Is Increased Echogenicity Related to a Decrease in Glomerular Filtration Rate? Objective Measurements in Pediatric Solitary Kidney Patients—A Retrospective Analysis

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Abstract

Quantitative measurements of renal echogenicity using a graphic program show close correlation with renal histology in adult patients, but this has neither been applied in pediatric patients nor correlated with glomerular filtration rate (GFR). To determine the direct relationship between echogenicity and GFR, we retrospectively analyzed 91 patients with a solitary functioning kidney under the age of 10, who underwent ultrasonography and serum cystatin C evaluation on a single day between January 2013 and December 2014. Echogenicity was quantified as previously reported. Echogenicity and kidney length were correlated with age-matched values of serum cystatin C-based GFR. Evaluation was performed at a median age of 17.1 months. GFR was low for age in eight of 54 right solitary kidney patients and four of 37 left solitary kidney patients. The right kidney-liver ratio was significantly elevated in the right decreased GFR group, while the left kidney-spleen ratio was not different in the left decreased GFR group. Age-matched longitudinal kidney length ratios were similar between the decreased and normal GFR groups for both sides. This is the first report to objectively prove the relationship between echogenicity and renal function in patients with a right solitary kidney. The right kidney-liver echogenicity ratio, measured objectively, showed feasibility in clinical practice as it showed a close relationship with decreased renal function when increased. However, absolute kidney echogenicity values, or the left kidney-spleen echogenicity ratio, were not independent markers for decreased renal function.

Introduction

Ultrasonography is the fundamental imaging modality in daily pediatric urological practice. The existence of hydronephrosis, longitudinal kidney length, corticomedullary differentiation, and cortical thickness are all known important parameters in the analysis of pediatric renal...
ultrasonography.[1] Renal echogenicity is another parameter as its increase is an important sign of renal parenchymal disease.[2, 3] However, it still has limited use, as it is too subjective to quantify. To overcome this issue, Manley and O’Neill first introduced quantitative measurement of renal echogenicity.[4] They scanned ultrasonographic images and measured the echogenicity of the right kidney by adjustment with the adjacent liver using graphic programs. Using a similar method, Moghazi et al. reported a close correlation between renal histopathology and echogenicity in adult patients.[5] However, this measurement has neither been correlated with glomerular filtration rate (GFR) nor applied in pediatric patients. To determine the direct relationship between echogenicity and GFR, only solitary kidney patients should be included in the analysis. We have applied this method in pediatric patients with a solitary kidney and correlated it with GFR for the first time in this study.

**Materials and Methods**

**Patients**

The database of the Department of Pediatric Urology at our institution was retrospectively queried for pediatric patients under the age of 10 years with a solitary functioning kidney. Because this study was performed retrospectively, the Institutional Review Board/Ethics Committee of Severance Hospital approved this study without the need for informed consent (approval number: 4-2014-0944). Patient records were anonymized and de-identified prior to analysis. Among these, 91 patients who underwent ultrasonography and serum cystatin C evaluation on the same day between January 2013 and December 2014 were included in this study. Exclusion criteria included: history of prematurity, grade 2 or higher hydronephrosis graded by the Society for Fetal Urology grading system,[6] known vesicoureteral reflux, the presence of another urinary tract anomaly or glomerular disease in the solitary functioning kidney, coexisting hepatobiliary or spleen disease, and systemic disease.

**Data collection**

Data regarding sex, laterality, age at evaluation, glomerular filtration rate, echogenicity of right kidney along with liver or left kidney with spleen, and longitudinal kidney length were collected. The glomerular filtration rate (GFR) was calculated based on the level of serum cystatin C using a previously reported formula by Grubb et al.: $84.69 \times \text{serum cystatin C (mg/L)}^{-1.680} \times 1.384$ (if child < 14 years).[7] The decreased GFR group was determined using recently reported reference GFR levels in Japanese children using the values of creatinine and cystatin C-based GFR.[8] When the calculated GFR was below the 2.5 percentile level for age, it was considered as ‘decreased GFR’. We additionally calculated cystatin C-based GFR Z-scores by age to analyze the correlation between renal function and echogenicity. The mean and standard deviation values were not shown in the original report; however, in contacting Uemura et al., we received these values and were thus able to calculate the Z-scores (Table 1).

