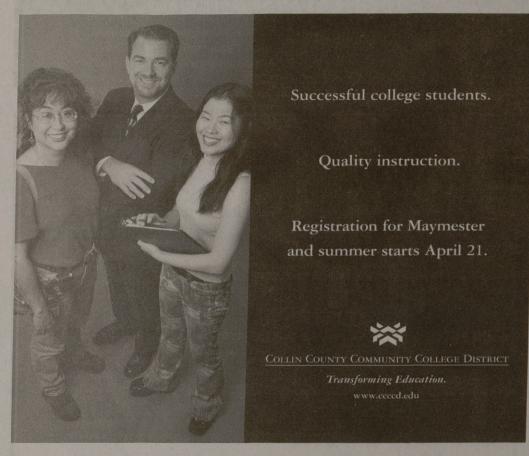
John D Huntley





Wednesday, April 2, 2003

Cancer drug prolongs Miss sickle cell patients' liviour

> By Lindsey Tanner THE ASSOCIATED PRESS

CHICAGO - A drug that reduces the disabling pain of sickle cell anemia can also significantly lower death rates and should be used by many more patients with the blood disorder, researchers say.

The drug, hydroxyurea, is more often associated with cancer and AIDS treatments and most sickle cell patients who need it aren't getting it, said Dr. Duane Bonds of the National Heart, Lung and Blood Institute.

In a government-funded study of adult sickle cell patients co-authored by Bonds, those who used hydroxyurea capsules at least periodically for nine years were 40 percent less likely to die during the study than those who never used the drug.

'Now there is new hope for these patients, who typically die 10 to 15 years earlier than patients with milder cases," said Dr. Claude Lenfant, the institute director.

The study appears in Wednesday's Journal of the American Medical Association. It covered 299 patients, but complete data was only recorded for 233 of them. Complete data were available for 233 of the 299 participants.

Sickle cell anemia, an inherited blood disorder, affects about 72,000 Americans, mostly blacks. Patients have defective hemoglobin, oxygen-carrying blood protein, which causes deformed red blood cells that can clog vessels. This deprives organs and tissues of adequate blood supply, causing severe pain.

There were 13 deaths among the 36 patients who never took hydroxyurea, compared with 21 deaths among the 106 patients who took the drug throughout the nine-year period. The 40 percent difference was calculated using an average of death rates for every three-month period in the study, Bonds said.

Follow-up research shows the benefits extend to prolonging life in moderately to severely affected patients, who make up some 30 percent of sickle cell patients, said Dr. Martin Steinberg of Boston University, the lead researcher. Most are

safe Drug offers promise for sickle cell ar

The drug costs about \$100 monthly, Thact since M possible side effects, including a pole2-mailed the increased risk of leukemia and a decrease ine cell counts, which could make patients pthey would be infections.

Bonds said the potent drug may be und because it's "more work for doctors," refrequent patient visits to adjust dosages and sure complications don't develop.

Patients are generally advised to take sules daily. Study participants took eitherhyurea or dummy pills in the first phase 1992-95. In the follow-up, from 1996 researchers observed results in patients wh tinued, stopped or started taking hydroxyun

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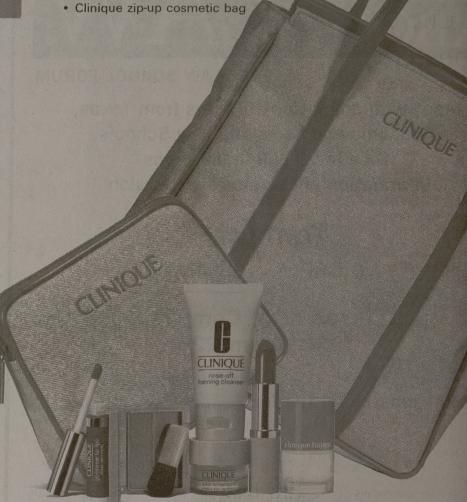
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