Pregnancy -check point for diagnosis of CKD in developing countries

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Introduction

Pregnancy is challenging among females with kidnev disease (CKD), and requires multidisciplinary approach comprising of obstetrician, nephrologists and neonatologist. Chronic kidney disease is often clinically and biochemically silent until renal impairment is advanced. Symptoms are unusual until the glomerular filtration rate declines to <25% of normal. Some women are found to have chronic kidney disease for the first time during pregnancy. Around 20% of women who develop early pre-eclampsia (≤30 gestation); especially those with heavy weeks' proteinuria, have previously unrecognized chronic kidney disease. (7,8,9) These diseased kidneys may be unable to adapt to the normal physiologic changes of pregnancy leading to poor maternal and fetal outcome. (4) Evidences suggest that females with mild renal impairment with normal BP and little or no proteinuria have good maternal and fetal outcomes, with little risk for accelerated progression toward ESRD or preterm delivery. However those pregnancies among CKD population have shown poor maternal and fetal outcomes. (1,2) Jones and Haslett (5) observed that CKD was associated with increased risk for adverse pregnancy outcomes especially among pregnancies with moderate or severe renal function impairment. More than one-third experience an irreversible decline in GFR and 10 percent progress to end-stage renal disease (ESRD) by 12 months postpartum. (4,6) These findings highlight the important need for early identification of CKD, pre-conception counseling, preconception review of medications and early referral of pregnant patients with CKD for specialist care.

Studies have suggested that the risk of deterioration in maternal renal function is increased mainly when conception has occurred at a plasma keratinize concentration in excess of 200 mol/L (equivalent to eGFR <25-30 mL/min/1.75 m²) or in the setting of poorly controlled hypertension. Whether such deterioration represents an acceleration of renal failure progression or merely the natural history of the underlying chronic kidney disease which has not been conclusively established needs to be evaluated. Women especially in developing countries like India conceive without preconception evaluation and have no regular follow up under an obstrtetician subjecting to increased risk to worsening of underlying basic disease with poor pregnancy and maternal outcome. The present study

looked into those aspects and emphasizing the need for evaluate all females before or during pregnancy for the risks to poor renal outcomes.

Aims and Objective

The present study looked into those females presenting with Chronic kidney disease (various stages) and assessing the impact of CKD detection at the time of pregnancy and its impact on fetal and maternal outcome, highlighting the need to address the influence of pregnancy on the underlying kidney disease and the kidney disease affecting the outcome of pregnancy.

Material and Methods

This retrospective observational study of female patients with CKD who were assessed about their pregnancy outcomes, their awareness of illness at the time of pregnancy and its outcome on renal as well as pregnancy outcomes conducted at tertiary care health care centre from northern part of India from January 2011 to January 2016.

CKD was defined as per standard definition of CKD by kidney disease outcomes quality initiative guideline. (10)

Hypertension was defined if there were 2 blood pressure measurements >140/90 mm Hg or there was history of antihypertensive drug use. If the hypertension was observed before 20 weeks of gestation with proteinuria or renal failure, hypertension was considered to be renal in origin.

Pre-eclampsia was defined as abrupt onset of hypertension after 20 weeks of gestation associated with the appearance of proteinuria with protein excretion >300 mg/day. Among females who had hypertension or proteinuria at the start of pregnancy if there was doubling of protein excretion associated with worsening of hypertension after 20 weeks of gestation was considered as preeclampsia. (11,12,13,14)

All female patients in child bearing age group (18-45 years) with first time diagnosis of "Chronic renal failure" or "Chronic kidney injury" either before or during pregnancy were identified and degree of awareness about the disease and its implication on maternal and fetal outcomes and management of pregnancy and future outcome were looked for.

Cases: Those pregnancies were the patient was not aware about the renal disease at the time of pregnancy

with no meticulous follow up during pregnancy by the obstetrician or nephrologists were considered cases.

Control: Those pregnancies were the obstetrician had explained about the illness and precautions and meticulous measures taken for successful pregnancy outcomes were identified as control.

Among females who were diagnosed with CKD at the time of conception and under regular follow up were assessed about progression of CKD using GFR assessment. Glomerular filtration rate (GFR) was estimated using the CKD epidemiology collaboration equation and expressed in ml/min/1.73 m² of body surface area. (15) Rate of progression was assessed from the first clinical or laboratory evidence of renal dysfunction assessed by GFR and decline during and after pregnancy in ml/min.

Patient data were retrieved from the computerized hospital based electronic information system and patient record files. All records of the patient's history, physical examination, laboratory investigations and discharge reports, their clinical presentation, etiological diagnosis and outcomes were evaluated and examined by two nephrologists. A clinical and histopathological diagnosis was made after scrutinizing all the details.

Their detailed history regarding symptomatology related to renal disease, pregnancy evaluation regarding renal disease and detection of renal disease from time from pregnancy was assessed and any association of worsening of progression was assessed following pregnancy.

Outcome criteria: The primary maternal outcome was decline in renal function after pregnancy at 6 weeks after delivery at completion of puerperium when the physiological changes related to pregnancy is settled. The other outcome criteria were doubling of the serum creatinine and 50% decline in GFR or ESRD at 1 year of follow-up. The fetal outcome was analyzed in terms of full-term and preterm delivery, stillbirth, and low birth weight. Among those females who were unaware of their illness at time of pregnancy was assessed for decline in renal function from the baseline at time of pregnancy to time the person becomes aware i.e., doubling of the serum creatinine and or 50% decline in GFR or ESRD.

The primary maternal outcome were assessed in terms of decline in renal function from the time of detection i.e. before or during pregnancy and the clinicians and females awareness of the illness and its implication on maternal management with its association to maternal outcome ie renal as well as overall outcome.

Fetal outcome was analyzed in terms of pre-term and full-term delivery, stillbirth and low birth weight. Stillbirth was defined as the occurrence of intrauterine fetal death after 24 weeks of gestation. Preterm delivery was defined as a live birth before the $37^{\rm th}$ week of gestation. Pregnancy was considered successful if it resulted in a live infant who was discharged from the hospital. Low birth weight is defined as a live-born infant weighing < 2,500 g.

Statistical analysis: All the data were analyzed using statistical package for the social sciences (SPSS) 16 statistical software (SPSS Inc., Chicago, Illinois, USA). All data are expressed as mean \pm standard deviation. Student's *t*-test was used to compare the mean values between two groups. One-way Analysis of Variance and Bonferroni test was used to compare the differences in mean values of >3 variables. The Chi-square test was used to compare the categorical variables between the groups. P < 0.05 was considered significant.

Results

A total of 465 female's patients presented in nephrology OPD from Jan 2014 -Jan 2016 with chronic renal failure (CKD) and retrospectively looked into their pregnancy outcomes and renal disease.

Demographic Profile (Table 1): Among 465 females, 85 pregnant ladies were not in follow up of any obstetrician and had home delivery while among the remaining 380 females, 172 of them despite being under obstetrician follow up had no antenatal or post natal evaluation about their renal illness i.e. serum creatinine, urine examination or ultrasound KUB region available.208 females who were aware about their renal disease during pregnancy were under regular obstetrical and nephrological follow up.

Detailed evaluation retrospectively identified 382 females giving history of edema, headache, recurrent urinary tract infections, and nocturia in them.

Pregnancy outcomes (Table 1): Among 380 patients, 172 females during pregnancy had awareness about their renal illness and were in regular follow up with the nephrologists/obstetrician during pregnancy. patients had in hospital delivery. According to GFR stage 1, 2 CKD was observed among 96 pregnancies while 47 were in stage 3 and 29 were in stage 4. GFR < 60ml/min which was similar to the unaware group i.e.... 44.1%. Nephritic range proteinuria was observed in 58 pregnant ladies while 125 females had hypertension during 1 or 2 nd trimester of pregnancy. Despite regular and optimized follow up, worsening edema was observed in 37%, urinary tract infections requiring admission was observed in 6% pregnancies. 34% of these ladies required admission during pregnancy for various reasons.

