Toronto ON
- 2018 Seminar Speaker, University of Ljubljana, Ljubljana, Slovenia
- 2018 Invited Seminar, University of Michigan, Ann Arbor, MI
- 2018 Invited Seminar, Department of Human Genetics, Univ. of Utah, UT
- 2018 Invited Speaker, Recursion Pharmaceuticals Systems Biology Minisymposium, UT

Peer-Reviewed Research Publications

1. B Floyd*, J Weile*, P Kannankeril, A Glazer, C Reuter, C MacRae, E Ashley*, D Roden*, *FP Roth**, V Parikh*[†]. Proactive variant effect mapping aids diagnosis in pediatric cardiac arrest. **Circulation: Genomic and Precision Medicine** (in press).
2. DK Kim*, B Weller*, CW Lin*, D Sheykhkarimli*, JJ Knapp*, G Dugied, A Zanzoni, C Pons, MJ Tofaute, SB Maseko, K Spirohn, F Laval, L Lambourne, N Kishore, A Rayhan, M Sauer, V Young, H Halder, N Marin-de la Rosa, O Pogoutse, A Strobel, P Schwehn, R Li, S Rothballer, M Altmann, P

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- Cassonnet, G Dugied, AG Cote, LE Vergara, I Hazelwood, BB Liu, M Nguyen, R Pandiarajan, B Dohai, PAR Coloma, J Poirson, P Giuliana, L Willems, M Taipale, Yves Jacob, Tong Hao, DE Hill, C Brun, JC Twizere, D Krapmann, M Heinig, C Falter, P Aloy, C Demeret[†], M Vidal[†], MA Calderwood[†], *FP Roth*[†], P Falter-Braun[†]. A proteome-scale map of the SARS-CoV-2 human contactome. **Nature Biotechnology** 41(1):140-149 (2023).
3. D Choi, N Khan, L Monterminib, N Tawil, B Meehan, D-K Kim, *FP Roth*, M Divangahi, J Rak[†]. Quantitative proteomics and biological activity of extracellular vesicles engineered to express SARS-CoV-2 spike protein. **Journal of Extracellular Biology** 2022;1:e58 (2022).
 4. D Kuang, J Weile, N Kishore, AF Rubin, S Fields, DM Fowler, *FP Roth*[†]. MaveRegistry: a collaboration platform for multiplexed assays of variant effect. 37(19):3382-3383 **Bioinformatics** (2021).
 5. D Kuang, R Truty, J Weile, B Johnson, K Nykamp, C Araya, RL Nussbaum, *FP Roth*[†]. Prioritizing genes for systematic variant effect mapping. **Bioinformatics** 36(22-23):5448-5455 (2021).
 6. J-H Kim, Y Seo, M Jo, H Jeon, Y-S Kim, E-J Kim, D Seo, W-H Lee, SR Kim, N Yachie, Q Zhong, M Vidal, *FP Roth*, K Suk[†]. A genetic interaction screen in yeast identifies genes relevant to human PAK1 function in glioma. **Journal of Biological Chemistry** 295(50):16906-16919 (2020).
 7. J van Leeuwen[†], C Pons, G Tan, JZ Wang, J Hou, J Weile, M Gebbia, W Liang, E Shuteriqi, Z Li, M Lopes, M Ušaj, A Dos Santos Lopes, N van Lieshout, CL Myers, *FP Roth*, P Aloy, BJ Andrews[†] & C Boone[†]. Systematic analysis of bypass suppression of essential genes. **Molecular Systems Biology** 16(9):e9828 (2020).
 8. MA Chiasson, NJ Rollins, JJ Stephany, KA Sitko, KA Matreyek, M Verby, S Sun, *FP Roth*, D DeSloover, D Marks, AE Rettie, DM Fowler[†]. Multiplexed measurement of variant abundance and activity reveals VKOR topology, active site and human variant impact. **eLife** 2020;9:e58026 (2020).
 9. D-K Kim, JJ Knapp, D Kuang, A Chawla, P Cassonnet, H Lee, D Sheykhkarimli, P Samavarchi-Tehrani, H Abdouni, A Rayhan, Li R, Pogoutse O, E Coyaud, S van der Werf, C Demeret, A-C Gingras, B Raught, Y Jacob[†], *FP Roth*[†]. A flexible genome-scale resource of SARS-CoV-2 coding sequence clones. **G3: Genes|Genomes|Genetics** 10(9):3399-3402 (2020).
