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I have CFIDS + am finally on pain meds. I have been on oxycontin for around 3 months. It is now not helping like it use to. My doctor (which understands CFIDS + stays on top of it) is referring me to a pain center. Guess what - my insurance won't pay for this with I have seen a orthopedic, rhumatologist, + neurologist. So this means I have to wait a couple of months in pain to see these other doctors. The dose of oxycontin does help but not like it should. I am a believer in pain control - not anti inflammatory meds which do not help. I do not want

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to live my life in pain. There is no quality of life & it affects not only me but my whole family, I will not live in pain. So please FDA let us have pain relief. It is the only humane thing to do. Stop the insurance companies from being able to control what the doctors need to do for us.

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