# Hermiston man deals with tough diagnosis

By KATHY ANEY STAFF WRITER

Hearing he had five years left to live jolted Jeff Sepulveda.

The Hermiston man and wife Rebecca got the grim prediction from his doctor in October at the Oregon Health & Science University Hospital in Portland. Sepulveda has scleroderma, an autoimmune disease that attacks healthy tissue. People with scleroderma often experience stiffening and tightening of the skin and connective tissue, the feeling of slowly turning to stone.

In Sepulveda's case, the disease affects his lungs. Pulmonary function has dropped to 46 percent as the fibrosis in his lungs worsens. The couple digested the devastating news as they drove home to Hermiston after the doctor's appointment.

"It hit me and my wife pretty hard," he said. "It was a quiet ride home."

They looked for options, but found few. Sepulveda takes a drug called mycophenolate to keep the disease at bay, but it's not a cure.

"It slows the process down," he said. "It's not a cure by any means."

Other treatments show promise but are still considered experimental by insurance companies.

Stem cell transplantation seemed to offer the most hope. Jeff's stem cells would be collected. He would undergo chemotherapy to destroy his malfunctioning immune system and then the collected stem cells would be returned to his body. The process takes several excruciating months and the outcome isn't assured. The added life expectancy could only be a few years, even if all goes well.

"It's not a for sure deal," he said. "It may take or it may not."

The procedure is expensive. Sepulveda will have to provide roughly \$200,000 out of pocket. Friends and family are working to raise the money.

The 37-year-old has a lot to live for — his wife, two young daughters, a close extended family, a good job

By JADE MCDOWELL

STAFF WRITER



Staff photo by E.J. Harris

Jeff Sepulveda of Hermiston was diagnosed with scleroderma, a fatal autoimmune disease that attacks a person's lungs, this summer and was told by his doctor he had five years to live.



Raynaud's disease.

unfinished dreams.

said.

West-end cities get grant

for housing study

The cities of Umatilla,

The cities received a grant

Stanfield and Echo are join-

ing forces to study housing

from the Oregon Department

of Land Conservation and Development to develop a

housing needs analysis. They

will work with consultant

release, while the city of

Hermiston has been partic-

ipating in discussions with

the other three cities about

addressing housing shortages

in the west end of Umatilla

County, Hermiston will not

be part of the project because

it already completed a very

similar housing analysis in

development director Tamra

Mabbott said the project

community

Umatilla

tions for the cities.

According to a news

Angelo Planning Group.

"My priority right now is

being here for my wife and kids as long as I can," he

according to the Sclero-

derma Foundation. The disease, which is difficult to

diagnose because it shares

symptoms with other auto-

immune diseases, typically

comes on between the ages

set for sometime in January,

where developers, homeless

advocates and other stake-

holders can give feedback

on housing needs in the area.

She said the report will likely

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be wrapped up by June.

Staff photo by E.J. Harris Poor circulation in the extremities was Jeff Sepulveda's first sign that something was wrong after he was diagnosed with

> Outwardly, Jeff looks fairly healthy. One can imag-

ine him hefting tires and fixing brakes at the Hermiston Les Schwab. Growing up in Boardman, he loved long-distance running and continued to run and bike for exercise as an adult.

So far, Jeff says life hasn't changed too much, although he's had to dial it down. He is taking off the winter from work because the cold air affects his lungs. It takes longer to complete physical tasks such as mowing the

A close look at his hands shows purple fingers, a symptom of Raynaud's disease, which he's had for the past 16 years. Raynaud's is

linked with scleroderma. Despite the grim reality of his condition, at least outwardly, Jeff's outlook seems as bright as the tree that glows in his living room. He and Rebecca are humbled by the supporters who are working to raise funds for the stem cell transplant or just offering words of encouragement.

"It has been amazing," Rebecca said. "People near and far have reached out to

Flyers hanging around town list several options for those interested in helping. Look for donation boxes at Les Schwab, Shell stations or Ace Hardware, deposit to the "Donations for Jeff Sepulveda" account at Bank of Eastern Oregon branches, to www.gofundme. com/jeffs-scleroderma-battle, pay five dollars for a "Fight for Jeff" wristband or attend a fundraising dinner at 5 p.m., Feb. 9 at the Port of Morrow Conference Center. To purchase tickets or donate an auction item, call Tia Sepulveda at 541-314-3509 or Dawn Coleman at 509-308-3666.

#### Rebecca is focused on "He's pretty quiet and super hard working. He's a great dad to our kids," she said. "This has all been a big shock to us." About 300,000 Americans have scleroderma,

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or at the Chamber Office at 501 South Main Street

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During this special time of year, encourage your loved ones who have served in the military to make an appointment with their local Veterans Service Representative to review their federal and state benefits. They may be entitled to Va Health care, Compensation, Pension, Education or Home loan benefits.

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for he careth for you. 1 Pet. 5:7

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# will include a look at housing needs now and 20 years down the road, an examination of residential land available for development and a study of barriers to that development. It will culminate in a set of recommenda-

Mabbott said those recommendations may be policy changes for the city council to implement, or zoning changes an expansion of the urban growth boundary or other strategies. 'It's important to have a

set of recommendations in place because it sets the stage for other work that needs to be done," she said. Mabbott said there will

be multiple public meetings