 10. D Kuang, J Weile, R Li, TW Ouellette, JA Barber, *FP Roth*[†]. MaveQuest: a web resource for planning experimental tests of human variant effects. **Bioinformatics** 36(12):3938-3940 (2020).
 11. K Luck^{*}, D-K Kim^{*}, L Lambourne^{*}, K Spirohn^{*}, BE Begg, W Bian, R Brignall, T Cafarelli, FJ Campos-Laborie, B Charloteaux, D Choi, AG Cote, M Daley, S Deimling, A Desbuleux, A Dricot, M Gebbia, MF Hardy, N Kishore, JJ Knapp, IA Kovács, I Lemmens, MW Mee, JC Mellor, C Pollis, C Pons, AD Richardson, S Schlabach, B Teeking, A Yadav, M Babor, D Balcha, O Basha, S-F Chin, SG Choi, C Colabella, G Coppin, C D'Amata, D De Ridder, S De Rouck, M Duran-Frigola, H Ennajaoui, F Goebels, A Gopal, G Haddad, M Helmy, Y Jacob, Y Kassa, R Li, N van Lieshout, A MacWilliams, D Markey, JN Paulson, S Rangarajan, J Rasla, A Rayhan, T Rolland, A San Miguel, Y Shen, D Sheykhkarimli, GM Sheynkman, E Simonovsky, M Tasan, A Tejada, J-C Twizere, Y Wang, R Weatheritt, J Weile, Y Xia, X Yang, E Yeger-Lotem, Q Zhong, P Aloy, GD Bader, J De Las Rivas, S Gaudet, T Hao, J Rak, J Tavernier, V Tropepe, DE Hill[†], M Vidal[†], *FP Roth*[†], MA Calderwood[†]. A reference map of the human protein interactome. **Nature** 580(7803):402-408 (2020).
 12. PM Muller^{*}, J Rademacher^{*}, RD Bagshaw^{*}, C Wortmann, C Barth, J van Unen, KM Alp, G Giudice, RL Eccles, LE Heinrich, P Pascual-Vargas, M Sanchez-Castro, L Brandenburg, G Mbamalu, M Tucholska, L Spatt, MT Czajkowski, RW Welke, S Zhang, V Nguyen, T Rustemi, P Trnka, K Freitag, B Larsen, O Popp, P Mertins, A-C Gingras, *FP Roth*, K Colwill, C Bakal, O Pertz, T Pawson, E Petsalaki[†], O Rocks[†]. Systems-analysis of RhoGEF/RhoGAP regulatory proteins reveals spatially organized Rac1 signaling from integrin adhesions. **Nature Cell Biology** 22(4):498-511 (2020).
 13. S Sun^{*}, J Weile^{*†}, M Verby, Y Wu, Wang, Y, AG Cote, I Fotiadou, J Kitaygorodsky, M Vidal, J Rine, P Ješina, V Kozich[†], *FP Roth*[†]. A proactive genotype-to-patient-phenotype map for cystathionine beta-synthase. **Genome Medicine** 12(1):13 (2020).
 14. JH Kim, Y Seo, M Jo, H Jeon, WH Lee, N Yachie, Q Zhong, M Vidal, *FP Roth*, K Suk. Yeast-based genetic interaction analysis of human kinome. **Cells**. 9(5):1156 (2020).

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15. A Celaj, M Gebbia, L Musa, AG Cote, J Snider, V Wong, M Ko, T Fong, P Bansal, JC Mellor, G Seesankar, M Nguyen, S Zhou, L Wang, N Kishore, I Stagljar, Y Suzuki, N Yachie[†] & *FP Roth*[†]. Highly combinatorial genetic interaction analysis reveals a multi-drug transporter influence network. **Cell Systems** 9:1-14 (2020).
16. D Esposito*, J Weile*, J Shendure, LM Starita, AT Papenfuss, *FP Roth*[†], DM Fowler[†] & AF Rubin[†]. MaveDB: an open-source platform to distribute and interpret data from multiplexed assays of variant effect. **Genome Biology** 20(1):223 (2019).
