

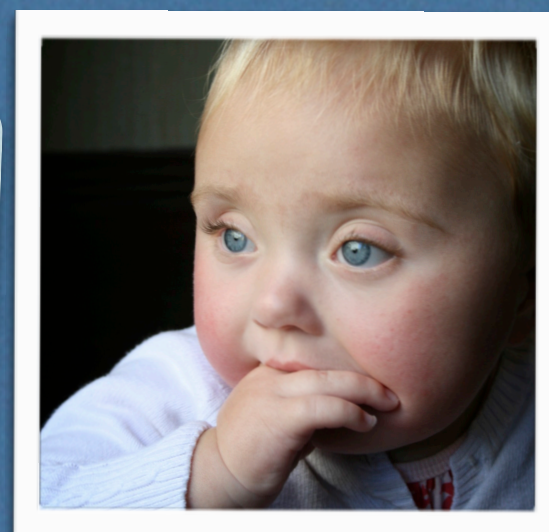
DOES YOUR PUZZLE PIECE JUST NOT FIT?

Smith Magenis Syndrome is an under diagnosed syndrome. Common characteristics are:

- Developmental Delay
- Flat midface, downturned mouth, prominent often rosy cheeks
- Low muscle tone (hypotonia)
- Early speech and motor delays
- Chronic sleep disturbance
- Maladaptive behaviors (hyperactivity, impulsivity, explosive outbursts, attention seeking, autistic-like/repetitive behaviors)
- Sensory integration issues
- May head bang, hit/bite self or pull fingernails off
- Happy disposition, sense of humor
- For more characteristics please visit www.prisms.org

SMS is associated with a small missing piece of chromosome 17p11.2

There may be thousands of people who have SMS and do not have a diagnosis. Estimated to occur in 1/25000 births, the syndrome is vastly under diagnosed.



Parents and Researchers Interested in
Smith Magenis Syndrome

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