



LUND UNIVERSITY

Generation of midbrain dopaminergic neurons in vivo and in vitro: the role of Neurogenin2

Andersson, Elin

2005

[Link to publication](#)

Citation for published version (APA):

Andersson, E. (2005). *Generation of midbrain dopaminergic neurons in vivo and in vitro: the role of Neurogenin2*. [Doctoral Thesis (compilation), Department of Experimental Medical Science]. Department of Experimental Medical Science, Lund University.

Total number of authors:

1

General rights

Unless other specific re-use rights are stated the following general rights apply:

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

Read more about Creative commons licenses: <https://creativecommons.org/licenses/>

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

LUND UNIVERSITY

PO Box 117
221 00 Lund
+46 46-222 00 00

*Sektionen för Neurovetenskap
Institutionen för Experimentell Medicinsk Vetenskap*

**Generation of midbrain dopaminergic neurons *in vivo* and *in vitro*:
the role of *Neurogenin2***

**Akademisk avhandling
av
Elin Andersson**

Som med vederbörligt tillstånd av Medicinska Fakulteten vid Lunds Universitet för
avläggande av doktorsexamen i medicinsk vetenskap kommer att offentligen försvaras i
Segefalksalen, Wallenberg Neurocentrum, Lund

Lördagen den 17 december 2005, kl 9.30

Fakultetsopponent:
Professor Thomas Perlmann
Institutionen för Cell- och Molekylärbiologi, Karolinska Institutet
Stockholm

Organization LUND UNIVERSITY Section for Neuroscience Department of Experimental Medical Science Wallenberg Neuroscience Center BMC A11 221 84 Lund	Document name DOCTORAL DISSERTATION	
Author(s) Elin Andersson	Date of issue 2005-12-17	
Sponsoring organization		
Title and subtitle Generation of midbrain dopaminergic neurons <i>in vivo</i> and <i>in vitro</i> : the role of Neurogenin2		
<p>Abstract Parkinsons disease (PD) is a neurodegenerative disorder where dopaminergic neurons of the substantia nigra (SNc) in the mesencephalon are progressively eliminated. The ensuing loss of dopaminergic innervation of the basal ganglia manifests itself as severe motor deficits in PD patients. Clinical trials have shown that cell replacement therapy, where dopaminergic neuroblasts derived from fetal ventral mesencephalon (VM) are transplanted to the striatum, may be an alternative to pharmacological treatment of PD patients. The limited access and ethical concerns with using fetal tissue have prompted the use of stem cells as a renewable and limitless source of dopaminergic neurons. However, the mechanisms of specification of mesDA neurons <i>in vivo</i> need to be elucidated for identification and generation of mesencephalic dopaminergic (mesDA) neurons from stem cells <i>in vitro</i>.</p> <p>In this thesis I have identified expression of the proneural gene <i>Neurogenin2</i> (<i>Ngn2</i>) in a restricted pattern in the embryonic VM during mesDA neurogenesis. The protein was expressed in the progenitor population in the ventricular zone but not in mature neurons in the mantle zone. When isolating the <i>Ngn2</i>-expressing cells and their direct descendants by FACS from an <i>Ngn2-GFP-KI</i> mouse, I found that the <i>Ngn2</i>-GFP-positive cell fraction contained dopaminergic neurons, in contrast to <i>Ngn2</i>-GFP-negative cells. This shows that <i>Ngn2</i> label early mesDA neuron precursors. Furthermore, when I analysed the <i>Ngn2</i> knockout mutants, I found that they displayed an early loss of mesDA neurons that was partially maintained at postnatal stages, showing that <i>Ngn2</i> has a role in the generation of the mesDA neurons. No other neuronal subtype in the VM was affected suggesting that this role for <i>Ngn2</i> is specific for the mesDA neurons.</p> <p>Using embryonic mouse tissue obtained at the stage of mesDA genesis, I was able to generate cultures of neural stem and progenitor cells, so called neurosphere cultures, that were neurogenic and maintained a ventral midbrain character over several passages. Although the neurospheres did not spontaneously give rise to dopaminergic neurons when differentiated, TH-positive cells were detected when <i>Nurr1</i> was over-expressed in the cultures. The frequency with which this occurred, and the morphology of the TH-positive cells, differed from the results obtained when over-expressing <i>Nurr1</i> in forebrain-derived expanded cells. This suggests that neurosphere expanded cells derived from VM specifically contain progenitors that can generate dopaminergic neurons under certain conditions. When over-expressing <i>Ngn2</i> together with <i>Nurr1</i> TH-positive cells were generated that displayed a mature neuronal morphology. Furthermore, I found that they expressed other dopaminergic markers which were not seen when either <i>Nurr1</i> or <i>Ngn2</i> were over-expressed alone. This suggests that <i>Nurr1</i> and <i>Ngn2</i> interact to specify a more mature dopaminergic phenotype.</p> <p>The results in this thesis have identified a new cellular marker of mesDA progenitors in the developing embryo and also provided new insight into the development of mesDA neurons.</p>		
Key words: stem cell, progenitor cell, neurospheres, bHLH, transcription factors, proneural genes, neuronal differentiation, TH, mesencephalon, development, neuronal specification, dopamine, Parkinson's, cell replacement		
Classification system and/or index terms (if any):		
Supplementary bibliographical information:	Language English	
ISSN and key title: 1652-8220	ISBN 91-85439-09-6	
Recipient s notes	Number of pages 176	Price
	Security classification	

Distribution by (name and address)

I, the undersigned, being the copyright owner of the abstract of the above-mentioned dissertation, hereby grant to all reference sources permission to publish and disseminate the abstract of the above-mentioned dissertation.

Signature



Date 2005-11-07

**Generation of midbrain dopaminergic
neurons *in vivo* and *in vitro*:**
the role of *Neurogenin2*

Elin Andersson, M.Sc



LUND
UNIVERSITY

Cover: TH-positive cells (red) with mature neuronal morphology and elaborate arborisation are generated from expanded fetal midbrain cells transduced simultaneously with retroviral constructs containing Nurr1 and Neurogenin2 (front). TH-positive cells with more immature morphology are generated after transduction with Nurr1 only (back). Cells also label for reporter gene GFP (green) which show that they are derived from transduced cells.

ISBN 91-85439-09-6

© 2005 Elin Andersson

Printed by Grahn's Tryckeri AB, Lund, Sweden

I made this!

*My soul, sit thou a patient looker-on;
Judge not the play before the play is done:
Her plot hath many changes; every day
Speaks a new scene; **the last act crowns the play.***

*Francis Quarles,
Epigram, Respice Finem*

LIST OF CONTENTS

ORIGINAL PAPERS.....	11
ABBREVIATIONS.....	12
SUMMARY.....	15
POPULÄRVETENSKAPLIG SAMMANFATTNING.....	16
INTRODUCTION.....	19
Dopaminergic neurons in the brain	19
Location of dopaminergic neurons in the brain.....	19
Projections and functions of midbrain dopaminergic neurons.....	20
Identification of dopaminergic neurons.....	21
Development of mesDA neurons	21
Patterning of the midbrain.....	21
Neurogenesis of mesDA neurons.....	23
Origin and organization of mesDA neurons.....	23
Genes involved in development of mesDA neurons.....	23
Cell replacement therapy	27
Stem cells	27
Stem cells – definitions and concepts.....	27
Neural stem cells.....	28
Neurospheres as a way to expand neural stem cells.....	30
Expansion of midbrain neural stem and progenitor cells.....	30
AIMS OF THIS THESIS.....	31
RESULTS AND COMMENTS.....	35
Identifying early midbrain neural progenitors	35
Sox-genes are differentially expressed in the midbrain.....	35
Neurogenin2 labels progenitor cells in the ventral midbrain (Paper I and II).....	36
Fate of Ngn2-expressing cells in the midbrain	38
Fate mapping of the Ngn2-positive cells reveals that they are mesDA precursors (Paper I).....	38
Ngn2-expressing precursors also give rise to other types of neurons (Paper I and II).....	39
The role of Ngn2 in mesDA neuron development	39
Absence of Ngn2 results in a dramatic loss of mesDA neurons (Paper II).....	39

