

BANCO NACIONAL DE LÍNEAS CELULARES (TRONCALES)

National Bank of Stem Cell Lines

IMPRESO DE SOLICITUD DE DEPÓSITO DE UNA LÍNEA iPS

Application Form to Deposit an iPS cell line of human origin

Documentos que se acompañan:

Attached documents:

☒ Copia de la autorización de derivación de la línea celular, junto con informe del Comité Ético del centro de procedencia.

A copy of the authorization for the derivation of the cell line, with the corresponding ethics committee approval

☒ Copia de cualquier publicación científica relacionada con la derivación y/o caracterización de la línea.

A copy of any relevant published scientific papers related to the derivation and/or characterization of the cell line

☒ C. V. del investigador principal (una página; formato libre).

A one page CV for the Principal Investigator

☒ Otros (especificar). Anexos 1-6

Others (specify) Annexes 1-6

SECCIÓN 1

Section 1

Información General

General Information

Nombre de la línea: SP02#1

Name of the line: SP02#1

Investigador principal: Angel Raya, Eduardo Tolosa

Principal Investigator: Angel Raya, Eduardo Tolosa

Tipo de célula de la que se obtiene la línea: Queratinocitos epidérmicos

Cell type origin of the cell line: Epidermal keratinocytes

¿El sujeto fuente tiene alguna patología?

Has the donor any pathological condition?

NO ☐

No

SÍ ☒

Yes

(especificar) Enfermedad de Parkinson
(specify) *Parkinson's Disease*

¿La patología es de origen genético?

Is the pathological condition of genetic origin?

NO ☒

No

SÍ ☐

Yes

(especificar)
(specify)

Identificación genética de la línea celular. Método y resultado

Genetic identity of the cell line. Method and result

Cariotipo/Karyotype (ver Anexo 2, see Annex 2)

Euploide/Euploid ☒

Anormal/Atypical ☐

(especificar/specify) 46, XY

SECCIÓN 2

Section 2

Datos del Depositante

Applicant Details

Investigador Principal: <i>Principal Investigator:</i> Angel Raya	Dirección Postal: <i>Postal address:</i> Baldiri Reixac 15-21, 08028 Barcelona
Centro de Trabajo: <i>Institution:</i> Institut de Bioenginyeria de Catalunya (IBEC)	Teléfono (phone): 93 402 0537 Fax: 93 402 0183 E-mail: araya@ibecbarcelona.eu

Investigador Principal: <i>Principal Investigator:</i> Eduardo Tolosa	Dirección Postal: <i>Postal address:</i>
Centro de Trabajo: <i>Institution:</i>	Teléfono (phone): Fax: E-mail:

SECCIÓN 3

Section 3

Datos de la Línea Celular

Details of Cell Line

Tejido de origen y localización anatómica de la muestra biológica <i>Tissue of origin and anatomic location of the biological sample</i> Biopsia de piel del brazo Skin biopsy from the arm	
Muestra biológica <i>Biological sample</i> <div>Fresco <input checked="" type="checkbox"/> Crioconservado <input type="checkbox"/> <i>Fresh</i> <i>Cryopreserved</i></div>	
Fecha de la donación del muestra biológica <i>Date of donation of the biological sample</i> July 04, 2011	Fecha del uso o descongelación <i>(si congelado)</i> <i>Date used or thawed (if frozen)</i> July 04, 2011

Origen del soporte celular o acelular utilizado para la derivación, así como de los componentes de los medios de cultivo (si se describen en publicación, indicar además referencia)

Origin of the cellular or cellular free support used in derivation in addition to the components of the culture mediums (if they are described in a publication, please indicate the reference).

Raya A et al. Nature protocols 2010; 5(4): 647-60

Sánchez-Danés A et al. EMBO Molecular Medicine 2012

Mantenimiento de la línea: Line maintenance en HFF / on HFF

Ratio de pase: Passage ratio 1:2- 1:4

Método de pase: Passage method mecánico / mechanical

Xenobióticos

Xenobiotics

si

Yes

no

No

Descripción de las características morfológicas de la línea en cultivo

(forma y tamaño colonias; forma y tamaño células; ratio núcleo/citoplasma; otros)

Description of the morphological characteristics of the line in culture (form and size of the colonies; form and size of the cells; nucleus/cytoplasm ratio; others)

Colonias grandes poligonales, ligeramente aplanadas, de un tamaño entre 1- 3 mm de diámetro de diversas formas con bordes lisos. Células de tamaño uniforme y una elevada relación núcleo/citoplasma.

Large and flat polygonal colonies of 1-3 mm of diameter, with well-defined edges, composed of uniformly sized cells with high nucleus/cytoplasm ratio.

Controles microbiológicos realizados (indicar detalladamente)

Microbiological controls carried out (indicate in detail)

Bacteriología

(Bacteriology)

Micología

(Mycology)

Micoplasma: PCR negativa

(Mycoplasma: by PCR) negative

Marcadores: ver Anexo 1*Markers: See Annex 1*

	Método (ARN/proteínas)	nº pase	resultado	comentarios
	<i>Method</i> <i>(RNA/proteins)</i>	<i>Passage n.</i>	<i>results</i>	<i>comments</i>
Oct 4	inmunofluorescencia	11	+	
Nanog	inmunofluorescencia	11	+	
Rex 1	-			
Sox 2	inmunofluorescencia	11	+	
SSEA3	inmunofluorescencia	11	+	
SSEA4	inmunofluorescencia	11	+	
TRA-1-60	inmunofluorescencia	11	+	
TRA-1-81	inmunofluorescencia	11	+	
Telomerasa	-			
Fosfatasa Alc.	Actividad	13	+	
Otros / Others				

Capacidad de diferenciación*Differentiation capacity*

	Ectodermo/ Ectoderm			Endodermo/Endoderm			Mesodermo/ Mesoderm		
	marcador	pase	resultado	marcador	pase	resultado	marcador	pase	resultado
	<i>marker</i>	<i>passage</i>	<i>result</i>	<i>marker</i>	<i>passage</i>	<i>result</i>	<i>marker</i>	<i>passage</i>	<i>result</i>
In Vitro	TUJ1	13	+	AFP /FOXA2	13	+ / +	SMA	13	+/+
Annexo 3									
<i>In vitro</i>	TUJ1	13	+	AFP /FOXA2	13	+ / +	SMA	13	+/+
<i>Annex 3</i>									
In vivo/ in vivo	(ver Anexo 4)			Método: formación de teratomas en ratones SCID			Resultado:+		
<i>pase/passage</i>				<i>Method: teratoma formation in SCID mice</i>			<i>Result:+</i>		

OPCIONAL/OPTIONAL:**Reprogramación del perfil de expresión génica (ver Anexo 5)***Reprogramming of gene expression profile (see Annex 5)***Reprogramación del perfil de metilación del ADN (ver Anexo 5)***Reprogramming of DNA methylation profile (see Annex 5)***Longitud telomérica***Telomere length*

<p>Descripción de las características de diferenciación <i>in vitro</i> <i>Description of the differentiation characteristics in vitro</i></p> <p>Mesodermo: cultivo de cuerpos embrioides en medio de cultivo suplementado con ácido ascórbico. Endodermo: cultivo de cuerpos embrioides. Ectodermo: cultivo de cuerpos embrioides en medio N2/B27. (ver Anexo 3) <i>Mesoderm: Embryoid bodies cultured in culture medium supplemented with ascorbic acid. Endoderm: Embryoid bodies culture. Ectoderm: Embryoid bodies culture in N2/B27 medium (see Annex 3)</i></p>
<p>Datos de la determinación de pluripotencialidad <i>in vivo</i> o formación de teratomas <i>Data of the pluripotentiality determination in vivo or teratoma formation</i></p> <p>Inyección intratesticular en ratones SCID de clumps de células indiferenciadas y tras aproximadamente 8 semanas, análisis de los teratomas producidos mediante técnicas de inmunohistoquímica para ectodermo, mesodermo y endodermo (Ver Anexo 4) <i>Clumps of undifferentiated cells were injected into the testis of SCID mice. Around 8 weeks later teratomas were analyzed by immunohistochemistry for ectoderm, endoderm and mesoderm (see Annex 4)</i></p>
<p>Datos de la tipificación HLA <i>HLA typification data</i></p>
<p>Integración de los transgenes de reprogramación: Southern blot (Ver Anexo 6) <i>Integration of reprogramming transgenes: Southern blot (See Annex 6)</i></p>
<p>Silenciamiento de los transgenes de reprogramación: RT-PCR o Q-RT-PCR (Ver Anexo 6) Positivo <i>Silencing of reprogramming transgenes: RT-PCR o Q-RT-PCR (See Annex 6) Positive</i></p>
<p>Mantenimiento a largo plazo en cultivo: > 20 pases <i>Long-term maintenance in culture:>20 passages</i></p>
<p>Pase en el momento del registro: pase 6 <i>Passage at the time of the recording: passage 6</i></p>

<p>¿Ha sido la línea modificada genéticamente? <i>Has the line been genetically modified?</i></p> <p>Sí Yes <input type="checkbox"/> No No <input checked="" type="checkbox"/></p> <p>Comentarios/ Comments:</p>	<p>¿Se llevó a cabo un análisis clonal? <i>Has a clonal analysis been carried out?</i></p> <p>Sí/ Yes <input type="checkbox"/> No <input checked="" type="checkbox"/> Resultado / Result</p>
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Otras observaciones o información relevantes (a juicio del Investigador Principal):
Other observations or relevant information (to the discretion of the Principal Investigator):

Otras observaciones o información relevantes (a rellenar por el BNLC):
Other comments or relevant information (to be completed by BNLC)

Seguimiento de la línea (a rellenar por el BNLC):
Follow up of the line (to be completed by BNLC)

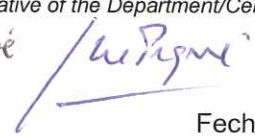


SECCIÓN 4

Declaración

Confirmo que la información contenida en estos impresos es cierta y asumo total responsabilidad sobre la misma.

I confirm that the information contained in this form is true and I assume total responsibility for it.

Firma en Representación del Centro / Signature in Representation of the Centre (Representante legal del Departamento/Centro) (Legal Representative of the Department/Centre) Josep A. Planell i Estany  Fecha/ Date: 21/05/2012	Firma del Investigador Principal Signature of the Principal Investigator Angel Raya  Fecha /Date 21/05/2012
Nombre y Cargo de la Persona Representante del Centro: Name and Position of the Person Representing the Centre: Director	
Dirección Postal: Postal Address: Institut de Bioenginyeria de Catalunya (IBEC) Baldiri Reixac 10-12 08028 Barcelona	Teléfono /Telephone: +34 934 039 706 Fax: +34 934 039 702 E-mail: japlanell@ibecbarcelona.eu

Firma en Representación del Centro / Signature in Representation of the Centre (Representante legal del Departamento/Centro) (Legal Representative of the Department/Centre) Josep M. Piqué   Fecha/ Date: 21/05/2012	Firma del Investigador Principal Signature of the Principal Investigator Eduardo Tolosa  Fecha /Date 21/05/2012
Nombre y Cargo de la Persona Representante del Centro: Name and Position of the Person Representing the Centre: Josep Maria Piqué, Director	
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Resolució per la qual s'autoritza al Servei de Neurologia del Hospital Clínic de Barcelona, per a realitzar activitats de recerca en relació amb el projecte *Generación de un modelo neuronal dopaminérgico a partir de células madre pluripotentes inducidas de pacientes con enfermedad de Parkinson asociada a mutaciones en el gen LRRK2*.

I. Antecedents.

El Servei de Neurologia del Hospital Clínic de Barcelona ha presentat a la Conselleria de Salut, per a la seva autorització, el projecte: *Generación de un modelo neuronal dopaminérgico a partir de células madre pluripotentes inducidas de pacientes con enfermedad de Parkinson asociada a mutaciones en el gen LRRK2*, del qual és investigador principal el Dr. Eduardo Tolosa Sarró.

II. Normativa d'aplicació.

- Llei 14//2006, de 26 de maig, sobre tècniques de reproducció humana assistida.
- Reial Decret 2132/2004, de 29 d'octubre, estableix els requisits i procediments per a sol·licitar el desenvolupament de projectes de recerca amb cèl·lules troncales obtingudes de preembrions sobrants.
- Llei 14/2007, de 3 de juliol, de recerca biomèdica.

III. Consideracions jurídiques.

Primera: El projecte d'investigació presentat s'acompanya de la documentació acreditativa de les condicions establertes a l'article 4 del Reial Decret 2132/2004, de 29 d'octubre precitat.

