ORIGINAL ARTICLE

Histopathological Pattern of Primary Glomerulonephritis -A Single Center Experience

Shaheen Sikder¹, Faijul Islam², Mahmudur Rahman³, Ehteshamul Hoque⁴

Abstract:

Glomerulonephritis is the most common cause of chronic kidney disease in our country like in other developing countries. The incidence and histological pattern of primary GN in our countare is inadequately described. This cross-sectional prospective study was conducted at the nephrology unit in Holy Family Red Crescent Medical College Hospital, a tertiary care teaching hospital in Dhaka, Bangladesh starting from January 2019 - December 2019 including all patients with suspected primary GN who underwent kidney biopsies. Total 35 biopsies were performed. M: F 1.3:1 .Mean age was 33.71+_12.2 yrs. Membranoproliferative GN (34.28%) was the most common cause followed by mesangioproliferative (22.85%), IgA nephropathy (17.14%), membranous (11.42%), FSGS (11.42%) and minimal change disease (2.8%). Among 35 cases, 20 had proliferative verity. In immunofluroscence study, 19 had mild to marked deposits of immunoglubulin. 5.71% had post-biopsy complication which was not significant enough.

As the sample size was small so needs more studies in large to get the specific epidemiological patterns of primary GN.

- 1. Assistant professor, department of Nephrology, Holy Family Red Crescent Medical College Hospital, Dhaka
- 2. Registrar, department of Nephrology, Holy Family Red Crescent Medical College Hospital, Dhaka
- 3. RMO, department of Nephrology, Holy Family Red Crescent Medical College Hospital, Dhaka
- 4. Professor and Head, department of Nephrology, Holy Family Red Crescent Medical College Hospital, Dhaka

Introduction:

Glomerulonephritis is a common cause of chronic kidney disease worldwide. It may be primary or secondary in etiology. Thus the pattern of glomerulonephritis varies from country to country, reflecting the effects of genetic, socioeconomic. and environmental factors¹. The disease spectrum is also been changing over the last few decades. IgA nephropathy is the commonest primary glomeruloneph in Asia. as well as in white Europeans and Americans²⁻⁵. In contrast, focal segmental glomerulosclerosis (FSGS) is the commonest glomerular disease among African - Americans, South Americans and in the middle east^{6,7}. Currently, we do not have a central biopsy registry in Bangladesh. Statistics of prevalence of glomerular disease in Bangladesh are limited.

In light of the paucity of published data from our country, this study was done to describe the histopathological pattern of primary glomerular disease at Holy Family Red Crescent Medical College Hospital.

Materials and method:

This was a hospital based cross-sectional observational study. Study period 01(one) year (January 2019- december 2019). Total 35 suspected cases of primary glomerulonephritis were included in this study and native kidney biopsy was done under local anesthesia at department of nephrology in HFRCMCH.

We recorded the demographic of cases, indication of renal biopsy, histopathological diagnosis, relevant laboratory investigations and post biopsy

complications.

Suspected case of primary glomerulonephritis with proteinuria > 1gm/day and/or persistent haematuria after excluding the possible causes of secondary glomerulonephritis.

Kidney biopsy was performed for all selected patients using 16-G automated biopsy needle. At least two cores of tissue were taken from each patient for light microscopy and direct immunofluroscence study.

Samples were fixed in 10% formalin solution and stained with hematoxylin and eosin and periodic acid schiff (PAS) for light microscopy. The other sample was preserved in normal saline for direct immunoflurescence (DIF)study.Immunoglobulins-IgG, IgM, IgA, also complement C1 and C3 were stained. Electron microscopy was not available for diagnostic purpose in our country.

Histopathological primary types of glumerulonephritis(PGN) were classified as follows-membranoproliferative glomerulonephritis (MPGN), messengio proliferative glomerulonephritis (MesPGN), Immunoglobulin nephropathy А (IgA), Focal segmental glomerulosclerosis (FSGS), Membranous Nephropathy (MN)and Minimal change disease (MCD).

Cases were further classified into proliferative GN and non proliferative GN .

All data were noted into a specially designed questionnaire and were analyzed using statistical package for social sciences (SPSS) version 20 computer software. Results were expressed as median or mean with standard deviation for continuous data and as frequencies with percentage for categorical data.

Results:

Out of 35 cases twenty cases were male (57.14%) and fifteen cases were female (42.85%). Male: Female was 1.3:1. Mean age33.71±12.2 (13-60). The majority of study subject were in the age group of 21 - 30 years.

Membranoproliferative glomerulonephritis (MPGN) was the commonest form of primary GN followed by messangioproliferative glomerulonephritis (MesPGN, Fig 1). No specific pathological pattern were found in any age group. (Table I)

The cases were also classified as proliferaitye (20) and non proliferative GN (15) (Table II)

There was male predominance in all disease categories except in membranoproliferative glomerulonephritis where eight out of twelve case (66.66%) were female.