Ngn2 is important for the mesDA neuron precursors specifically (Paper II).....	39
Over-expression of Ngn2 does not induce dopaminergic differentiation (Paper II).	40
Potential of the <i>Ngn2</i>-GFP positive cells.....	40
<i>In vitro</i> differentiation (Paper I).....	40
<i>In vivo</i> differentiation (Paper I).....	40
Expanding neural stem and progenitor cells from the ventral midbrain.....	42
Neurosphere cultures (Paper III).....	42
Differentiation potential of progenitors from ventral midbrain expanded as neurospheres (Paper III).....	42
Neurospheres from E11.5 ventral midbrain are regionally specified (Paper III).....	43
Over-expression of Ngn2 and Nurr1 together generate a more complete mesDA phenotype (Paper III).....	43
DISCUSSION.....	47
Phenotype of the mitotic progenitor in the ventral midbrain.....	47
Dynamic expression of Ngn2 within the VZ.....	47
Possible interaction of Ngn2 with Sox2 in neurogenesis of ventral midbrain cells..	48
Ngn2 involved in acquiring full neuronal phenotype in mesDA neuron precursors..	49
Does Ngn2 have a role in both differentiation and specification of mesDA neurons?...	49
Possible mechanisms for Ngn2 fate specification.....	50
Interaction of Ngn2 and homeodomain proteins.....	50
Induction of mesDA phenotype <i>in vitro</i> by over-expression of Ngn2 and Nurr1....	51
What about the lateral population?.....	52
Induction of Ngn2 in ventral midbrain.....	52
Why is it so hard to get a dopaminergic neuron <i>in vitro</i>?.....	53
Are we expanding the correct progenitor?.....	54
MATERIAL AND METHODS.....	57
Experimental animals.....	57
PCR genotyping.....	57
Histological analysis.....	57
Preparation of tissue.....	57
Immunohistochemistry.....	57
<i>In situ</i> hybridization.....	57

Tissue culture	59
Dissection.....	59
Neurosphere cultures.....	59
Differentiation of neurospheres.....	59
Primary cultures and co-cultures.....	60
Immunocytochemistry.....	60
RT-PCR.....	60
FACS procedure	61
Transplantation	61
BrdU labeling of embryonic tissue and primary cells	61
Retroviral transduction	61
List of mediums	62
PCR primers and programs	63
List of antibodies	64
REFERENCES	67
ACKNOWLEDGEMENTS	77
APPENDIX	81
Paper I	85
Paper II	111
Paper III	135
Colour plates	161

ORIGINAL PAPERS

- I. Lachlan H. Thompson*, Elin Andersson*, Josephine B. Jensen, Perrine Barraud, Francois Guillemot, Malin Parmar, and Anders Björklund
Neurogenin2 identifies a transplantable dopamine neuron precursor in the developing ventral mesencephalon
* equal contribution

Submitted to Experimental Neurology

- II. Elin Andersson*, Josephine B. Jensen*, Malin Parmar, Francois Guillemot, and Anders Björklund
Development of the mesencephalic dopaminergic neuron system is compromised in the absence of Neurogenin2
* equal contribution

Accepted in Development

- III. Elin KI Andersson, Dwain K Irvin, Jessica Ahlsjö, Emeli Nilsson, and Malin Parmar
Ngn2 and Nurr1 facilitates dopaminergic neuron differentiation from neurosphere expanded ventral mesencephalic cells

Submitted to Molecular and Cellular Neuroscience

ABBREVIATIONS

bFGF	Basic fibroblast growth factor
bHLH	Basic helix loop helix
BMP	Bone morphogenetic protein
BrdU	5-bromo-2-deoxyuridine
cDNA	Complementary DNA
CKO	Conditional knock-out
CNS	Central nervous system
DA	Dopamine
DAPI	4'-6-Diamidino-2-phenylindole
E	Embryonic day
EGF	Epidermal growth factor
ES	Embryonic stem (cell)
FACS	Fluorescence activated cell sorting
FGF8	Fibroblast growth factor 8
GABA	Gamma-aminobutyric acid
GFP	Green fluorescent protein
HD	Homeodomain (protein)
HEPES	4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid
IHC	Immunohistochemistry
IZ	Intermediate zone
L-DOPA	levodopa, 3,4-dihydroxy-L-phenylalanine
MesDA	Mesencephalic dopamine (neuron)
MHO	Mid-hindbrain organizer
MZ	Mantle zone
PCR	Polymerase chain reaction
PD	Parkinson's disease
PNS	Peripheral nervous system
RRF	Retrorubral field
SHH	Sonic hedgehog
SNc	Substantia nigra pars compacta
SVZ	Sub-ventricular zone
TH	Tyrosine hydroxylase
TUNEL	Terminal deoxynucleotidyl Transferase Biotin-dUTP Nick End Labeling
VM	Ventral midbrain
VTA	Ventral tegmental area
VZ	Ventricular zone

