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The Effect of Antihypertensive Dosing on Hypertension in Children with Chronic Kidney Disease School of Medicine





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Background

- · Hypertension (HTN) is a highly prevalent and major risk factor for poor cardiovascular and renal outcomes in chronic kidney disease (CKD).
- · Previous research suggests that HTN is underdiagnosed and undertreated in children with CKD (Hypertension 2018;71:1-7).
- . To our knowledge no studies have investigated the effect of antihypertensive dose on blood pressure control in this population.

Objective

- To determine the effect of antihypertensive dose on HTN status in children with CKD.
- · Hypothesis: Uncontrolled HTN is associated with lower antihypertensive dose.

Methods

Study population: 255 participants studied in the Chronic Kidney Disease in Children (CKiD) study at their third visit.

Inclusion criteria: Age 1-16 years with estimated GFR 30-90 mL/min/1.73m², taking at least one antihypertensive medication, successful 24h ABPM study.

Exclusion: White-coat hypertension (n=5).

Study variables:

- Cumulative Drug Dose Index (cDDI): We developed a new quantitative tool, Drug Dose Index (DDI, Fig 1) which expresses dose as a ratio between the current dose and the maximum potential dose, accounting for age, weight and if indicated, renal dose adjustments. cDDI is the sum DDI for all antihypertensive agents taken by the subject.
- Primary outcome: HTN status classified into controlled HTN (cHTN) or uncontrolled HTN (uHTN = masked HTN or ambulatory HTN) based on 2017 AAP guidelines and 24h ABPM study.
- Secondary outcome: Left ventricular hypertrophy (LVH = LVMI>38g/m²).

Statistical analysis:

Univariate analysis: t-test and ANOVA (continuous variables) and chi-square test (categorical variables) used to compare cDDI between the outcome groups (Fig 4): a) cHTN vs. uHTN, b) cHTN, MH vs. AH, and c) LVH vs. no LVH.

Multivariate Logistic Regression Analysis:

• Two outcomes: Table 2) uHTN and Table 3) LVH.

Results

Drug Dose Index (DDI): Figure 1

A measure of drug dose relative to its maximum dose that accounts for:

- pediatric dosing
- renal dose adjustments

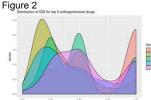
Weight (kg) x $dose_{mg/kg} > Max dose_{mg}$ (based on max $dose_{mg/kg}$) or dose_{ma}>Max dose_{ma}

 $DDI = dose_{mg}/Max dose_{mg}$ (up to maximum of 1)

 $DDI = dose_{mq/kq}/Max dose_{mq/kq}$ (up to maximum of 1)

WCH MH

105





 0.40 ± 1.13

 51.29 ± 21.02

 0.50 ± 0.50

 0.42 ± 1.15

58 (44%)

 0.61 ± 0.52

13 (9%)

0.19

pharmacological factors

BMI z-score

estimated GFR

Family history of

 0.57 ± 0.52

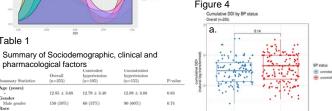
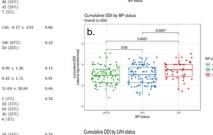
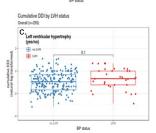


Figure 3

AAP 2017





Tables 2 and 3 Logistic Regression Analysis Model 1: Predictors of Uncontrolled Hypertension





Discussion

- No differences in cDDI between uHTN vs. cHTN. or LVH vs. no LVH, but higher cDDI was associated with AH on univariate
- · Non-Caucasian race, absence of RAASi and diuretic agents, and higher number of agents were associated with greater odds of
- Female gender, higher BMI z-score, lower eGFR, higher number of agents and lower cDDI were associated with greater odds of LVH.

Conclusions

- This was the first quantitative analysis of antihypertensive dose expressed as a newly developed measure, cDDI, and its relationship with hypertension status in children with CKD.
- · Lower dose (cDDI) is not a significant predictor of uHTN, however it may be associated with LVH.
- · RAASi and diuretic use were associated with lower odds of
- Further research is needed to validate the use of cDDI and the the roles of medication class and dose on blood pressure control in children with CKD.