PURDUE EMPLOYEES WHO AUTHORED, REVISED OR APPROVED THE TEXT OF PACKAGE INSERTS FOR OXYCONTIN		
Original OxyContin Package Insert Approved December 12, 1995		
NAME	**CURRENT TITLE**	**CURRENT JOB DESCRIPTION**
James Conover	Executive Director, Regulatory Affairs: Resigned May 12, 2000	Former Purdue employee-oversaw regulatory department
Michael Friedman	Executive Vice President, Chief Operating Officer	Chief Operating Officer
Paul Goldenheim	Executive Vice President, Worldwide R&D	Responsible for Purdue's research and development and medical affairs
William Herlihy	Associate General Counsel	Legal counsel responsible for compliance with FDA & DEA law and regulations
Ellen Ingber	Executive Director, Project Management	Leads product development teams involved in clinical studies and product registration issues
Michael Innaurato	Senior Director, Hospital Specialty Division	Oversees the hospital specialty sales force
Robert Kaiko	Vice President, Clinical Research	Oversees clinical development of Purdue's products
Robert Reder	Vice President, Medical Affairs and Worldwide Drug Safety	Oversees Purdue's US medical affairs department and worldwide drug safety and pharmacovigilence function
Lee Ann Storey	Assistant Director, Drug Safety Assurance	Evaluates safety assurance processes to assure compliance with adverse event reporting regulations
Howard Udell	Executive Vice President, General Counsel	Chief Legal Officer

All individuals above can be reached at (203) 588-8000, with the exception of James Conover

PURDUE EMPLOYEES WHO AUTHORED, REVISED OR APPROVED THE TEXT OF PACKAGE INSERTS FOR OXYCONTIN		
OxyContin Package Insert Approved July 18, 2001		
NAME	**CURRENT TITLE**	**CURRENT JOB DESCRIPTION**
Mark Alfonso	Vice President, Marketing	Oversees Purdue's marketing department
Elizabeth Connelly	Manager, Regulatory Affairs	Assists in ensuring that governmental requirements necessary to obtain approval of new products and to continue production of existing products are met
Michael Friedman	Executive Vice President, Chief Operating Officer	Chief Operating Officer
Paul Goldenheim	Executive Vice President, Worldwide R&D	Responsible for Purdue's research and development and medical affairs
William Herlihy	Associate General Counsel	Legal counsel responsible for compliance with FDA & DEA law and regulations
Ellen Ingber	Executive Director, Project Management	Leads product development teams involved in clinical studies and product registration issues
Christopher Prue	Senior Director, U.S. Regulatory Affairs	Ensures that governmental requirements necessary to obtain approval of new products and to continue production of existing products are met
Robert Reder	Vice President, Medical Affairs and Worldwide Drug Safety	Oversees Purdue's US medical affairs department and worldwide drug safety and pharmacovigilence function
Patricia Richards	Associate Medical Director	Plans, designs, directs and executes assigned clinical drug development programs for NDA submissions
Sally Riddle	Sr. Director, Product Management	Establishes, directs, and coordinates marketing activities for all assigned products
Anthony Santopolo	Vice President, US Regulatory Affairs	Purdue's Senior US Regulatory Officer; responsible for ensuring that governmental requirements necessary to obtain approval of new products and to continue production of existing products are met
Howard Udell	Executive Vice President, General Counsel	Chief Legal Officer

Individuals above can be reached at: (203) 588-8000

PURDUE EMPLOYEES WHO AUTHORED, REVISED OR APPROVED THE TEXT OF PACKAGE INSERTS FOR OXYCONTIN		
OxyContin Patient Package Insert Approved January 16, 2002		
NAME	**CURRENT TITLE**	**CURRENT JOB DESCRIPTION**
Mark Alfonso	Vice President, Marketing	Oversees Purdue's marketing department
Elizabeth Connelly	Manager, Regulatory Affairs	Assists in ensuring that governmental requirements necessary to obtain approval of new products and to continue production of existing products are met
Michael Friedman	Executive Vice President, Chief Operating Officer	Chief Operating Officer
Paul Goldenheim	Executive Vice President, Worldwide R&D	Responsible for Purdue's research and development and medical affairs
William Herlihy	Associate General Counsel	Legal counsel responsible for compliance with FDA & DEA law and regulations
Ellen Ingber	Executive Director, Project Management	Leads product development teams involved in clinical studies and product registration issues
Christopher Prue	Senior Director, U.S. Regulatory Affairs	Ensures that governmental requirements necessary to obtain approval of new products and to continue production of existing products are met
Robert Reder	Vice President, Medical Affairs and Worldwide Drug Safety	Oversees Purdue's US medical affairs department and worldwide drug safety and pharmacovigilence function
Patricia Richards	Associate Medical Director	Plans, designs, directs and executes assigned clinical drug development programs for NDA submissions
Sally Riddle	Sr. Director, Product Management	Establishes, directs, and coordinates marketing activities for all assigned products
Anthony Santopolo	Vice President, US Regulatory Affairs	Purdue's Senior US Regulatory Officer; responsible for ensuring that governmental requirements necessary to obtain approval of new products and to continue production of existing products are met
Howard Udell	Executive Vice President, General Counsel	Chief Legal Officer

Individuals above can be reached at: (203) 588-8000