Summary

SUMMARY

Parkinsons disease (PD) is a neurodegenerative disorder where dopaminergic neurons of the substantia nigra (SNc) in the mesencephalon are progressively eliminated. The ensuing loss of dopaminergic innervation of the basal ganglia manifests itself as severe motor deficits in PD patients. Clinical trials have shown that cell replacement therapy, where dopaminergic neuroblasts derived from fetal ventral mesencephalon (VM) are transplanted to the striatum, may be an alternative to pharmacological treatment of PD patients. The limited access and ethical concerns with using fetal tissue have prompted the use of stem cells as a renewable and limitless source of dopaminergic neurons. However, the mechanisms of specification of mesDA neurons *in vivo* need to be elucidated for identification and generation of mesencephalic dopaminergic (mesDA) neurons from stem cells *in vitro*.

In this thesis I have identified expression of the proneural gene *Neurogenin2* (*Ngn2*) in a restricted pattern in the embryonic VM during mesDA neurogenesis. The protein was expressed in the progenitor population in the ventricular zone but not in mature neurons in the mantle zone. When isolating the *Ngn2*-expressing cells and their direct descendants by FACS from an *Ngn2-GFP*-KI mouse, I found that the *Ngn2-GFP*-positive cell fraction contained dopaminergic neurons, in contrast to *Ngn2-GFP*-negative cells. This shows that *Ngn2* label early mesDA neuron precursors. Furthermore, when I analysed the *Ngn2* knockout mutants, I found that they displayed an early loss of mesDA neurons that was partially maintained at postnatal stages, showing that *Ngn2* has a role in the generation of the mesDA neurons. No other neuronal subtype in the VM was affected suggesting that this role for *Ngn2* is specific for the mesDA neurons.

Using embryonic mouse tissue obtained at the stage of mesDA genesis, I was able to generate cultures of neural stem and progenitor cells, so called neurosphere cultures, that were neurogenic and maintained a ventral midbrain character over several passages. Although the neurospheres did not spontaneously give rise to dopaminergic neurons when differentiated, TH-positive cells were detected when *Nurr1* was over-expressed in the cultures. The frequency with which this occurred, and the morphology of the TH-positive cells, differed from the results obtained when over-expressing *Nurr1* in forebrain-derived expanded cells. This suggests that neurosphere expanded cells derived from VM specifically contain progenitors that can generate dopaminergic neurons under certain conditions. When over-expressing *Ngn2* together with *Nurr1* TH-positive cells were generated that displayed a mature neuronal morphology. Furthermore, I found that they expressed other dopaminergic markers which were not seen when either *Nurr1* or *Ngn2* were over-expressed alone. This suggests that *Nurr1* and *Ngn2* interact to specify a more mature dopaminergic phenotype.

The results in this thesis have identified a new cellular marker of mesDA progenitors in the developing embryo and also provided new insight into the development of mesDA neurons.