| Age          | N  | 2.5% tile | 50.0% tile | 97.5% tile | Mean   | Standard deviation |
|--------------|----|-----------|------------|------------|--------|-------------------|
| 3–5 months   | 17 | 76.6      | 91.7       | 106.7      | 91.7   | 9.5               |
| 6–11 months  | 47 | 75.7      | 98.5       | 133.0      | 100.8  | 15.8              |
| 12–17 months | 31 | 83.3      | 106.3      | 132.6      | 106.6  | 13.7              |
| 18 months–16 years | 1042 | 83.5 | 113.1 | 156.7 | 115.2 | 18.3 |

Table 1. The reference values of GFR for each age group. The median, 2.5 percentile, and 97.5 percentile values were cited from a report by Uemura et al.[8] The mean and standard deviation values were cited from a report by Uemura et al. The mean and standard deviation values were not shown in their original report; however, in contacting them, we received these values and calculated the Z-scores (Table 1).
Renal ultrasonography and quantification of echogenicity

All patients were instructed to increase their water intake with their usual diet, without intravenous hydration or diuretics, and ultrasounds were performed after oral hydration according to the previous recommendation of the Society for Fetal Urology.[6] Parents and older children with communication skills were instructed to increase water intake before evaluation for hours. In the case of infants or younger children, parents were instructed to increase the water intake of the patient to the extent possible. Two experienced pediatric radiologists (MJK and MJL) performed all the ultrasonographic evaluations using an iU22 ultrasound unit (Philips Ultrasound, Bothell, WA, USA) with a 5–8 or 1–5 MHz convex transducer. Gain was adjusted for the optimization of imaging by the radiologists during the evaluation. Echogenicity was quantified as previously reported.[4, 5, 9] In brief, among pictures in same study, longitudinal images of the right kidney with the adjacent liver, or the left kidney with the spleen, were chosen and captured (Fig 1). In cases of a left solitary kidney, an image of the liver was additionally captured, although it was not located in the same image as the left kidney. In each picture, the region of interest (ROI) was outlined around the whole kidney and around the liver or spleen with ImageJ software, version 1.48v (National Institutes of Health, USA). Then, renal echogenicity was measured in 256-degree grayscale numerically from 0 to 255. The mean pixel density of the right kidney and liver were 70.3 and 67.3 respectively in the former patient with a right kidney-liver echogenicity ratio of 1.04, and 119.6 and 70.4 respectively in the latter patient with a ratio of 1.70.

Fig 1. Longitudinal images of the right solitary kidneys with adjacent livers were captured in a 6-month-old female infant with an estimated glomerular filtration rate of 117.2 mL/min/1.73 m² (A), (B) and in a 7-month-old male infant with an estimated glomerular filtration rate of 51.6 mL/min/1.73 m² (C), (D). In each picture, the region of interest was outlined around the entire kidneys (A), (C) and around the livers (B), (D) with ImageJ software, version 1.48v (National Institutes of Health, USA). Renal echogenicity was measured in 256-degree grayscale numerically from 0 to 255. The mean pixel density of the right kidney and liver were 70.3 and 67.3 respectively in the former patient with a right kidney-liver echogenicity ratio of 1.04, and 119.6 and 70.4 respectively in the latter patient with a ratio of 1.70.

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measured numerically according to the 256 degrees of grayscale labeled by the software, from 0 (black) to 255 (white), for each pixel within the ROI. The inverse ratio of the mean pixel densities of the kidney and adjacent organ (liver in case of the right kidney and spleen in case of the left kidney) was calculated. In cases of a left solitary kidney, the calculation was performed again between the left kidney and the liver. Along with echogenicity, longitudinal kidney length was measured and adjusted with mean length of age-matched normal children, as reported by Kim et al.[10] Echogenicity and kidney length were correlated with age-matched values of serum cystatin C-based GFR. Each sonographic parameter was measured by a single investigator (YSL) to avoid interobserver variability, as in the study by Moghazi et al.[5]

**Statistical Analysis**

Univariate analyses were performed using the Fisher’s exact test and Mann-Whitney U-test in each kidney. Spearman correlation analyses were performed to analyze the eGFR Z-scores for age and the right kidney-liver echogenicity ratio. In addition, to analyze the change in renal echogenicity with age, correlation analysis was also performed between the echogenicity ratio and age in the patients with normal GFR in each kidney group. SPSS software, version 18.0 (SPSS Inc., Chicago, IL) was used. P-values <0.05 were considered statistically significant.

