DO YOU KNOW WHAT THE ICD-10 CODES ARE FOR THESE RARE GLOMERULAR DISEASES?

The use of these ICD-10 codes aims to help address gaps in diagnosis, surveillance, and research of some rare glomerular diseases[1-3]

Between 2020 and 2023, new ICD-10 codes have been launched for C3G (including C3GN and DDD), IgA nephropathy, and aHUS, allowing for greater specificity in coding the unique clinical features that can present with these glomerular diseases[4]

aHUS, atypical hemolytic uremic syndrome; C3G, complement 3 glomerulopathy; C3GN, complement 3 glomerulonephritis; DDD, dense deposit disease; HUS, hemolytic uremic syndrome; ICD-10, International Statistical Classification of Diseases and Related Health Problems 10th Revision; IgA, immunoglobulin A.

*This asterisk denotes a variable character; a complete list of codes can be found in the panel on the right.

1. Sun AZ et al. *Perm J*. 2020;24(2):19.126. doi:10.7812/TPP/19.126; 2. Dendooven A et al. *BMC Nephrol*. 2021;22(1):193. doi:10.1186/s12882-021-02365-3; 3. Kwon CS et al. *J Health Econ Outcomes Res*. 2021;8(2):36-45. doi:10.36469/001c.26129; 4. Centers for Medicare & Medicaid Services. Accessed November 27, 2023. https://www.cms.gov/medicare/coding/icd10

ICD-10 CODES FOR GLOMERULAR DISEASES[4]

C3GN

N00.A: acute nephritic syndrome with C3GN

N01.A: rapidly progressive nephritic syndrome with C3GN

N02.A: recurrent and persistent hematuria with C3GN

N03.A: chronic nephritic syndrome with C3GN

N04.A: nephrotic syndrome with C3GN

N05.A: unspecified nephritic syndrome with C3GN

N06.A: isolated proteinuria with C3GN

N07.A: hereditary nephropathy, not elsewhere classified with C3GN

DDD

N00.6: acute nephritic syndrome with DDD

N01.6: rapidly progressive nephritic syndrome with DDD

N02.6: recurrent and persistent hematuria with DDD

N03.6: chronic nephritic syndrome with DDD

N04.6: nephrotic syndrome with DDD

N05.6: unspecified nephritic syndrome with DDD

N06.6: isolated proteinuria with DDD

N07.6: hereditary nephropathy, not elsewhere classified with DDD

IgA nephropathy

N02.B: recurrent and persistent IgA nephropathy

N02.B1: with glomerular lesion

N02.B2: with focal and segmental glomerular lesion

N02.B3: with diffuse membranoproliferative glomerulonephritis

N02.B4: with diffuse membranous glomerulonephritis

N02.B5: with diffuse mesangial proliferative glomerulonephritis

N02.B6: with diffuse mesangiocapillary glomerulonephritis

N02.B9: other recurrent and persistent IgA nephropathy

aHUS

D59.30: HUS unspecified

D59.31: infection-associated HUS

D59.32: aHUS with an identified genetic cause

D59.39: aHUS (nongenetic)

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