**Table 1. Cases in which microdeletion has been detected in the 8q22.1 region and the characteristic features of the syndrome, reported since 2000.**

|  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | Teebi, 2000Raas- Rothschild *et al.*, 2009 1 | Allanson, 2002 Son | Allanson, 2002 Mother | Salpietro, Briuglia, Rigoli, Merlino, & Dallapiccola, 2003 Shieh *et al.*, 2006 2, Mazziotti *et al.*, 2014 | Shieh *et al.*, 2006 1 | Barber *et al.*, 2008 Family 2, Son | Barber *et al.*, 2008 Family 2, mother | Raas- Rothschild *et al.*, 2009 2 | Overhoff *et al.*, 2014 1 | Overhoff *et al.*, 2014 2 | Jamuar *et al.*, 2015 1 | Jamuar *et al.*,2015 2 | Our Patient. |
| Facial features |
|  | Age at evaluation | 4 yr | 4 yr | Adult | 14 mon | 21 mon | 22 mon | Adult | 9 mon | 12 yr | 11 wk | 14 yr | 4 yr | 7 mon |
| Gender | M | M | F | F | M | M | F | M | M | M | M | M | M |
| Tight or thick skin  | + | + | + | + | + | + | + | + | + | + | + | + | + |
| Frontal upsweep | + | + | + | + | U | U | + | + | + | U | + | + | + |
| Telecanthus | + | + | + | + | + | + | + | + | + | + | + | + | + |
| Blepharophimosis | + | + | + | + | + | + | + | + | + | + | + | + | + |
| Sparse eyebrows | + | + | + | + | + | + | + | + | + | + | + | + | + |
| Bulbous nose | + | + | + | + | + | + | + | + | + | + | + | + | + |
| Long philtrum | + | + | + | + | + | + | + | + | + | + | + | + | + |
| Cheek dimple | + | + | + | + | + | U | U | U | + | − | + | − | + |
| Abnormal ear configuration | + | + | + | + | + | + | U | + | + | + | + | + | + |
| Hearing loss | − | − | − | − | − | − | − | − | − | − | + | − | + |
| Intelligence and Developmental Disability | − | + | − | + | + | + | + | + | + | U | + | + | + |
| Cleft (submucous) palate | − | + | + | − | + | − | + | − | + | + | + | − | + |
| Dental Anomaly | − | U | + | + | + | U | U | − | − | U | U | U | + |

U:unspecified; +: the finding was identified; **−**: the finding wasn’t identified.

M: Male; F: Female.