

Supporting Children & Young People with CFS/ME

Information for Schools & Colleges

This leaflet contains information and guidance designed to help professionals in educational settings to understand the symptoms experienced by children and young people with Chronic Fatigue Syndrome/Myalgic Encephalitis (CFS/ME) and to support their continued attendance and success.

What is CFS/ME?

CFS/ME is relatively common in children, affecting at least 1% of teenagers. It is probably the largest cause of long-term absence from school. CFS/ME symptoms may include:

- Extreme fatigue after normal everyday activities
- Pain (including headaches)
- Impaired mental functioning (poor concentration, distractibility, difficulty focussing, poor memory, 'brain fogging')
- Malaise and dizziness, nausea
- Sleep disturbance

We usually find that children and young people with CFS/ME have only a finite amount of energy capacity and the normal "recharging of batteries" is impaired. If the amount of available energy is exceeded, then the young person experiences a 'payback' over the next day or two where they may feel 'fluey' and wiped out. This is also known as a "Boom and Bust" cycle. In our experience, young people with CFS/ME may also be more likely than those without fatigue to experience anxiety and low mood. The management of CFS/ME involves finding out how much high energy activity time a young person can spend every day, without experiencing this payback. This is called the baseline.

Energy Expenditure

There are three types of energy expenditure: physical, cognitive and emotional.

- **Physical** energy expenditure occurs for any activities involving mobility.
- **Cognitive** energy is anything to do with thinking tasks – all classroom work is high energy cognitive activity. Other high energy cognitive activity would be watching television or chatting with friends.
- **Emotional** energy is often easy to overlook. This includes any emotional distress, rows, anxiety and 'over-thinking' the young person experiences. Although this can be difficult to control, it needs to be factored into the available amount of energy.



Treatment Programmes

The Paediatric CFS/ME service at the Royal United Hospitals, Bath offers a range of treatment programmes, each of which are tailored to the individuals needs and agreed as part of our regular consultations with them and their families. They may include:

Adjustments to sleep-wake cycle

Changing sleep-wake cycles can lead to an increase in symptoms in the short term, and this may well have an impact on their school attendance.

Changing activity patterns

We usually expect to see a **reduction** in levels of activity at the start of treatment. Students will often be advised to reduce their planned school attendance in the short term. It is better to attend for less time, consistently, rather than trying to attend full time and missing random school days/weeks due to payback. Once they are achieving a manageable amount of school attendance every day, without experiencing payback symptoms, they will then work on building their attendance back up in a gradual and sustainable way.

Ways to help manage CFS/ME at school:

Rest breaks during school time

Some children/young people with CFS/ME benefit from regular rest breaks. These breaks should be in a quiet place, ideally away from the classroom. It would be helpful if breaks could be scheduled, especially for younger children. Other students prefer a medical timeout card which they can use in lessons if they feel they need a break. They should be allowed to leave the classroom without having to explain why. In a quiet space, e.g. the library, they should spend the time reading or listening to music or doing a downtime activity. They can be expected to return to the lesson after 10-15 minutes.

Concentration

We have found that young people with CFS/ME benefit from working in chunks of no more than 30-45 minutes; severely affected students may only be able to sustain concentration for 15 – 20 minutes. Students often have increased sensitivity to everyday sounds and are likely to be more easily distracted in the classroom environment. Students may experience 'brain fog' i.e. difficulty processing information – it can be helpful to provide written instructions and memory aids.

Exams

Young people with CFS/ME benefit from being allowed to sit exams in a room on their own, or with few other students. They should be allowed to get up and move around for timed rest breaks. We would also advise that they are allowed 25% extra time.

The above allowances will really help children and young people with CFS/ME to reach their full academic potential.

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Please contact the Patient Advice and Liaison Service (PALS) if you require this leaflet in a different format, or would like to feedback your experience of the hospital.

Email ruh-tr.pals@nhs.net or telephone 01225 825656.