## Case information

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| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| Patient ID | Age | Gender | Family history | Gene | Country  | Mutation site/SV | Disease name | Phenotype  | Symptom |
| 1 | 2y3m | M | no | SCAMP5 | China | Chr15:75018813; NM\_001178111.2: c.538G>T p. (Gly180Trp)  | - | Global developmental delay; epilepsy | Global developmental delay; epilepsy |
| 2 | 8y4m | M | no | SCAMP5 | * Italy
 | - | Global developmental delay; epilepsy;ASD;Dysmorphism | Global developmental delay; epilepsy;ASD;Dysmorphism |
| 3 | 2y6m | F | no | SCAMP5 | America | - | Global developmental delay;epilepsy | Global developmental delay;epilepsy |
| 4 | 32y | F | Don’t know | SCAMP5 | America | - | Global developmental delay;epilepsy; ;Dysmorphism | Global developmental delay;epilepsy; ;Dysmorphism |

Family history: Yes, No or don’t know;

家族病史：是否有家族病史；

Country: The patient's nationality or the patient's race;

国家：病人的国籍或人种；

Mutation site/SV: Nucleotide and protein change information or genotype;

突变/结构变异：核苷酸、蛋白质的改变或基因型信息/结构变异的位置；

Disease name: The exact disease the patient was diagnosed;

疾病名称：病人被确切诊断的疾病名称；

Phenotype: The phenotype of the disease;

表型：疾病的表型信息；

Symptom: The clinical symptom;

症状：临床症状信息