POPULÄRVETENSKAPLIG SAMMANFATTNING

I hjärnan finns många olika typer av nervceller. De använder sig av olika signalsubstanser, kallade *neurotransmittorer*, för att kommunicera med andra nervceller. En viss typ av nervceller använder neurotransmittorn *dopamin*. Dopaminceller finns på många ställen i hjärnan men de flesta ligger i mellanhjärnan i ett par olika cellgrupper som var och en skickar signaler till sina specifika områden i andra delar av hjärnan. En av dessa cellgrupper kallas *substantia nigra* och signalerar till ställen som styr en människas motorik. Hos patienter med Parkinsons sjukdom, dör cellerna i denna grupp och då försvinner även dopaminsignalerna till de delar som styr motoriska förmågor. Därför har Parkinson-patienter typiska symptom, som problem med motoriken och svårighet att sätta igång rörelser. För att lindra dessa symptom kan Parkinson-patienter ta medicin som ska ersätta dopaminet. Man har också testat andra behandlingsmetoder som går ut på att ersätta dopamincellerna inuti hjärnan. Genom att ta dopaminceller från fostervävnad och transplantera till hjärnan har man lyckats återskapa dopaminsignalleringen utan mediciner. Tyvärr kan denna teknik ännu inte tillämpas på många patienter eftersom det är svårt att få tag på tillräckligt mycket vävnad. Man har därför börjat undersöka hur man kan generera dopaminceller på annat sätt. En metod är att använda stamceller, celler som kan förökas i kultur och som kan utvecklas till vilka sorters celler som helst. För att få stamcellerna att bli dopaminceller så måste man veta vad det är som gör att just den sortens nervceller bildas. Vi måste förstå vilka de bakomliggande faktorerna är som styr cellutvecklingen mot dopaminceller.

I min avhandling har jag undersökt vilka signaler och gener som är viktiga för att dopaminceller ska bildas. För att ta reda på det har jag tittat på dopaminceller under fosterutvecklingen. I mina studier använde jag möss som en modell för vad som händer i människan. Jag fann att en gen, *Neurogenin2*, var påslagen (uttryckt) i precis de celler som skulle bli dopaminceller hos mössfoster. När jag sedan undersökte muterade möss där denna gen var borttagen såg jag att dopamincellerna i mellanhjärnan inte bildades som de skulle. Detta visar att *Neurogenin2* är viktig för bildandet av dopaminceller. Jag försökte också påverka odlade stamceller att utvecklas till dopaminceller genom att se till att *Neurogenin2* uttrycktes i cellerna. När jag uttryckte *Neurogenin2* tillsammans med en annan gen, *Nurr1*, som också är viktig för att det ska bli dopaminceller, gav det bättre resultat än att använda dem var och en för sig och jag såg att det bildades dopaminceller i cellkulturerna.

Med resultaten som presenteras i den här avhandlingen har vi kommit ännu en bit på väg för att veta hur dopaminceller genereras. Mina resultat kan användas bl.a för att identifiera celler som ska bli dopaminceller. I ett längre perspektiv kan mina resultat bidra till att man kan generera dopaminceller från stamceller och därmed ge de patienter som lider av Parkinsons sjukdom en alternativ behandling.

Introduction

INTRODUCTION

Dopamine is a neurotransmitter employed by specific neurons in the central nervous system (CNS). Dopaminergic neurons are involved in neural processes as diverse as neuroendocrine hormonal release, cognition, emotion, reward and initiation of motor responses, however it is for their role in the neurodegenerative disorder Parkinson's disease (PD) that so much interest has been placed upon this subtype of CNS neuron. The neuropathology of PD is a gradual loss of mesencephalic (midbrain) dopaminergic neurons and their innervation of the basal ganglia in the ventral forebrain (for review see Lang and Lozano, 1998a; Lang and Lozano, 1998b). It is predominantly the dopaminergic neurons in the substantia nigra, one of the nuclei in the midbrain, that are affected in PD. Attempts to replace lost dopaminergic transmission in the basal ganglia by transplantation of immature dopaminergic neuroblasts from fetal tissue have proved to be a viable approach, in animal models of PD but also in PD patients (Lindvall et al., 1990; for review see Dunnett and Bjorklund, 1999; Winkler et al., 2005). Since cell replacement therapy requires large numbers of transplantable dopaminergic neurons and the yield from fetal tissue is limited, stem cells have been suggested and explored as an option to meet this need. Advances within the field of stem cell research in early 1990s saw the emergence of protocols how to grow and maintain neural stem cells in culture (Reynolds et al., 1992; Reynolds and Weiss, 1992; Reynolds and Weiss, 1996). This fuelled research on the specification of dopaminergic neurons during embryogenesis and how this could be applied to the *in vitro* generation of dopaminergic neurons from stem cells.

The aim of this thesis work has been to further elucidate mechanisms and events important for the development of the midbrain dopaminergic neurons *in vivo* and explore ways to generate this kind of neuron *in vitro*.