Post biopsy bleeding occurred in two cases (5.71%). No case required blood transfusion or nephrectomy.





| Diagnosis | 10 -20 | 21 - 30 | 31 - 40 | 41 - 50 | 51 - 60 |
|---------------|-----------|-----------|-------------|-----------|-----------|
| (no of cases) | yrs(n =5) | yrs(n=15) | yrs (n = 7) | yrs (n=4) | yrs (n=4) |
| MPGN(12) | 2 | 4 | 2 | 2 | 2 |
| MesPGN (8) | 1 | 3 | 2 | 1 | 1 |
| IgA (6) | 2 | 2 | 1 | 1 | 0 |
| FSGS (4) | 0 | 2 | 1 | 0 | 1 |
| MN (4) | 0 | 3 | 1 | 0 | 0 |
| MCD (1) | 0 | 1 | 0 | 0 | 0 |

Table I: pathological pattern of PGN in different age group (n-35)

Table II: proliferative and non proliferative classification of PGN

| Proliferative (n=20) | Non prolifentive (n=15) | | |
|----------------------|-------------------------|--|--|
| MPGN (12) | MN (4) | | |
| MesPGN (8) | IgA (6) | | |
| | FSGS(4) | | |
| | MCD(1) | | |

Discussion:

The majority of study subject who underwent renal biopsy were in the age group of 20-30 years but no distinct pathological pattern observed. in any age group. Mundi I et al⁸ found most of cases were in 21 - 40 yrs age range and distinct pattern of PGN was found in different age group. The age range of our cases was 16-60 years with a slight male predominance (1.3:1) except in cases of MPGN. Most of the studies shows male predominance 9,10,11 with exception of Habib M A¹² from Bangladesh.

In the current study MPGN is the commonest primary GN seen. In most studies from Bangladesh MesPGN was found to be the commonest primary ¹⁴ and proliferative GN¹³ though another study from Bangladesh found Focal segmental proliferative GN (29.47%) as the commonest entity.¹²

In our study MesPGNwas found second most common cause of primary GN . A study on global evolutionary trend of GN done in Singapore for three decades stated that in the first decade most Asians countries had mesPGN and MPGN are the common pattern of primary GN and still it is prevalent in some Asain countries like China,

25

Japan and Thailand.1 Apart from geographical, genetic and socioeconomic factor, one facto which may influence the pattern of glomerulonephritis in various countries could be the hygeine hypothesis ¹⁵ The hygeine hypothesis proposes that bacterial and other infections occuring in less developed and developing countries leads to development of some type of human glomerulonephritis, including MPGN and MesPGN. This would be true in Asian countries like China, 16 Indonesia 27, Malaysia 18, Thailand 19, and Singapore14 which have a high prevalence of MPGN and MesPGN. In some countries like Malaysia 18 and Singapure14 prevalence of MPGN and MesPGN is already decreasing in keeping the urbanization and better housing and other amenities in these countries. Bangladesh is a rising country in context of urbanization and other fields of development, i.e, this hypothesis can explain the majority of MPGN and MesPGN in our study. Chugh KS also found high prevalence of MPGN and MesPGN in India ²⁰ but Golay V et al found lower incidence (0.6%)in a recant study ²¹ which does not match our findings and we could not explain that.Our study findings regarding IgA Nephropathy (17.14%) is compatible with finding in other Asian countries¹¹. It is the most common from of primary GN in Asia, accounting for up to 30 - 40 % of all biopsies, for 20% in Europe and for 10% in north America. 22

Our study finding regarding focal segmental glumerulosclerosis (FSGS) (11.42%) and membranous nephropathy (MN) (11.42%) nearly identical with Habib MA¹² which showed FSGS was (11.53%) and MN (7.37%). Mundi I et al and Mannan R et al^{8,11} demonstrated. FSGS, MN and mininal change (MCD) are the commonest from of primary GN in India. Data from Singapore and other countries also showed that the prevalence of FSGS and MN have become increased in recent years.

The overall frequency of important complications after renal biopsy varied from 5% to 13% in previous reports.²³ which mainly included hematuria as in our study. No death or even nephrectomy was observed. This complications may be minimized in future by performing biopsy under USG/CT guidance.

Limitation of the study:

Small sample size of a single centre is the key limitations. A large number of study subjects from multi centre and availability of electron microscope could make our study more representative.

Conclusion:

To conclude, from the study and data analyzed the prevalence of GN is different all over the world due to various factors. Membranoproliferative and mesangioproliferative GN are more common in our country. whereas MN, MCD, FSGS and IgA nephropathy is not very prevalent. Renal biopsy is a safe procedure in expert hand. It also been realized that it is essential and necessary to maintain a control biopsy registry with as increased participation of many more nephrology centre. That will give us more accurate information about incidence, prevalence and distribution of glomerulonephritis in our country.

