Central precocious puberty (CPP) is a rare disorder in children characterized by premature reactivation of the hypothalamic-pituitary-gonadal (HPG) axis, causing early puberty and the development of secondary sexual characteristics, accelerated growth, and accelerated bone maturation. A small body of research links CPP with increased risk for central nervous system and congenital disorders, but little is known about the association of CPP with other medical conditions.

We explored a large-scale, retrospective claims database (Florida Medicaid) to examine potential associations of precocious puberty with other medical conditions.

**OBJECTIVES**

Among 720,931 children, 1,644 (0.23%) were diagnosed with precocious puberty. There was a significant difference in disease burden between groups (p<0.0001).

After adjustments, compared to Medicaid-enrolled children without the disorder, those with precocious puberty were:

- 6 times more likely to be diagnosed with hemiplegia (OR 6.3, 95% CI 3.8-10.4, p=0.0001);
- 5 times more likely to be diagnosed with cerebrovascular disease (OR 5.1, 95% CI 3.2-8.1, p=0.0001) or moderate or severe renal disease (OR 4.7, 95% CI 2.2-9.9, p=0.0001);
- 4 times more likely to be diagnosed with congestive heart failure (OR 3.9, 95% CI 2.1-7.0, p=0.0001), connective tissue disease (OR 3.7, 95% CI 1.4-9.8, p=0.009), or diabetes (OR 3.5, 95% CI 2.0-6.2, p=0.0001); and
- 3 times more likely to be diagnosed with chronic pulmonary disease (OR 3.1, 95% CI 2.2-4.3, p<0.0001) or AIDS (OR 2.8, 95% CI 1.6-4.9, p=0.0002).

**METHODS**

Data were obtained from Florida Medicaid computerized claims records, which contain basic demographic information; ICD and CPT diagnosis and treatments codes; and payment data. Information is patient de-identified and fully compliant with the HIPPA Privacy Rule. Subjects were identified from enrollees in the Florida Medicaid program who had a paid claim from July 1997 through June 2004.

We compared children aged 3-7 years with precocious puberty (ICD-9 258.1, which may include premature adrenarche in addition to CPP) to those aged 3-7 years without this diagnosis. Patients were age-, sex-, race-, and year-matched to controls in the year of precocious puberty diagnosis (one-to-many).

ICD-9 diagnoses were used to classify disease groups and disease burden (Charlson Index). The Charlson index is a weighted index of comorbidity that is widely used in retrospective claims analysis. Fisher's exact test was used to examine differences between groups and logistic regression to examine likelihood estimates.

**RESULTS**

**CONCLUSIONS**

This is the first large-scale analysis to show a strong association between ICD-9 claims-based diagnoses of precocious puberty (which may include premature adrenarche in addition to CPP) and medical disorders. Although researchers have examined the psychosocial consequences of precocious puberty, this study found that precocious puberty significantly increased the likelihood of children developing comorbid medical conditions, including hemiplegia, renal disease, congenital heart failure, cardiopulmonary disease, connective tissue disorders, diabetes, and AIDS. The mechanisms by which children with this hormonal disorder become at increased risk for these particular medical conditions should be explored.

**REFERENCES**