Segona: En data 22 de setembre de 2010, el projecte ha estat avaluat favorablement pel Comitè Ètic d'Investigació Clínica del Centre de Medicina Regenerativa de Barcelona.

Tercera: La Comissió de Seguiment i Control de la Donació i Utilització de Cèl·lules i Teixits Humans, en sessió de 11 de febrer de 2011, ha informat favorablement el projecte. Els informes d'aquesta Comissió són preceptius i vinculants de conformitat amb l'article 3 del Reial Decret 2132/2004, de 29 d'octubre, precitat.

D'acord amb lo exposat, en exercici de les competències que tinc atribuïdes,

RESOLC:

Autoritzar al Servei de Neurologia del Hospital Clínic de Barcelona Fundació per a realitzar activitats de recerca en relació amb el projecte *Generación de un modelo neuronal dopaminérgico a partir de células madre pluripotentes inducidas de pacientes con enfermedad de Parkinson asociada a mutaciones en el gen LRRK2*, amb subjecció a les condicions acreditades en l'expedient.



Boi Ruiz García
Conseller

Barcelona, 18 d'abril de 2011

D. Ramon Gomis de Barbarà, Director de Investigación del
Hospital Clínic de Barcelona, y Presidente del CEIC

CERTIFICA:

Que el Comité de Investigación del Hospital Clínic, en la sesión
celebrada en el día de hoy, ha analizado el proyecto de investigación
titulado:

*Generation of induced pluripotent stem cells (iPS)-derived dopaminergic
neurons from patients with familial and sporadic Parkinson's disease.*

cuyo investigador principal es el Dr. **Tolosa, Eduard**

del Servicio de **Neurología**

entendiendo que dicho estudio se incluye en una de las líneas de
investigación biomédica acreditadas en este centro, cumpliendo
los requisitos metodológicos necesarios, y que es viable en todos
sus términos, por lo que lo ha considerado adecuado y ha decidido
su aprobación.

Lo que firmo en Barcelona, a 27/11/2008



CLÍNIC
BARCELONA
Hospital Universitari
COMITÉ ÈTIC
INVESTIGACIÓ CLÍNICA

Registro: 2008 / 4625

Dña. Begoña Gómez Pérez , del Servicio de Farmacia del Hospital
Clínic de Barcelona y Secretaria del Comité Ético de Investigación
Clínica (CEIC)

CERTIFICA:


Que el Comité Ético de Investigación Clínica, según consta en el acta
de la reunión celebrada en el día de hoy, ha analizado el proyecto de
investigación titulado:

*Generation of induced pluripotent stem cells (iPS)-derived dopaminergic
neurons from patients with familial and sporadic Parkinson's disease.*

cuyo investigador principal es el Dr. **Tolosa, Eduard**
del Servicio de **Neurología**

entendiendo que dicho estudio se ajusta a las normas éticas esenciales
y criterios deontológicos que rigen en este Centro, y, por tanto, ha
decidido su aprobación.

Lo que firmo en Barcelona, a 27/11/2008



CLÍNIC
BARCELONA
Hospital Universitari
COMITÉ ÈTIC
INVESTIGACIÓ CLÍNICA

Registro: 2008 / 4625

Disease-specific phenotypes in dopamine neurons from human iPS-based models of genetic and sporadic Parkinson's disease

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Keywords: autophagy; disease modeling; LRRK2 mutation; neurodegeneration; pluripotent stem cells

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Accepted January 10, 2012

Induced pluripotent stem cells (iPSC) offer an unprecedented opportunity to model human disease in relevant cell types, but it is unclear whether they could successfully model age-related diseases such as Parkinson's disease (PD). Here, we generated iPSC lines from seven patients with idiopathic PD (ID-PD), four patients with familial PD associated to the G2019S mutation in the *Leucine-Rich Repeat Kinase 2* (LRRK2) gene (LRRK2-PD) and four age- and sex-matched healthy individuals (Ctrl). Over long-time culture, dopaminergic neurons (DAn) differentiated from either ID-PD- or LRRK2-PD-iPSC showed morphological alterations, including reduced numbers of neurites and neurite arborization, as well as accumulation of autophagic vacuoles, which were not evident in DAn differentiated from Ctrl-iPSC. Further induction of autophagy and/or inhibition of lysosomal proteolysis greatly exacerbated the DAn morphological alterations, indicating autophagic compromise in DAn from ID-PD- and LRRK2-PD-iPSC, which we demonstrate occurs at the level of autophagosome clearance. Our study provides an iPSC-based *in vitro* model that captures the patients' genetic complexity and allows investigation of the pathogenesis of both sporadic and familial PD cases in a disease-relevant cell type.

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[†]These authors contributed equally to this work.

INTRODUCTION

Parkinson's disease (PD) is a common and devastating neurodegenerative disorder characterized by motor clinical manifestations, although non-motor features are also important in later stages of the disease (Lees et al, 2009; Obeso et al, 2010; Schapira & Tolosa, 2010). Movement alterations in PD stem from the selective loss of dopaminergic neurons (DAn) from the pars compacta of the substantia nigra, a specific subtype of neurons patterned as ventral midbrain DAn (vmDAn). Neuronal loss is typically progressive and accompanied by α -synuclein (SNCA)-containing intraneuronal inclusions known as Lewy bodies and Lewy neurites. While the majority of PD cases are sporadic, likely resulting from complex interactions among gene susceptibility and environmental factors, around 10% of patients present monogenic forms of the disease (Lesage & Brice, 2009). Pathogenic mutations in the *Leucine-Rich Repeat Kinase 2* (*LRRK2*) and *SNCA* genes have been identified and associated to PD, while mutations in four genes (*Parkin*, *DJ-1*, *PINK1* and *ATP13A2*) cause early onset parkinsonism (Lees et al, 2009). In particular, *LRRK2* mutations cause an autosomal dominant, late-onset familial PD, whose clinical and pathological features are indistinguishable from those of the common, sporadic form of PD (Paisan-Ruiz et al, 2004; Zimprich et al, 2004). While over 50 variants have been identified throughout the different *LRRK2* domains in PD patients, the mutation *G2019S* has been regarded as the most common cause of dominant familial PD and accounts for up to 2% of sporadic PD cases (Cookson, 2010).

The pathogenic mechanisms that lead to neurodegeneration in PD are not well understood, since current experimental PD models do not recapitulate key neuropathological features of the disease (Dawson et al, 2010). In particular, the special susceptibility of DAn to neurodegeneration and the progressive nature of this process in PD, together with the presence of Lewy bodies, have proven especially difficult to model in animal models of PD. The advent of induced pluripotent stem cell (iPSC) technology (Takahashi & Yamanaka, 2006) has made it possible to reprogram human somatic cells to pluripotency (Park et al, 2008b; Takahashi et al, 2007; Yu et al, 2007), thus enabling the generation of disease-specific iPSC (Lee & Studer, 2010). In this way, disease-relevant cell types have been generated from iPSC representing a variety of early onset diseases, and shown to display specific features of the disease *in vitro* (Carvajal-Vergara et al, 2010; Ebert et al, 2009; Ku et al, 2010; Lee et al, 2009; Marchetto et al, 2010; Moretti et al, 2010; Rashid et al, 2010; Raya et al, 2009; Zhang et al, 2011). Disease-specific iPSC have also been generated from sporadic cases of PD, but DAn differentiated from these cells did not show any conspicuous phenotype (Soldner et al, 2009), prompting uncertainty as to whether iPSC technology might be useful to model diseases of non-monogenic, complex etiology and/or late onset. Here, we generated iPSC lines from both idiopathic PD (ID-PD) and familial PD associated to the *G2019S* mutation in the *LRRK2* gene (the most frequent cause of familial PD, Cookson, 2010), as well as from age- and sex-matched healthy individuals. We show that, over long-time culture, DAn

differentiated from PD-iPSC display specific PD pathology related phenotypes and, thus, provide proof-of-concept for iPSC-based cellular models that capture PD patients' genetic complexity and allow investigation of the PD pathogenesis in disease-relevant cell types.

RESULTS

Generation of PD-specific iPSC cell lines

We recruited a total of 15 individuals into this study: 7 patients diagnosed with ID-PD, with no family history of disease and no known PD-related mutations; 4 unrelated patients diagnosed with familial PD carrying the *G2019S* mutation in the *LRRK2* gene (*LRRK2*-PD); and 4 healthy individuals with no history of neurological disease (Ctrl). Individuals in each group were also selected so that no significant bias in gender or age occurred (Table 1). Primary cultures of epidermal keratinocytes and dermal fibroblasts were established from all individuals and were used for reprogramming at passages 2–4. We used retroviral delivery of OCT4, KLF4 and SOX2 to generate 2–6 independent iPSC lines per individual, totalling 50 iPSC lines. Of those, 2 lines per patient were thoroughly characterized and shown to be fully reprogrammed to pluripotency, as judged by colony morphology and growth dynamics, sustained long-term passaging (>20 passages), karyotype stability, alkaline phosphatase (AP) staining, expression of pluripotency associated transcription factors (OCT4, SOX2, NANOG, CRIPTO and REX1) and surface markers (SSEA3, SSEA4, TRA1-60 and TRA1-81), silencing of retroviral transgenes, demethylation of *OCT4* and *NANOG* promoters, *in vitro* pluripotent differentiation ability and generation of teratomas comprising derivatives of the three main embryo germ layers (Table 1 and Fig 1A–N).

The efficiency of iPSC generation varied among different individuals, but did not depend on the presence or type of disease, nor on the age of donors. Whereas, most of the iPSC lines analyzed met our criteria for *bona fide* pluripotent stem cells, those that failed to silence the reprogramming transgenes, did not differentiate appropriately *in vitro*, or presented karyotype alterations were identified and excluded from further studies (Table 1). Overall, iPSC generated from PD patients or from healthy individuals were indistinguishable in all tests performed, with the exception that *LRRK2*-PD iPSC lines carried the *LRRK2*^{*G2019S*} mutation (Fig 1O).

Generation of PD-specific vmDA neurons

For the directed differentiation of iPSC towards vmDA neurons (the cell type most affected in PD), we used a 30-day protocol recently implemented in our laboratory that relies on lentiviral-mediated forced expression of the ventral midbrain determinant LMX1A, together with DA neuron patterning factors (Sánchez-Danés et al, 2012). All iPSC lines tested generated DAn using this differentiation protocol, as judged by co-staining with the neuron-specific class III- β -tubulin (TUJ1) and tyrosine hydroxylase (TH; Fig 2A–C). Moreover, DAn differentiated in this way were mature and mostly of the A9 subtype, as revealed by expression of dopamine transporter (DAT), G protein-activated

Table 1. Summary of iPSC lines generated

	Patient			Disease					iPSC						
	Code	Sex	Age ^a	Age onset	Family history	Mutation	Initial symptoms ^b	LDopa response	# of Lines	Clones selected	Karyotype	Transgene silencing ^c	Pluripotency markers ^c	In vitro differentiation ^c	Teratoma assay ^c
CONTROL	SP09	M	66						4	SP09.2	46,XY	Passed	Passed	Passed	N/P
										SP09.4	46,XY	Passed	Passed	Passed	Passed
	SP11	F	48						3	SP11.1	46,XX	Passed	Passed	Passed	Passed
										SP11.4	46,XX	Passed	Passed	Passed	N/P
	SP15	F	47						4	SP15.2	46,XX	Passed	Passed	Passed	Passed
										SP15.3	46,XX	Passed	Passed	Passed	N/P
	SP17	M	52						3	SP15.4	47,XX + 20	Passed	Passed	N/P	N/P
										SP17.1	47,XY + 20	Passed	Passed	Passed	N/P
										SP17.2	46,XY	Passed	Passed	Passed	Passed
										SP17.3	46,XY	Passed	Passed	Failed	N/P
ID-PD	SP01	F	63	58	No	No	T and B	N/A	4	SP01.1	46,XX	Passed	Passed	Passed	Passed
										SP01.4	46,XX	Passed	Passed	Passed	N/P
	SP02	M	55	48	No	No	T	N/A	2	SP02.1	46,XY	Passed	Passed	Passed	Passed
										SP02.2	46,XY	Passed	Passed	Passed	N/P
	SP04	M	46	40	No	No	B	Good	2	SP04.1	46,XY	Passed	Passed	Passed	N/P
										SP04.2	46,XY	Passed	Passed	Passed	Passed
	SP08	F	66	60	No	No	T	Good	4	SP08.1	46,XX	Passed	Passed	Passed	Passed
										SP08.2	46,XX	Passed	Passed	Passed	N/P
										SP08.3	46,XX	Failed	N/P	N/P	N/P
	SP10	M	58	50	No	No	D	Good	2	SP10.1	46,XY	Passed	Passed	Passed	N/P
										SP10.2	46,XY	Passed	Passed	Passed	Passed
	SP14	M	55	51	No	No	B	Good	2	SP14.1	46,XY	Passed	Passed	Passed	Passed
										SP14.2	46,XY	Passed	Passed	Passed	N/P
	SP16	F	51	48	No	No	B	N/A	4	SP16.2	46,XX	Passed	Passed	Passed	Passed
										SP16.3	46,XX	Passed	Passed	Passed	N/P
LRRK2-PD	SP05	M	66	52	Yes	LRRK2	B	Good	2	SP05.1	46,XY	Passed	Passed	Passed	Passed
										SP05.2	46,XY	Passed	Passed	Passed	N/P
	SP06	M	44	33	Yes	LRRK2	T	Good	6	SP06.1	46,XY	Passed	Passed	N/P	N/P
										SP06.2	46,XY	Passed	Passed	Passed	Passed
	SP12	F	63	49	Yes	LRRK2	T	Good	4	SP12.3	46,XX	Passed	Passed	Passed	Passed
										SP12.4	46,XX	Passed	Passed	Passed	N/P
	SP13	F	68	57	Yes	LRRK2	T	Good	4	SP13.2	46,XX	Passed	Passed	Passed	N/P
										SP13.4	46,XX	Passed	Passed	Passed	Passed

N/A, information not available; N/P, test not performed.

