COMMUNITY STUDY OF PREVALENCE AND RISK FACTORS FOR CHRONIC KIDNEY DISEASE AMONG PAEDIATRIC AGE GROUP IN ONDO WEST LOCAL GOVERNMENT AREA OF ONDO STATE NIGERIA

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INTRODUCTION AND

LITERATURE

The NKF/KDOQI in 2002 defined CKD as kidney damage manifested by structural or functional abnormalities lasting three or more months with or without decreased glomerular filtration rates (GFR) or a GFR< 60mls/min/ 1.73 m²

- Little is known about the epidemiology of chronic kidney disease (CKD) among the paediatric population especially in Sub-Sahara Africa.
- Due to the asymptomatic nature of early CKD
- A prevalence of 12.1 to 74.7 cases per million children has been reported previously.

- Nigerian data so far:
- ≻main source: major tertiary in-hospital data
- ≻Tip of the iceberg (not truly representative).
- Examples: 4.0% (Enugu); 4.5% (Mid-west); 1.6% (Ilorin); 3.1% (Uyo)

Okoro et al, 1999; Ibadin et al, 2003; Adedoyin et al, 2012; Ikpeme et al, 2014

- The highest percentage of paediatric CKD cases are potentially reversible (congenital)
- In Port-Harcourt, 28.9% of CKD were due to congenital disorders

- CKD in children is compounded by one or a combination of growth problems, nutrition, electrolyte imbalance, anaemia and hypertension.
- The child's body system copes less with uraemia resulting in high levels of mortality among them.
- Early detection and management of kidney malfunction is crucial to delay or prevent progression of CKD to ESRD.

OBJECTIVES

•We therefore set out to determine

prevalence of CKD

risk factors for CKD among children in

Ondo State.



- 114 school children whose parents/guardians gave consent were studied
- Children outside 2-17 years and those who were acutely ill were excluded
- Their bio data was recorded on a proforma
- Their weights and heights were obtained with a standard stadiometer (*RGZ-160 Lincon* Mark Medical England)

- BMI was calculated using wt/ht²
- BP was measured using *Accossons Mercury* Spyhgmomanometer with appropriate cuff for age on the right upper arm after 5 minutes rest to the nearest 2mmHg

- Blood samples were collected for Serum chemistry, haemogram, fasing lipids and albumin.
- eGFR was calculated using *Schwartz formula*
- Urine samples for analysis were obtained after adequate counsel of the parents/guardians
- Combi 10 Unistrip[®] was employed for urinalysis

- Height, BMI and BP percentiles were determined using the appropriate charts
- Written consent was obtained from the school authority
- Data was analyzed with SPSS 17.



Gender distribution of subjects





Distribution of subjects by class



Clinical and Lab parameters of subjects

Parameter	Mean	Standard deviation
Age (years)	8.99	4.26
eGFR (ml/min/1.73m ² BMI (kg/m ²)	86.59 16.80	27.6 3.09
SBP (mmHg)	97.88	16.29
DBP (mmHg)	57.84	11.66
PCV (%)	37.23	4.34
Serum creatinine (umol/L)	75.14	16.72
Total cholesterol (mmol/L)	4.20	0.83
Triglyceride (mmol/L)	1.85	0.29
HDL-cholesterol (mmol/L)	1.24	0.21
Albumin (g/L)	40.60	6.23

KDOQI STAGING OF CKD (n=104)

KDOQI STAGE	FREQUENCY	PERCENT
Stage 1	36	34.6%
Stage 2	60	57.7%
Stage 3	8	7.7%
Stage 4	0	0%
Stage 5	0	0%

CKD BY GENDER

KDOQI STAGE	Male	Female
Stage 1	20 (55.6%)	16 (44.4%)
Stage 2	29 (48.3%)	31 (51.7%)
Stage 3	5 (62.5%)	3 (37.5%)
Stage 4	0	0
Stage 5	0	0

eGFR by various criteria for CKD in our paediatric age-group

• In our study

≻(8) 7.7% (cut off <60ml/min/1.73m²)

>(37) 35.6% (cut off <75ml/min/1.73m²)

Hans Pottel et al. *Pediatric Nephrology: Journal of the International Pediatric Nephrology Association (2015), vol. 30, pp. 821-828*

Prevalence of risk factors

Risk factor	Percentage	Male	Female
		Freq (%)	Freq (%)
Pre- hypertension	12.4%	5 (35.7)	9 (64.3%)
Hypertension	12.8%	1 (10.0%)	9 (90%)
Overweight	7.9%	5 (55.6%)	4 (44.4%)
Obesity	5.3%	1 (16.7%)	5 (83.3%)

BMI vs eGFR



Inverse relationship between TC and eGFR



SBP vs eGFR



DBP vs eGFR



Conclusions

- There is an apparent high prevalence of CKD among paediatric population of Ondo State
- There is a high prevalence of risk factors among them
- Hypertension, obesity and dyslipidaemia showed a significant relationship to eGFR.

Recommendations

- Higher nos of subjects needed;
- Follow-up of subjects with established risk factors;
- Follow-up of subjects with reduced eGFR;
- Identify causes of reduced eGFR in them;
- Screening of siblings of subjects with risk factors and/or reduced eGFR.



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