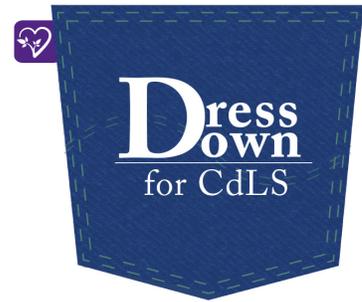


DRESS DOWN FOR CdLS

The Cornelia de Lange Syndrome Foundation, a national family support organization located in Avon, CT, exists to ensure early and accurate diagnosis of CdLS, promote research into the causes of the syndrome, and help people with CdLS and their families make informed decisions. An estimated 20,000 people in the U.S. have CdLS but remain undiagnosed or without support services.



Individuals with CdLS range from mildly to severely affected. Most have similar physical characteristics including small size, hands and feet; thin eyebrows that meet in the middle; long eyelashes; an upturned nose; and thin, downturned lips. Some individuals have limb differences, including missing fingers or arms.

DATE: _____

DONATION: _____

CONTACT PERSON: _____

All funds benefit the Cornelia de Lange Syndrome Foundation,
a 501(c)(3) nonprofit organization, Tax ID # 06-1057497, founded in 1981 by parents of children with CdLS.
It's the only organization in the country devoted to the syndrome.



REACHING OUT. PROVIDING HELP. GIVING HOPE.

www.CdLSusa.org

