



CLINICAL TRIALS OF TEXAS, INC.

Promoting Health Through Research

Curriculum Vitae

Bruce Richard Lantry, MD, FAAP

7940 Floyd Curl Drive, Suite 700
San Antonio, Texas 78229

2040 Babcock Road, Suite 301
San Antonio, Texas 78229

EDUCATION

- 1995-1997 Pediatrics Residency, University of Texas Health Science Center San Antonio, San Antonio, Texas
- 1994-1995 Pediatrics Internship, University of Texas Health Science Center San Antonio, San Antonio, Texas
- 1990-1994 Doctor of Medicine, University of Texas Health Science Center San Antonio Medical School, San Antonio, Texas
- 1987-1990 Bachelor of Arts in Biology: with honors, University of Texas at Austin, Austin, Texas

LICENSE(S)

- 2010 Department of Public Safety
- 2009 DEA Permit
- 1997 Texas Medical Board, # K2944

CERTIFICATION(S)

- 2005 Board Certified in Pediatrics

PROFESSIONAL EXPERIENCE

- 1999-2003 Pediatric Section Chief, Christus Santa Rosa Hospital – Medical Center, San Antonio, Texas
- 1998-Present Practice, First Steps Pediatrics, San Antonio, Texas
- 1998-2003 Medical Director/Committee Chairperson, Columbia HCA/Methodist Children's Hospital, San Antonio, Texas
- 1997-1998 Medical Center Pediatrics, San Antonio, Texas

PROFESSIONAL MEMBERSHIP(S)

- American Academy of Pediatrics (Fellow)
- Pediatric Telephone Nurse Triage

HOSPITAL AFFILIATION(S)

- Columbia HCA/Methodist Children's Hospital, San Antonio, Texas
- Christus Santa Rosa Hospital – Medical Center, San Antonio, Texas
- Southwest Texas Methodist, San Antonio, Texas

RECENT PRESENTATIONS AND/OR PUBLICATIONS

Breastfeeding Education for Medical Professionals, Texas Breastfeeding Summit, 2002

Gomez, J., Lantry, B. "Current Use of Adequate Pre-participation History Forms For Heart Disease Screening of High School Athletes." *Archives of Pediatrics & Adolescent Medicine* Vol:153, July 1999

Lecture Presentation, IPV vs. OPV, Connaught Vaccines, 1998

RESEARCH AFFILIATION(S)

2008-Present Investigator, Clinical Trials of Texas, Inc, San Antonio, Texas

RESEARCH EXPERIENCE

2010-Present: A Phase 1/2a, Randomized, Double-Blind, Placebo-Controlled, Dose-Escalation Study To Evaluate The Safety, Tolerability, Immunogenicity And Vaccine-Like Viral Shedding Of "Study Drug", A Live, Attenuated Intranasal Vaccine Against Respiratory Syncytial Virus (RSV) And Parainfluenza Virus Type 3 (PIV3), In Healthy 6 to < 24 Month-Old Children And In 2 Month-Old Infants (CTT-000215)

2009-Present: An Observational Prospective Study to Assess Respiratory Syncytial Virus (RSV) Respiratory Events Among Premature Infants (32 to 35 Week Gestational Age) - Outcomes and Risk Tracking Study (The Report Study) (CTT-000198)

2009-Present: A Phase 1/2a, Randomized, Double-Blind, Placebo-Controlled Study To Evaluate The Safety, Tolerability, Immunogenicity, And Viral Shedding Of "Study Drug", A Live Attenuated Intranasal Vaccine Against Respiratory Syncytial Virus In Healthy 1 To <12 Month-Old Children (CTT-000180)

Signature and Date