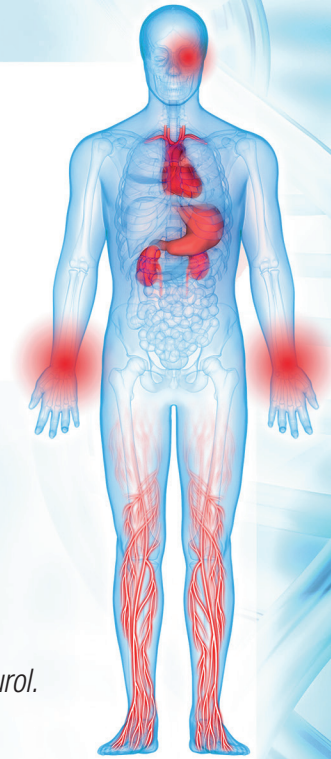


# Identifying Signs and Symptoms of Hereditary Transthyretin (hATTR) Amyloidosis in Pharmacy Practice



## Suggested Readings

Adams D, Ando Y, Beirão JM, et al. Expert consensus recommendations to improve diagnosis of ATTR amyloidosis with polyneuropathy [published online ahead of print, 2020 Jan 6]. *J Neurol*. 2020;10.1007/s00415-019-09688-0. doi:10.1007/s00415-019-09688-0

Adams D, Gonzalez-Duarte A, O'Riordan WD, et al. Patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. *N Engl J Med*. 2018;379(1):11-21. doi:10.1056/NEJMoa1716153

Benson MD, Waddington-Cruz M, Berk JL, et al. Inotersen Treatment for Patients with Hereditary Transthyretin Amyloidosis. *N Engl J Med*. 2018;379(1):22-31. doi:10.1056/NEJMoa1716793

Conceição I, Damy T, Romero M, et al. Early diagnosis of ATTR amyloidosis through targeted follow-up of identified carriers of TTR gene mutations. *Amyloid*. 2019;26(1):3-9. doi:10.1080/13506129.2018.1556156

Gertz MA, Mauermann ML, Grogan M, Coelho T. Advances in the treatment of hereditary transthyretin amyloidosis: A review. *Brain Behav*. 2019;9(9):e01371. doi:10.1002/brb3.1371

Kachur, E. Novel therapies for transthyretin amyloidosis. *US Pharm*. 2019;44(8):8-10. Available: <https://www.uspharmacist.com/article/novel-therapies-for-transthyretin-amyloidosis>. Accessed October 27, 2020.

Kapoor M, Rossor AM, Laura M, Reilly MM. Clinical Presentation, Diagnosis and Treatment of TTR Amyloidosis. *J Neuromuscul Dis*. 2019;6(2):189-199. doi:10.3233/JND-180371

Maurer MS, Schwartz JH, Gundapaneni B, et al. Tafamidis Treatment for Patients with Transthyretin Amyloid Cardiomyopathy. *N Engl J Med*. 2018;379(11):1007-1016. doi:10.1056/NEJMoa1805689

Witteles RM, Bokhari S, Damy T, et al. Screening for Transthyretin Amyloid Cardiomyopathy in Everyday Practice. *JACC Heart Fail*. 2019;7(8):709-716. doi:10.1016/j.jchf.2019.04.010