

FDA FIDDLES 27,000 MEN DIE

4/30/08 Care To Live brings you week #5 of our Rockville area weekly plea from patients, physicians or families of patients, who suffer from advanced prostate cancer, which will continue until the FDA acts in a humane manner.

On behalf of our members we filed a Citizens Petition on July 27, 2007 asking the FDA to reconsider the failure to approve Provenge and by law they should have properly responded in January. To date they have not properly responded.

The FDA ignores our Petition while our members continue to die painful deaths.

Why can't the FDA Commissioner consider the patients and take some action? We want better, safer treatments NOW!

To read our past, present, and future ads visit our website. See you next week in Rockville.

Dear Reader,

Some 220,000 American men are diagnosed with prostate cancer each year. Unfortunately, 27,000 of them go on to die from the disease yearly. Those unfortunate individuals often suffer a lingering, painful death.

Unlike breast cancer where

there are many options for treatment for advanced disease, prostate cancer has few treatment options. The patients suffer with the side effects of hormone treatment. These include weight gain, loss of libido and muscle mass, depression, exacerbation of heart disease and fatigue.



Dr. Robert Rostock

When that fails there is only one effective chemotherapy drug which many patients refuse or can't complete due to considerable side effects.

Last year the FDA failed to approve a promising new drug called Provenge for the treatment of prostate cancer.

The FDA convened a panel of experts to advise them whether the drug should be ap-

proved. The panel voted 17-0 that the drug was safe and 13-4 that it was effective. In an unprecedented move, the FDA did not approve the drug.

The decision was marred with conflicts of interest and back room politics. The FDA has neither disciplined or investigated the individuals involved.

As a result of the decision, many physicians have become critical and cynical of the FDA.

It is difficult to watch patients suffer and die when they have run out of options and there is a promising treatment such as Provenge.

The FDA could have made the drug available while monitoring the results of an additional trial. Instead they chose to wait for more data and allow more deaths to occur from this disease.

*Sincerely,
Robert A Rostock MD*

Chairman of Oncology, Geisinger, South Wilkes Barre, PA; Chairman, Radiation Oncology, Wyoming Valley Health Care, and Chair, Institutional Review Board; former faculty member and specialist in immunology research, Johns Hopkins Hospital

CareToLive is a not for profit corporation

This is OUR FDA

We need to take it back from certain FDA individuals with their own ambitions

FDA Approve Provenge Now!

More on this sordid affair at

www.CareToLive.com

HEY HEY FDA HOW MANY MEN DIED TODAY ?

5/14/08 Care To Live brings you week #7 of our published plea from patients, physicians or families of patients, who suffer from advanced prostate cancer, which will continue until the FDA acts in a humane manner.

On behalf of our members we filed a Citizens Petition on July 27, 2007 asking the FDA to reconsider the failure to approve Provenge and by law they should have properly responded in January. To date they have not properly responded.

The FDA ignores our Petition while our members continue to die painful deaths. 1 in 6 men get prostate cancer. 75 American men die of it every day which is over 27,000 men a year.

Why can't the FDA Commissioner consider the patients and take some action? We want better, safer treatments NOW!

Join our May 30 nationwide demonstration protesting the FDA delay of Provenge. To learn more about us and to see all our current, past and future ads, please visit our website.

Dear Reader,

My family has been closely following the Provenge saga as a treatment for prostate cancer. My two brothers and a brother-in-law are victims of this disease, as well as me. As a survivor of prostate cancer, I

know full well that the disease has not been totally eradicated from my body; and, I also know

ical/medical specialties group called Physicians for Provenge) actively voicing their support for this vaccine.

The action by the FDA to delay approval of Provenge, even though it was approved by its own expert advisory panel as "safe and effective", is deplorable.

I am most concerned that the delay of the availability of Provenge has now become a political issue, losing sight of the tens of thousands of men who could benefit from Provenge as another approach to treatment.

Does anyone care that men and their families are suffering when they could have had additional, quality time with their loved ones?

My fondest hope is to be able to see my grandchildren grow and be a part of their lives and I am certain I speak for countless others.

We have been told by several physicians that, due to family history, our own two sons may be faced with this disease unless safe and effective treatments such as Provenge are provided to them. The question remains, "How many more men will die as a result of not having Provenge available to them?"

Sincerely,

*C. Richard Wick
Psychologist, B.S., M.A.
St. Joseph, Michigan*



CareToLive Member C. Richard Wick, his wife and 3 Grandchildren

that the use of radiation or surgery will not be a treatment option for me should the cancer reappear.

So, I along with 50+ members of the Man-To-Man prostate group here in Berrien County, Michigan, meet monthly to hear what other options are out there for us.

Recently, we had Dr. Mark Moyad from the University of Michigan present current research related to treatment of prostate cancer. He stated that he supports use of Provenge for this devastating disease and there are a number of groups (i.e. a broad diversity geograph-

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