Table 1: Clinical and demographic profile of CKD patients during pregnancy

Parameters	Awareness at the time of Pregnancy	Unaware about CKD at time of
	(n=172)	pregnancy (n= 208)
Average age at time of	26.7+/- 5 years	27.4 +/- 4.7 years
Pregnancy		
Average number of pregnancy	3+/-1	4/+/-1
GRF at presentation		
Stage 1-2(>60ml/min)	96	117
Stage 3 (30-50ml/min)	47	57
Stage 4(15-30 ml/min)	29	34
Stage 3+4 (< 60ml/min)	76	91
Proteinuria		
Subnephrotic	114	138
Nephrotic	58	70
Urinary tract infection	64	68
Hypertension	125	146
Preclampsia	30	78
Basic Disease		
Chronic Glomerulonephtitis	58	72
Chronic Interstitial Nephritis	72	79
ADPKD	7	5
Stone Disease	5	8
Others	30	44

Among the 208 females who were the unaware group, were not in regular follow up under a proper obstetrician and or nephrologists as they were unaware about their renal illness. This population was not judiciously managed during pregnancy due to lack of awareness by the patient and or the obstetrician about the underlying renal disease.

On retrospectively observing their baseline values, 37.5% had presented with preeclampsia and 2 patients had ecalmpsia at 32 weeks of gestation. 46 patients required early admission before termination of pregnancy for worsening hypertension, worsening edema, recurrent infections. i.e. 12% had recurrent urinary tract infection with early fetal loss in 4 of them. In the unaware group, 142 underwent normal delivery, while early termination was required in 46.7% of them. Ante-partum /post partum bleeding was observed in 12.6% while infection was encountered in 15.2%.

In the planned pregnancy group, precalmpsia was observed in 17.6% of planned pregnancies, more evident among females with GRF < 60ml/min i.e. 12.6% as compared to 29.5% respectively among the 2 groups. 78 females underwent caesarian section while 132 had normal delivery. Those pregnancies which were in regular follow up of the obstetrician were the ones who had caesarian section while the rest had presented with worsening hypertension, breathlessness and worsening edema requiring early termination in about 24% of them. Ante - partum /post- partum bleeding was observed in about 4.5% of pregnancy and. Renal outcomes (Table 2): Decline in GFR was assessed from time of first detection of renal failure to time after or any time when patient became aware about her renal illness.

Table 2: Renal outcomes during or and following delivery or at time patient becomes aware of renal disease

Parameters	Awareness at the time of Pregnancy (n=172)	Unaware about CKD at time of pregnancy (n= 208)
Average fall in GFR at		
follow up after delivery		
or at the time the patient		
is aware		
Stage 1-2	86	71
Stage 3	48	63
Stage 4	26	41
Stage 3+4	84	101
Nephrotic range	69	96

proteinuria at follow up		
ESRD	12	31
Worsening of GFR	51	87
Average time to dialysis	5.1+/-2.5 years	2.3+/1.2 years
from pregnancy		
Worsening of	29	65
Hypertension		
Delivery	131	111
Full Term	41	97
Pre term		
Foetal outcome		
Alive	147	90
Still birth	74	118
Low birth	45	98

Those pregnancies which were planned and monitored judiciously had better outcome. Worsening of renal failure during and following pregnancy was observed among all stages of CKD but was more evident when gofer <60ml/min. This decline in GFR was much less as compared to those pregnancies in unaware group where meticulous follow up was not done. There was greater decline in GRF among all stages with progression to ESRD earlier as compared to aware group. Those females who were having presumably glomerular illness faired poorly than ones with interstitial illness among both the groups.

