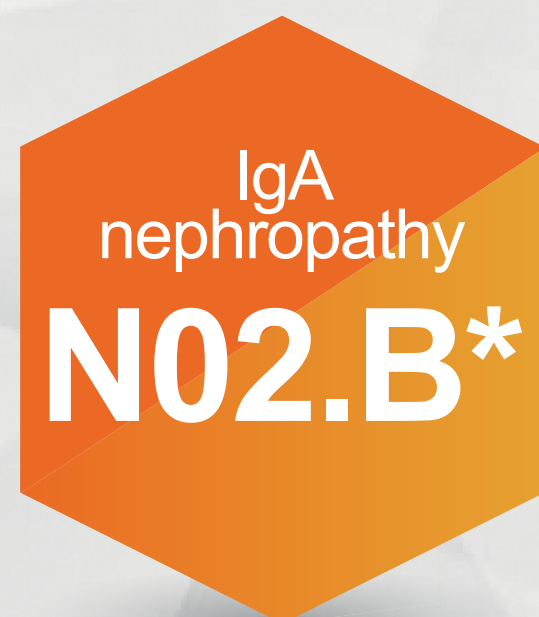
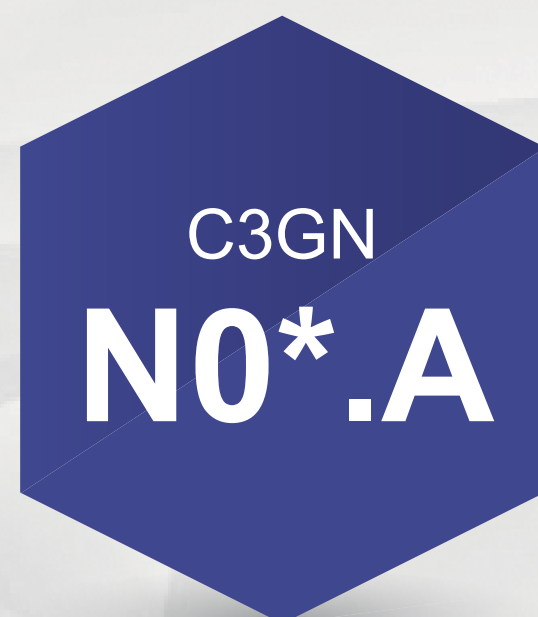


DO YOU KNOW WHAT THE ICD-10 CODES ARE FOR THESE RARE GLOMERULAR DISEASES?



The use of these ICD-10 codes aims to help address gaps in diagnosis, surveillance, and research of some rare glomerular diseases¹⁻³



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 diseases⁴

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 DISEASES⁴

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)

