

Name		Position Title	
Mitchell Drumm		Associate Professor	
Education/Training			
Institution and Location	Degree	Year(s)	Field of Study
The Ohio State University	B.S.	1979-1983	Genetics
University of Michigan	Ph.D.	1984-1990	Human Genetics
University of Michigan	Postdoc	1990-1992	Human Genetics

- **Research and Professional Experience.**

1992 **Research Investigator**, Department of Internal Medicine, University of Michigan
1992-1999 **Assistant Professor**, Departments of Pediatrics and Genetics, Case
1992-2000 **Assistant Professor**, Department of Genetics, Case Western Reserve University
1999-present **Associate Professor**, Department of Pediatrics, Case Western Reserve University
2000-present **Associate Professor**, Department of Genetics, Case Western Reserve University

- **Professional Publications.**

1. F.S. Collins, M.L. Drumm, J.L. Cole, W.K. Lockwood, G.F. Vande Woude and M.C. Iannuzzi. Construction of a General Human Chromosome Jumping Library, with Application to Cystic Fibrosis. *Science* 235, 1046 (1987).
2. M.L. Drumm, C.L. Smith, M. Dean, J.L. Cole, M.C. Iannuzzi and F.S. Collins. Physical Mapping of the Cystic Fibrosis Region by Pulsed Field Gel Electrophoresis. *Genomics* 2, 346 (1988).
3. J.E. Richards, T.C. Gilliam, J.L. Cole, M.L. Drumm, J.J. Wasmuth, J.F. Gusella, F.S. Collins. Chromosome Jumping from D4S10 (G8) toward the Huntington Disease Gene. *Proc. Natl. Acad. Sci. USA* 85, 6437 (1988).
4. R.P. Erickson, C.E. Ross, J.L. Gorski and M.L. Drumm. Bkm sequences from the human X chromosome contain large clusters of GATA/GACA repeats. *Ann. Hum. Genet.* 52, 167 (1988).
5. M.C. Iannuzzi, M. Dean, M.L. Drumm, N. Hidaka, J.L. Cole, A. Perry, C. Stewart, B. Gerrard, F.S. Collins. Isolation of Additional Polymorphic Clones from the Cystic Fibrosis Region, using Chromosome Jumping from D7S8. *Am. J. Hum. Genet.* 44, 695 (1989).
6. J.M. Rommens, M.C. Iannuzzi, B. Kerem, M.L. Drumm, G. Melmer, M. Dean, R. Rozmahel, J.L. Cole, D. Kennedy, N. Hidaka, et al. Identification of the Cystic Fibrosis Gene: Chromosome Walking and Jumping. *Science* 245, 1059 (1989).
7. J.R. Riordan, J.M. Rommens, B. Kerem, N. Alon, R. Rozmahel, Z. Grzelczak, J. Aielensky, S. Lok, N. Plavsic, J. Chou, M.L. Drumm, M.C. Iannuzzi, F.S. Collins, L. Tsui. Identification of the Cystic Fibrosis Gene: Cloning and Characterization of Complementary DNA. *Science* 245, 1066 (1989).
8. M. Dean, M.L. Drumm, C. Stewart, B. Gerrard, A. Perry, N. Hidaka, J.L. Cole, F.S. Collins, M.C. Iannuzzi. Approaches to Localizing Disease Genes as Applied to Cystic Fibrosis. *Nucleic Acids Res.* 18, 345 (1990).
9. M. Dean, C. Stewart, A. Perry, B. Gerrard, T. Beck, U. Rapp, M. Drumm, M. Iannuzzi, F. Collins and S.O'Brien. Genetic Markers for Oncogenes, Growth Factors, and Cystic Fibrosis in *Modern Trends in Human Leukemia VIII*, Springer-Verlag 1989. p 360-365.
10. M.C. Iannuzzi, R.C. Stern, F.S. Collins, C. Tom Han, N. Hidaka, T.V. Strong, L. Becker, M.L. Drumm, M.B. White, B. Gerrard and M. Dean. Two Frameshift mutations in the Cystic Fibrosis gene associated with mild disease. *Am. J. Hum. Genet.* 48, 227 (1991).
11. M.L. Drumm, H.A. Pope, W.H. Cliff, J.M. Rommens, S.A. Marvin, L.-C. Tsui, F.S. Collins, R.A. Frizzell and J.M. Wilson. Corection of the Cystic Fibrosis Defect In Vitro by Retrovirus-Mediated Gene Transfer. *Cell* 62, 1227 (1990).
12. D. Marchuk, M. Drumm, A. Saulino and F.S. Collins. Construction of T-vectors, a rapid and general system for direct cloning of unmodified PCR products. *Nucleic Acids Res.* 19, 1154 (1991).