Fetal outcome (**Table 1**): Among the planned pregnancy group, had better fetal outcome with 147 had live births as compared to 74 still births among pregnancies which were not regularly monitored. 24% had preterm delivery among planned pregnancy group while 26% had babies <2500 gm. However in the unaware group, 56.7% had still births while preterm delivery was observed in 46.7% pregnancies. 96 pregnancies had low birth babies weighing <2500gm.

Full term pregnancy was higher among ladies with > 60ml/min GFR in both the groups.

Discussion

The prevalence of CKD in pregnancy is reported rarely especially in developing countries. This discrepancy may be due to several reasons besides underreporting. The prominent ones could be 1. Pregnancy happens in young population and are expected to be healthy. 2. Pregnant females are not routinely screened for renal dysfunction. 3. Females with substantial renal insufficiency or renal failure are either beyond child-bearing age or infertile. Hence it is essential to screen all females during pregnancy for CKD to prevent progression of renal disease besides poor maternal and fetal outcome.

The kidney undergoes physiological changes during pregnancy However outcomes. In such circumstances, multidisciplinary process is intimidated which can help in careful management of maternal and fetal outcome.

Every pregnancy should be thoroughly investigated and evaluated and those females with CKD, prepregnancy counseling is recommended. Hypertension, proteinuria and teratogenic medications eg ACE inhibitors usage is prevalent among these females and needs to be optimized before conception. There is increase risk of maternal and fetal outcomes among pregnant CKD patients and needs to be communicated and understood well by them before conception. (20) In our study population 18.2% had home delivery with no ante- natal checks even. Despite being under obstetrician follow up, 44.7% were not aware of the renal disease at the time of pregnancy, suggesting that pre-pregnancy evaluation as well as counseling is indeed essential for better fetal and maternal outcome. Pregnancies which were planned had better outcome in terms of fetal and maternal outcomes. Worsening of GFR was observed in 29.6% of pregnant females while it was 41.85% among females who were unaware. Preeclampsia was observed in 17.2% among females who were aware as compared to unaware females which was 37.5%. Higher preterm a well as low birth babies were born to females unaware about their renal illness. Time to ESRD was faster among unaware females as compared to those ladies who were aware about their illness.

Clinical marker of renal disease during Pregnancy:

Hypertension during pregnancy has greater impact on GFR deterioration, more so among females with deckling GFR during pregnancy. (21) This observation was evident in our study where females unaware about their renal disease had worsening of hypertension in 31.8% as compared to aware females i.e. 17.05% with worsening of GFR following delivery.

34.1% had nephritic range proteinuria at the baseline among the aware group while it was 34.3% among the unaware group, how ever there was worsening of proteinuria during and following pregnancy in both groups but more marked in unaware group to 47.15% with higher number developing preeclampsia and worsening renal function. Literature

suggests asymptomatic pregnant women with proteinuria >500 mg/day, not previously known renal disease and no evidence of preeclampsia, 20% of them progressed to ESRD at a median time of 5 year. When proteinuria exceeds 1 g/day, there is a greater tendency for accelerated GFR-decline and nearly a 2-fold higher incidence of ESRD.

Renal failure at the time of presentation has great impact on future maternal and fetal outcome as observed in our study females with advanced renal disease fairing poorly in terms of fetal and maternal outcome. Our study observed that among planned pregnancy group, 24% had preterm delivery while 26% babies born were weighing <2500 gm. However in the unaware group, 56.7% had still birth, preterm delivery was observed in 46.7% pregnancies. 96 pregnancies had low birth babies weighing <2500gm. Among systemic review of pregnancy outcomes in CKD it was observed that the incidence of premature birth in women with CKD was consistently higher along with the risk of IUGR observed to be five times higher. The risk of neonatal mortality was fivefold higher and a nine fold risk for still births. Studies observed a fivefold increase in risk for low birth weight. (25)

