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"Uncertain Suffering will have a unique place in medical anthropology, public health scholarship, and the social sciences of health. It involves a layered and deeply philosophical approach to the limits of the role/responsibility of modern American medicine to address the suffering of African ...

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In *Uncertain Suffering*, anthropologist Carolyn Rouse digs beneath the surface of sickle disease care among African Americans to interpret the cultural barriers to [End Page 519] overcoming racial health care disparities in the United States.

Project MUSE - Uncertain Suffering: Racial Health Care

On average, black Americans are sicker and die earlier than white Americans *Uncertain Suffering* provides a richly nuanced examination of what this fact means for health care in the United States through the lens of sickle cell anemia, a disease that primarily affects blacks In a wide ranging analysis that moves from individual patient cases to ...

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Uncertain Suffering leads the reader through a complex set of arguments and evidence to build up a vivid picture of how patients with sickle cell disease struggle in their quest for health care. The reward is insight to the troubling connection between the disjointed weakness of the American health-care system and racial disparities in health and health care.

Uncertain Suffering: Racial Health Care Disparities and

Uncertain Suffering: Racial Health Care Disparities and Sickle Cell Disease, by Carolyn Moxley Rouse. Berkeley, CA: University of California Press, 2009. 328 pp. I am not a sickle cell disease (SCD) patient, but Carolyn Moxley Rouse in *Uncertain Suffering* made me feel as if I were walking in the

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