Dopaminergic neurons in the brain

Location of dopaminergic neurons in the brain

The dopaminergic neurons in the brain are organized into ten nuclei ranging from the caudalmost cell group A8, the retro-rubral field, to the rostralmost A17, a group of amacrine interneurons in the retina (Björklund and Lindvall, 1984; Dahlstrom and Fuxe, 1964). Small groups of dopaminergic neurons are located in e.g the olfactory bulb and the diencephalon (part of forebrain), however, the vast majority of dopaminergic neurons, around 75%, reside in nuclei in the ventral mesencephalon (VM): the substantia nigra pars compacta (SNc, A9), the ventral tegmental area (VTA, A10) and the retro-rubral field (RRF, A8) (Björklund and Lindvall, 1984; Dahlstrom and Fuxe, 1964) (Fig 1a). The dopaminergic neurons of the mesencephalic nuclei (the mesDA neurons) are often investigated together since they develop from the same progenitor location.

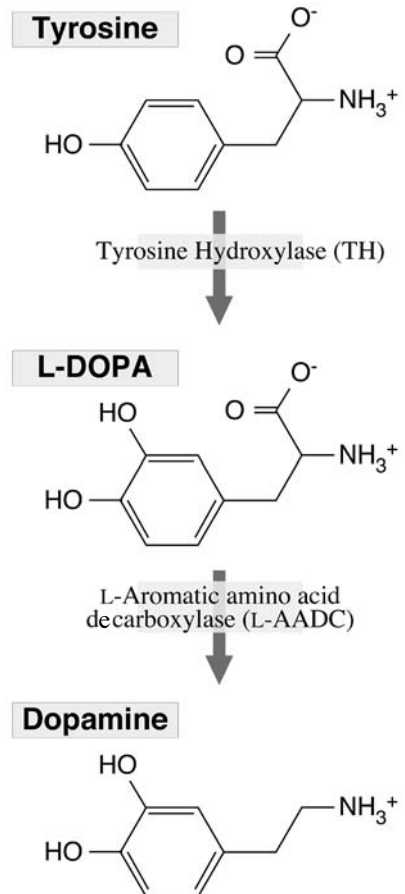
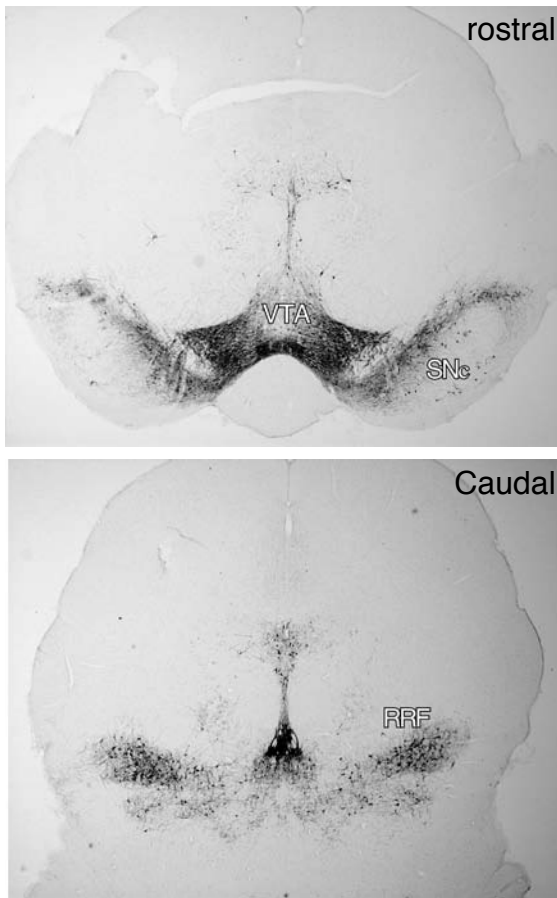


Fig. 1A) Coronal section of P18 mouse midbrain showing the localization and distribution of the midbrain dopaminergic nuclei ventral tegmental area (VTA), substantia nigra (SNc) and retro-rubral field (RRF) B) Enzymatic pathway of dopamine synthesis