Worsening of renal failure during and following pregnancy was observed among all stages of CKD but was more evident when gfr <60 ml/min. This decline in GFR was much less as compared to the unaware population with no proper management for CKD. In the aware group, worsening of GFR was observed among 30% patients while it was 39.7% in the unaware group. 6.9% became ESRD after pregnancy in aware group while 16.8% became so in the unaware group focusing the need for careful evaluation and management of CKD among pregnant females. Average time to dialysis from the last pregnancy was faster among the unaware population. Those females who were presumably glomerular illness faired poorly than ones with interstitial illness. This is evident from literature wherein approximately 25-38% of pregnant women with moderate CKD show decline in renal function evident by an increase in S creatinine during pregnancy(23,24) which may persist in one third of the women for 6 months postpartum and in 10% of the total cohort reach ESRD. Those women with a more severe renal functional impairment (GFR<40 mL/min/1.73m²) and proteinuria exceeding 1 g/day had poorer outcomes, the combination resulting in worse outcomes than either factor alone. (22) This was evident in our study population, 48.8% had progressed to stage 3+4 in the aware group and 48.9% in the unaware population with a larger percentage progressing to ESRD in the unaware population. There was decline in GFR even among those in stage 1or 2 yet it was more evident in the unaware group suggesting that judicious monitoring of pregnancy is essential among these females. Complications are even higher in women with more severe kidney disease at conception Our observation

was supported by literature evidences that 82% of women with severe CKD had chronic hypertension and 64% developed preeclampsia⁽²⁴⁾ with significant detoriation in maternal kidney function in over 25% of women. The risk of accelerated progression to ESRD was highest when serum creatinine was found to be greater than 1.9 mg/dl at the beginning of the pregnancy. In our study population preeclampsia was an important complication of pregnancy evident in both study population though higher in unaware group i.e. 37.5% pregnancies.

Hence it is essential to screen all ladies before pregnancy for proteinuria as well as renal dysfunction to avoid maternal and fetal complication. In our study population those pregnancies where antenatal checks up were available and were intensively monitored by the obstetrician with multidisciplinary support of a nephrologists and neonatologist faired better then unmonitored pregnancies. Higher preterm a well as low birth babies were born to females unaware about their renal illness as compared to better monitored female.

Conclusions

Chronic kidney disease is often clinically and biochemically silent until renal impairment is advanced. Some women are found to have chronic kidney disease for the first time during pregnancy. Around 20% of women who develop early pre-eclampsia (≤30 weeks' gestation); especially those with heavy proteinuria, have previously unrecognized chronic kidney disease. Hence pregnancy can be a check point in females to pick up for any evidence of renal dysfunction and appropriately managed for better maternal and fetal outcome.

References

- Williams D and Davison J. Chronic kidney disease in pregnancy. BMJ 2008;336:211-215.
- Hou S, Historical perspective of pregnancy in chronic kidney disease. Adv Chronic Kidney Dis.2007;14:116-118.
- Munkhaugen J, Lydersen S, Romundstad PR et al. Kidney function and future risk for adverse pregnancy outcomes: a population-based study from HUNT II, Norway. Nephrology Dialysis Transplantation. 2009;24:3744-50.
- Katz AI, Lindheimer MD. Does pregnancy aggravate primary glomerular disease? Am J Kidney Dis 1985;6:261-5.
- Jones DC, Hayslett JP. Outcome of pregnancy in women with moderate or severe renal insufficiency. N Engl J Med 1996;335:226-32.
- Nevis IF, Reitsma A, Dominic A, McDonald S, Thabane L, Akl EA, et al. Pregnancy outcomes in women with chronic kidney disease: A systematic review. Clin J Am Soc Nephrol 2011;6:2587-98.
- Jungers P, Chauveau D, Choukroun G, Moynot A, Skhiri H, Houillier P, et al. Pregnancy in women with impaired renal function. Clin Nephrol 1997;47:281-8. [PubMed]
- Davison J, Baylis C. Renal disease. In: De Swiet M, ed. Medical disorders in obstetric practice. 3rd ed. Oxford: Blackwell Science, 1995:226-305.

