Fechtner-Epstein syndrome

It is caused by a mutation in the myosin heavy chain gene **MYH9**. Hereditary nephritis, deafness, cataracts, and May–Hegglin anomaly appear in this syndrome. Inheritance is autosomal dominant. It was once classified as Alport syndrome.

Links

References

- KASHTAN, Clifford E. *Collagen IV-Related Nephropathies* [online]. ©28, August, 2001. The last revision 15, July,2010, [cit. 17-03-2023]. <https://www.ncbi.nlm.nih.gov/books/NBK1207/>.