**Results**

Fifty-four patients with right solitary functioning kidneys and 37 patients with left solitary functioning kidneys were enrolled in this study (Table 2). Ultrasonography was performed and serum cystatin C evaluated at a median age of 17.1 months. The following were the causes leading to the condition of solitary kidney: multicystic dysplastic kidneys in 82 (90.1%) patients, and renal agenesis in the other 9 (9.9%) patients. The mean pixel density of the right kidney was 68.6 (interquartile range [IQR]: 56.8–82.5), significantly different from that of the left kidney (52.9; IQR: 41.4–76.5; p = 0.005).

**Right kidney-liver echogenicity ratio**

Among the 54 patients with right solitary kidneys, GFR was decreased in eight right solitary kidney patients (14.8%) for their age (Table 3). Although the median kidney echogenicity was not different between the groups (p = 0.658), the right kidney-liver ratio was significantly elevated in the decreased GFR group (p = 0.029). The age-matched longitudinal kidney length ratio was not different between the groups (p = 0.422). Correlation analysis showed weak correlation between eGFR Z-scores for age and the right kidney-liver echogenicity ratio. The Spearman correlation coefficient was –0.227 (p = 0.098). A scatter diagram is shown in Fig 2.

To analyze the change in renal echogenicity with age, Spearman correlation analysis was performed between the right kidney-liver echogenicity ratio and age in 46 patients with normal GFR. It revealed a correlation coefficient of 0.361 with a p-value of 0.014 (Fig 3A).

**Left kidney-spleen echogenicity ratio**

GFR was decreased in four left solitary kidney patients (10.8%) for their age (Table 4). The left kidney-spleen ratio was not different in these four left solitary kidney patients with decreased GFR compared with the left solitary kidney patients with normal GFR (p = 0.493). The left kidney-liver ratio also showed no significant difference between groups (p = 0.114).

Correlation analysis was performed between the left kidney-spleen echogenicity ratio and age in 33 patients with normal GFR. The correlation coefficient was 0.015 with a p-value of 0.935 (Fig 3B).
Manley and O’Neill first introduced quantitative measurement of renal echogenicity in 2001. [4] They scanned ultrasonographic images and measured the echogenicity of the right kidney by adjustment with the liver using graphic programs. Moghazi et al. used a similar method and correlated the ultrasonographic findings with renal histological parameters. [5] They revealed

Table 2. Characteristics of 91 pediatric patients with solitary kidney.

| Variable                                      | Number (%) | or median (interquartile range) |
|-----------------------------------------------|------------|---------------------------------|
| Gender (male:female)                          | 41:50      |                                 |
| Laterality (right:left)                       | 54:37      |                                 |
| Cause of non-functioning kidney               |            |                                 |
| MCDK                                          | 82 (90.1)  |                                 |
| Renal agenesis                                | 9 (9.9)    |                                 |
| Mode of presentation                          |            |                                 |
| Prenatal diagnosis                            | 88 (96.7)  |                                 |
| Urinary tract infection                       | 2 (2.2)    |                                 |
| Palpable mass                                 | 1 (1.1)    |                                 |
| Median age at evaluation (months)             | 17.1 (8.1–27.5) |                             |
| Existence of proteinuria on urinalysis        | 0 (0.0)    |                                 |
| Serum cystatin C (mg/L)                       | 1.01 (0.90–1.16) |                           |
| Median glomerular filtration rate (mL/min/1.73m²) | 115.3 (91.3–139.9) |                      |
| Decreased glomerular filtration rate for age  | 12 (13.2)  |                                 |
| Median kidney echogenicity                    |            |                                 |
| Right kidney<sup>a</sup>                      | 68.6 (56.8–82.5) |                        |
| Left kidney<sup>b</sup>                       | 52.9 (41.4–76.5) |                        |
| Total                                         | 65.0 (51.5–79.2) |                       |
| Median liver echogenicity<sup>a</sup>         | 63.5 (51.9–77.6) |                        |
| Median spleen echogenicity<sup>b</sup>        | 62.3 (47.4–79.5) |                        |
| Median kidney size                            | 6.8 (6.3–7.9) |                          |
| MCDK: multicystic dysplastic kidney.          |            |                                 |
| <sup>a</sup> 54 patients with right solitary kidney. |
| <sup>b</sup> 37 patients with left solitary kidney. |

