

	A	B	C	D	E	F	G	H	I
	AG Trade Name, if any	Brand-Name	Active Ingredient	Dosage Form	NDA #	Dosage Strength	14-Digit Generic Product Identifier	AG - 9 digit NDC # (Labeler Code - Product Code)	AG Labeler/Entity Name
1									
2		Painfree	phenylthicodone	tablet	18444	5 mg	15-27-63-84-91-22-05	00001-*005	Reliapill
3		Painfree	phenylthicodone	tablet	18444	10 mg	15-27-63-84-91-22-10	00001-*010	Reliapill
4		Sumadex	sumacoxib	capsule	19106	80 mg	14-27-32-77-93-81-80	00001-7080	Reliapill
5		Sumadex	sumacoxib	capsule	19106	240 mg	14-27-32-77-93-81-24	00001-7240	Reliapill

	H	J	K	L	M	N
1	AG - 9 digit NDC # (Labeler Code - Product Code)	NDC Date of Launch	NDC Date of Discontinuance, if any	Date of first announcement of AG marketing	Has Company marketed this drug as an ANDA generic? (Yes/No)	QUESTION 6 AG marketed pursuant to settlement? (Yes/No)
2	00001-*005	1/30/2004	1/30/2005	1/18/2004	No	No
3	00001-*010	1/30/2004	1/30/2005	1/18/2004	No	No
4	00001-7080	2/22/2005		2/20/2005	Yes	Yes
5	00001-7240	2/22/2005		2/20/2005	Yes	Yes

	A	B	C	D	E	F	G	H	I	J	K
1	Brand-Name of Drug Subject to ¶ 4 or Drug with AG marketed by Any Company	NDA # of RLD	Active Ingredient	Dosage Form	ANDA #	Dosage Strength	9 digit NDC # (Labeler Code - Product Code)	Labeler/Entity Name	NDC Date of Launch	NDC Date of Discontinuance, if any	180-day exclusivity? (Yes/No)
2	Benetan	17823	lactotannate	tablet	71116	40 mg	00001-2340	Reliapill	7/28/2001		Yes
3	Benetan	17823	lactotannate	tablet	71116	80 mg	00001-2380	Reliapill	7/28/2001		Yes
4	Benetan	17823	lactotannate	tablet	71116	200 mg	00001-2320	Reliapill	1/29/2004		No
5	Benetan	17823	lactotannate	tablet	71116	40 mg	23232-2340	Reliapill of Atlantis	7/29/2001		Yes
6	Benetan	17823	lactotannate	tablet	71116	80 mg	23232-2380	Reliapill of Atlantis	7/29/2001		Yes
7	Marvacid	18998	trichloroprazole	capsule	74555	15 mg	00001-6015	Reliapill	10/9/2003		Yes
8	Marvacid	18998	trichloroprazole	capsule	74555	30 mg	00001-6030	Reliapill	10/9/2003		Yes
9	Virostop XR	57983	dicyclovir	capsule XR	65833	100 mg	00001-1100	Reliapill	6/1/2002		No
10	Virostop XR	57983	dicyclovir	capsule XR	65833	200 mg	00001-1200	Reliapill	6/1/2002		No
11	Virostop XR	57983	dicyclovir	capsule XR	65833	400 mg	00001-1400	Reliapill	6/1/2002		No
12	Virostop XR	57983	dicyclovir	capsule XR	65833	800 mg	00001-1800	Reliapill	12/23/2004		No
13	Virostop XR	57983	dicyclovir	capsule XR	65833	100 mg	23232-1100	Reliapill of Atlantis	6/3/2002		No
14	Virostop XR	57983	dicyclovir	capsule XR	65833	200 mg	23232-1200	Reliapill of Atlantis	6/3/2002		No
15	Virostop XR	57983	dicyclovir	capsule XR	65833	400 mg	23232-1400	Reliapill of Atlantis	6/3/2002		No
16	Sucrastat	17907	acetotryptamide	tablet	70234	2 mg	00001-*232	Reliapill	10/10/2004		No
17	Sucrastat	17907	acetotryptamide	tablet	70234	5 mg	00001-*235	Reliapill	10/10/2004		No
18	Sucrastat	17907	acetotryptamide	tablet	70234	10 mg	00001-*231	Reliapill	10/10/2004		No
19	Hemolite	20554	bicardipine	tablet XR	79666	10 mg	00001-4210	Reliapill	11/20/2005		No
20	Hemolite	20554	bicardipine	tablet XR	79666	20 mg	00001-4220	Reliapill	11/20/2005		No
21	Sumadex	19106	sumacoxib	capsule	77777	80 mg	00001-7080	Reliapill	2/23/2006		No
22	Sumadex	19106	sumacoxib	capsule	77777	240 mg	00001-7240	Reliapill	2/23/2006		No
23	Sumadex	19106	sumacoxib	capsule	77777	80 mg	23232-7080	Reliapill of Atlantis	2/25/2006		No
24	Sumadex	19106	sumacoxib	capsule	77777	240 mg	23232-7240	Reliapill of Atlantis	2/25/2006		No