References:

1. Woo KT, Chan CM, Chin YM, Choong HL, Tan HK, Foo M et al. Global evolutionary trend of the prevalence of Primary glomerulonephitis over the past three decades. Nephron Clin Pract 2010; 116: c337-46.

2. Chang JH, Kim DK, Kim W. Changing prevalence of glomerular diseases in Korean adults: A review of 20 years of experience. Nephrol Dial Transplant. 2009;24:2406-10.

3. Utsonomiya Y, Koda T, Kado T, Okada S, Hayashi A, Kanzaki S et al. Incidence of pediatric IgA nephropathy Pediatr Nephrol 2003;18:511 - 15.

4. Gesualdo L, Di Palma AM, Morrone LF, Stripolli GF, Schena FP. The Italian experience of the national registry of renal biopsies. Kidney Int 2004; 66:890-94.

5. Hanko JB, Mullan RN, Rourke DM, McNamee PT, Maxwell AP, Courtney AE. The changing pattern of adult primary glomerular disease. Nephrol Dial Transplant 2009;24:3050-54.

6. Barden GL, Mulhern JG, O'Shea MH, Nash SV, Ucci AA Jr, Germain MJ. Changing incidence of glomerular disease in adults. Am J Kidney Dis 2000;35:878-83.

7. Mitwalli AH, Al Wakeel J, Abu-Aisha H, Alma A, AlSohaibani M, Tarif N, et al. Prevalence of glomerular disease: King Khalid University Hospital, Saudi Arabia. Saudi J Kidney Dis Transpl. 2000;11:442-48.

8. Ahmed P I, Zaman S U, Jahan F, Gupto R D, Chowdhury M N, et al. Pattern of primary glomerulonephritis in Dhaka Medical College Hospital, Bangladesh. Bangladesh J Medicine. 2014;25:42-46.

9. Mannan R, Bhasin TS, Singh PA, Misra V, Manjari M. The pattern of glomerulonephritis in the North Indian Gangetic Plain - a 13 year epidemiological study. Journal of clinical and diagnostic research. 2012;6(5):855-58.

10. Das U, Dakshinamurty KV, Prayaga A. Pattern of biopsy proven renal disease in a single center of south India: 19 years experience. Indian J Nephrol.

2011;21(4):250-56.

11. HabibMA, Badruddoza SM. Pattern of glomerular disease among adults in Rajshahi, the northern region of Bangladesh. Saudi J kidney Dis Transpl 2012;23(4):876-80.

12. Abdullah A, Khanam A, Biswas S, Niloy A, Shahin M, Murshed K, et al. Medical causes and histological pattern of glomerulonephritis. Mymensingh Med. J. 2008;17(1):38-41.

13. Hurtado A, Johnson RJ: Hygiene hypothesis and prevalence of glomerulonephritis. Kidney Int 2005;68:s62-67.

14. Balakrishnan N, John Jt, Korula A, et al. Spectrum of biopsy proven renal disease and changing trends at a tertiary care centre 1990-2001. Indian J nephrol 2003;13:29-35.

15.Mitwali Ah. Glomerulonephritis in Saudi Arabia: a review. Saudl J Kidney dis Transpl 2000;11:567-76.

16. Narasimhan B, Chacko B, John GT, Korula A, Kirubakaram MG, Jacob CK. Characterization of Kidney lesions in Indian adults: Towards a renal biopsy registry. Nephrol 2006;19:205.

17. Rahman T, Islam N, Rashid HU, Rahman M. Morphological syndrome-an experience based on light microsocopy of needle Biopsy. Bangladesh Renal J 1984;3:1-5. 18. McKenzie LM, Hendrickson SL, Briggs Wa, et al. NPHS2 variation insporadic focal segmental glomerulosclerosis. J Am Soc Nephrol 2007;18:2987-95.

19. Mubarak M, Kazi JL, Naqvi R, Ahmed E, Akhter F, Naqvi SA, et al. Pattern of renal disease observed in native renal biopsies in adult in a single center in Pakistan. Nephrology 2011;16:87-92.

20. Falk RJ, Jennette C, Nachman PH. Primary glomerular disease. In: Brennwr BM, ed. The kidney. 7th ed. Philadelphia: W.B Saunders Company; 2004.p. 1293-54.

21. Rashid HU, Sharmin S, Khanam A, Islam A, Ahmed J. Clinical presentation, histological diagnosis and management of primary glomerulonephritis in Bangladeshi adult population. Bangladesh Renal J 2003;22:35-58.

22. Mendelssohn DC, Cole EH, Outcomes of percutaneous kidney biopsy, including those of solitary native kidneys. Am J kidney Dis 1995; 26:580-85.

23. Whittier WL, Korbet SM. Timing of complication in percutaneous renal biopsy. J AM Soc Nephrol 2004; 15: 142-47.