

Do you know that... CHILDREN CAN HAVE CFS/ME TOO?

Young Faces of ME



SAM:

"ME took away my young life, my education and my friends. I was worthy of better treatment and of being believed."

"I loved school. I wanted to be there in the company of my peers."

ANSTICE:

"When I was a child I was bed-bound for 10 months, at times unable to speak or eat solid foods."



"I was unable to do any school work. ME robbed me of education and qualifications."

ZOE:

"Going in the garden is an outing. I hoped to be a musician; my brother is – I've never been able to go to his gigs."

"As a child a month into the illness I had to lie down a lot - two decades later I still do."



**OMEGA is a support group for children, adults and carers affected by CFS/ME too.
We are campaigning to extend the community services
to cover children under 14 years old.**

"I feel so sorry for the many young kids who are suffering the isolation and loneliness that I have suffered. It's not right and it shouldn't happen." (Sam)

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