ALPHA 1-ANTITRYPSIN DEFICIENCY REGISTRY ADVERSE REACTION FORM

Form Completion Instructions:

This form should be completed for each individual where an adverse reaction has occurred from receipt of augmentation therapy. Complete one form per reaction.

Be sure to record the Prolastin Lot Number and the dosage administered.

Refer to question 38a of Form #05A as well as the "Problem" column on Form 10 to gather information to complete this form.

If further information is required, contact the local caregiver.

This form is completed for <u>each</u> date therapy given where at least one adverse reaction has occurred. If no problems have arisen, do not complete form.

1	Date form completed: FIIQOL-F2d (fuzzed)
2.	•
3.	Patient name code: namecode (censored)
4.	Clinical Center: Clinic (censored)
5.	Date of augmentation therapy:Aug Date_f2d (fuzzed) / /
	a. Location of therapy: FII Q05a
	(1)Registry Clinical Center(4)Home
	(2)Local physician's office
	(3)Other (Specify) F11Q05a3(9)Unknown
-6.	Prolastin lot number: FIIQØ6
7.	a. Additional Prolastin lot number used: FIIQ06a
	a. Frequency of therapy:(1)Weekly(2)Monthly(3)Other (specify): Never entered
	Note: Coded by the CCC as the interval (days) between infusions. Code '98' represents "intermittent" or occasional use.
8a.	Type of therapy:(1)Infusion(2)Other (specify): Never entered
b.	When did the reaction occur? FIIQ8B
	(1)During the infusion(4)Other (Specify) F11Q8B4
	(2)Immediately after the infusion(9)Unknown but within 24 hours
	(3)Greater than 24 hours after the infusion
c.	Was hospitalization or emergency room treatment required?: FIIQ8C (1)Yes(2)No
ď.	Was medication and/or contact with physician required?: FUQSD(1)Yes(2)No
e.	Was therapy discontinued permanently?:F.\\Q&E(1)Yes(2)No
f.	Date therapy permanently discontinued: FII08F f2d (fyzzed) //
9.	what frequency did reaction occur? F1108G month day year what frequency did reaction occur? F1108G month day year specify F1108G5 pages following was patient pre-medicated? F1108H specify F1108H1 This form
()	what trequency and reaction see explanation

White/Yellow: Clinical Coordinating Center, Pink: Clinical Center

	A1AD	Adverse	Reaction	#1	1
Patient Registry ID:			Rev. 3	2/9	14
Date of Therapy:/			Page 2	of	2
month day year					

Check each sign/symptom that occurred at the time of this therapy. Use the following codes for "Action Taken":

1=Flow rate decreased 2=Infusion temporarily ceased, then resumed 3=Infusion ceased, not resumed 4=Infusion continued, no change 5=Other (specify in comments)

			REACTIC (check)		ACTION TAKEN (indicate code from above)
i	a.	Anaphylactic Shock:	F11Q09A1		FUQØ9A2
ı	b.	Anxiety:			F11009B2
(c.	Chest Tightness:	FUQ09CI		F11009C2
(d.	Chills:	F11009.D1		FUQØ9D2
(e.	Dizziness/Fainting:	FIIQ09EI_		FILQ09E2
í	f.	Dyspnea/Shortness of Breath:	FILOOPFI		F11009F2
(g.	Emesis:	FILQUAGL		F1100992
ı	'n.	Fever:			FILQ109H2
i		Flushing:	F1100911		FUQ09I2
j		Headache:	FILQUAJI		F11009J2
ļ	ζ.	Hives:	F11009K1		F11009 K-2
l	•	Hypotension:	FUQ09L1		FUQ09L2
r	n.	Muscle Cramps:	FIIQ09MI		F11Q09M2
r	١.	Nausea:	FIIQO9NI		FILQO9N2
C).	Pain:	F1100901		FUQ0902
Ç) .	Rash:	FIIQØ9PI_		FUQ 09P2
	٦.	Tachycardia:			F1100992
r		Wheezing:			FUQUIR2
S	;.	Other			FUQ0952
		(Specify): <u>F1\00953</u>	(see next	page)	
C	Cor	nments: COMMENTS			
		COMMENTS		_	
_					
_	-				
				••	
F	orr	m Completed By (Name): <u>Y)C\</u>	ier entered		
P	'nv	sician Signature: <u>Nevev</u> e	ntered		

White/Yellow: Clinical Coordinating Center, Pink: Clinical Center

Notes to Form 11, Question 9

Based on the value in F11Q09S3 ["Other(Specify)"] the following variables were also coded into the dataset.