^aAge at biopsy.

^bT, tremor; B, bradykinesia; D, foot dystonia.

^cTests performed as exemplified in Fig 1.

inward rectifier potassium channel 2 (GIRK2) and FOXA2, respectively (Fig S1 of Supporting information). We found a large variability in the numbers of DAn obtained from different iPSC lines, ranging from 9 to 29% of all differentiated cells, which did not depend on the presence or type of disease (Fig 2D), nor on the efficiency of lentiviral infection (Fig S2 of Supporting information), but rather appeared to depend on the specific iPSC clone used (Fig S3 of Supporting information), consistent with previous results on differentiation to other neuronal lineages (Hu et al, 2010). Thus, our results so far confirm previous data on the similar ability of control and ID-PD iPSC to give rise to DAn (Soldner et al, 2009, 2011) and extend these findings to iPSC derived from LRRK2-PD patients.

Spontaneous phenotypes of PD-specific vmDA neurons

Because SNCA is the main component of Lewy bodies, and immunolabelling with antibodies against SNCA has become the standard and most sensitive immunohistochemical method for the neuropathological diagnosis of PD (Lees et al, 2009), we first investigated whether SNCA could be detected in DAn differentiated from PD-specific iPSC. The majority of DAn differ-

entiated from Ctrl-iPSC and ID-PD iPSC exhibited barely detectable levels of endogenous SNCA in their cytoplasm, while ~20% of DAn showed cytoplasmic staining (Fig 3A, B and D). However, most DAn derived from LRRK2-PD iPSC exhibited diffuse cytoplasmic accumulations of SNCA (Fig 3C and D), which did not form obvious aggregates or inclusions. This finding is consistent with results from animal studies that reported post-translational regulatory interactions of mutant LRRK2 and SNCA (reviewed in Cookson, 2010). In this line, the accumulation of SNCA in our model of LRRK2-PD iPSC-derived DAn did not appear to depend on transcriptional regulation of SNCA expression, which was upregulated during DAn differentiation to a similar extent compared with Ctrl-iPSC- and ID-PD iPSC-derived cells (Fig S4 of Supporting information), which also showed similar LRRK2 levels (Fig S5 of Supporting information).

Since neuronal degeneration in PD patients takes decades to occur, we next attempted to maintain iPSC-derived DAn over longer culture times. For this purpose, we co-cultured them over a monolayer of mouse post-natal cortical astrocytes (Johnson et al, 2007), which supported viable cultures of vmDA neurons

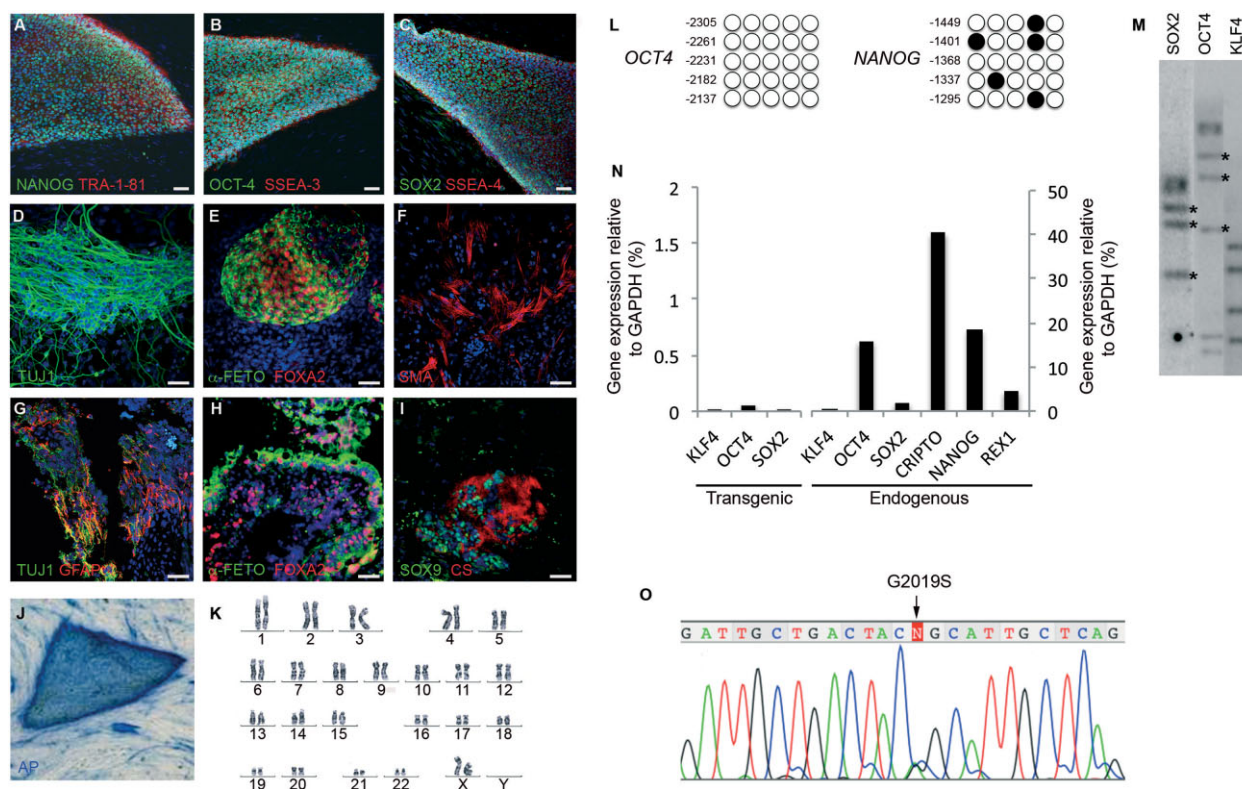


Figure 1. Generation and characterization of PD patient-specific iPSC lines.

- A–C.** Representative colonies of passage-20 LRRK2-PD-iPSC (cell line SP13.4) stained positive for the pluripotency-associated markers NANOG, OCT4 and SOX2 (green), TRA-1-81, SSEA3 and SSEA4 (red).
- D–F.** Immunofluorescence analyses of LRRK2-PD-iPSC (cell line SP13.4) differentiated *in vitro* show the potential to generate cell derivatives of all three primary germ cell layers including ectoderm (D, stained for TUJ1, green), endoderm (E, stained for α -fetoprotein, green, and FOXA2, red) and mesoderm (F, stained for smooth muscle actin, SMA, red).
- G–I.** Immunofluorescence analyses of sections from a teratoma induced by injecting LRRK2-PD-iPSC (cell line SP13.4), showing derivatives of the three main embryo germ layers: ectoderm (G, stained for TUJ1, green, and GFAP, red), endoderm (H, stained for α -fetoprotein, green, and FOXA2, red) and mesoderm (I, stained for SOX9, green, and chondroitin sulphate, CS, red). In (A–I) nuclei are counterstained with DAPI, shown in blue. Scale bars, 50 μ m.
- J.** LRRK2-PD-iPSC (cell line SP13.4) stained for alkaline phosphatase (AP) activity.
- K.** Normal karyotype of LRRK2-PD-iPSC (cell line SP13.4) at passage 20.
- L.** Bisulphite genomic sequencing of the *OCT4* and *NANOG* promoters showing demethylation in LRRK2-PD-iPSC (cell line SP13.4).
- M.** Southern blot analysis of LRRK2-PD-iPSC (cell line SP13.4) showing genomic integrations (asterisk) of the indicated retroviruses.
- N.** RT-qPCR analyses of the expression levels of retroviral-derived reprogramming factors (transgenic) and endogenous expression levels (endogenous) of the indicated genes in LRRK2-PD-iPSC (cell line SP13.4).
- O.** Direct sequence of genomic DNA from LRRK2-PD-iPSC (cell line SP13.4) identifying the *LRRK2*^{G2019S} mutation.

for up to 75 days (latest time point analysed, see Fig S6 of Supporting information). Under these conditions, Ctrl-iPSC gave rise to DAN that were morphologically homogeneous and showed the expected features of mature DAN, including complex dendritic arborizations (Fig 4A and F). However, DAN differentiated from ID-PD or LRRK2-PD iPSC developed a range of altered morphologies over long-term culture that were not evident at shorter time points (Fig S7 of Supporting information). After detailed analysis, we categorized the morphologies of individual DAN in these cultures in three types: type 1 were neurons with mature morphology and long neurites with complex arborization, similar to those differentiated from Ctrl-iPSC; type 2 DAN showed fewer and simpler

processes, reminiscent of immature neurons; and type 3 were DAN with clear signs of degeneration, including very short or absent neurites, vacuolated soma, fragmented nucleus and positive staining for cleaved caspase-3 (Fig 4A–D and Fig S7 of Supporting information). In aged (~75 days) cultures, most of DAN differentiated from ID-PD or LRRK2-PD iPSC were of altered type 2 or type 3 morphologies (Fig S7 of Supporting information), with a significant percentage of them undergoing apoptosis (Fig 4E).

To rule out any subjectivity that might have occurred in the ascription of different morphologies to iPSC-derived DAN, we also directly measured the number and length of neurites. For this, we used high-power confocal images of TH-stained single

