

primary studies - published, non RCT

# Pubertal Height Growth and Adult Height in Cystic Fibrosis After Newborn Screening.

Code: PM27244789 Year: 2016 Date: 2016 Author: Zhang Z

Study design (if review, criteria of inclusion for studies)

Historical cohort study

## **Participants**

107 children born in 1985-1994 and followed through 2012

#### Interventions

Newborn screening (NBS)

### **Outcome measures**

Adolescent peak height velocity (PHV), and adult height. PHV was estimated by a semiparametric growth curve model and compared with Tanner reference.

#### Main results

Meconium ileus (MI; n = 25) was associated with the worst pubertal growth and adult height, including 1 child who did not experience apparent PHV; children with pancreatic sufficiency (n = 18) achieved the best growth (normal PHV and adult height). In children with pancreatic insufficiency without meconium ileus (n = 64), the subgroup most likely to benefit from NBS, screened children had similar PHV but better adult height compared with controls. Specifically, in boys, the screened group (n = 22) achieved normal PHV (9.5 cm at 13.5 years); the control group (n = 19) had similar onset age (13.6 years) but 0.6-cm lower magnitude (P = .08). In girls, the screened group (n = 10) had somewhat later (12.5 years vs 11.7 years, P = .12) and lower PHV (7.3 cm vs 7.9 cm, P = .33) than the controls (n = 13), coinciding with later menarche (13.6 years vs 12.2 years, P = .10). Adult height was taller in the screened than the control (50th vs 29th percentile, P = .02), even after adjusted for genetic potential (32nd vs15th percentile, P = .006). Differences in adult height were primarily attributable to NBS and better prepubertal growth.

#### **Authors' conclusions**

Early linear growth benefits of NBS were sustained through puberty, leading to better adult height in cystic fibrosis.

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## See also

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## Keywords

Neonatal Screening; Newborn; non pharmacological intervention - diagn; screening; diagnostic procedures;