



Every picture tells a story.

Emma, diagnosed at 3 years



Anthony, diagnosed at 3 years



Rayven, diagnosed at 1 year



Zachary, diagnosed at 5 months

For Emma, Anthony, Rayven, and Zachary, their stories begin with a diagnosis of Cornelia de Lange Syndrome (CdLS). Though not the storyline their parents expected, diagnosis helps these children and their families get appropriate medical and educational interventions as they go through each chapter of their lives.

But not all stories begin this way. CdLS remains undiagnosed in thousands of children. That's because CdLS is not a "one-size-fits-all" genetic condition. While many children have the classic signs—small size, eyebrows that meet, thin upper lip, long eyelashes, missing arms or fingers—others are so mildly affected, the condition goes unnoticed.

Help change a child's story by learning more about CdLS.

Go to www.CdLSusa.org or call 800-753-2357.



CdLS Foundation

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