17. J Zhang, LN Kinch, Q Cong, P Katsonis, O Lichtarge, C Savojardo, G Babbi, PL Martelli, E Capriotti, R Casadio, A Garg, D Pal, J Weile, S Sun, M Verby, *FP Roth* & NV Grishin[†]. Assessing predictions on fitness effects of missense variants in calmodulin. **Human Mutation** 40(9):1463-1473 (2019).
18. F Yang, D-K Kim, H Nakagawa, S Hayashi, S Imoto, L Stein, *FP Roth*[†]. Quantifying immune-based counterselection of somatic mutations. **PLoS Genetics** 15(7):e1008227 (2019).
19. Y Wu, J Weile, AG Cote, S Sun, J Knapp, M Verby, *FP Roth*[†]. A web application and service for imputing and visualizing missense variant effect maps. **Bioinformatics** 35(17):3191-3193 (2019).
20. E Deneault, M Faheem, SH White, DC Rodrigues, S Sun, W Wei, A Piekna, T Thompson, JL Howe, L Chalil, V Kwan, S Walker, P Pasceri. *FP Roth*, RKC Yuen, KK Singh, J Ellis[†] & SW Scherer[†]. CNTN5-/+ or EHMT2-/+ iPSC-derived neurons from individuals with autism develop hyperactive neuronal networks. **eLife** 8:e40092 (2019).
21. A Sokolov, S Ashenden, N Sahin, R Lewis, N Erdem, E Ozaltan, A Bender, *FP Roth*, M Cokol[†]. Characterizing ABC-transporter substrate-likeness using a clean-slate genetic background. **Frontiers in Pharmacology** 10:448 (2019).
22. ZB Weinstein, N Kuru, S Kiriakov, AC Palmer, AS Khalil, PA Clemons, MH Zaman, *FP Roth*, M Cokol[†]. Modeling the impact of drug interactions on therapeutic selectivity. **Nature Communications** 9(1):3452 (2018).
23. D Choi, L Montermini, DK Kim, B Meehan, *FP Roth*, J Rak[†]. The impact of oncogenic EGFRvIII on the proteome of extracellular vesicles released from glioblastoma cells. **Molecular and Cellular Proteomics**. 17(10):1948-1964 (2018).
24. JJ Diaz-Mejia, A Celaj, JC Mellor, A Cote, A Balint, B Ho, P Bansal, F Shaeri, M Gebbia, J Weile, M Verby, A Karkhanina, Y Zhang, C Wong, J Rich, D Prendergast, G Gupta, S Ozturk, D Durocher, GW Brown, *FP Roth*[†]. Mapping DNA-damage-dependent genetic interactions in yeast via party mating and barcode fusion genetics. **Molecular Systems Biology** (2018).
25. J Weile, S Sun, AG Cote, J Knapp, M Verby, JC Mellor, Y Wu, C Pons, C Wong, N van Lieshout, F Yang, M Tasan, G Tan, S Yang, DM Fowler, R Nussbaum, JD Bloom, M Vidal, DE Hill, P Aloy & *FP Roth*[†]. A framework for exhaustively mapping functional missense variants. **Molecular Systems Biology** 13(12):957 (2017).
26. I Vigentini[†], M Gebbia, A Belotti, R Foschino, *FP Roth*. CRISPR/Cas9 system as a valuable genome editing tool for wine yeasts with application to decrease urea production. **Frontiers in Microbiology** 8(2194):1-11 (2017).
27. M Jo, AY Chung, N Yachie, M Seo, H Jeon, Y Nam, Y Seo, E Kim, Q Zhong, M Vidal, HC Park[†], *FP Roth*[†], K Suk[†]. Yeast genetic interaction screen of human genes associated with amyotrophic lateral sclerosis: identification of MAP2K5 kinase as a potential drug target. **Genome Research** 27(9):1487-1500 (2017).
28. J Zhang, LN Kinch, Q Cong, J Weile, S Sun, AG Cote, *FP Roth*, NV Grishin[†]. Assessing predictions of fitness effects of missense mutations in SUMO-conjugating enzyme UBE2I. **Human Mutation**. 38(9):1051-1063 (2017).
29. A Celaj, U Schlecht, J Smith, W Xu, S Suresh, M Miranda, AM Aparicio, M Proctor, RW Davis, *FP Roth*[†] & RP St.Onge[†]. Quantitative analysis of protein interaction network dynamics in yeast. **Molecular Systems Biology** 13(7):934 (2017).