- Williams D. Renal disorders. In: James DK, Steer PJ, Weiner CP, Gonik B, eds. High risk pregnancy. Management options. 3rd ed. Philadelphia: Elsevier Saunders, 2006:1098-124.
- Levey AS, Coresh J, Balk E, Kausz AT, Levin A, Steffes MW, et al. National Kidney Foundation practice guidelines for chronic kidney disease: Evaluation, classification, and stratification. Ann Intern Med 2003;139:137-47.
- Sibai BM, Stella CL. Diagnosis and management of atypical preeclampsia-eclampsia. Am J Obstet Gynecol. 2009;200:481e1–7.
- 12. Report of the national high blood pressure education program working group on high blood pressure in pregnancy. Am J Obstet Gynecol. 2000;183:S1–22.
- 13. Harrington K, Fayyad A, Thakur V, Aquilina J. The value of uterine artery Doppler in the prediction of uteroplacental complications in multiparous women. Ultrasound Obstet Gynecol. 2004;23:50–5.
- Espinoza J, Romero R, Nien JK, Gomez R, Kusanovic JP, Gonçalves LF, et al. Identification of patients at risk for early onset and/or severe preeclampsia with the use of uterine artery Doppler velocimetry and placental growth factor. Am J Obstet Gynecol. 2007;196:326e1–13.
- Levey AS, Stevens LA, Schmid CH, Zhang YL, Castro AF, 3rd, Feldman HI, et al. A new equation to estimate glomerular filtration rate. Ann Intern Med. 2009;150:604–12.
- Davison J, Baylis C. Renal disease. In: De Swiet M, ed. Medical disorders in obstetric practice. 3rd ed. Oxford: Blackwell Science, 1995:226-305.
- Williams D. Renal disorders. In: James DK, Steer PJ, Weiner CP, Gonik B, eds. High risk pregnancy.

- Management options. 3rd ed. Philadelphia: Elsevier Saunders, 2006:1098-124.
- Cunningham FG, Cox SM, Harstad TW, Mason RA, Pritchard JA. Chronic renal disease and pregnancy outcome. Am J Obstet Gynecol 1990;163:453-9. [PubMed]
- Jungers P, Houillier P, Chauveau D, Choukroun G, Moynot A, Skhiri H, et al. Pregnancy in women with reflux nephropathy. Kidney Int 1996;50:593-9.
- Kate S Wiles, Kate Bramham, Alina Vais, Kate R Harding, Paramit Chowdhury, Cath J Taylor, Catherine Nelson-Piercy. Pre-pregnancy counselling for women with chronic kidney disease: a retrospective analysis of nine years' experience BMC Nephrology (2015) 16:28.
- Piccoli GB, Conijn A, Attini R, Biolcati M, Bossotti C, Consiglio V, et al. Pregnancy in chronic kidney disease: need for a common language. J Nephrol. 2011;24:282– 299
- Imbasciati E, Gregorini G, Cabiddu G, Gammaro L, Ambrosso G, Del Giudice A, et al. Pregnancy in CKD stages 3 to 5: fetal and maternal outcomes. Am J Kidney Dis. 2007;49:753–762.
- Jones DC, Hayslett JP, et al. Outcome of pregnancy in women with moderate or severe renal insufficiency. N Engl J Med. 1996;335:226–232.
- Cunningham FG, Cox SM, Harstad TW, Mason RA, Pritchard JA. Chronic renal disease and pregnancy outcome. Am J Obstet Gynecol. 1990;163:453–459.
- Immaculate Nevis et al. Pregnancy Outcomes in Women with Chronic Kidney Disease: A Systematic Review. CJASN. 2011; 6:2587-2598.