

	A	B	C	D	E	F	G	H	I
1	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

	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)

	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)

	G	L	M	N	O	P	Q
1	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 ▶▶

	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												

	C	M	N	O	P	Q	R
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▶▶