	А	В	С	D	Е	F	G	Н	
	AG Trade	Brand-Name	Active Ingredient	Dosage	NDA#	Dosage	14-Digit Generic	AG - 9 digit	AG Labeler/Entity
	Name, if			Form		Strength	Product	NDC #	Name
	any						Identifier	(Labeler	
								Code -	
								Product	
1								Code)	

	Н	J	K	L	M	N
	AG - 9 digit	<b>NDC Date</b>	NDC Date of	Date of first	Has Company	<b>QUESTION 6</b>
	NDC #	of	Discontinuance, if	announcement	marketed this	AG marketed
	(Labeler	Launch	any	of AG marketing	drug as an ANDA	pursuant to
	Code -		_		generic?	settlement?
	Product				(Yes/No)	(Yes/No)
1	Code)				,	,

	Α	В	С	D	Е	F	G	Н	I	J	K
	Brand-Name of	NDA#	Active	Dosage	<b>ANDA</b>	Dosage	9 digit	Labeler/Entity Name	NDC Date	NDC Date of	180-day
	Drug Subject to	of RLD	Ingredient	Form	#	Strength	NDC #		of Launch	Discontinuance, if	exclusivity?
	¶ 4 or Drug with						(Labeler			any	(Yes/No)
	AG marketed by						Code -				
	Any Company						Product				
1							Code)				

	G	L	M	N	0	Р	Q
	9 digit	Date 180-day	Date 180-day	Name of ANDA-	Name of ANDA-	Name of	Enter columns
	NDC#	exclusivity	exclusivity	Generic	Generic	ANDA-	for additional
	(Labeler	began	ended	Company #2	Company #3	Generic	companies
	Code -	-		During	During	Company #4	here▶▶
	Product			Exclusivity	Exclusivity	During	
1	Code)					Exclusivity	

	Α	В	С	D	Е	F	G	Н	I	J	K	L
	Active Ingredient	Dosage	ANDA#	Dosage	Date of	Date of	14-Digit	Patent	Paragraph #	Date of Patent	Paragraph #	Date of
		Form		Strength	ANDA	ANDA	Generic	Number	of	Certification	of Amended	Amended
					Filing	Approval	Product		Certification		Patent	Certification
						for	Identifier				Certification	
						Dosage						
1						Strength						

	С	M	N	0	Р	Q	R
	ANDA#	Patent	Paragraph #	Date of	Paragraph #	Date of	Enter sets
		Number	of	Patent	of Amended	Amended	of columns
			Certification	Certification	Patent	Certification	for
					Certification		additional
							patents
1							here▶▶