	G	L	M	N	O	P	Q
	9 digit NDC # (Labeler Code - Product Code)	Date 180-day exclusivity began	Date 180-day exclusivity ended	Name of ANDA- Generic Company #2 During Exclusivity	Name of ANDA- Generic Company #3 During Exclusivity	Name of ANDA- Generic Company #4 During Exclusivity	Enter columns for additional companies here ►►
1							
2	00001-2340	7/28/2001	1/28/2002				
3	00001-2380	7/28/2001	1/28/2002				
4	00001-2320						
5	23232-2340						
6	23232-2380						
7	00001-6015	10/8/2003	4/8/2004	Dependapill			
8	00001-6030	10/8/2003	4/8/2004	Dependapill			
9	00001-1100						
10	00001-1200						
11	00001-1400						
12	00001-1800						
13	23232-1100						
14	23232-1200						
15	23232-1400						
16	00001-*232						
17	00001-*235						
18	00001-*231						
19	00001-4210						
20	00001-4220						
21	00001-7080						
22	00001-7240						
23	23232-7080						
24	23232-7240						

	A	B	C	D	E	F	G	H	I	J	K	L
	Active Ingredient	Dosage Form	ANDA #	Dosage Strength	Date of ANDA Filing	Date of ANDA Approval for Dosage Strength	14-Digit Generic Product Identifier	Patent Number	Paragraph # of Certification	Date of Patent Certification	Paragraph # of Amended Patent Certification	Date of Amended Certification
1												
2	lactotannate	tablet	71116	40 mg	8/27/1997	7/25/2001	43-29-44-12-55-21-40	5,293,001	4	8/27/1997		
3	lactotannate	tablet	71116	80 mg	8/27/1997	7/25/2001	43-29-44-12-55-21-80	5,293,001	4	8/27/1997		
4	lactotannate	tablet	71116	200 mg	8/27/1997	1/28/2004	43-29-44-12-55-21-25	5,293,001	4	8/27/1997		
5	trichloroprazole	capsule	74555	15 mg	4/7/2001	10/8/2003	74-12-49-33-56-82-15	6,000,444	4	4/7/2001		
6	trichloroprazole	capsule	74555	30 mg	4/7/2001	10/8/2003	74-12-49-33-56-82-30	6,000,444	4	4/7/2001		
7	dicyclovir	capsule XR	65833	100 mg	4/9/1999	5/25/2002	26-15-23-72-01-12-10		Not applicable			
8	dicyclovir	capsule XR	65833	200 mg	4/9/1999	5/25/2002	26-15-23-72-01-12-20		Not applicable			
9	dicyclovir	capsule XR	65833	400 mg	4/9/1999	5/25/2002	26-15-23-72-01-12-40		Not applicable			
10	dicyclovir	capsule XR	65833	800 mg	4/9/1999	12/21/2004	26-15-23-72-01-12-80		Not applicable			
11	acetotryptamide	tablet	72234	2 mg	12/29/1998	10/8/2004	65-22-46-87-26-33-02	5,488,293	4	12/29/1998		
12	acetotryptamide	tablet	72234	5 mg	12/29/1998	10/8/2004	65-22-46-87-26-33-05	5,488,293	4	12/29/1998		
13	acetotryptamide	tablet	72234	10 mg	12/29/1998	10/8/2004	65-22-46-87-26-33-10	5,488,293	4	12/29/1998		
14	bicardipine	tablet XR	79666	10 mg	3/5/2002	11/17/2005	38-72-09-14-28-16-10	5,988,290	4	3/5/2002		
15	bicardipine	tablet XR	79666	20 mg	3/5/2002	11/17/2005	38-72-09-14-28-16-20	5,988,290	4	3/5/2002		
16	sumacoxib	capsule	77777	80 mg	8/29/2001	8/23/2005	14-27-32-77-93-81-80	5,900,999	4	8/29/2001		
17	sumacoxib	capsule	77777	240 mg	8/29/2001	8/23/2005	14-27-32-77-93-81-24	5,900,999	4	8/29/2001		

	C	M	N	O	P	Q	R	S	T	U	V
	ANDA #	Patent Number	Paragraph # of Certification	Date of Patent Certification	Paragraph # of Amended Patent Certification	Date of Amended Certification	Patent Number	Paragraph # of Certification	Date of Patent Certification	Paragraph # of Amended Patent Certification	Date of Amended Certification
1											
2	71116	4,486,200	3	8/27/1997			4,576,066	3	8/27/1997		
3	71116	4,486,200	3	8/27/1997			4,576,066	3	8/27/1997		
4	71116	4,486,200	3	8/27/1997			4,576,066	3	8/27/1997		
5	74555	5,892,333	3	4/7/2001							
6	74555	5,892,333	3	4/7/2001							
7	65833										
8	65833										
9	65833										
10	65833										
11	72234	4,644,521	3	12/29/1998			4,151,212	2	12/29/1998		
12	72234	4,644,521	3	12/29/1998			4,151,212	2	12/29/1998		
13	72234	4,644,521	3	12/29/1998			4,151,212	2	12/29/1998		
14	79666	4,600,001	3	3/5/2002							
15	79666	4,600,001	3	3/5/2002							
16	77777	5,833,333	4	8/29/2001			4,655,121	3	8/29/2001		
17	77777	5,833,333	4	8/29/2001			4,655,121	3	8/29/2001		

	C	W	X	Y	Z	AA	AB
1	ANDA #	Patent Number	Paragraph # of Certification	Date of Patent Certification	Paragraph # of Amended Patent Certification	Date of Amended Certification	Enter sets of columns for additional patents here ►►
2	71116						
3	71116						
4	71116	4,701,299	3	8/27/1997	4	12/12/1998	
5	74555						
6	74555						
7	65833						
8	65833						
9	65833						
10	65833						
11	72234						
12	72234						
13	72234						
14	79666						
15	79666						
16	77777						
17	77777						