

her life. Everyone called her "Grandma Margaret."

Margaret leaves behind five daughters, Tillie Rodriguez, Ruby Ramirez, Mary Ramirez, Lorraine Chavez, JoAnn Beckman; and five sons, Pete Sierra Jr., Charlie Sierra, Amador Sierra, Johnny Sierra, and Joe Madrigal; sixty-five grandchildren; and four great-grandchildren.

I extend to the family my condolences and wish blessings to them in their time of mourning. We say, "goodbye, we miss you, God bless."

IN SUPPORT OF NATIONAL
WOMEN'S HEALTH WEEK

HON. CARRIE P. MEEK

OF FLORIDA

IN THE HOUSE OF REPRESENTATIVES

Wednesday, May 16, 2001

Mrs. MEEK of Florida. Mr. Speaker, I rise today in support of National Women's Health Week, to speak of a topic near and dear to me which is Lupus.

I know firsthand the heartache that lupus causes. I lost a sister to lupus and have seen many others suffer from this incurable disease. I know all too well the difficulties persons with lupus face to maintain employment and lead normal lives. I have seen the often-devastating side effects of current treatment regimens. I also know the profound impact that my sister's disease had on me and that lupus often has on the family and friends of lupus patients.

More people have lupus than AIDS, cerebral palsy, multiple sclerosis, sickle-cell anemia and cystic fibrosis combined. Yet I believe that much of the public does not yet have this awareness. The Lupus Foundation of America estimates that between 1,400,000 and 2,000,000 people have been diagnosed with lupus. Many others have the disease, but have not even been diagnosed because of the insidious way in which lupus "masks" itself, thereby often making it difficult to diagnose. Many lupus victims are mis-diagnosed, and some victims even die, without even knowing that they have this disease.

Lupus is a wide-spread and devastating autoimmune disease that causes the immune system to attack the body's own tissue and organs, including the kidneys, heart, lungs, brain, blood, or skin. It afflicts women nine times more than it does men, and is three times more prevalent in women of color than Caucasian women. Lupus has its most significant impact on young women during their childbearing years (ages 15-44).

Lupus patients from poor or rural areas often cannot access the level of specialty care required to manage such a varied and complex disease. When first presenting symptoms of the disease, lupus patients usually contact their family physician. It is not unusual for people to have lupus for three to five years and to visit up to five doctors before they receive a correct diagnosis. Unfortunately, medical schools do not provide family physicians with sufficient training to recognize lupus.

I am sure that increased public awareness of the pervasiveness of lupus will substantially assist our efforts to increase funding not only for research, but also for the treatment and support services that the Congress authorized last November when it passed my lupus bill,

H.R. 762, as part of the Public Health Improvement Act of 2000 (P.L. 106-505). Passage of H.R. 762 was an important step in the fight against lupus, one of which I am extremely proud. But it is not enough. It is time to take the next step this year by funding the research, treatment and support services that the Congress authorized last year when it passed my lupus bill.

Lupus affects multiple organ systems and can be an expensive disease to manage. Treatment requires the participation of many different medical specialists and expensive specialized testing and procedures. The average annual cost of medical treatment for a lupus patient is between \$6,000 and \$10,000. However, for some people with lupus, medical costs may exceed several thousand dollars every month. Lupus can be financially devastating for many families.

It was these human factors that caused me to offer H.R. 762 and to work so hard for so many years with all of you for its passage. The case management and comprehensive treatment services that we authorized in H.R. 762 for individuals with lupus, and the support services that we authorized for their families, will be tremendously helpful, but only if we adequately fund them. We need a coordinated, targeted, well-executed appropriations strategy to make the promise of these programs a reality.

My lupus bill that the Congress passed last year authorizes appropriations of such funds as are necessary for FY 2001 through FY 2003 for lupus research, education, and treatment, including a grant program to expand the availability of lupus services. It also empowers the Secretary of the Department of Health and Human Services to protect the poor and the uninsured from financial devastation by limiting charges to individuals receiving lupus services pursuant to the grant program, the way that we do under the Ryan White Care Act, should the Secretary deem it appropriate to adopt such limitations.

H.R. 762 authorizes research to determine the reasons underlying the increased prevalence of lupus in women, including African-American women; basic research concerning the etiology and causes of the disease; epidemiological studies to address, among other things, the differences among the sexes and among racial and ethnic groups with respect to the frequency of the disease; the development of improved diagnostic techniques; clinical research for the development and evaluation of new treatments, including new biological agents; and information and education programs for health care professionals and the public.

The bill also authorizes appropriations of such sums as are necessary for FY 2001 through FY 2003 for a grant program. This program would support a wide range of services for the diagnosis and disease management of lupus for lupus patients, as well as a broad range of support services for lupus patients and their families, including transportation services, attendant care, homemaker services, day or respite care, counseling on financial assistance and insurance, and other support services.

I think it is appropriate during National Women's Health Week, that Congress fully fund research and treatment programs such as this.

TRIBUTE TO THE LATE
FRANCISCA GARMON

HON. JOE BACA

OF CALIFORNIA

IN THE HOUSE OF REPRESENTATIVES

Wednesday, May 16, 2001

Mr. BACA. Mr. Speaker, it is with great sadness that I rise in memory of Francisca Garmon, of my district, who passed away on Mother's Day, May 13, 2001.

Francisca served as vice president of the local union, United Steelworkers of America Local 7600, which represents approximately 4,000 Kaiser Permanente Medical Center Employees in San Bernardino and Riverside counties. A woman of great faith, Francisca was known for her resilience and tenacity. A gifted communicator, she was asked by the union to serve as a spokeswoman because of her speaking abilities. A talented singer, she made a recording last year at the request of the Steelworkers International.

Francisca is survived by her husband, James Garmon, a physician's assistant at Kaiser Permanente's San Bernardino Clinic. She is also survived by her mother Virginia; Children Johnny, Troy and Anna (Sey), who is a customer service representative at Kaiser's Corona Call Center; grandchildren Dana, Kaleb and Jacob; brother, Richard; sisters Evelyn, Jeannie and Rosie; and many other relatives.

Francisca had worked for Kaiser Permanente for 18 years. Prior to becoming a grievance officer she served as an assistant grievance officer. In the year 2000, Fran became vice president of USWA Local 7600. She served as co-chair of the Legislative and Education Committee, was active in the Labor Management Partnership and was a political activist for State and Federal labor laws.

Francisca's presence, along with her efforts and hard work, will be missed tremendously by all her Brothers and Sisters of Local 7600, and indeed, all in our community, but our comforting memories of her will live on.

Francisca also served her country in the armed forces, in the Air Force from April 4, 1970, through September 11, 1970, being honorably discharged due to pregnancy; and also in the Army for two years, being honorably discharged on April 17, 1979.

Francisca made a lasting difference in her community. Our hearts go out to her family and loved ones. With God's grace we know she will have peace.

THE COST OF HIGH ENERGY
PRICES ON OUR NATION'S AGRICULTURE PRODUCERS

HON. JERRY MORAN

OF KANSAS

IN THE HOUSE OF REPRESENTATIVES

Wednesday, May 16, 2001

Mr. MORAN of Kansas. Mr. Speaker, I rise today to call attention to the energy crisis that is draining the farm economy. My district, like many rural areas across the country, has suffered greatly as a result of high energy prices. Agricultural producers in particular have been hit hard as higher diesel and natural gas prices increase fuel, irrigation energy, and fertilizer costs.