13. B. Kluge-Beckerman, M.L. Drumm and M.D. Benson. Nonexpression of the human serumamyloid A three (SAA3) gene. *DNA and Cell Biology* 10, 651 (1991)
 14. M.L. Drumm, D.J. Wilkinson, L.S. Smit, R.T. Worrell, T.V.Strong, R.A. Frizzell, D.C. Dawson and F.S. Collins. Chloride Conductance Expressed by Δ F508 and Other Mutant CFTRs in *Xenopus* Oocytes. *Science* 254, 1797 (1991).
 15. R.D. Krauss, J.K. Bubien, M.L. Drumm, T. Zheng, S.C. Peiper, F.S. Collins, K.L. Kirk, R.A. Frizzell and T.A. Rado. Transfection of wild type CFTR into cystic fibrosis lymphocytes restores chloride conductance at G₁ of the cell cycle. *EMBO J.* 11, 875-883 (1992).
 16. T.R. Flotte, S.A. Afione, R. Solow, M.L. Drumm, D. Markakis, W.B. Guggino, P.L. Zeitlin and B.J. Carter. Expression of the Cystic Fibrosis Transmembrane Conductance Regulator from a Novel Adeno-associated Virus Promoter. *J. Biol. Chem.* 268, 3781-3790 (1993).
 17. J.L. Overholt, A. Saulino, M.L. Drumm, R.D. Harvey. Rectification of whole-cell CFTR chloride current. *Am. J. Physiol.* 268, C636-C646 (1995).
 18. R.C. Stern, C.F. Doershuk and M.L. Drumm. 3849+10kb C->T mutation and disease serveryity in cystic fibrosis. *Lancet* 346:274-276 (1995).
- Drumm, M.L.
19. T.J Kelley, L. Al-Nakkash and M.L. Drumm. CFTR-mediated chloride permeability is regulated by type III phosphodiesterases in airway epithelial cells. *Am. J. Resp. Cell Mol. Biol.* 13, 657-664 (1995)
 20. J. Xie, M.L. Drumm, J. Ma and P.B. Davis. Intracellular loop between TM4 and TM5 of CFTR is involved in regulation of chloride channel conductance state. *J. Biol. Chem.* 270, 28084-28091 (1995).
 21. T. Tao, J. Xie, M.L. Drumm, J. Zhao, P.B. Davis and J. Ma. Slow conversions among subconductance states of cystic fibrosis transmembrane conductance regulator chloride channel. *Biophysical J.* 70, 743-753 (1996).
 22. J. Ma, J.E. Tasch, T. Tao, J. Zhao, J. Xie, M.L. Drumm and P.B. Davis. Phosphorylation-dependent block of the cystic fibrosis transmembrane conductance regulator chloride channel by exogenous R domain protein. *J. Biol. Chem.* 271, 7351-7356 (1996).
 23. T.J. Kelley, L. Al-Nakkash, C.U. Cotton and M.L. Drumm. Activation of Endogenous Δ F508 CFTR by Phosphodiesterase Inhibition. *J. Clin. Invest.* 98, 513-520 (1996).
 24. J. Zhao, B Zerhusen, J. Xie, M.L. Drumm, P.B. Davis and J. Ma. Rectification of CFTR chloride channels mediated by extracellular divalent cations *Biophysical J.* 71, 2458-2466 (1996).
 25. J. Xie, M.L. Drumm, J. Zhao, J. Ma and P.B. Davis. Human epithelial CFTR without exon 5 maintains partial chloride channel function in intracellular membranes. *Biophysical J.* 71, 3148-3156 (1996).
 26. T.J. Kelley, L. Al-Nakkash and M.L. Drumm. C-natriuretic peptide increases chloride permeability in normal and CF airway cells. *Am. J. Resp. Cell Mol. Biol.* 16, 464-470 (1997).
 27. T.J. Kelley, K.R. Thomas, Laura J.H. Milgram and M.L. Drumm. *In vivo* activation of Δ F508 CFTR in murine nasal epithelium. *Proc. Natl. Acad. Sci. U.S.A.* 94, 2604-2608 (1997).
 28. T.J. Kelley, C.U. Cotton and M.L. Drumm. *In vivo* activation of CFTR-dependent chloride transport in murine airway epithelium by C-type natriuretic peptide. *Am. J. Physiol.* 273, L1065-L1072 (1997).
 29. J. Ma, J. Zhao, M.L. Drumm, J. Xie and P.B. Davis. Function of the R domain in the CFTR chloride channel. *J. Biol. Chem.* 272, 28133-28141 (1997).
 30. W.K. Steagall, R.J. Marsick, T.J. Kelley, and M.L. Drumm. Type II protein kinase A regulates CFTR in airway, pancreatic, and intestinal cells. *Am. J. Phys.* 274, C819-C826 (1998).
 31. M.K. Mansoura, S.S. Smith, A.D. Choi, N.W. Richards, T.V. Strong, M.L. Drumm, F.S. Collins and D.C. Dawson. Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) anion binding as a probe of the pore. *Biophysical Journal* 74, 1320-1332 (1998).
 32. T.J. Kelley, C.U. Cotton and M.L. Drumm. Regulation of amiloride-sensitive sodium absorption in murine airway epithelium by C-type natriuretic peptide. *Am. J. Physiol.* 274, L990-L996 (1998).

33. T.J. Kelley and M.L. Drumm. Inducible nitric oxide synthase expression is reduced in cystic fibrosis murine and human airway epithelial cells: Possible involvement in CF-related sodium hyperabsorption and susceptibility to bacterial infection. *J. Clin. Invest.* 102, 1200-1207 (1998).
34. H.L. Elmer, K.G. Brady, M.L. Drumm and T.J. Kelley. Nitric oxide-mediated regulation of transepithelial sodium and chloride transport in murine nasal epithelium. *Am. J. Phys.* 276, L466-L473 (1999)
35. J.F. Chmiel, M.L. Drumm, M.W. Konstan, T.W. Ferkol, and C.M. Kerckmar. Pitfall in the use of genotype analysis as the sole diagnostic criterion for cystic fibrosis. *Pediatrics* 103, 823-826 (1999).
36. W.K. Steagall and M.L. Drumm. Stimulation of cystic fibrosis transmembrane conductance regulator-dependent short circuit currents across \square F508 murine intestines. *Gastroenterology*, 116, 1379-1388 (1999).
37. K. G. Brady, T. J. Kelley, and M. L. Drumm. Examining basal chloride transport using the nasal potential difference response in a murine model. *Am. J. Phys.* 281, L1173-L1179 (2001)
38. L.M. Ulatowski, K.L. Whitmore, T. Romigh, A.S. VanderWyden, S.M. Satinover and M.L. Drumm. Strain-specific variants of the mouse *Cftr* promoter region reveal transcriptional regulatory elements. *Hum. Mol. Genet.* *in press*
39. A.M. van Heeckeren, M.D. Schluchter, M.L. Drumm, and P.B. Davis. Role of *Cftr* Genotype in the Response to Chronic *Pseudomonas aeruginosa* Lung Infection in Mice *Am J Physiol Lung Cell Mol Physiol.* *in press*
40. Drumm ML and Davis PB. Some like it hot: curcumin and CFTR. *Trends in Molecular Medicine*, 2004 In press.

- **Expertise Summary**

Dr. Drumm's areas of expertise are in genetics and molecular genetics. Over the past 20 years, he has been applying this expertise to cystic fibrosis. In addition to genetics, his expertise has expanded to the use of mouse models to study human genetic disease, focusing on the genetic regulation of ion transport and endocrine processes in cystic fibrosis.