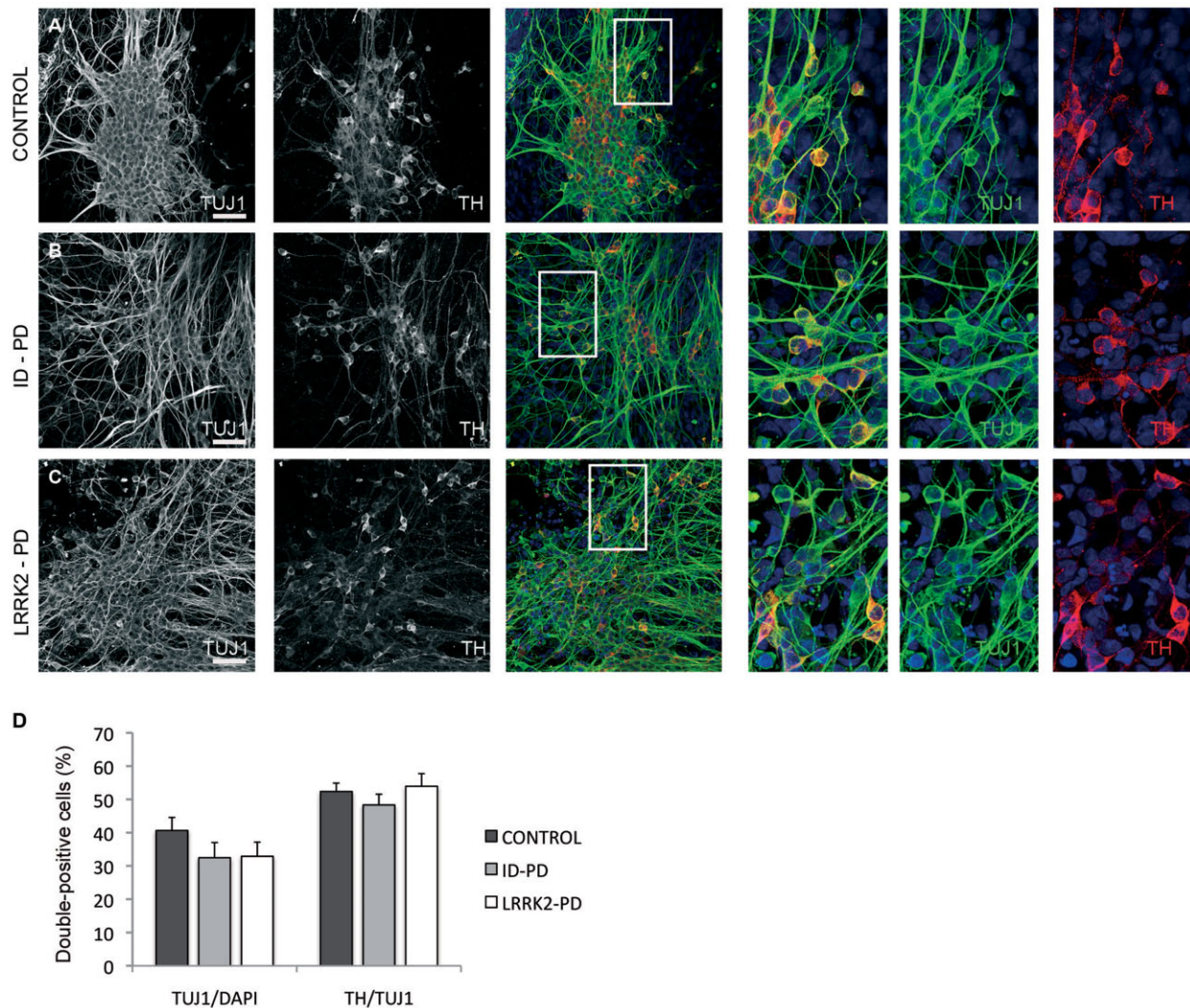


Figure 2. Differentiation of DA neurons from PD patient-specific iPSC. Ctrl-, ID-PD and LRRK2-PD iPSC were differentiated into DA neurons and analysed by immunofluorescence for expression of TUJ1 (green) and TH (red) at the end of the 30-day differentiation protocol.

A-C. All iPSC-derived neurons express TUJ1 and TH. Shown are representative images of differentiation experiments from Ctrl-iPSC (cell line SP11.1), ID-PD iPSC (cell line SP08.1) and LRRK2-PD iPSC (cell line SP06.2). Nuclei are counterstained with DAPI, shown in blue. Scale bars, 50 μ m.

D. Quantitative analyses of cells stained positive for TUJ1 (left bars) or TUJ1 and TH (right bars). TUJ1-positive cells are represented as the percentage of total number of cells (stained with DAPI). Bars represent average with SEM as error bars. Data for CONTROL is the average of 4 iPSC lines, for ID-PD from 7 iPSC lines and LRRK2-PD from 4 iPSC. No significant differences were found in the ability of iPSC to generate neurons [$F(2,12) = 0.953$; $p = 0.413$] or DA neurons [$F(2,12) = 0.679$; $p = 0.526$] after 30 days of differentiation.

neurons, randomly chosen, differentiated from all the PD patient-specific and control iPSC lines and used software-assisted analysis (Fig 4F, see also Chu et al, 2009). These analyses confirmed that both the number and length of neurites of Ctrl-iPSC-derived DAN were significantly higher than those of DAN differentiated from ID-PD or LRRK2-PD iPSC (Fig 4G and H). The causative role of the genetic background of patient-specific cells was further investigated by ectopically expressing mutated LRRK2^{G2019S} in DAN differentiated from Ctrl-iPSC. In these experiments, DAN differentiated for 65 days were co-transfected with plasmids encoding wild-type (WT) LRRK2 or LRRK2^{G2019S} and GFP, and cells were analysed after 10 days.

Even though the transfection efficiency (as evaluated by GFP/TH double positive cells) was low in all cases, we could readily detect DAN transfected with WT LRRK2, which were morphologically indistinguishable from untransfected DAN (Fig 4I-K and M). In contrast, few DAN transfected with mutant LRRK2^{G2019S} were found after 10 days, suggesting that the expression of pathogenic LRRK2 is deleterious for these cells (Fig 4M). Indeed, surviving DAN expressing mutated LRRK2^{G2019S} displayed evident morphological alterations, including fewer and shorter neurites compared to untransfected DAN or DAN expressing WT LRRK2 (Fig 4I-L and N). So far, our data show that, even though PD-specific iPSC differentiate

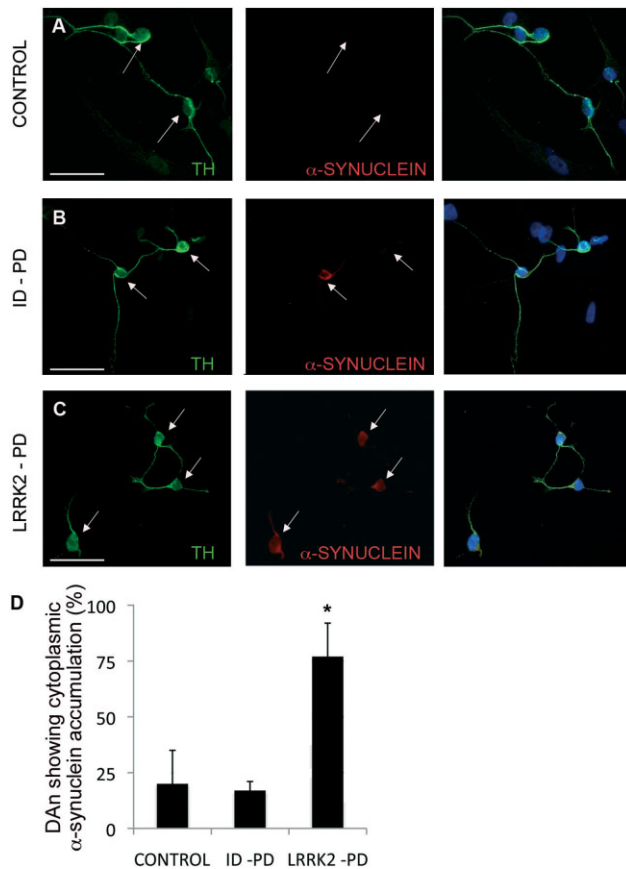


Figure 3. Abnormal SNCA accumulation in DA neurons from LRRK2-PD iPSC.

- A-C.** Immunofluorescence analyses of DA neurons after 30 days of differentiation from Ctrl-iPSC (A), ID-PD iPSC (B) and LRRK2-PD iPSC (C), co-stained for TH (green) and SNCA (red). Arrows point to DA neurons, as identified by TH positive staining.
- D.** Quantitative analyses of DA neurons showing cytoplasmic accumulation of SNCA. Bars represent average with SEM as error bars. Data is the average of at least two-independent experiments using 3 Ctrl-iPSC lines, 4 ID-PD iPSC lines and 3 LRRK2-PD iPSC lines. The differences in the number of DA neurons showing accumulation of SNCA are statistically significant (asterisk) when comparing those derived from LRRK2-PD iPSC with either those from Ctrl-iPSC ($p=0.019$) or from ID-PD iPSC ($p=0.002$) [$F(2,16)=6.888$; $p=0.007$].

normally into vmDA neurons, they display evident signs of morphological alterations upon long-time culture.

Impaired autophagy of PD-specific vmDA neurons

We next investigated the bases for the altered morphology and shorter/fewer neurites of PD iPSC-derived DAn compared to those differentiated from Ctrl-iPSC. Given that autophagy has been shown to play key roles for maintaining neurite length (reviewed in Chu et al, 2009) and that several lines of evidence indicate that dysregulated autophagy may be a pathogenic mechanism in PD (Menzies et al, 2011), we first analysed the autophagosome content in iPSC-derived DAn. For this purpose, we used immunofluorescence labelling with antibodies to the light chain type 3 protein (LC3), a marker of autophagosomes

(Kabeya et al, 2000). After 75 days of culture, DAn differentiated from Ctrl-iPSC showed a diffuse cytoplasmic LC3 staining with very few autophagosomes (LC3-positive puncta, Fig 5A). In contrast, a marked increase in LC3-positive puncta was evident in untreated DAn differentiated from ID-PD or LRRK2-PD iPSC (Fig 5B–E). The fact that autophagosomes were evident even in the absence of inhibitors of lysosomal proteolysis in these neurons suggests either massive upregulation of autophagosome formation, or a compromise in the clearance of autophagic compartments in these cells. To distinguish between these possibilities, we compared levels of p62 in the three groups of cells. A large fraction of p62 undergoes degradation through autophagy because it serves as a cargo-recognizing protein that it is sequestered in autophagosomes along the cytosolic cargo. Immunostaining for p62 revealed a marked increase in the levels of this protein in DAn differentiated from ID-PD or LRRK2-PD iPSC, where it showed a punctuate pattern reminiscent of the one observed for LC3 (Fig 5F–J).

To directly analyse autophagic clearance we compared the rates of degradation of LC3-II in the different cells by immunoblot (LC3-flux assay), even though we were aware that neural-specific changes could be underestimated in this type of analysis, as DAn represent only a fraction of the cells present in whole-culture lysates. Part of the LC3 located in the inner membrane of the autophagosome is degraded along with the cargo when these compartments fuse to lysosomes, consequently the increase in levels of LC3-II upon blockage of lysosomal proteolysis provides information on autophagic clearance. As expected, immunoblot for LC3 confirmed an increase in basal steady-state levels of LC3-II in the PD-derived cells (Fig 5K and L). Blockage of lysosomal degradation resulted in an increase in LC3-II levels in all cells but, despite some level of individual variability, the autophagic flux was significantly reduced in PD-derived cells (Fig 5K and M). To separately analyse autophagosome formation independently of clearance, we also compared changes in LC3-II at two different times after blockage of lysosomal degradation (under these conditions, any increase in LC3-II would directly result from autophagosome formation as degradation is no longer occurring). Although we found a consistent trend for increased autophagosome formation in DAn differentiated from ID-PD or LRRK2-PD, when compared to Ctrl-iPSC (Fig 5K and N), this increase was not statistically significant and clearly less pronounced than the changes observed on clearance. Our findings thus support that the marked increase in the abundance of these compartments in PD-iPSC-derived neurons when compared with control was due, for the most part, to blockage on their clearance.

Ultrastructural analysis by electron microscopy confirmed marked differences in the autophagic system between DAn differentiated from Ctrl-iPSC and those of PD origin. As shown in Fig 6A–C, autophagic vacuoles were in general more abundant in the PD groups. In control cells, most of the autophagic profiles corresponded to autophagolysosomes (autophagic vacuoles already fused to lysosomal compartments; Fig 6A and D). In contrast, in the case of the ID-PD and LRRK2-PD, autophagosomes (prior to lysosomal fusion) were predominant (Fig 6B–D), supporting that the increase in LC3-positive structures is

