

## Projekta Izp-2020/1-0269 rezultāti

### Primārie (pārsvarā) antivielu deficīti pieaugušajiem: meklējot slimības cēloņus un klīniskās izpausmes

*Oriģināli zinātniskie raksti, kas publicēti zinātniskos žurnālos, rakstu krājumos vai konferenču rakstu krājumos, kuri ir indeksēti datu bāzēs Web of Science Core Collection, SCOPUS vai ERIH PLUS*

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2. Nartisa, I.; Kirsteina, R.; Neiburga, K.D.; Zigure, S.; Ozola, L.; Grantina, I.; Micule, I.; Murmane, D.; Slisere, B.; Gailite, L.; Vilne, B.; Rots, D.; Taurina, G.; Kurjane, N. Clinical and genetic characterization of Netherton syndrome due to SPINK5 founder variant in Latvian population. - Pediatr Allergy Immunol, 2023, <https://doi.org/10.1111/pai.13937>
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6. Lucane Z, Davidsone Z, Micule I, Auzenbaha M, Kurjane N. A novel frameshift variant in the ADA2 gene of a patient with a neurological phenotype: a case report. - Pediatr Rheumatol Online. 2022, <https://doi.org/10.1186/s12969-022-00781-9>
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*Zinātniskās datubāzes un datu kopas*

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