Adele Hall’s personal and professional record exhibits her spirit of commitment to others. Her entire life has exemplified the core values that we all strive for: commitment to the work and family responsibilities. Adele Hall’s various roles and achievements throughout the history of the Central Exchange have demonstrated that she has succeeded in fulfilling her dream of increasing the visibility and effectiveness of Kansas City’s most reputable women’s organizations. She has served as a board member for the Trust Fund of the Library of Congress, the George Bush Presidential Library Center, the American Academy of Pediatrics, and the Salvation Army. Currently, she is serving as Co-Chairman of a $175 million capital campaign for the Nelson-Atkins Museum of Art. She is the Vice-Chairman of the United Negro College Fund and the Youth Corps of America.

As a founding member of one of Kansas City’s most reputable women’s organizations, the Central Exchange, she has worked tirelessly to promote the advancement of women in all sectors of society. For the past 20 years the Central Exchange has worked to bring people of diverse backgrounds together to encourage the personal and professional growth of women. Today the Central Exchange boasts nearly 900 members from all over the Kansas City metropolitan area. The astounding membership can be attributed to what members of the Central Exchange value the most, creating opportunities to meet and learn from other women. This is an extremely difficult goal when many women are busy with work and family responsibilities. Adele Hall’s various roles and achievements throughout the history of the Central Exchange have demonstrated that she has succeeded in fulfilling her dream of increasing the visibility and effectiveness of Kansas City’s women’s organizations.

As a leader of our community, Adele Hall has been particularly committed to an issue I know well: AIDS. As many of my colleagues know, since the moment I first stepped foot in Washington, I have fought for increased funding for critical HIV and AIDS education, treatment and research programs. I have also worked to expand our current programs to areas that are still in need of our help. Africa, India, the Caribbean, and Central and Eastern Europe in particular need our help and Senator MOYNIHAN has been particularly committed to an issue I know well: AIDS. Unfortunately, it seems the rise in hospice care the 1990s brought about an increase in fraud and abuse of the Medicare system, which in turn sparked a misguided crackdown on terminally ill patients.

HCFA officials discovered roughly $83 million in such abuse and began pushing their intermediaries to crack down on the problem. In 1997, the Inspector General of the Department of Health and Human Services warned HCFA officials to do a better job enforcing their 6-month reimbursement guideline. While HCFA’s plans may have been well-intentioned, its intermediaries’ attempt to enforce the rule was disastrous. For example, the Wall Street Journal reported that UGS, a subsidiary of Blue Cross Blue Shield in Wisconsin and a Medicare intermediary, sent letters to five terminally ill patients which declared that they were not eligible for Medicare hospice and, adding insult to injury, requested these patients to pay $450,000 for the care they received.

Outrage from several hospices and Federal legislators has led to a small change in HCFA’s aggressive crackdown on its 6-month rule. Last week, HCFA’s administrator, Nancy-Ann Min DeParle, wrote to thousands of hospices to explain that there has been a “dis- turbing misperception” about HCFA’s efforts to enforce its 6-month regulation. However, she never specifically declared that reimbursement for care of hospice patients will continue for as long as they receive treatment. She only offered to create a “voluntary” case-by-case review of patients who remain in hospice care longer than 6 months.

Regardless of Administrator DeParle’s change in position, we must clarify the law so that there is no question about HCFA’s responsibility to provide care for the terminally ill. It is the right and moral thing to do. More importantly, it will let hospice patients live out their final days in dignity. I urge my colleagues to cosponsor my bill and I submit the Wall Street Journal article of June 5th to be printed in the RECORD.