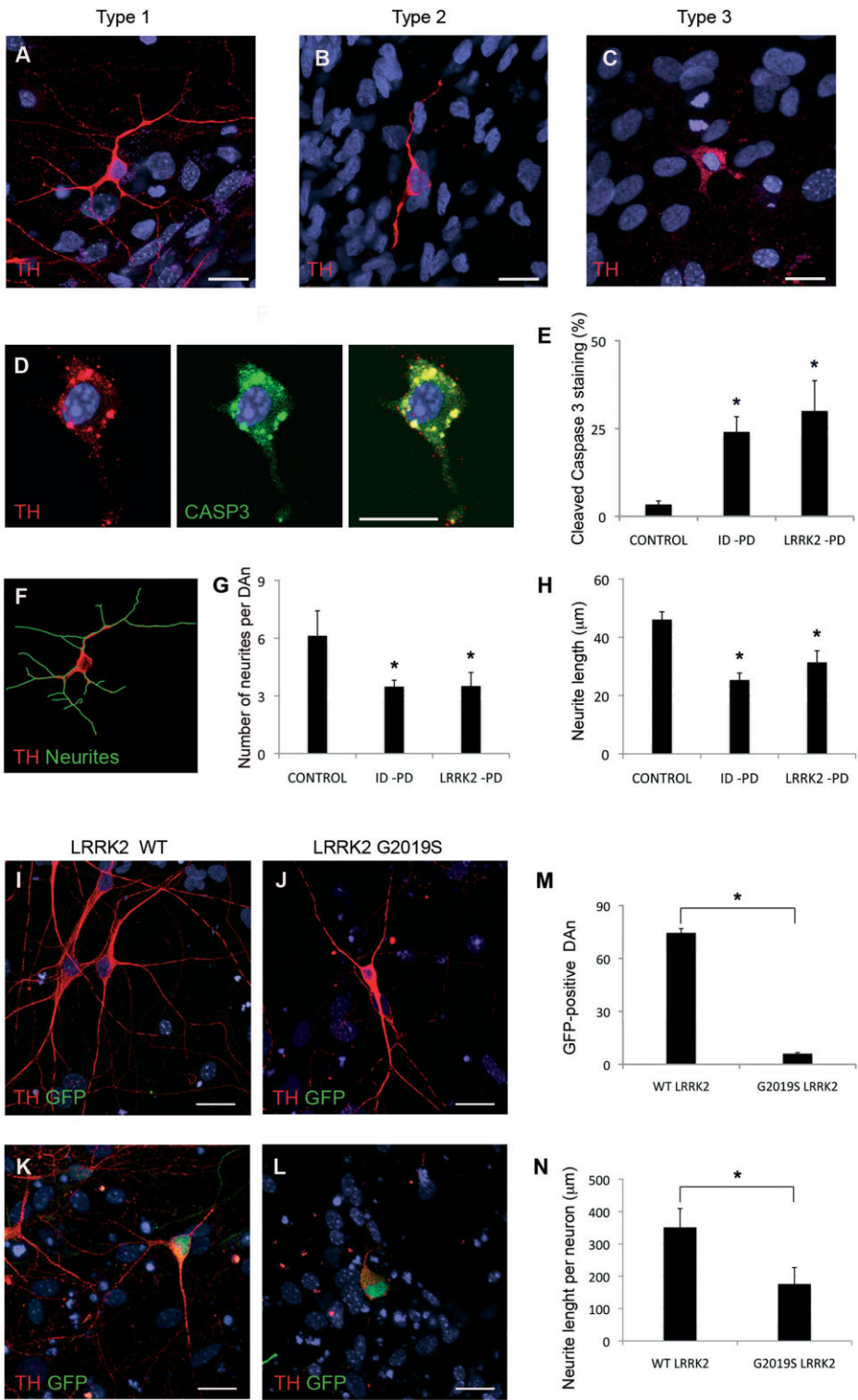


Figure 4.

likely due to problems in the maturation of autophagosomes. An increase in intracellular lipid droplets, previously shown to undergo degradation by macroautophagy (Singh et al, 2009), and the presence of dilated ER in the PD cells are also common features of cells with compromised autophagy (Fig 6B, C and E).

Lastly, defective autophagosome clearance was also confirmed analyzing the co-localization of autophagosome (LC3) and lysosomal (LAMP-1) markers in DAn differentiated from ID-PD, LRRK2-PD or Ctrl-iPSC. The very low levels of LC3 puncta detected in basal conditions in the control cells (Fig S8 of Supporting information) precluded evaluating LC3/LAMP-1 co-localization in these cells. To be able to compare the autophagic flux among the three groups of cells, we reduced degradation of LC3 in lysosomes by treatment with leupeptin (which does not modify autophagosome/lysosome fusion). Under these conditions, we detected a significantly lower level of LC3/LAMP-1 co-localization in both ID-PD and LRRK2-PD iPSC-derived DAn, when compared to control cells (Fig 7A–D). These results confirm compromised maturation of autophagosomes into autophagolysosomes in these cells.

To evaluate whether morphological alterations and autophagy defects in DAn were mechanistically linked, we next reproduced in Ctrl-iPSC-derived DAn the blockage of autophagic clearance observed in ID-PD and LRRK2-PD iPSC, by inducing autophagosome formation through mTOR inhibition with rapamycin and, at the same time, blocking lysosomal degradation with leupeptin. This treatment resulted in ~20% of Ctrl-iPSC-derived DAn showing higher number of autophagosomes, which stained positive for LC3 as well as for LAMP-1 (Fig S9 of Supporting information). This combined treatment had an even more marked effect in DAn differentiated from ID-PD or LRRK2-PD iPSC. In either case, rapamycin/leupeptin treatment induced a significant shortening in the average total neurite length of DAn (Fig 7E). Importantly, the increased susceptibility of ID-PD and LRRK2-PD iPSC-derived DAn to rapamycin/leupeptin

treatment appeared to be cell type-specific, since patients' fibroblasts showed an increase of LC3-positive puncta and changes in LC3 levels comparable to control cells upon this treatment (Fig S10 of Supporting information), further supporting that the compromise in autophagosome clearance may be specific for DAn. These results indicate that DAn differentiated from either ID-PD or LRRK2-PD iPSC have a deficient competence for autophagic clearance compared with Ctrl-iPSC-derived DAn, and suggest that the PD pathology related phenotypes revealed in our *in vitro* model are caused, at least in part, by an impairment in the autophagy pathway.

DISCUSSION

In this study, we describe the generation and characterization of a collection of iPSC lines representing age- and sex-matched patients of sporadic and LRRK2-associated PD, as well as control individuals. Consistent with previous studies describing the generation of patient-specific iPSC in the context of PD (Park et al, 2008a; Soldner et al, 2009, 2011) or other conditions (Dimos et al, 2008; Park et al, 2008a; Raya et al, 2010), the efficiency of iPSC generation varied among different individuals, but did not depend on the presence or type of disease, nor on the donor age. We chose retroviral delivery of reprogramming transgenes to secure the derivation of iPSC lines from valuable biopsy material, because of the higher efficiency of this system in our hands compared to that of lentivirus-based systems. A clear advantage of the latter is the possibility of using inducible excisable lentivirus (Soldner et al, 2009), so that iPSC are free of reprogramming transgenes. However, we reasoned that a careful analysis of the silencing of integrated transgenes in our iPSC lines, together with the generation of independent iPSC lines from several patients representing each condition, would rule out any confounding effects due to residual transgene

Figure 4. Morphological alterations of PD-iPSC-derived DA neurons after long-term culture.

- A–C.** Immunofluorescence analyses of Ctrl-iPSC (cell line SP11.1, **A**), ID-PD iPSC (cell line SP08.1, **B**) and LRRK2-PD iPSC (cell line SP06.2, **C**), differentiated towards DA neurons and cultured for 75 days on cortical astrocytes, stained for TH (red). Images are representative of the three morphologies of DA neurons found under these conditions.
- D.** Immunofluorescence analysis of DA neurons differentiated from LRRK2-PD iPSC (cell line SP06.2) for 75 days, co-stained for TH (red) and cleaved caspase-3 (green). In **A–D** nuclei are counterstained with DAPI, shown in blue. Scale bar, 12.5 μ m.
- E.** Quantitative analyses of the percentage of DA neurons differentiated for 75 days showing cleaved caspase-3 staining. Data for CONTROL is the average of 4 iPSC lines, for ID-PD from 7 iPSC lines and LRRK2-PD from 4 iPSC. Asterisks denote statistically significant differences between CONTROL and ID-PD ($p = 0.004$) or LRRK2-PD ($p = 0.011$) [$F(2,12) = 5.668$; $p = 0.018$].
- F.** Image of the same DA neuron shown in **(A)**, with neurites outlined by green traces as processed by NIH ImageJ software.
- G.** Quantitative analyses of the number of neurites in DAn differentiated for 75 days. Data for CONTROL is the average of 87 DAn from 4 iPSC lines, for ID-PD is from 119 DAn from 7 iPSC lines and for LRRK2-PD is from 91 DAn from 4 iPSC. Asterisks denote statistically significant differences between CONTROL and ID-PD ($p < 0.001$) or LRRK2-PD ($p < 0.001$) [$F(2,294) = 21.096$; $p < 0.001$].
- H.** Quantitative analyses of the neurite length in DA neurons differentiated for 75 days. Data for CONTROL is the average of 540 neurites from 4 iPSC lines, for ID-PD is from 409 neurites from 7 iPSC lines and for LRRK2-PD is from 303 neurites from 4 iPSC. Asterisks denote statistically significant differences between CONTROL and ID-PD ($p < 0.001$) or LRRK2-PD ($p < 0.001$) [$F(2,1249) = 42.161$; $p < 0.001$]. (**E**, **G** and **H**) bars represent average with SEM as error bars.
- I–L.** DAn (TH positive, red) and GFP (green) from Ctrl-iPSC (SP11.1) co-transfected with plasmids encoding WT LRRK2 (WT, **I** and **K**) or mutant LRRK2^{G2019S} (**J** and **L**) and GFP. (**I** and **J**) Images showing GFP negative DA neurons. (**K** and **L**) Co-transfected DA neurons being GFP positive. Nuclei are counterstained with DAPI, shown in blue. Scale bar, 12.5 μ m.
- M.** Quantitative analysis of the number of GFP positive neurons present in co-transfected cells.
- N.** Quantitative analyses of the neurite length per neuron in co-transfected DAn. For (**M** and **N**) data are average of two-independent experiments using two different Ctrl-iPSC lines. Bars represent average with SEM as error bars. Statistically significant differences are indicated with asterisks ($p = 0.003$ in **M**; $p = 0.006$ in **N**).

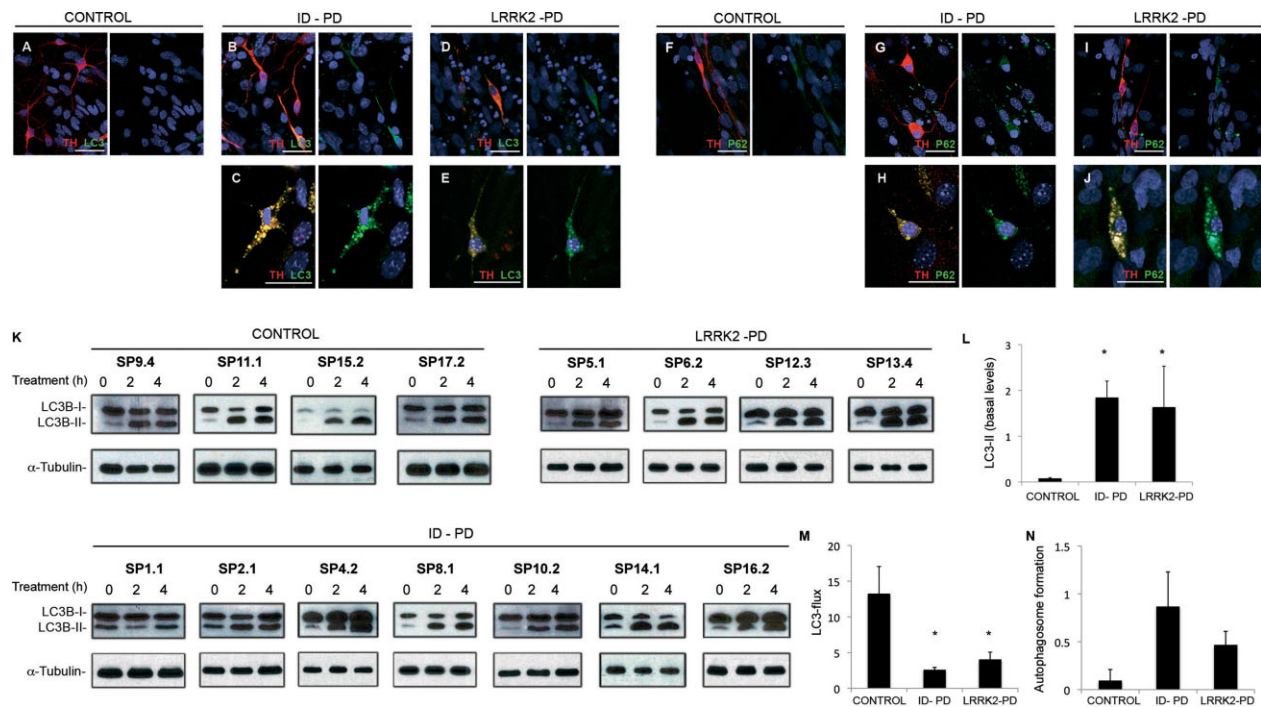


Figure 5. Alterations in autophagic clearance in DA neurons from PD patient-specific iPSC.

