

**ANALYTE:**  
**UBE3A**

<b>NAME:</b>	ubiquitin protein ligase E3A
<b>SYMBOL:</b>	UBE3A
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNOMYS:</b>	ANCR AS Angelman syndrome E6-AP FLJ26981
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">Reactome</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Angelman / Prader Willi Syndrome](#)
- [Autism \(gene panel\)](#)
- [Cerebral palsy \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Early onset epileptic encephalopathy \(gene panel - 845 genes\)](#)
- [Epilepsy \(gene panel\)](#)
- [Epilepsy gene panel](#)
- [Epilepsy, seizures \(gene panel\)](#)
- [Intellectual Disability \(gene panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Malformations of cortical development \(235 genes\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)

### Related Diseases

- [15q11q13 microduplication syndrome](#)
- [Angelman syndrome due to a point mutation](#)
- [Angelman syndrome due to imprinting defect in 15q11-q13](#)
- [Angelman syndrome due to maternal 15q11q13 deletion](#)
- [Angelman syndrome due to paternal uniparental disomy of chromosome 15](#)

## Related Gene Panels

- [Autism \(57 genes\) - IPG](#)
- [Cerebral palsy \(212 genes\) - UZA](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
- [Early onset epileptic encephalopathy \(845 genes\) - ULB](#)
- [Epilepsy gene panel - VUB](#)
- [Epilepsy, seizures \(196 genes\) - IPG](#)
- [Intellectual Disability \(104 genes\) - IPG](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability/Epilepsy \(1091 genes\) - ULG](#)
- [Malformations of cortical development \(235 genes\) - VUB](#)
- [Neurodevelopmental disorders \(1300 genes\) - ULB](#)
- [Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders \(1162 genes\) - VUB](#)
- [Rare epilepsy with developmental delay \(> 240 genes\) - UZA](#)
- [test](#)

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