Projections and functions of midbrain dopaminergic neurons

The dopaminergic neurons of the midbrain nuclei have distinct functions in the brain and consequently innervate separate structures. The dopaminergic neurons whose cellbodies are located in the SNc, and which are the neurons most affected in PD, project to the dorso-lateral striatum and caudate putamen forming the so-called nigrostriatal pathway (Ungerstedt, 1971). The nigrostriatal pathway modulates the output of these basal ganglia structures that, together with cortical areas control initiation of voluntary movement, posture etc. Since the nigrostriatal innervation is lost in PD patients, they have characteristic symptoms of motor dysfunction such as rigidity and slowness of movements that is accompanied by tremor (Lang and Lozano, 1998a; Lang and Lozano, 1998b). The VTA neurons innervate limbic areas in the ventro-medial

striatum (mesolimbic system) and pre-frontal cortex (mesocortical system) and are involved in emotion, cognitive processes and reward behaviours (Björklund and Lindvall, 1984; Ungerstedt, 1971). The RRF neurons send axons along the same pathways as VTA and SN.

Identification of dopaminergic neurons

Dopaminergic neurons are generally identified by their expression of the rate-limiting enzyme in the dopamine pathway, tyrosine hydroxylase (TH). TH modifies the amino acid tyrosine to dopamine precursor L-DOPA, which is in turn converted to dopamine by aromatic amino acid decarboxylase (AADC) (Fig 1b). Dopaminergic neurons share expression of proteins involved in production, storage and release of dopamine, such as TH, AADC, dopamine transporter (DAT) and vesicular monoamine transporter (VMAT2). Other proteins are differently expressed in different dopaminergic nuclei. These proteins are likely to play a role in the function of the particular dopaminergic neuron subtype but can be used simply as markers to distinguish them. The mesDA neurons have very similar expression profiles, more so during development than at adult stages. They can obviously be identified by location and projections in intact adult tissue, and their morphologies differ slightly (Thompson et al., 2005). At adult stages, mesDA neurons in SNc and VTA can also be distinguished by presence of the markers *Girk2* and *calbindin*, respectively (Liang et al., 1996; Schein et al., 1998; Thompson et al., 2005). Additionally, progenitor marker *Aldh1* (see below) is maintained preferentially in SNc neurons in adult (McCaffery and Drager, 1994). However, since mesDA neurons develop from the same cells, at approximately the same time, and no early marker has been reported to be expressed specifically by either, it is at present not possible to distinguish them during development.

Development of mesDA neurons

Patterning of the midbrain

Development of the CNS is initiated by the formation of neural ectoderm, the so-called *neural plate*. The neural plate will invaginate into a neural fold and subsequently close along the dorsal midline forming the neural tube (*neurulation*). The neural tube consists at this stage of one layer of dividing cells, the *neuroepithelial* cells. They will divide to give rise to all cells of the CNS. A rostro-caudal and dorsal-ventral patterning of the neural tube is established early which provides positional information to the dividing neuroepithelial cells and ensures that the neurons generated in a specific position is of the correct subtype. Local organizing centers are involved in defining such developmental compartments by secreting factors that influence surrounding tissue (Jessell and Sanes, 2000b; for review of patterning of the neural tube see Lumsden and Krumlauf, 1996).

MesDA neurons develop in close proximity to two organizing centers, the floorplate and the isthmus which control the dorso-ventral and anterior-posterior patterning, respectively. Floorplate cells were shown to ectopically induce dopaminergic neurons in dorsal midbrain (Hynes et al., 1995b). This inductive effect of floorplate cells is mediated by the protein Sonic hedgehog, SHH (Hynes et al., 1995a; Wang et al., 1995; Ye et al., 1998) which is secreted by the