- A–E.** Immunofluorescence analyses of Ctrl-iPSC (cell line SP17.2, **A**), ID-PD iPSC (cell line SP16.2, **B** and **C**) and LRRK2-PD iPSC (cell line SP12.3, **D** and **E**), differentiated towards DA neurons and cultured for 75 days on cortical astrocytes, co-stained for LC3 (green) and TH (red). Images are representative of the most abundant morphologies of DA neurons found under these conditions.
- F–J.** Immunofluorescence analyses of Ctrl-iPSC (cell line SP17.2, **F**), ID-PD iPSC (cell line SP10.2, **G** and **H**) and LRRK2-PD iPSC (cell line SP06.2, **I** and **J**), differentiated towards DA neurons and cultured for 75 days on cortical astrocytes, co-stained for p62 (green) and TH (red). Images are representative of the morphologies of DA neurons found under these conditions. In **A–J**, nuclei are counterstained with DAPI, shown in blue. Scale bars, 35 μ m.
- K.** Western blot analysis for LC3 in DA neuron cultures at 75 days of differentiation from the indicated iPSC lines, treated with leupeptin and NH_4Cl during the indicated period of time. α -tubulin is used as a loading control.
- L.** Quantification analyses of the basal levels (treatment time = 0) of LC3-II relative to α -tubulin. Asterisks denote statistically significant differences between CONTROL and ID-PD ($p = 0.003$) and between CONTROL and LRRK2-PD ($p = 0.046$) [$F(2,12) = 4.253$; $p = 0.046$].
- M.** Quantification of LC3 flux normalized to α -tubulin. Asterisks denote statistically significant differences between CONTROL and ID-PD ($p = 0.002$) and between CONTROL and LRRK2-PD ($p = 0.029$) [$F(2,12) = 9.614$; $p = 0.003$].
- N.** Quantification of the autophagosome formation normalized to α -tubulin. No significant differences were found among groups [$F(2,12) = 1.570$; $p = 0.248$]. (**L–N**) Data per each group is the average of the blots shown in **K**. Bars represent average with SEM as error bars.

expression in individual iPSC lines, while still allowing a high efficiency of iPSC generation. In any case, we omitted c-MYC from the reprogramming cocktail due to the possibility of its interaction with LRRK2 in the context of eukaryotic initiation factor 4E (eIF4E) function regulation (Imai et al, 2008; Ruggero et al, 2004).

All the iPSC lines tested in our study were competent to give rise to DAN, although the efficiency at which they did so varied among lines. Similar findings have been reported for iPSC in a variety of differentiation protocols, including neuronal lineages other than DAN (Hu et al, 2010), cardiomyocytes (Zhang et al, 2009) and haematopoietic cells (Woods et al, 2011). Importantly, the variability in differentiation efficiency of our iPSC lines was independent of the presence or type of disease, indicating inter-line variation, rather than a result of the disease. Moreover, DAN differentiated from Ctrl- or PD-iPSC appeared morphologically and phenotypically indistinguishable after

30 days in culture, in agreement with a previous report that analysed the DAN differentiation ability of IP-PD iPSC (Soldner et al, 2009), and extending these findings to iPSC derived from LRRK2-PD patients. Despite this similarity, we found anomalous accumulation of SNCA in DAN differentiated from LRRK2-PD iPSC, compared to those from Ctrl- or ID-PD iPSC. The identification of this PD-related phenotype in LRRK2 mutant DAN is consistent with the notion that LRRK2 and SNCA may participate in intersecting pathways (reviewed in Cookson, 2010), and with the fact that LRRK2 can accelerate mutant SNCA-induced neuropathology in mice in a dose-dependent manner (Lin et al, 2009). Moreover, these results demonstrate the ability of iPSC-based cellular systems to recapitulate PD-related pathology and to model a monogenic form of PD.

Probably the most significant finding of our work is the identification of spontaneous phenotypes in long-term cultures of DAN from both idiopathic and LRRK2-associated PD. To our

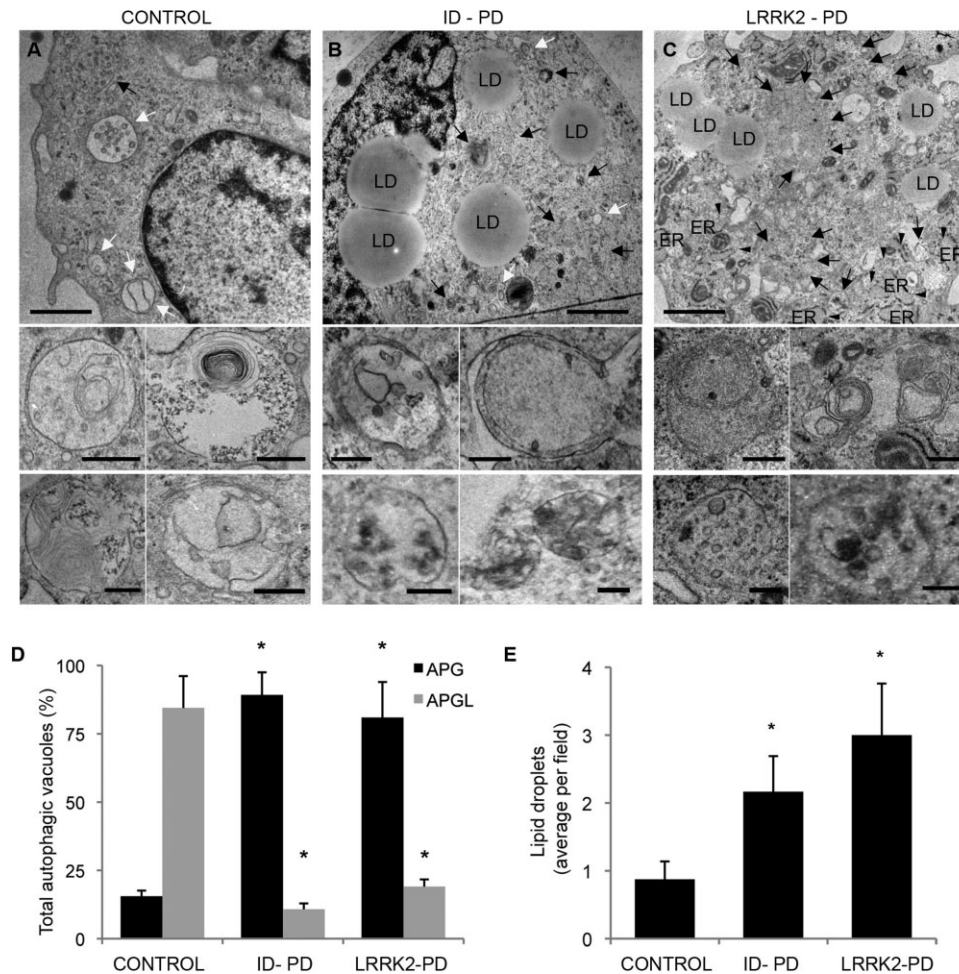


Figure 6. Autophagic system in iPSC-derived neurons.

- A-C.** Ultrastructure of DA neurons at 75 days of differentiation from Ctrl-iPSC (A), ID-PD iPSC (B) and LRRK2-PD (C) iPSC. Lower inserts show higher magnification images to illustrate individual examples of autophagic vacuoles. Black arrows: autophagosomes; white arrows: autophagolysosomes; arrowheads: dilated ER; LD: lipid droplets. Scale bars: 0.5 μ m (top) and 0.2 μ m (bottom).
- D.** Percentage of autophagosomes (APG) and autophagolysosomes (APGL) relative to the total amount of autophagic vacuoles per field. Asterisks denote statistically significant differences in the percentage of autophagosomes [$F(2,16) = 28.274$; $p < 0.001$] and of autophagolysosomes [$F(2,16) = 8.894$; $p = 0.003$] between CONTROL and ID-PD (autophagosomes, $p < 0.001$; autophagolysosomes, $p = 0.003$), and between CONTROL and LRRK2-PD (autophagosomes, $p = 0.003$; autophagolysosomes, $p = 0.010$).
- E.** Content of lipid droplets per field. Asterisks denote statistically significant differences [$F(2,16) = 28.274$; $p < 0.001$] between CONTROL and ID-PD ($p = 0.010$), and between CONTROL and LRRK2-PD ($p = 0.013$). Bars represent average with SEM as error bars ($n = 4-8$).

knowledge, this is the first time such phenotypes have been described. Indeed, Soldner and colleagues did not find any differences between DAn differentiated from ID-PD iPSC and from control iPSC (Soldner et al, 2009), a fact that they attributed to the short time span of cultured neurons (32–42 days). Moreover, these authors suggested that additional manipulations might be necessary to accelerate PD-pathology related phenotypes in iPSC-derived DAn *in vitro*, such as increasing oxidative stress, challenge with neurotoxins or overexpressing PD-related genes (Soldner et al, 2009). Thus, it may not appear entirely unexpected that DAn differentiated from our LRRK2-PD iPSC displayed PD-related alterations *in vitro*. However, during the writing of this manuscript, a study was published that found

no alterations in DAn differentiated from iPSC from one LRRK2-PD patient, unless these were challenged with H_2O_2 , 6-hydroxydopamine (a PD-related neurotoxin) or MG-132 (proteasome inhibitor; Nguyen et al, 2011). We believe that the lack of spontaneous PD pathology-related phenotypes in previous iPSC-based models of ID-PD (Soldner et al, 2009) and LRRK2-PD (Nguyen et al, 2011), in contrast with our results presented here, may be due to at least two reasons: First, the longer time span of cultured neurons (up to 75 days) in our experiments, which may have induced culture-related stress conditions mimicking *in vivo* aging in PD patients, and thus accelerated the development of PD-related phenotypes *in vitro*. In support of this, we found that DAn differentiated for 30 days from ID-PD- or LRRK2-PD iPSC, a

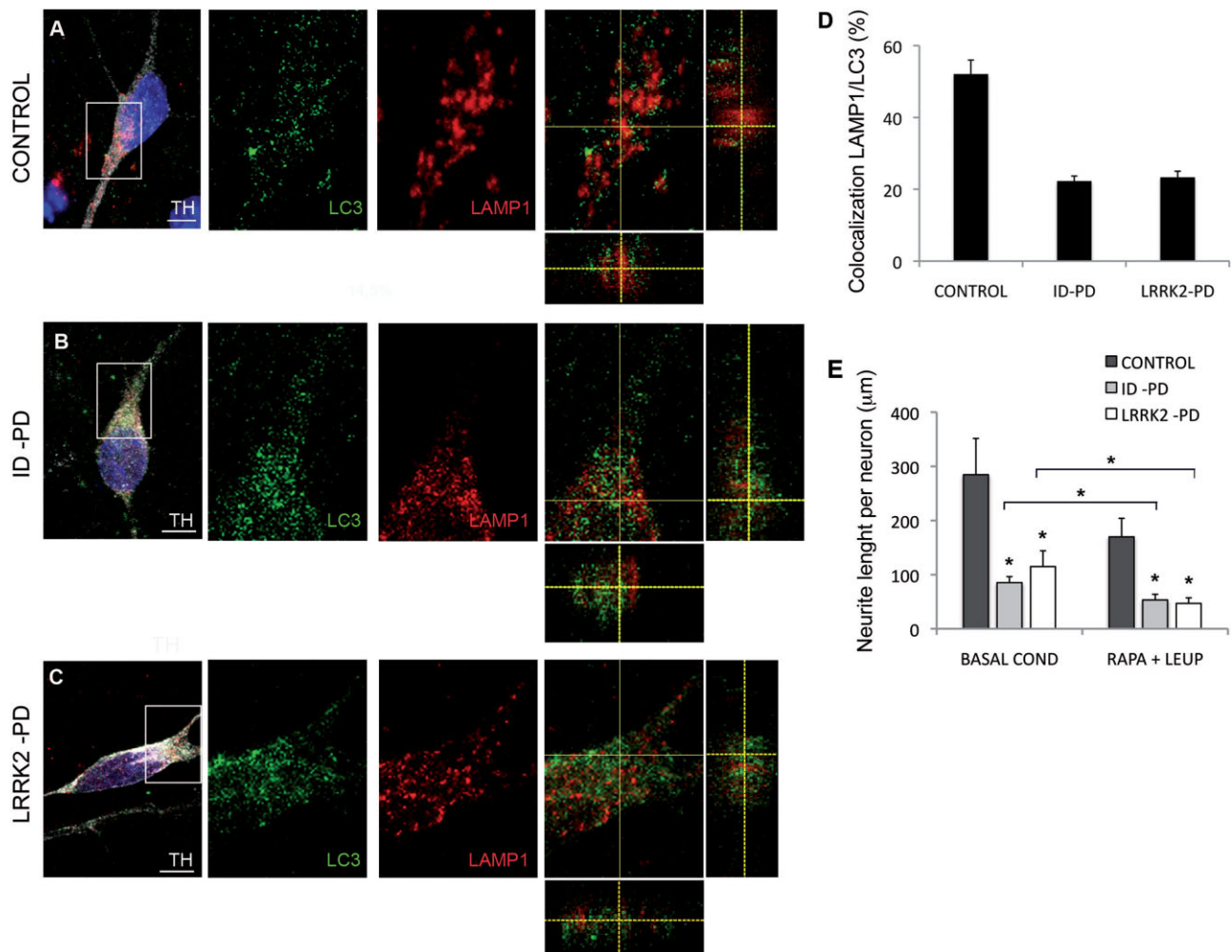


Figure 7. Defective autophagosome clearance in PD-iPSC-derived DA neurons.