### Discussion

Manley and O’Neill first introduced quantitative measurement of renal echogenicity in 2001. [4] They scanned ultrasonographic images and measured the echogenicity of the right kidney by adjustment with the liver using graphic programs. Moghazi et al. used a similar method and correlated the ultrasonographic findings with renal histological parameters. [5] They revealed

Table 3. Comparison of clinical parameters between the non-decreased and decreased glomerular filtration rate groups among 54 patients with right solitary kidney.

| Variables                                      | Non-decreased (n = 46) | Decreased (n = 8) | p-value |
|------------------------------------------------|------------------------|-------------------|---------|
| Gender (male:female)                           | 18:28                  | 4:4               | 0.702   |
| Age at evaluation (months)                     | 16.6 (IQR: 7.5–26.7)   | 10.5 (IQR: 7.1–23.4) | 0.450   |
| Median glomerular filtration rate (mL/min/1.73 m²) | 119.2 (IQR: 101.4–140.6) | 63.2 (IQR: 53.6–78.8) | <0.001  |
| Median kidney echogenicity                     | 68.4 (IQR: 58.0–82.4)  | 59.5 (IQR: 49.3–109.4) | 0.658   |
| Median liver echogenicity                      | 65.4 (IQR: 53.2–77.6)  | 49.9 (IQR: 31.6–80.0) | 0.263   |
| Median kidney-liver echogenicity ratio         | 1.09 (IQR: 0.93–1.26)  | 1.50 (IQR: 1.02–2.07) | 0.029   |
| Median kidney size                             | 6.9 (IQR: 6.1–7.9)     | 6.3 (IQR: 5.7–7.2)  | 0.311   |
| Median kidney size-age matched ratio           | 1.06 (IQR: 0.93–1.18)  | 0.99 (IQR: 0.89–1.13) | 0.422   |

IQR: interquartile range.

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that, compared to renal size, cortical thickness, or parenchymal thickness, renal echogenicity showed the strongest correlation with renal histological parameters, such as glomerular sclerosis, tubular atrophy, interstitial fibrosis, and interstitial inflammation. Recently, Hershkovitz et al. reported changes in fetal renal echogenicity during the fetal period using similar method. [9] They gathered pictures of the longitudinal kidney from ultrasonography and converted them into 256 degrees of grayscale using a common graphic program. Then they also normalized the difference in gain using liver echogenicity. For this adjustment with the liver, Manley and O’Neill, and Moghazi et al. could analyze right kidneys only. In this study, we used a similar method in pediatric patients with a solitary functioning kidney. In our study, decreased GFR was related to an increased right kidney-liver echogenicity ratio. However, correlation analysis between the eGFR Z-scores and renal echogenicity revealed only a weak correlation. Therefore, it seems that decreased echogenicity does not imply increased renal function, although increased echogenicity is closely correlated with decreased renal function. Previous reports include patients with bilateral kidneys. However, it was not possible to obtain the relationship between echogenicity and GFR for each kidney, as neither the information on the differential renal function of each kidney nor the total GFR was assessed. To overcome this

Fig 2. Correlation analysis showed weak correlation between eGFR Z-scores for age and the right kidney-liver echogenicity ratio. The Spearman correlation coefficient was \(-0.227\) (\(p = 0.098\)).

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problem, we performed this study including only solitary kidney patients. To our knowledge, this is the first report to objectively prove the relationship between increased echogenicity and decreased renal function.

There have been a few reports of the subjective measurement of renal echogenicity in pediatric patients. Spira et al. reported that 83.3% of chronic kidney disease patients with a solitary functioning kidney showed increased echogenicity. [3] Chi et al. revealed increased echogenicity to be a poor prognostic factor in pediatric hydronephrosis patients. [2] In addition, it does not always reflect irreversible renal injury: Wiersma et al. reported that in children with acute illness, renal echogenicity increases for weeks, even without renal disease. [11] Peerboccus et al. reported that a change in echogenicity can be observed, even over the course of a single day, depending on the hydration status. [12] Nevertheless, there has been no report of an objective

Fig 3. To analyze the change in echogenicity with age, Spearman correlation analysis was performed in the normal glomerular filtration rate group for each kidney. (A) Among 46 patients with right solitary kidneys and a normal glomerular filtration rate, the age and right kidney-liver echogenicity ratio revealed a correlation coefficient of 0.361 with a p-value of 0.014. (B) In the analysis of the correlation between age and left kidney-spleen echogenicity ratio in 33 patients with left solitary kidneys and a normal glomerular filtration rate, the correlation coefficient was 0.015 with a p-value of 0.935.

