SECTION 1: CONTACT INFORMATION Principal Investigator (Individual who derived the cell lin	ne):				
Submitting Institution (where cell line was derived):					
Laboratory Contact Information Name:					
Phone:					
Email:					
Member of NINDS GO iPSC ConsortiumYI	ESNO				
SECTION 2: CELL LINE IDENTIFICATION AND D					
iPS Cell Line Name/Identifier:					
SUBJECT INFORMATION					
If subject banked at NINDS Repository (fibroblast, blood). Submitter Name	, please list:				
Submitter ID					
Repository Number ND					
Diagnosis:ALSHDPD	FTDControl				
If the biopsy, fibroblast or blood is from a subject not bank contact ninds@coriell.org prior to submission.	xed at the NINDS Repository, please				
SECTION 3: INVENTORY INFORMATION					
Number of Vials:					
Label on Vial:					
Passage Number at Freeze:					

SECTION 4: REPROGRAMMING INFORMATION

Starting Material Cell Type	Fibroblast	Other:	
Reprogramming MethodRetroviral vector	Lentiviral vector	Sendai virus	vector
Episomal	mRNA	Other:	
Reprogramming Factors			
OCT4 SOX2	KLF4	cMYC NAM	NOGLIN28
Date of Establishment:			
Has the original cell line and/or il	PS line been describ	ed in a publication	? Yes No
If yes, please list reference(s) and	PubMed PMID:		
SECTION 5: CULTURING INFO	ORMATION		
Is this line grown on a feeder layer	er?	Yes	No
Source of feeder cells:			
Is this line grown on substrate-co	ated dishes?	Yes	No
If yes, please specify:			
Growth Media (please list base me by commercial name and supplier)	edia, serum and addit		-
Concentration of FGF:			
Special Instructions for Recovery			
Do you use ROCK Inhibitor in re			
Concentration of ROCK Inhibito	r:	_	

Passaging M	Iethod (C	choose one):									
Trypsin	C	ollagenase	D	ispase		Tryp	1E		EZ To	ol	
_ Versene,	_C	Other:									
Passaging fi5 days				days		Othe	r:				
					•						
Split Ratio (1	:5		1:6		_	_Other	::	
Do you use	ROCK In	hibitor in p	assage n	nedia?		Yes		_No			
	If Yes,	what is the c	concentra	tion of	ROCK	Inhibito	or?				
Freeze Med	ia:										
Do you use	ROCK In	hibitor in f	reeze me	edia?		Yes		_No			
	If Yes,	what is the c	concentra	ition of l	ROCK	Inhibito	or?				
SECTION 6	6: CHAR	ACTERIZA	TION (F UND	TEFE	RENTL	ATED	CEL	L LIN	E	
	4	3	1	90	31	לז					
	SSEA-4	SSEA-3	SSEA-1	1-6	FRA 1-8	NANOG	OCT4	SOX2	KLF4	MYC	Other
	SSI	SSI	SSI	TRA 1-60	TRA 1-81	NA	Ŏ	Š	K	Σ	Õ
Immuno- staining											
RT-PCR											
	<u> </u>				<u> </u>						
If other, ple	ase specif	iy:									

SECTION 7: CHARACTERIZATION OF DIFFERENTIATION POTENTIAL

	Teratoma Formation Performed at
	Pluritest (name of facility)
	Score card
	Other:
_	In Vitro Differentiation; please identify method (check all that apply):
	Embryoid body formation
	Neural differentiation (Retinoic acid)
	Neural differentiation (EZ Sphere method)
	Dopaminergic neurons (co-culture with PA6 cells)
	Cardiomyocyte differentiation
	Hematopoietic differentiation
	Adipocyte differentiation
	Endodermal differentiation
	Pancreatic differentiation
	BMP4-induced differentiation
	Other:
	ion of the cell line was assessed by mmunostaining RT-PCR Both Other
SECTION (3: ADDITIONAL CHARACTERIZATION
ECTION	S. ADDITIONAL CHARACTERIZATION
	been karyotyped? ssage number:)No Please provide the ISCN summary.

Has this line been characterized for a specific mutation (ex. point mutation, CAG repeat)?

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Yes	No	If yes, please describe the results:
Has this line l	been genotyped (e No	ex. Affymetrix 6.0 GeneChip)? If yes, please describe the platform and provide link to data:
	110	if yes, prease desertee the platform and provide him to data.
Has this line l	had other charact	erization (transcriptome, epigenome etc.)
Yes	No	If yes, please describe the platform and provide link to data:
Was this iPS(T line tested for g	enomic integration of reprogramming vectors?
_Yes	_No	If yes, please describe the method used to test for integration:
Was there gen	nomic integration No	of reprogramming vectors in this iPSC line? If yes, what was the copy number of any integrated vector(s):
Please provid	e any other releva	ant information regarding this cell line: