Fig. S1. Smith-Magenis syndrome cases from infancy to adult. All cases shown either have a documented 17p11.2 deletion (a, b, d-g, i-k, m-o) or a de novo mutation (c, h, l) in the RAI1 gene. The same individuals at different ages are indicated by the same corresponding letter.
Supplemental Figure 2.

**Fig. S2. Self-hugging and hand-clasping behavior.** Male child, 5 y of age. Same child is shown in Fig. S1, panel d.