Table 4. Comparison of clinical parameters between patients with normal and decreased glomerular filtration rates among 37 patients with left solitary kidney.

| Variables                                    | Non-decreased (n = 33) | Decreased (n = 4) | p-value |
|----------------------------------------------|------------------------|-------------------|---------|
| Gender (male:female)                         | 18:15                  | 1:3               | 0.340   |
| Age at evaluation (months)                   | 19.8 (IQR: 8.8–33.2)   | 7.0 (IQR: 4.4–29.3) | 0.140   |
| Median glomerular filtration rate (mL/min/1.73m²) | 117.2 (IQR: 97.0–151.0) | 72.2 (IQR: 69.3–77.4) | <0.001 |
| Median kidney echogenicity                   | 52.2 (IQR: 41.4–69.2)   | 83.8 (IQR: 48.9–108.3) | 0.092   |
| Median spleen echogenicity                   | 61.4 (IQR: 46.2–77.8)   | 77.6 (IQR: 56.7–94.7) | 0.186   |
| Median liver echogenicity                    | 59.5 (IQR: 44.3–74.0)   | 56.0 (IQR: 41.2–73.5) | 0.906   |
| Median kidney-spleen echogenicity ratio      | 0.88 (IQR: 0.76–1.06)   | 0.92 (IQR: 0.80–1.45) | 0.493   |
| Median kidney-liver echogenicity ratio       | 0.90 (IQR: 0.75–1.12)   | 1.35 (IQR: 0.98–1.98) | 0.114   |
| Median kidney size                           | 6.9 (IQR: 6.6–7.9)      | 6.6 (IQR: 5.8–7.4)  | 0.285   |
| Median kidney size-age matched ratio         | 1.05 (IQR: 1.00–1.16)   | 1.04 (IQR: 0.91–1.13) | 0.620   |

IQR: interquartile range.
measurement of renal echogenicity in pediatric patients. Our study is the first report to apply objective measurements of renal echogenicity in pediatric patients.

Just as the echogenicity of the right kidney has been frequently compared with that of the liver, the echogenicity of the left kidney has been compared with that of the spleen, clinically.[13] Yet, the objectively measured echogenicity of the left kidney adjusted by that of the spleen has not been previously reported. In this study, we analyzed the echogenicity of the left kidney adjusted by that of the spleen. However, unlike our right kidney-liver analysis, we did not observe clinical significance. During the ultrasonographic evaluation, the right kidney is usually covered by the liver while the left kidney is partially covered by the spleen. This difference could lead to the difference in gain. Although the grayscale density of the spleen showed similar values as that of liver, those of the right kidney and left kidney were significantly different. This result appears to be due to differences in the echogenicity of the liver and spleen, causing a gain difference, rather than any real differences between the right kidney and left kidney. In addition, as none of the patients in the study had a disease of the liver or spleen, the difference in liver echogenicity between the non-decreased GFR group (65.4) and the decreased GFR group (49.9) in right solitary kidney patients appears to be caused by gain adjustment. Similarly, the difference in spleen echogenicity between the non-decreased GFR group (61.4) and the decreased GFR group (77.6) in the left solitary kidney patients also appears to be caused by gain adjustment. Although it also showed no statistical significance, the left kidney-liver echogenicity ratio was higher in the decreased GFR group. These results may indicate that the liver is more feasible than the spleen as a controlled organ. Using the liver as the reference organ for both kidneys might have produced a different result; however, this option was not available in our retrospective study. Further prospective studies that control for gain would yield useful information on the relationships among the liver, spleen, and kidney.

The echogenicity of the right kidney showed a statistically significant correlation with age. This contradicts a previous report: Han and Bobcock reported that the echogenicity of the right kidney was lower than that of the liver after 3 years of age in normal children.[13] They also revealed the echogenicity of the left kidney to be lower than that of the spleen after 6 months of age. Although there is a difference between the objective measurement in our study and the subjective measurement in Han and Bobcock’s study, there might be other reasons. First, there could be a difference in hydration status during ultrasonography. Echogenicity in pediatric patients is influenced largely by hydration.[12] After hydration, the echogenicity of the kidney increases due to the significant expansion of tubules that then provide two distinct acoustic surfaces.[4] Although we did not use intravenous hydration or diuretics for ultrasonography, patients were instructed to increase water intake before evaluation, which would appear to have greater validity in older children with better communication skills. In addition, the fact that only patients with solitary kidney were included in this study could be another reason, due to the possible effect of compensatory renal growth. In the case of the fetal kidney, the kidney has been known to show different echogenicities according to the trimester.[9] The echogenicity decreased in the second and third trimesters when the collecting system is stable and new nephrons emerge rapidly. In patients with a solitary kidney, compensatory renal growth is observed from the fetal period to 1–2 years of age.[14–16] During this period, echogenicity should decrease, as has been observed during the second and third trimester. This study, performed with solitary kidney patients under the age of 10 years, cannot be taken to represent the renal echogenicity of the general population. As the normal change in renal echogenicity during childhood constitutes important clinical information, further investigation performed with objective measurements in the normal control group should be mandatory.

