

Epilepsy Etiquette

a simple etiquette guide for people who don't have epilepsy



EPILEPSY: DOS AND DON'TS

Don't refer to me as an epileptic or label me.

I am an individual with many interesting things about me. I also have epilepsy.

Don't assume that epilepsy is a mental condition.

Epilepsy is a physical disorder that has nothing to do with mental health or illness.

Don't offer me advice on my epilepsy.

I know my condition best. Unsolicited advice and tips are not helpful.

Don't assume that everyone with epilepsy has the same condition.

There are many different forms of epilepsy. Seizures can differ between people.

Don't assume that living with epilepsy is easy.

It's extremely hard physically and emotionally living with epilepsy.

Don't patronise me because I have epilepsy.

Treat me with respect and don't talk down to me. I want to be understood without judgement or preconceived assumptions.

SEIZURES

Do talk to me about my needs and how you can help.

Feel free to discuss my epilepsy with me and find out how you can help. Listen to any instructions I have.

Don't use words like fits, spells, epis or attacks

Just call them seizures. Ignorance and discrimination only make it harder so stick to the facts.

Do give me time alone after a seizure.

I may be confused, disorientated and embarrassed so just leave me alone to gather myself.

DOS



Guide me from danger - remove harmful objects from nearby & cushion my head.



Stay with me until I am recovered.



Call an ambulance if the seizure lasts more than 5 minutes.



Aid my breathing by gently placing me in the recovery position once the seizure has finished.



Be calmly reassuring.

DON'TS



Restrain my movements



Put anything in my mouth



Move me unless I am in danger



Attempt to bring me round



Give me anything to eat or drink until I am recovered

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Understanding Epilepsy What It Is And How It Can Affect Your Life

Humberto Foyaca-Sibat



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