

Taking Cystic Fibrosis To School

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Advice from the UKCFMA The UKCFMA has advised that the vast majority of parents should plan for their child or children with cystic fibrosis (CF) to return to school. There may be a very small number of individual children with CF who might be advised not to go back to school yet because of the severity of their cystic fibrosis.

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Even though Jessie has cystic fibrosis, she can still attend school and do many of the same things as her classmates. Written from Jessie's perspective, this book explains and educates her classmates about her condition. Jessie explains to her classmates what cystic fibrosis is and tells them that it feels like "elephants dancing on her chest."

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do many of the same things as her classmates. Written from Jessie's perspective, this book explains and educates her classmates about her condition.

~~Taking Cystic Fibrosis to School by Cynthia S. Henry~~

Whether you are the parent of a child who is starting preschool, the teacher of a student with cystic fibrosis, or a college student leaving home for the first time, preparation is key. The following resources are intended to help you manage CF at school.

~~CF and School—Cystic Fibrosis Foundation~~

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A student with cystic fibrosis may fatigue easily. This could be due to chronic infections, early waking for morning respiratory treatments, poor lung function and malnutrition. You may notice reduced endurance in comparison to other students, and long school days can be difficult for some.

~~Cystic Fibrosis in the Classroom—CFRI~~

Cystic fibrosis shouldn't stop your child enjoying a full and rewarding school experience. Compromises may need to be found, and minor adjustments made, but working in partnership with the school and your CF team will help ensure your child's education is not limited by cystic fibrosis.

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