30. F Yang, S Sun, G Tan, M Costanzo, DE Hill, M Vidal, BJ Andrews, C Boone & *FP Roth*[†]. Identifying pathogenicity of human variants via paralog-based yeast complementation. **PLOS Genetics** 13(5):e1006779 (2017).
31. MJ Betts, O Wichmann, M Utz, T Andre, E Petsalaki, P Minguéz, L Parca, *FP Roth*, A-C Gavin, P Bork & RB Russell[†]. Systematic identification of phosphorylation-mediated protein interaction switches. **PLOS Computational Biology** 13(3):e1005462 (2017).
32. C Cenik, HN Chua, G Singh, A Akef, MP Snyder, AF Palazzo, MJ Moore[†] & *FP Roth*[†]. A common class of transcripts with 5'-intron depletion, distinct early coding sequence features, and N1-methyladenosine modification. **RNA** 23(3):270-283 (2017).
33. J van Leeuwen^{*}, C Pons^{*}, JC Mellor, TN Yamaguchi, H Friesen, J Koschwanez, MM Ušaj, M Pechlaner, M Takar, M Ušaj, B VanderSluis, K Andrusiak, P Bansal, A Baryshnikova, C Boone, J Cao, A Cote, M Gebbia, G Horecka, I Horecka, E Kuzmin, N Legro, W Liang, N van Lieshout, M McNee, B-J San Luis, F Shaeri, E Shuteriqi, S Sun, Lu Yang, J-Y Youn, M Yuen, M Costanzo, A-C Gingras, P Aloy, C Oostenbrink, A Murray, TR Graham, CL Myers[†], BJ Andrews[†], *FP Roth*[†] & C Boone[†]. Exploring genetic suppression interactions on a global scale. **Science** 354(6312):pii:aag0839 (2016).
34. T Shekhar-Guturja, GM Gunaherath, EM Wijeratne, JP Lambert, AF Averette, SC Lee, T Kim, YS Bahn, F Tripodi, R Ammar, K Döhl, K Niewola-Staszewska, L Schmitt, RJ Loewith, *FP Roth*, D Sanglard, D Andes, C Nislow, P Coccetti, AC Gingras, J Heitman, AA Gunatilaka & LE Cowen[†]. Dual action antifungal small molecule modulates multidrug efflux and TOR signaling. **Nature Chemical Biology** 12(10):867-875 (2016).
35. *FP Roth*[†] & J Wakeley[†]. Taking exception to human eugenics. (Letter to the Editor) **Genetics** 206(2):821-823 (2016).
36. Q Zhong^{*†}, SJ Pevzner^{*}, T Hao, Y Wang, R Mosca, J Menche, M Taipale, M Tašan, C Fan, X Yang, P Haley, RR Murray, F Mer, F Gebreab, S Tam, A MacWilliams, A Dricot, P Reichert, B Santhanam, L Ghamsari, MA Calderwood, T Rolland, B Charloteaux, S Lindquist, AL Barabási, DE Hill, P Aloy, ME Cusick, Y Xia, *FP Roth*[†] & M Vidal[†]. An inter-species protein-protein interaction network across vast evolutionary distance. **Molecular Systems Biology** 12(4):865 (2016).
37. N Yachie^{*†}, E Petsalaki^{*}, JC Mellor, J Weile, Y Jacob, M Verby, SB Ozturk, S Li, AG Cote, R Mosca, JJ Knapp, M Ko, A Yu, M Gebbia, N Sahni, S Yi, T Tyagi, D Sheykhkarimli, JF Roth, C Wong, L Musa, J Snider, Y-C Liu, H Yu, P Braun, I Stagljar, T Hao, MA Calderwood, L Pelletier, P Aloy, DE Hill, M Vidal & *FP Roth*[†]. Pooled-matrix protein interaction screens using Barcode Fusion Genetics. **Molecular Systems Biology** 12(4):863 (2016).
38. S Sun, F Yang, G Tan, M Costanzo, R Oughtred, J Hirschman, C Theesfeld, P Bansal, N Sahni, S Yi, A Yu, T Tyagi, C Tie, DE Hill, M Vidal, BJ Andrews, C Boone, K Dolinski & *FP Roth*[†]. An extended set of yeast-based functional assays accurately identifies human disease mutations. **Genome Research** 26(5):670-80 (2016).