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	REACTION (check)	ACTION TAKEN (indicate code from above)				
9. (con	9. (continued)					
t.	Asthma attack <u>F11Q09T1</u>	F11Q09T2				
u.	Arthralgia, body ache <u>F11Q09U1</u>	F11Q09U2				
v.	Flu-like symptoms <u>F11Q09V1</u>	F11Q09V2				
w.	Itching <u>F11Q09W1</u>	F11Q09W2				
x.	Cyanosis <u>F11Q09X1</u>	F11Q09X2				
y.	Indigestion/diarrhea <u>F11Q09Y1</u>	F11Q09Y2				
Z.	Warmth/sweating <u>F11Q09Z1</u>	F11Q09Z2				
aa.	Tremors-shaking <u>F11Q9AA1</u>	F11Q9AA2				
ab.	Edema-fluid retention F11Q9AB1	F11Q9AB2				
ac.	Weakness-malaise <u>F11Q9AC1</u>	F11Q9AC2				
	miscellaneous <u>F11Q9AD1</u>	F11Q9AD2				

Variable: SEVERITY

This is an overall severity score:

(1) Severe	Presence of one or more of the following: anaphylactic shock, dyspnea, shortness of breath, wheezing, hypotension. Also included were asthma attack and cyanosis, both subject to physician review.
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(2) Moderate Absence of any "severe" symptoms, and presence of one or more of the following: chest tightness, chills, dizziness/fainting, emesis, fever, headache, hives, rash, tachycardia.

(3) Mild Absence of any "severe" or "moderate" symptoms.

Notes on Coding:

Question 5a. Location of Therapy.

The codes shown on the form are those used to enter data. Codes appearing on previous versions of the forms differed from these.

Question 7a. Frequency of Therapy:

The interval in days between infusions is entered, e.g., 7, 14, 30. Code "98" is used to represent "intermittent" or "occasional" use.

Question 9a-s.

For each question, there are two variables, coded to capture information on whether the reaction occurred (1=yes, 0=blank), and what the action taken was (using codes indicated on the form, with 9=unknown).

9a. Anaphylactic shock:

F11Q09A1 - Anaphylactic shock, reaction

F11Q09A2 - Anaphylactic shock, action

9s. Other

F11Q09S1 - Other, reaction F11Q09S2 - Other, action

Additional variables coded at the CCC:

Additional Signs and Symptoms:

The following additional variables were created to code other specific adverse reaction signs and symptoms not included as specific questions on the original form:

	Variable	<u>Names</u>
Adverse Reaction	Reaction	<u>Action</u>
Asthma attack	F11Q09T1	F11Q09T2
Arthralgia, body ache	F11Q09U1	F11Q09U2
Flu-like symptoms	F11Q09V1	F11Q09V2
Itching	F11Q09W1	F11Q09W2
Cyanosis	F11Q09X1	F11Q09X2
Indigestion-diarrhea	F11Q09Y1	F11Q09Y2
Warmth-sweating	F11Q09Z1	F11Q09Z2
Tremors-shaking	F11Q9AA1	F11Q9AA2
Edema-fluid retention	F11Q9AB1	F11Q9AB2
Weakness-malaise	F11Q9AC1	F11Q9AC2
Miscellaneous	F11Q9AD1	F11Q9AD2

Additional variables coded at the CCC (continued):

Severity of Reaction was coded at the CCC using the following algorithm:

SEVERITY = 1 Severe -

Defined as the presence of one or more of the

following "severe" symptoms: dyspnea, shortness of

breath, wheezing, hypotension

2 Moderate - Absence of any "severe" symptom, and presence of

one or more "moderate" symptoms

3. Mild -

Absence of any "severe" or "moderate" symptoms

Classification of symptoms in Question 9a:

Question a. Anaphylactic shock b. Anxiety c. Chest tightness d. Chills e. Dizziness/fainting f. Dyspnea/SOB g. Emesis h. Fever i. Flushing j. Headache k. Hives l. Hypotension m. Muscle cramps n. Nausea o. Pain p. Rash q. Tachycardia r. Wheezing t. Asthma attack u. Arthralgia, body ache v. Flu-like symptoms w. Itching x. Cyanosis y. Indigestion/diarrhea z. Warmth/sweating aa. Tremors-shaking ab. Edema-fluid retention ac. Weakness-malaise	Severity Severe - none in database Mild Moderate Moderate Moderate Severe Moderate Mild Moderate Moderate Severe Mild Mild Mild Mild Moderate Severe Severe Mild Mild Moderate Severe Mild Mild Moderate Severe Severe Mild Mild Mild Mild Mild Mild Mild Mild
ad. Miscellaneous	

Other symptoms classified as "mild": "Flaky spells", "Mild depression", "Cold sensation", "Nasal congestion", "Gas", "Numbness around mouth".

These symptoms were classified as indicated, subject to physician review

Additional variables coded at the CCC (continued):

<u>Variable</u>	Description	
F11Q8G	With what frequency did the reaction(s) occur?	
	Codes used:	
	Single episode two or three times greater than 3 times with every infusion Other Unknown	
F11Q8G5	If other frequency, specify	
	Format: character \$25	
F11Q8H	Was patient pre-medicated?	
	Codes used:	
	1 Yes 2 No	
F11Q8H1	Was patient premedicated? Specify:	
	Format: character \$25	
BADLOT	Was the adverse reaction associated with a known bad lot of Prolastin?	
	Codes used	
	1 Yes 0 No	