Nevertheless, the study has limitations. Its retrospective design introduces the potential for selection bias. Although we used a similar method in previous studies, the gain was not
controlled due to the retrospective nature, and this could affect the renal echogenicity value among images even in the same study. However, Manley and O’Neill have previously reported a difference of less than 2.8% in renal echogenicity among pictures in the same study.[4] Moreover, the fact that even the images of uncontrolled gain showed clinical significance paradoxically reveals the high feasibility of this method in daily practice. In this study, for each picture, the ROI was outlined around the whole kidney rather than a specific region. Although, Manley and O’Neill first introduced the partial measurement of renal parenchyma,[4] Hershkovitz et al. measured the echogenicity of the whole kidney.[9] We chose whole kidney measurement as it is simple and easy, as well as free from selection bias. However, the medulla, renal pelvis, and sinus fat were included in the analysis. To minimize the effect of collecting system inclusion, we excluded patients with hydronephrosis. In addition, sinus fat is known to be less prominent in this age group. However, this could still affect the echogenicity value. Another limitation is that the hydration status was not controlled in all patients. This could lead to an overestimation of cystatin C-based GFR, and also affect the echogenicity of the kidney. Peer-boccus et al. reported a change in echogenicity after hydration.[12] In this study, however, analysis of the time interval between hydration and ultrasonography was not available. In addition, we used a single measurement of cystatin C to correlate it with the ultrasonography performed on the same day. Although the intra-individual variability of serum cystatin C has been reported to be less than that of serum creatinine in pediatric patients with decreased GFR,[17] serial follow up of both cystatin C and ultrasonography might yield better results.

Conclusions
This is the first report to objectively prove the relationship between echogenicity and renal function in patients with a right solitary kidney. The right kidney-liver echogenicity ratio, measured objectively, showed feasibility in clinical practice as it showed a close relationship with decreased renal function when increased. Nevertheless, the absolute values of kidney echogenicity alone, or the left kidney-spleen echogenicity ratio, were not independent markers for decreased renal function.

Acknowledgments
We appreciate the help of Dr. Osamu Uemura and his collaborators. In this study, we used cystatin C-based GFR Z-scores for age to analyze the correlation between renal function and echogenicity. The mean and standard deviation values were not shown in Uemura et al.’s original report. At our request, they additionally derived the mean and standard deviation values for this study.

Author Contributions
Conceived and designed the experiments: YSL SWH. Performed the experiments: YSL MJL. Analyzed the data: YSL MJL NLL. Contributed reagents/materials/analysis tools: YSL MJK YJI SWK. Wrote the paper: YSL MJL NLL SWH.

References
1. Lee YS, Jung HJ, Im YJ, Hong CH, Han SW. The significance of detrusor wall thickness as a prognostic factor in pediatric bladder outlet obstruction. Journal of pediatric surgery. 2012; 47(9):1682–7. Epub 2012/09/15. doi: 10.1016/j.jpedsurg.2012.03.051 PMID: 22974606.
2. Chi T, Feldstein VA, Nguyen HT. Increased echogenicity as a predictor of poor renal function in children with grade 3 to 4 hydronephrosis. The Journal of urology. 2006; 175(5):1898–901. Epub 2006/04/08. doi: 10.1016/j.juro.2006.02.028 PMID: 16600795.
3. Spira EM, Jacobi C, Frankenschmidt A, Pohl M, von Schnakenburg C. Sonographic long-term study: paediatric growth charts for single kidneys. Archives of disease in childhood. 2009; 94(9):693–8. Epub 2009/06/24. doi:10.1136/adc.2008.153601 PMID: 19546100.

