

**GENETIC TEST:****Polyarteritis nodosa, childhood-onset / Vasculitis, autoinflammation, immunodeficiency, and hematologic defects syndrome**

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|-----------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>FULL NAME:</b>                 | Polyarteritis nodosa, childhood-onset / Vasculitis, autoinflammation, immunodeficiency, and hematologic defects syndrome                          |
| <b>TEST TYPE:</b>                 | Clinical                                                                                                                                          |
| <b>TEST SPECIALTY:</b>            | Molecular Genetics                                                                                                                                |
| <b>TEST PURPOSE:</b>              | Post-natal Diagnosis                                                                                                                              |
| <b>SPECIMEN:</b>                  | Peripheral (whole) blood on EDTA                                                                                                                  |
| <b>METHOD CATEGORY:</b>           | Sequence analysis: entire coding region                                                                                                           |
| <b>METHOD TECHNIQUE:</b>          | Next Generation Sequencing (NGS)                                                                                                                  |
| <b>RIZIV CODE:</b>                | 565471-565482                                                                                                                                     |
| <b>TURNAROUND TIME (MAXIMUM):</b> | 4 - 6 months                                                                                                                                      |
| <b>CREATED:</b>                   | 23 Jul 2019 - 09:53                                                                                                                               |
| <b>CHANGED:</b>                   | 19 Oct 2021 - 10:06                                                                                                                               |
| <b>URL:</b>                       | <a href="https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/14676">https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/14676</a> |

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## RELATED CONTENT

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### Related Diseases

- [Diamond-Blackfan anemia](#)
- [Sneddon syndrome](#)
- [Vasculitis due to ADA2 deficiency](#)

### Related Laboratories

- [Centrum Menselijke Erfelijkheid - KUL](#)

### Related Analytes

- [adenosine deaminase 2](#)

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