A–C. DAN (TH positive, white) cultured on astrocytes for 75 days stained for LC3 (green) and LAMP1 (red) after 6 h of treatment with leupeptin from CONTROL (cell line SP11.1, **A**), ID-PD (cell line SP10.2, **B**) and LRRK2-PD (cell line SP05.1, **C**) iPSC. Z-stack views are shown on the merged images (right panels). Nuclei are counterstained with DAPI, shown in blue. Scale bars, 5 μm.

D. Quantitative analyses of the colocalization of LAMP1 and LC3. Data for each group is the average of 15, 15 and 14 DAN representing three different Ctrl-, ID-PD and LRRK2-PD iPSC lines, respectively. Asterisks denote statistically significant differences between CONTROL and ID-PD ($p < 0.001$) or LRRK2-PD ($p < 0.001$) [$F(2,41) = 41.628$; $p < 0.001$].

E. Quantitative analyses of the total neurite length per neuron with or without treatment with rapamycin and leupeptin. Under basal conditions (without treatment), data for CONTROL is the average of 97 DAN from 4 iPSC lines, for ID-PD is from 127 DAN from 7 iPSC lines and for LRRK2-PD is from 80 DAN from 4 iPSC lines. After treatment with rapamycin and leupeptin (RAPA + LEUP), data for CONTROL is the average of 60 DAN from 4 iPSC lines, for ID-PD is from 112 DAN from 7 iPSC lines and for LRRK2-PD is from 74 DAN from 4 iPSC lines. Asterisks above data bars denote statistically significant differences under basal conditions [$F(2,301) = 5.886$; $p = 0.003$] between CONTROL and ID-PD ($p = 0.003$), and between CONTROL and LRRK2-PD ($p = 0.022$), and after treatment with rapamycin and leupeptin [$F(2,243) = 48.812$; $p < 0.001$] between CONTROL and ID-PD ($p < 0.001$), and between CONTROL and LRRK2-PD ($p < 0.001$). Asterisks comparing data sets denote statistically significant differences in total neurite length after treatment with rapamycin and leupeptin in DAN from ID-PD ($p = 0.001$) and LRRK2-PD ($p < 0.001$), but not from CONTROL ($p = 0.090$). In (**D–E**) bars represent average with SEM as error bars.

time when no alterations (other than SNCA accumulation in LRRK2-PD iPSC-derived DAN) were evident, showed increased susceptibility to sub-lethal concentrations of the PD-related toxin MPP⁺ (Fig S11 of Supporting information). Thus, it is likely that the prolonged culture of these cells in our 75-day differentiation protocol served to amplify the intrinsic susceptibility of PD-iPSC-derived DAN to undergo neurodegeneration in response to aging, the most important PD-related susceptibility

factor. Second, it should be noted that the majority of iPSC-derived neurons in our experiments were of vmDA neuron phenotype, whereas, this cell population was much rarer in previous studies, with Soldner and colleagues reporting ~10% of DAN (TH+/TUJ1+ cells) of a phenotype that was not examined in detail (Soldner et al, 2009), and Nguyen and colleagues reporting less than 1% of neurons of a vmDA phenotype (Nguyen et al, 2011). We believe that the enrichment

of vmDA neuron in our experiments facilitated the identification of PD pathology-related phenotypes, which are known to affect preferentially this specific subtype of DAN.

Our iPSC-based PD model should help investigating the pathogenic mechanisms that underlie PD neurodegeneration. Thus, although several lines of evidence suggest a causal link between impaired autophagy and the development of PD-related pathology, definitive proof for this has been elusive (reviewed in Yang & Mao, 2010). Our findings of defective autophagosome clearance in DAN from PD patient-specific iPSC, together with the positive correlation between the expansion of the autophagic compartment and the degree of morphological alterations in these cells, provide strong support for this hypothesis. In addition, the iPSC-based PD model described here should also prove an invaluable tool to investigate early functional alterations that predate the onset of neurodegeneration, which surely will help identifying potentially new therapeutic targets for the prevention, rather than rescue, of PD-related neurodegeneration.

Overall, our studies demonstrate the potential of iPSC-based technology to experimentally model the pathogenic mechanisms of late-onset diseases such as PD. Not only did we find PD pathology related phenotypes in long-term cultures of DAN representing a monogenic form of PD, but also in those from patients of ID-PD. This critical point indicates that the cause for the increased susceptibility of ID-PD-derived DAN to undergo degeneration *in vitro* after long-time culture, albeit complex, is encoded in the genome of ID-PD patients, or at least of those tested in our study. Therefore, intrinsic cell-autonomous factors, rather than environmental influences, appear to be sufficient to trigger neurodegeneration of DAN from PD patients, in a process that requires time to manifest itself, but that can be modelled within the time-frame of *in vitro* experiments. We believe that key aspects to the success of our strategy were the efficient differentiation of the PD relevant cell type, the ability to maintain DAN cultures over a long-term culture span and the use of multiple patients per condition, which allowed controlling the inherent variability of human pluripotent stem cell lines. In this way, we have identified *in vitro* phenotypes associated with ID and/or LRRK2 PD, which could be harnessed as readouts for drug screening studies. Our findings provide important conceptual and technical advances for the understanding of PD pathogenesis and, eventually, for the identification of novel therapeutic strategies in this disease.

MATERIALS AND METHODS

Description of PD patients

Studies were approved by the authors' Institutional Review Board and conducted under the Declaration of Helsinki. Patients were encoded to protect their confidentiality, and written informed consent obtained. The generation of human iPS cells was done following a protocol approved by the Spanish competent authorities (Commission on Guarantees concerning the Donation and Use of Human Tissues and Cells of the Carlos III Health Institute). PD was diagnosed according to UK Brain Bank criteria (Hughes et al, 1992). We identified subjects

participating in the study among patients attending the outpatient Movement Disorder Unit at the Hospital Clinic of Barcelona (Barcelona, Spain). Four patients were known from previous studies to have LRRK2-associated PD. The remaining seven patients had a sporadic form of PD and LRRK2 mutation was excluded by direct DNA sequencing. Healthy controls were recruited among patients' spouses.

Generation of iPSC

Epidermal keratinocytes were used for iPSC generation, except for patients SP05 and SP09, for whom dermal fibroblasts were used. Keratinocytes were cultured in serum-free low calcium medium (Epilife, Invitrogen) and reprogrammed using a 1:1:1 mix of retroviruses encoding FLAG-tagged OCT4, SOX2 and KLF4, essentially as previously described (Aasen et al, 2008). Fibroblasts were reprogrammed using a 1:1:1 mix of retroviruses encoding FLAG-tagged OCT4, SOX2 and KLF4, following a previously described protocol (Raya et al, 2010). For selecting reprogrammed cells, the medium was changed to human ES cell (hESC) medium, consisting of KO-DMEM (Invitrogen) supplemented with 20% KO-Serum Replacement (Invitrogen), 2 mM Glutamax (Invitrogen), 50 μ M 2-mercaptoethanol (Invitrogen), non-essential aminoacids (Cambrex) and 10 ng/ml bFGF (Peprotech). Cultures were maintained at 37°C, 5% CO₂, with media changes every other day. Colonies were picked based on morphology 45–60 days after the initial infection and plated onto fresh feeders. Lines of patient-specific iPS cells were maintained by mechanical dissociation of colonies and splitting 1:3 onto feeder cells in hESC medium or by limited trypsin digestion and passaging onto Matrigel-coated plates with hESC medium pre-conditioned by mouse embryonic fibroblasts. The iPSC lines generated and characterized in this study are currently in the process of being registered and deposited in the Spanish Stem Cell Bank, through which they will be made available to interested researchers.

Characterization of iPSC lines

Expression of retroviral transgenes and endogenous pluripotency-associated transcription factors by quantitative Polymerase Chain Reaction (after reverse transcription) (RT-PCR), integration of retroviral transgenes by Southern blot and methylation status of *OCT4* and *NANOG* promoters were assessed as previously reported (Aasen et al, 2008). *In vitro* differentiation towards endoderm, mesoderm and neuroectoderm was carried out essentially as described (Raya et al, 2008). Severe combined immunodeficient (SCID) beige mice (Charles River Laboratories) were used to test the teratoma induction capacity of patient-specific iPS cells essentially as described (Aasen et al, 2008). All animal experiments were conducted following experimental protocols previously approved by the Institutional Ethics Committee on Experimental Animals, in full compliance with Spanish and European laws and regulations.

iPSC differentiation to DA neurons

For DAN differentiation, iPSC were transduced with LV.NES.LMX1A.GFP and processed as previously described (Sánchez-Danés et al, 2011). For DAN yield analysis cells were co-cultured with PA6 for 3 weeks in N2B27 medium. For SNCA and LRRK2 staining, DAN generated on the top of PA6 for 3 weeks were trypsinized and cultured for 3 days on Matrigel-coated dishes. For long-term culture, neural progenitor cells

were seeded onto mouse primary cortical astrocytes, prepared as described elsewhere (Giralt et al, 2010), and maintained in N2B27 medium. After 9 weeks, some wells were incubated with 4 μ M rapamycin (LC Labs) and/or 100 μ M leupeptin (Sigma) for 6 h prior to fixation. For LC3 flux studies in DAn, neurons differentiated on murine astrocytes for 9 weeks were treated for 2 and 4 h with lysosomal inhibitors NH₄Cl (20 mM, Sigma) and leupeptin (200 μ M) in N2B27 media. After the treatment a pellet was obtained using a cell scraper (BD).

Expression of wild-type or mutant LRRK2 in DAn

DAn differentiated on top of murine astrocytes were grown on coverslips and transfected at day 65 of the differentiation protocol. The transfection was done as described previously (Kaech & Banker, 2006). Briefly, transfection media consisting on 200 μ l of MEM media (Invitrogen), 5 μ l lipofectamine 2000 (Invitrogen), 1 μ g of plasmids encoding WT LRRK2 or mutant LRRK2^{G2019S} (kindly provided by Giorgio Rovelli, Novartis Pharma Ltd, Basel, Switzerland, and Philipp Kahle, Hertie Institute for Clinical Brain Research, University Clinics Tübingen, Germany) and 0.25 μ g pmaxGFP plasmid (Lonza) was prepared. The media was aspirated from the wells and in each coverslip containing the DAn culture 200 μ l of transfection media was added and incubated for 4 h at 37°C followed by replacement (two times) with fresh N2B27 media. GFP positive cells started to appear 2 days after transfection.

Immunohistochemistry

Cells were fixed with 4% paraformaldehyde in PBS at 4°C for 10 min and permeabilized for 15 min in 0.5% Triton in PBS. Cells were then blocked in Triton-X100 with 3% donkey serum for 2 h. The following antibodies were used: mouse anti-Tra-1-81 (Chemicon, MAB4381, 1:100), rabbit anti-Sox2 (Chemicon, AB5603, 1:500), mouse anti-SSEA-4 (MC-813-70, 1:2) and mouse anti-SSEA-3 (MC-631, 1:2) from the Developmental Studies Hybridoma Bank at the University of Iowa, mouse anti-SMA (Sigma, 1:400), mouse anti-Tuj1 (Covance, 1:500), mouse Oct-3/4 (Santa Cruz, 1:100), rabbit anti- α -fetoprotein (Dako, 1:400), goat anti-Nanog (Everest Biotech; 1:100), goat anti-FoxA2 (R&D Biosystems, 1:100), rabbit anti-TH (Sigma, 1:1000), mouse anti-TH (Chemicon, MAB5280, 1:1000), rabbit anti-Girk2 (Sigma, 1:40), rat anti-DAT (Chemicon, MAB369, 1:400), rabbit anti-cleaved Caspase-3 (Cell Signalling, 1:400), rabbit anti-LC3 (Cell Signalling, 1:100), mouse anti-LAMP1 (U. Iowa Developmental Hybridoma Bank, 1:100), rabbit anti-p62 (Enzo Life Sciences, 1:200), chicken anti-GFP (Aves Labs, 1: 500), rabbit anti-LRRK2 (MJFF C41-2, kindly provided by MJFF, 1:100), mouse anti-chondroitin sulphate (Sigma, 1:400), rabbit anti-Nurr1 (Santa Cruz, 1: 50) and mouse anti-SNCA (Ab-2, Thermo Scientific, 1:1000). Secondary antibodies used were all the Alexa Fluor Series from Invitrogen (all 1:500). Images were taken using Leica SP5 confocal microscope. For quantification of the DAn yield randomly 300 cells per differentiated aggregate were counted (average 6–8 differentiated aggregates per experiment). Data points represent the average of at least three-independent experiments. To visualize nuclei, slides were stained with 0.5 μ g/ml DAPI (4',6-diamidino-2-phenylindole) and then mounted with PVADABCO. Direct AP activity was analysed using an AP blue/red membrane substrate solution kit (Sigma) according to the manufacturer's guidelines.

