

Exhibit E

**CHRONOLOGY OF SIGNIFICANT ACTIVITIES
IND 54,307 and NDA 21-229**

Application #	Date	Description
IND 54,307	Oct. 15, 1997	Submitted original IND (and Study No. 171)
IND 54,307	Dec. 15, 1997	Submitted Study Nos. 1997092 and 1997095
IND 54,307	Jan. 27, 1998	Submitted Study No. 183
IND 54,307	Jan. 30, 1998	Astra Merck Inc. notified FDA of change of responsibilities for IND 54,307, identifying The Procter & Gamble Company as its agent.
IND 54,307	Feb. 27, 1998	Submitted Study No. 1998003
IND 54,307	Jun. 4, 1998	Submitted Study Nos. 1998005 and 1998006
IND 54,307	Jul. 28, 1998	Submitted Study No. 1998014
IND 54,307	Dec. 14, 1998	Submitted Study No. 1998067
NDA 21-229	Jan. 27, 2000	Submitted original NDA
NDA 21-229	Oct. 20, 2000	Joint FDA Meeting of Gastrointestinal Drugs Advisory Committee and Nonprescription Drugs Advisory Committee
NDA 21-229	Nov. 27, 2000	FDA issued not approvable Action Letter, citing need for additional information on several issues
NDA 21-229	Nov. 31, 2000	Notified FDA of intent to file additional information in relation to NDA 21-229
NDA 21-229	Jan. 30, 2001	Met with FDA to discuss product labeling
NDA 21-229	Jul. 19, 2001	Submitted proposed Actual Use Study 2001007
NDA 21-229	Feb. 12, 2002	Resubmitted NDA in response to deficiencies cited in Nov. 27, 2000 FDA Action Letter with revised product labeling and directions for use, modified indication, and data
NDA 21-229	Jun. 21, 2002	Joint FDA Meeting of Gastrointestinal Drugs Advisory Committee and Nonprescription Drugs Advisory Committee
NDA 21-229	Aug. 8, 2002	Received FDA approvable letter recommending changes in product labeling
NDA 21-229	Oct. 9, 2002	Submitted label comprehension study (Study No. 22103)
NDA 21-229	Dec. 20, 2002	Resubmitted NDA in response to Aug. 8, 2002 FDA letter with revised product labeling
NDA 21-229	Feb. 24, 2003	Submitted updated labeling
NDA 21-229	Jun. 12, 2003	Submitted sample package labeling, plastic overlay for the 42-count package, and unit-dose blister labeling
NDA 21-229	Jun. 19, 2003	Submitted proposed labeling, carton labeling and consumer information leaflet
NDA 21-229	Jun. 20, 2003	FDA issued approval letter for NDA 21-229