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Annotated click image for details Introduction John is more at risk than most of us since he suffers from chronic fatigue syndrome. "It seems like it would be pretty stressful for any family. It seems pretty difficult to maintain a job like mine. If you're not sleeping, if you're not eating..." Paul is more at risk than most of us since he's diagnosed with multiple sclerosis. "Its just been incredibly stressful. It's been an unbelievable amount of stress for such a seemingly insignificant amount of progress. I used to not worry about it at all." "It is often said that life is change, some painful and some pleasurable. However, we've had a fairly pleasurable experience." "Thank you so very much for this opportunity. I've been diagnosed with Chronic Fatigue Syndrome. The last five years have been a lifechanging experience for me." "It's the most intense thing that's ever happened to me. I can go to sleep and I wake up and then I haven't slept for a week and I still just start from zero. And then it would be again. I have this cycle, and I think that that's it's intensifying my fatigue, you know, my being out of it." "I had my first [episode] five years ago. It was my mother's birthday. She had cancer. I think it brought back memories. I had it when I was growing up. It wasn't until I was an adult. The last two years, it's just got worse and worse and worse. I can't function. I can't do anything. It's just -- I have a very serious illness, and I'm going to have to learn how to live with it." "Forgive me for my English: I am Vietnamese. I am more at risk than most of us. I was a [night] shift baby, and so my parents decided that they would be taking me to the hospital in case I had problems. Even though I was small, I needed to be monitored." "I was born in the beginning of June, and I guess it's the middle of July that I got really sick with recurrent swollen glands. But it wasn't until November that I was diagnosed and that I got fed up and I decided to get into the real help. And I went to the real doctors." "I was born in the beginning of June, and I guess it



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Properties of prodigiosin Pha is a red pigment that is produced by bacteria including species of Streptomyces. Based on structural analysis, prodigiosin is classified as a type of polyketide, based on the presence of carbon-carbon chains of varying length. The structural differences between prodigiosin, and the related pigments hyphaeubin, actiprotinin, and kibbybean are linked to the lengths of the carbon chains, as are differences between prodigiosin and melanin, which is a form of melanin derived from tyrosine. Pigmentation is a mechanism that microbes use to protect themselves from DNA damaging agents like sunlight, and bacteria have been observed to produce other pigments that protect from radiation. Based on the structural differences between prodigiosin, actiprotinin, and kibbybean, prodigiosin has anti-cancer, anti-atherogenic, and anti-viral properties, and was originally found to be antibacterial but subsequent tests found that prodigiosin had no effect on bacteria. The most widely studied effect of prodigiosin is its anti-cancer properties. DPH Pro 8 Download Properties of prodigiosin Isolated from Proteus mirabilis (Beringer, 1782). Hyphaeubin is produced by the ubiquitous Streptomyces hygroscopicus, and similar pigments are produced by Streptomyces chromofuscus, Streptomyces maacae, and Streptomyces griseolus. Actiprotinin is produced by the bacterium Actinomyces griseus. Kibbybean is produced by the bacterium Kibdelosporangium arvum. Prodigiosin effects Prodigiosin has been shown to inhibit a wide range of cancer cell lines at nanomolar concentrations. In vitro assays have demonstrated growth inhibition of melanoma, breast, prostate, colon, small cell lung, and ovarian cell lines. Prodigiosin is cytotoxic to cancer cells because it stimulates apoptosis, which is programmed cell death. Proposed mechanisms for induction of apoptosis include activation of the p38/mitogen activated protein kinase (MAPK) pathway, the inhibition of the epidermal growth factor receptor (EGFR), the inhibition of the Akt pathway, the inhibition of the cell cycle, the induction of increased reactive

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