4. Manley JA, O’Neill WC. How echogenic is echogenic? Quantitative acoustics of the renal cortex. American journal of kidney diseases: the official journal of the National Kidney Foundation. 2001; 37(4):706–11. Epub 2001/03/29. PMID: 11273869.

5. Moghazzi S, Jones E, Schroepple J, Arya K, McClellan W, Hennigar RA, et al. Correlation of renal histopathology with sonographic findings. Kidney international. 2005; 67(4):1515–20. Epub 2005/03/23. doi:10.1111/j.1523-1755.2005.00230.x PMID: 15780105.

6. Fernbach SK, Maizels M, Conway JJ. Ultrasound grading of hydronephrosis: introduction to the system used by the Society for Fetal Urology. Pediatric radiology. 1993; 23(6):478–80. Epub 1993/01/01. PMID: 8255658.

7. Grubb A, Nyman U, Bjork J, Lindstrom V, Rippe B, Sterner G, et al. Simple cystatin C-based prediction equations for glomerular filtration rate compared with the modification of diet in renal disease prediction equation for adults and the Schwartz and the Counahan-Barratt prediction equations for children. Clinical chemistry. 2005; 51(8):1420–31. Epub 2005/06/18. doi:10.1373/clinchem.2005.051557 PMID: 15961546.

8. Uemura O, Nagai T, Ishikura K, Ito S, Hataya H, Gotoh Y, et al. Reference glomerular filtration rate levels in Japanese children: using the creatinine and cystatin C based estimated glomerular filtration rate. Clinical and experimental nephrology. 2014. Epub 2014/10/20. doi:10.1007/s10157-014-1042-6 PMID: 25326724.

9. Hershkovitz R, Amichay K, Stein GY, Tepper R. The echogenicity of the normal fetal kidneys during different stages of pregnancy determined objectively. Archives of gynecology and obstetrics. 2011; 284(4):807–11. Epub 2010/11/11. doi: 10.1007/s00404-010-1738-0 PMID: 21063717.

10. Kim SS, Bang WJ, Seo JW, Cho KS, Han SW. Discrepancy of Measured Renal Length between Ultrasonography and Dimercaptosuccinic Acid (DMSA) Scintigraphy. 대한비뇨기과학회지. 2007; 48(1):77–81. doi: 10.4111/kju.2007.48.1.77

11. Wiersma F, Toorenvliet BR, Ruige M, Holscher HC. Increased echogenicity of renal cortex: a transient feature in acutely ill children. AJR American journal of roentgenology. 2008; 190(1):240–3. Epub 2007/12/21. doi: 10.2214/ajr.07.2606 PMID: 18094318.

12. Peerboccus M, Damry N, Pather S, Devriendt A, Avni F. The impact of hydration on renal measurements and on cortical echogenicity in children. Pediatric radiology. 2013; 43(12):1557–65. Epub 2013/08/06. doi: 10.1007/s00247-013-2748-4 PMID: 23913159.

13. Han BK, Babcock DS. Sonographic measurements and appearance of normal kidneys in children. AJR American journal of roentgenology. 1985; 145(3):611–6. Epub 1985/09/01. doi: 10.2214/ajr.145.3.611 PMID: 3895872.

14. Peters CA, Gaertner RC, Carr MC, Mandell J. Fetal compensatory renal growth due to unilateral ureteral obstruction. The Journal of urology. 1993; 150(2 Pt 2):597–600. Epub 1993/08/01. PMID: 7686988.

15. Laufer I, Griscom NT. Compensatory renal hypertrophy. Absence in utero and development in early life. The American journal of roentgenology, radium therapy, and nuclear medicine. 1971; 113(3):464–7. Epub 1971/11/01. PMID: 5127707.

16. van Vuuren SH, van der Doef R, Cohen-Overbeek TE, Goldschmeding R, Pistorius LR, de Jong TP. Compensatory enlargement of a solitary functioning kidney during fetal development. Ultrasound in obstetrics & gynecology: the official journal of the International Society of Ultrasound in Obstetrics and Gynecology. 2012; 40(6):665–8. Epub 2012/05/15. doi: 10.1002/uog.11168 PMID: 22581658.

17. Sambasivan AS, Lepage N, Filler G. Cystatin C intrapatient variability in children with chronic kidney disease is less than serum creatinine. Clinical chemistry. 2005; 51(11):2215–6. Epub 2005/10/26. doi:10.1373/clinchem.2005.056150 PMID: 16244312.