

# Wisconsin Newborn Screening Study

## Specimen Information

### Status: Specimens are Available

Specimens were collected associated with clinical care over a longitudinal time period of at least 17 years. Left over specimen (plasma) was saved when available; thus the timeframe for specimen collection is variable. In the table below Visit 1 (baseline) represents the first sample taken and Visit 2 represents all other potential time points during this longitudinal study.

Visit #	Time From Baseline	Specimens Collected
1	+0 Days	EDTA plasma
2	+17 Years	EDTA plasma

### Study Design

**Study Type?** Interventional

**Randomized Study?** No

**Placebo Controlled?** No

**Length of Participation**

**Number of Study Visits?**

### Additional Information

**Phase?** Not Applicable

**Study Sponsor?** Farrell, Philip

**Study Drugs?** N/A

### Eligibility

**Age** 1 Years to 21 Years

**Mutation(s)** No Mutation Requirement

**FEV1% Predicated** No FEV1 Limit

**PA Status** Not Applicable

**Other** Other key inclusion criteria include: must have been born in the State of Wisconsin, must have been born between April 15, 1985 and June 30, 1994, must have had a valid newborn screening test for cystic fibrosis in the first 28 days of life and must have a sweat chloride test greater or equal to 60 mmol/Liter

# Study Results

## **CITATION:**

Pediatrics 2001,Jan(107(1)):1-13

For more information about the results of this study and where it was conducted, visit [ClinicalTrials.gov](https://clinicaltrials.gov).