

**ANALYTE:**  
**ADAR**

|                             |                                                                                                                           |
|-----------------------------|---------------------------------------------------------------------------------------------------------------------------|
| <b>NAME:</b>                | adenosine deaminase RNA specific                                                                                          |
| <b>SYMBOL:</b>              | ADAR                                                                                                                      |
| <b>VERSION OF ORPHANET:</b> | 2023-06-22 14:14:43                                                                                                       |
| <b>SYNONYMS:</b>            | ADAR1                                                                                                                     |
| <b>XREF(S):</b>             | <u>Orphanet</u><br><u>Ensembl</u><br><u>Genatlas</u><br><u>HGNC</u><br><u>OMIM</u><br><u>Reactome</u><br><u>SwissProt</u> |
| <b>CREATED:</b>             | 13 May 2019 - 01:01                                                                                                       |
| <b>CHANGED:</b>             | 22 Jun 2023 - 16:14                                                                                                       |

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### Related Genetic Tests

- [Ataxia Spasticity \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
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- [Early onset epileptic encephalopathy \(gene panel - 845 genes\)](#)
- [Epilepsy \(gene panel\)](#)
- [Epilepsy gene panel](#)
- [Hereditary Spastic Paraplegia \(94 genes\)](#)
- [Hereditary spastic paraplegia \(gene panel - 249 genes\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Leukodystrophy \(gene panel\)](#)
- [Movement Disorders \(gene panel\)](#)
- [Movement Disorders \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Neuromuscular disorders \(548 genes\)](#)
- [Periodic Fever \(88 genes\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
- [Skin disorders \(gene panel\)](#)
- [Spastic Paraplegia \(gene panel\)](#)

## Related Diseases

- [Aicardi-Goutières syndrome](#)
- [Dyschromatosis symmetrica hereditaria](#)
- [Familial infantile bilateral striatal necrosis](#)

## Related Gene Panels

- [Ataxia Spasticity - UGent](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
- [Dermatogenetic / severe, rare and hereditary genodermatoses \(394 genes\) - ULB](#)
- [Early onset epileptic encephalopathy \(845 genes\) - ULB](#)
- [Epilepsy gene panel - VUB](#)
- [Hereditary Spastic Paraplegia \(94 genes\) - KUL](#)
- [Hereditary spastic paraplegia \(188 genes\) - ULB](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability/Epilepsy \(1091 genes\) - ULG](#)
- [Leukodystrophy - UGent](#)
- [Movement Disorders - UGent](#)
- [Movement Disorders - ULG](#)
- [Neurodevelopmental disorders \(1300 genes\) - ULB](#)
- [Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders \(1162 genes\) - VUB](#)
- [Neuromuscular disorders \(548 genes\) - ULB](#)
- [Periodic Fever \(88 genes\) - ULB](#)
- [Primary immune deficiencies \(444 genes\) - KUL](#)
- [Primary immune deficiencies - UGent](#)
- [Rare epilepsy with developmental delay \(> 240 genes\) - UZA](#)
- [Skin disorders - UGent](#)
- [Spastic Paraplegia \(89 genes\) - IPG](#)

- test

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