

Lechowicz, U., Pollak, A., Frączak, A., Rydzanicz, M., Stawiński, P., Lorens, A., Skarżyński, P.H., Skarżyński, H., Płoski, R., Ołdak, M. (2018), *Application of next-generation sequencing to identify mitochondrial mutations: Study on m.7511T>C in patients with hearing loss*. *Molecular Medicine Reports*, 17 (1), pp. 1782-1790

Cited: 2 times

Cited by:

Cui, Y., He, D.-J. (2018), Mitochondrial tRNA^{Ile} A4317G mutation may be associated with hearing impairment in a Han Chinese family. *Molecular Medicine Reports*, 18 (6), pp. 5159-5165. DOI: 10.3892/mmr.2018.9519

Lechowicz, U., Pollak, A., Raj-Koziak, D., Dziendziel, B., Skarżyński, P.H., Skarżyński, H., Ołdak, M. (2018), Tinnitus in patients with hearing loss due to mitochondrial DNA pathogenic variants. *European Archives of Oto-Rhino-Laryngology*, 275 (8), pp. 1979-1985. DOI: 10.1007/s00405-018-5028-y