

flammatory diseases. Sales breakdown for 2008: Thalomid (23%), Revlimid (59%), Vidaza (9%), Alkeran (4%), and royalties/other (5%). R&D pipeline includes immune modulators, selective cytokine inhibitory drugs, estrogen receptor modulators, and kinase in-

9.3%; Janus Cap'l Mgmt., 9.0%; AXA Fin'l, 4.6% (4/08 Proxy). Chrmn. & CEO: Sol J. Barer. Inc.: DE. Addr.: 86 Morris Ave., Summit, NJ 07901. Tel.: 908-673-9000. Internet: www.celgene.com. cal malignancies. Celgene also plans to ex-

239.7 527.2 432.9 Past Est'd '06-'08 5 Yrs. 45.5% to '12-'14 19.5% 73.0% 117.5% 33.0% 34.0% 14.0% 78.5% 47.5% QUARTERLY SALES (\$ mill.) 275.0 898.9 414.6 1405.8

2841.4

53.9

473.3

Cal-Mar.31 Jun. 30 Sep. 30 Dec. 31 endar 2006 181.8 197.2 244.9 2007 347.9 349.9 293.4 461.0 566.6 586.7 623.5 2237.8 2008 600 625 675 700 2600 2009 3200 2010 750 775 825 850 EARNINGS PER SHARE A Mar.31 Jun. 30 Sep. 30 Dec. 31 endar Year 2006 .04 .03 .05 .06 .18 2007 .14 .13 .09 .18 .54 2008 .32 .31 .34 .38 1.37 .38 .45 .50 .52 1.85 2009 .70 2010 .60 .65 2.60 .65 QUARTERLY DIVIDENDS PAID Cal-Full endar Mar.31 Jun.30 Sep.30 Dec.31 2005 NO CASH DIVIDENDS 2006 2007 **BEING PAID** 2008

2311.0

Past

10 Yrs.

24.4

3084.4

37.9

395.0

Current Assets

Accts Payable Debt Due

Current Liab.

Sales "Cash Flow"

Dividends Book Value

Earnings

2009

ANNUAL RATES

of change (per sh)

We look for Celgene's bottom line to advance 35% in 2009, thanks largely to increased adoption of Revlimid Vidaza. Revlimid sales should rise 30% this year, to \$1.7 billion, driven by an expanding geographic footprint, marketshare gains, and longer treatment durations. Indeed, we look for Revlimid to gain approval in Japan during 2009. Japan is the second-largest oncology market after the U.S. That said, about 95% of the *Rev*limid product mix in the U.S. is for multimyeloma and myelodysplastic syndromes (MDS). Vidaza sales will probably double, reaching \$400 million.

Use of Revlimid to treat a wider range of malignancies should be a key longterm earnings driver. Leveraging the success of the drug as a second-line treatment for multiple myeloma, Celgene has initiated several studies aimed at expanding the existing myeloma label to include newly diagnosed patients. Indeed, we look for regulatory action on that front in both the U.S. and EU this year. There are multiple studies into other unserved indications, such as long-term maintenance therapy in myeloma and other hematologiplore the use of Revlimid as a treatment for solid tumors.

The company has a more potent immunomodulatory drug (IMiD) in the **pipeline.** Celgene recently completed Phase II studies of *Actimid*. This IMiD is an analogue to Relimid that has shown clinical activity in multiple uniaue myeloma and myelofibrosis. Indeed, 29% of patients that had not responded to Revlimid saw improvement with Actimid. The drug has been found to be generally well tolerated by most patients, with manageable adverse events. Further study is necessary, however, and Actimid is not likely to be available before 2012-2014. This treatment represents the next generation in Celgene's hematological franchise. These timely shares offer wide appreciation potential to 2012-2014. Celgene's cancer drugs, particularly Revlimid, offer strong growth prospects for both the near and long terms. Further, the ongoing global recession should have little effect on the company's performance, as these drugs are potentially life-saving. Jerry W. Gray Jr. April 17, 2009

(A) Diluted earnings. Excludes nonrecurring gains (losses): '93, (10¢); '97, (5¢); '98, 14¢; '00, (10¢); '01, 1¢; '02, (\$1.13); '08, (\$4.83). Next earnings report due early May.

(B) In millions, adjusted for stock splits.

(C) As of 12/31/08, had federal and state net operating loss carryforwards of \$63.0 million and tax credit carryforwards of \$65.2 million.

Company's Financial Strength Stock's Price Stability Price Growth Persistence **Earnings Predictability**

65

80

40