39. X Yang, J Coulombe-Huntington, S Kang, GM Sheynkman, T Hao, A Richardson, S Sun, F Yang, YA Shen, RR Murray, K Spirohn, BE Begg, M Duran-Frigola, A MacWilliams, SJ Pevzner, Q Zhong, SA Trigg, S Tam, L Ghamsari, N Sahni, S Yi, MD Rodriguez, D Balcha, G Tan, M Costanzo, B Andrews, C Boone, XJ Zhou, K SalehiAshtiani, B Charloteaux, AA Chen, MA Calderwood, P Aloy, *FP Roth*[†], DE Hill[‡], LM Iakoucheva[†], Y Xia[†] & Marc Vidal[†]. Widespread expansion of protein interaction capabilities by alternative splicing. **Cell** 164(4):805-817 (2016).
40. TV Vo, J Das, MJ Meyer, NA Cordero, N Akturk, X Wei, BJ Fair, AG Degatano, R Fragoza, LG. Liu, A Matsuyama, M Trickey, S Horibata, A Grimson, H Yamano, M Yoshida, *FP Roth*, JA Pleiss, Y Xia & H Yu[†]. A proteome-wide fission yeast interactome reveals network evolution principles from yeasts to human. **Cell** 164(1-2):310-23 (2016).
41. T Hart^{*}, MC Chandrashekar^{*}, M Aregger^{*}, Z Steinhart, KR Brown, G MacLeod, M Mis, M Zimmermann, A Fradet-Turcotte, S Sun, P Mero, P Dirks, S Sidhu, *FP Roth*, OS Rissland, D Durocher, S Angers & J Moffat[†]. High-resolution CRISPR screens reveal fitness genes and genotype-specific cancer liabilities. **Cell** 163(6):1515-26 (2015).

42. E Petsalaki[†], AO Helbig, A Gopal, A Pasculescu, *FP Roth*[†] & T Pawson. SELPHI: Correlation-based identification of kinase-associated networks from global phospho-proteomics datasets. **Nucleic Acids Research** 43(W1):W276-82 (2015).
43. N Sahni, S Yi, M Taipale, JI Fuxman Bass, J Coulombe-Huntington, F Yang, J Peng, J Weile, GI Karras, Y Wang, IA Kovács, A Kamburov, I Krykbaeva, MH Lam, G Tucker, V Khurana, A Sharma, Y-Y Liu, N Yachie, Q Zhong, Y Shen, A Palagi, A San-Miguel, C Fan, D Balcha, A Dricot, DM Jordan, JM Walsh, AA Shah, X Yang, A Stoyanova, A Leighton, MA Calderwood, Y Jacob, ME Cusick, K Salehi-Ashtiani, LJ Whitesell, S Sunyaev, B Berger, A-L Barabási, B Charloteaux, DE Hill, T Hao, *FP Roth*[‡], Y Xia[‡], AJM Walhout[‡], S Lindquist^{†‡} & M Vidal^{†‡}. Widespread macromolecular interaction perturbations in human genetic disorders. **Cell** 161(3):647-660 (2015).
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- 2022 – 2026 NIH/NHLBI R01HL164675 (Roden, Roth, Ashley). “Generating missense variant effect maps for cardiovascular disease genes”
- 2021 – 2026 NIH/NHGRI UM1 HG011989 (Vidal). “Molecular phenotyping of ~100,000 coding variants across Mendelian disease genes”
- 2021 – 2023 Beam Therapeutics (Roth). “Variant effect maps to enable alpha-1 antitrypsin therapy”
- 2021 – 2023 NIH R21 HG011523 (Roth). “Technology for ten-minute resolution protein interaction mapping at proteome scale”
- 2021 – 2023 Biogen, Inc. (Roth). “Systematically and proactively testing variant effects for SOD1”
- 2019 – 2024 NIH/NIGMS R01 GM133185 (Vidal, Roth, Calderwood)
“Incomplete penetrance via edgetic suppression”
- 2019 – 2023 NIH/NHGRI RM1 HG010461 Center for Excellence in Genomic Sciences (Fowler).
"Center for the multiplexed assessment of phenotype"
- 2018 – 2025 CIHR Foundation Grant (Roth). “Systematically mapping context-dependent protein interactions and functional human sequence variation”

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