Neurite analysis

Neurite analysis was performed at the indicated time-points on iPSC-derived neurons differentiated on top of cortical mouse astrocytes fixed and stained for TH. We randomly selected a minimum of 20 DAn per iPSC line (on the only condition that were isolated from surrounding DAn, so that we could unambiguously ascribe neurites to a single DAn), using a Leica SP5 confocal microscope, and analysed with the ImageJ plugin NeuronJ to determine the number and length of neurites per cell.

Fibroblast culture and treatment for autophagy studies

Fibroblasts from all patients were cultured in IMDM (Invitrogen) supplemented with 10% FBS (Gibco) and 2 mM Glutamax (Invitrogen). Fibroblasts were grown until reaching 65–75% confluence, when they were used as controls or treated for 6 h with rapamycin (0.2 or 2 μ M) and 100 μ M leupeptin. After the treatment, some cells were fixed and analysed by immunofluorescence for LC3 and the rest was processed as a pellet for Western blot analysis for LC3. For LC3 flux studies in fibroblasts, cells were treated for 2 and 4 h in supplemented IMDM containing lysosomal inhibitors, consisting in 20 mM NH₄Cl (Sigma) and 200 μ M leupeptin, after the treatment a pellet was obtained using 0.25% trypsin-EDTA.

Protein extraction and Western blotting

Proteins were extracted using RIPA protein extraction buffer (Sigma) supplemented with protease inhibitor (Sigma) and phosphatase inhibitor cocktails (Sigma). In brief, cells were detached from culture dishes, centrifuged and washed once with cold PBS. Cell pellets were then resuspended and incubated in RIPA buffer for 20 min on ice and sonicated in one pulse of 5 s at 10% amplitude on ultrasonicator (Branson Digital Sonifier, Branson Ultrasonics Corporation, Danbury, CT, USA) to ensure high efficiency of lysis. After centrifugation, supernatants of total protein extracts were denatured in loading buffer for 10 min at 98°C. Proteins were separated using SDS-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to PVDF membrane. The membrane was probed with anti- α -tubulin (Sigma T6074; 1:10,000), anti-LC3 (Cell Signalling; 1:1000) and anti-LRRK2 (MJFF2 c41-2; 1:50).

Electron microscopy and morphometric analysis

Cells were fixed in 2.5% glutaraldehyde/2% paraformaldehyde in 100 mM sodium cacodylate, pH 7.43 (SC) and post-fixed in 1% osmium tetroxide in SC followed by 1% uranyl acetate. After ethanol dehydration and embedding in LX112 resin (LADD Research Industries), ultrathin sections were stained with uranyl acetate followed by lead citrate. All grids were viewed on a JEOL 100CX II transmission electron microscope at 80 kV. Morphometric analysis was performed using ImageJ in 15–20 different micrographs for each condition after thresholding. Autophagic vacuoles were identified using previously established criteria (Dunn, 1990; Nixon et al, 2005). Autophagic vacuoles (vesicles <0.5 μ m) were classified as autophagosomes when they met two or more of the following criteria: double membranes (complete or at least partially visible), absence of ribosomes attached to the cytosolic side of the membrane, luminal density similar to cytosol and identifiable organelles or regions of organelles in their lumen. Vesicles of similar size but with a single membrane (or less than 40% of the membrane visible as double), luminal density lower

The paper explained

PROBLEM:

Parkinson's disease (PD) has a lifetime risk of 2%, making it the second most common neurodegenerative disease after Alzheimer's disease. It is associated to a selective loss of dopaminergic neurons (DAn) from the substantia nigra pars compacta responsible for controlling body movement. Around 90% of PD cases are sporadic, while 10% have a familial or genetic origin, of which mutations in the LRRK2 gene are the most frequent cause. The pathogenic mechanisms leading to PD are not well understood, to a large extent owing to the lack of suitable genetic models of this disease in animals. Therefore, there is an urgent need for developing reliable experimental models that recapitulate the salient features of PD. Recently, induced pluripotent stem cells (iPSC) have been used to model human diseases, although it is currently unclear if this approach would be able to model age-related diseases.

RESULTS:

We generated iPSC from sporadic PD patients, PD patients with LRRK2 mutation and healthy controls. All of these iPSC lines were comparable in their ability to generate DAn, and no major differences were found in these neurons under conventional

culture conditions, except for increased accumulation of SNCA in neurons from LRRK2 mutant iPSC. However, when cultured for over 2.5 months (a condition mimicking aging *in vitro*), DAn differentiated from patient-specific iPSC developed evident signs of neurodegeneration, including fewer and shorter neurites and a significant increase in apoptotic cells. Most importantly, this degeneration occurred in neurons from either sporadic PD or mutant PD, but not from healthy individuals. We then show that these alterations are caused by a deficient autophagic machinery.

IMPACT:

We have generated an iPSC-based model for PD that recapitulates key pathological features of PD such as neurodegeneration and SNCA accumulation in DAn generated from PD-patients. Our findings indicate that the bases of the increased susceptibility of DAn to degenerate upon aging in sporadic PD patients should be encoded in their genome. Moreover, our iPSC-based PD model provides a valuable tool to investigate the pathogenic mechanisms of PD, such as altered autophagy, and to screen for drugs that may prevent or rescue neurodegeneration in PD.

than the surrounding cytosol, multiple single membrane-limited vesicles containing light or dense amorphous material were classified as autophagolysosomes.

RT-qPCR analyses

Total mRNA was isolated by guanidinium thiocyanate–phenol–chloroform extraction (TRIzol, Invitrogen) and treated with DNaseI. One microgram was used to synthesize cDNA with the SuperScript III Reverse Transcriptase Synthesis Kit (Invitrogen). Quantitative PCR analyses were done in triplicate on 50 ng with Platinum Syber Green qPCR Super Mix (Invitrogen) in an ABI Prism 7000 thermocycler (Applied Biosystems). All results were normalized to HPRT and β -2-microglobulin. The primers used were: B2micro-F, 5'-GCCGTGTGAAC-CATGTGACT-3'; B2micro-R, 5'-GCTTACATGTCT CGATCCCACTT-3'; HPRT-F, 5'-TTATGGACAGGACTGAACGTCTTG-3' HPRT-R, 5'-GCACACAGAGGGC-TACAATGTG-3'; NANOG-F, 5'-ACAACCTGGCCGAAGAATA GCA-3'; NANOG-R, 5'-GGTTCCTCCAGTCGGGTTTCCAC-3'; FOXA2-F, 5'-AGACTCCTGC TTCTCAAGCACCT-3'; FOXA2-R, 5'-ACTTCCTGCAACAACAGCAATGG-3'; LRRK2-F, 5'-GGGAGCAGATGCCAATCAAGCAA-3'; LRRK2-R, 5'-TCCAC-CAATTT GGGACTGTCTCT-3'; SNCA-F, 5'-AGCGGACCTCCACAAGTAAC-GAAT-3'; SNCA-R, 5'-TTGGCATCTGTCTTCTCCCAAGT-3'; TAU-F, 5'-ACGAAGCGCTGAAGCACA GGATTA-3'; TAU-R, 5'-TCCTTTGGGACTGCCAT-GAGACTT-3'.

Statistical analysis

Differences among groups were evaluated by one-way ANOVA, and comparisons between two groups by Student's *t*-test, using the StatPlus build for Microsoft Excel.

Author contributions

ASD differentiated DAn from iPSC, performed neurodegeneration and autophagy studies and analyzed data; YRP and SJD generated and characterized iPSC lines; ICC and MV analysed SNCA and MPP+ susceptibility; CC, ME and ET identified and recruited the PD patients, and obtained authorization for the study; SM performed LC3 flux studies; CDG helped with iPSC cultures; BP performed electron microscopy studies; AMC designed the autophagy studies, analysed data and edited the manuscript; AG, JMC, MM, JA and JLB provided materials; AC conceived, designed and supervised the study, and edited the manuscript; AR generated iPSC lines, conceived, designed and supervised the study and wrote the manuscript.

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Supporting information is available at EMBO Molecular Medicine online.

The authors declare that they have no conflict of interest.

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– CURRICULUM VITÆ –

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Education & Training

1984-1990:	MD	School of Medicine, University of Valencia, Spain.
1990-1995:	PhD	Department of Physiology. University of Valencia, Spain.
1995-2000:	Postdoc	Biochemistry & Molecular Biology, IIC/FVIB, Valencia, Spain
2000-2004	Postdoc	Developmental Biology, Salk Institute, La Jolla, CA

Employment History

1991-1995:	Predoctoral Fellow. Department of Physiology, University of Valencia, Spain. Supervisor: Joaquín Romá.
1994,1995:	Special Project Associate. Peripheral Nerve Center, Department of Neurology, Mayo Clinic, Rochester, MN, USA. Supervisor: Phillip A. Low.
1995-1999:	Postdoctoral Research Fellow. Molecular Pathology Unit. Instituto de Investigaciones Citológicas, FVIB. Valencia, Spain. Supervisor: Juan Saus.
1996:	Assistant Professor. Department of Physiology, University of Valencia, Spain.
1999-2000:	Senior Research Associate. Molecular Pathology Unit. Instituto de Investigaciones Citológicas, FVIB. Valencia, Spain. Supervisor: Juan Saus.
2000-2004:	Research Associate. Gene Expression Laboratory. Salk Institute for Biological Studies, La Jolla, CA, USA. Supervisor: Juan Carlos Izpisúa-Belmonte.
2004-2006:	Senior Research Associate. Gene Expression Laboratory. Salk Institute for Biological Studies, La Jolla, CA, USA. Supervisor: Juan Carlos Izpisúa-Belmonte.
2006-present:	ICREA Research Professor
2006-2009:	Scientific Coordinator, Center for Regenerative Medicine in Barcelona, Barcelona, Spain.
2008-present:	Group Leader, Networking Center of Biomedical Research in Bioengineering, Biomaterials and Nanomedicine (CIBER-BBN)
2009-present:	Group Leader, Institute for Bioengineering of Catalonia, Barcelona Science Park, Barcelona, Spain.

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BIOGRAPHICAL SKETCH

Provide the following information for the key personnel and other significant contributors in the order listed on Form Page 2.
Follow this format for each person. **DO NOT EXCEED FOUR PAGES.**

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EDUCATION/TRAINING (Begin with baccalaureate or other initial professional education, such as nursing, and include postdoctoral training.)			
INSTITUTION AND LOCATION	DEGREE (if applicable)	YEAR(s)	FIELD OF STUDY
- University of Barcelona	MD	1963-1967	Medicine
- Hartford Hospital, Hartford, Conn., USA		1968-1969	Internship
- Dept. Neurol, U Minnesota Hospital, Univ. Minnesota, USA		1969-1972	Residency in Neurology
- University of Barcelona	PhD	1984	Ph.D. in Medicine

A. Positions

1974-1975 Visiting Assistance Scientist Brookhaven National Laboratory, Upton LI, New York, USA

1976-1979 Staff, Dept. Neurology University Minnesota Hospital, Mpls., Minn. USA

1981-1982 Chief, Neurology Service, Instituto Neurológico Municipal, Barcelona, Spain

1982-2007 Chief, Neurology Service, Hospital Clínic Universitari,

1993-2007 Director, University of Barcelona Brain Bank, Universitat de Barcelona, Spain

Since January 2007 Senior Consultant , Neurology Service, Hospital Clínic Universitari, Barcelona, Spain

Since 1984 .- 2009 Director, Movement Disorders Centre, Hospital Clínic Universitari, Barcelona.

Since January 2010 Director Parkinson Disease Research IDIBAPS, University of Barcelona.

Since 2010 – Steering Committee of CIBERNED (Centro Investigación Biomédica en Red Enfermedades Neurodegenerativas)