## **OHSU CYSTIC FIBROSIS RESEARCH TEAM**

707 S.W. Gaines Road, CDRC-P Portland, Oregon 97239-2091 503-494-8023 (Dial option 5)

#### **ABOUT OUR TEAM**

The Cystic Fibrosis research team is composed of doctors and research coordinators who work closely with the rest of the adult and pediatric cystic fibrosis teams. In addition to conducting research, we like to consider ourselves an extra pair of eyes that help ensure our patients receive the best care possible!

#### **RESEARCH COORDINATORS**



Brendan Klein, M.P.H., CCRP	Annie Hanson, B.S.	Jenna Bucher, B.S., CCRC	Pierce Nusbaum, B.S.
503-418-8108	503-494-5487	503-494-7807	503-418-1169
kleinb@ohsu.edu	hansoan@ohsu.edu	bucherj@ohsu.edu	nusbaum@ohsu.edu

#### **RESEARCH DOCTORS/INVESTIGATORS**

Michael Powers, M.D. Kelvin MacDonald, M.D. Anne Stone, M.D. Gopal Allada, M.D. Aaron Trimble, M.D. Amy Garcia, M.D.

# **INTERESTED IN CF RESEARCH?** HERE'S HOW YOU CAN LEARN MORE AND GET INVOLVED:

- Ask your provider about research opportunities at your next CF appointment.
- Contact the research team using the information above.
- Contact the CF Foundation's Clinical Trials Hotline at 1-800-FIGHT-CF, or visit the Clinical Trial Finder website: <u>www.cff.org/Find</u>

### **CURRENT STUDIES AT OHSU**

Please contact the research team if you have questions about what studies you or your child may be eligible for.

Anti-Infecti TEACH: Tes tobramycin This study v inhaled tob Age 12 Years and Older	ve   Enrolling ting the effect o in people with o vill look at the ef ramycin and will <b>Mutations</b> No Mutation Requirement	f adding oral CF fect of adding use a placeb FEV1% Predicted 25-100%	azithromycin g oral azithron o control. Number of Visits 5	to inhaled hycin to Length of Participation 14 Weeks	Anti-Infective Aerovanc: Pha children 6 yea This study will vancomycin in placebo contro Age 22 years and younger	Enrolling ase 3 study of inl rs and older wit look at the effect adults and child ol. Mutations No Mutation Requirement	haled vancor h CF ctiveness of t lren 6 years a FEV1% Predicted 30-90%	nycin in adu he inhaled o ind older an Number of Visits 13	Ilts and Irug d will use a Length of Participation 12 months	
Observational   Enrollment on hold temporarily CHEC-SC: Sweat chloride observational study				Observational   Enrolling CF Activity Study: Inpatient Exercise Program during Hospitalization						
This study will look at sweat chloride concentration in people who are currently taking CFTR modulators.				program for su	This study will examine the potential benefits of an inpatient exercise program for subjects with CF.					
Age 4 Months	Mutations No Mutation	FEV1% Predicted No FEV1	Number of Visits 1	Length of Participation 1 day	<b>Age</b> Pediatric	Mutations No Mutation	FEV1% Predicted No FEV1	Number of Visits During	Length of Participation g Admission	
<b>Restore CFTR Function</b>   Enrolling TranslateBio: Phase 1/2 study of MRT5005, combined single and multiple ascending dose, administered by nebulization to adult subjects with CF					Anti-Inflammatory I Enrolling Corbus: Phase 2 study to evaluate efficacy and safety of Lenabasum in Cystic Fibrosis					
This study will examine the safety, tolerability and biological activity of MRT5005 administered by nebulization and will use a placebo control.				This study will look at the safety and effectiveness of the anti- pulmonary inflammation drug Lenabasum and will use a placebo control.						
Age	Mutations	FEV1% Predicted	Number of Visits	Length of Participation	Age	Mutations	FEV1% Predicted	Number of Visits	Length of Participation	
18 Years and Older	Both in Class I or II	50-90%	Part A: 10 Parts B/C: 15	Part A: 49 weeks Parts B/C: 53 Weeks	12 Years and Older	No Mutation Requirement	40-90%	10	32 weeks	