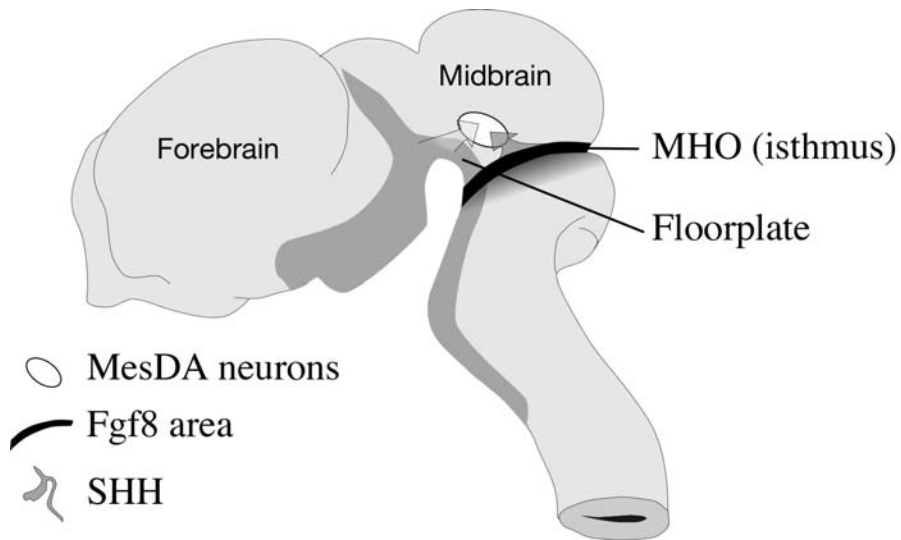


Fig. 2 Schematic drawing of a developing mouse brain, viewed from the side. MesDA neurons are located in the ventral midbrain, above the mesencephalic flexure. They develop at the site where signaling molecules FGF8, secreted from the mid-hindbrain organizer (MHO), and SHH from floorplate cells intersect.

floorplate cells all along the ventral neural tube (Fig 2). SHH is instrumental in defining ventral cell identities throughout the neuraxis, but its effect is most studied in the spinal cord (for review see Jessell, 2000). Interestingly, forebrain dopaminergic neurons are also generated under the influence of SHH (Ye et al., 1998) implicating SHH as a crucial factor for dopaminergic neuron development.

The isthmus, or the mid-hindbrain organizer (MHO) as it is also called, is a constriction of the neural tube separating the midbrain and the hindbrain. The MHO is established at the site of interaction between two transcription factors: *Otx2*, expressed in the early rostral neural tube that will later develop into the forebrain and midbrain, and *Gbx2* whose expression domain covers the presumptive hindbrain and spinal cord. *Otx2* and *Gbx2* interact by suppressing expression of the other to position the MHO (Broccoli et al., 1999; Wurst and Bally-Cuif, 2001). The MHO can be shifted caudally by ectopic expression of *Otx2* in the rostral hindbrain or rostrally by loss of *Otx* alleles, which expands the *Gbx2* expressing domain. Shifting the MHO caudally and rostrally increases and decreases the number of mesDA neurons, respectively (Broccoli et al., 1999; Brodski et al., 2003). The secreted factor Fibroblast growth factor 8 (FGF8) is responsible for the patterning effects of the MHO and FGF8 soaked beads were shown to induce an ectopic midbrain (Crossley et al., 1996). Dopaminergic neurons develop in the ventral neural tube just rostral to the MHO at a site where SHH and FGF8 signals intersect, indicating that the combined action of SHH and FGF8 controls the precise location of mesDA neurons and is essential for their formation (Ye et al., 1998) (Fig 2).

A number of transcription factors, *Lmx1b*, *Pax2*, *Pax5*, *En1* and *En2* and secreted signalling molecule *Wnt1* are expressed in or around the isthmus following its establishment (reviewed by Joyner, 1996; Liu and Joyner, 2001; Wurst and Bally-Cuif, 2001). A few of these, *En1*, *En2* and *Lmx1b* will continue to be expressed in maturing mesDA neurons and will then exercise functions other than general patterning of the mid-hindbrain region (see below).

Neurogenesis of mesDA neurons

Immunohistochemistry (IHC) for TH has shown that dopaminergic neurons start to appear at embryonic day E11 in mice (Foster et al., 1988) and E12.5 in rats (Specht et al., 1981). However, depending on detection method and staging of embryos TH-positive cells have been reported as early as E9.5 in mice (Di Porzio et al., 1990). Labelling studies in rats showed that mesDA neurons incorporated [³H]thymidine during E11-E15 with a peak at E13 (Altman and Bayer, 1981). Similar studies on mice showed that most mesDA neurons are born on E12 (E11.5 if the morning of the plug is E0.5), however neurons of the SNc and RRF are born slightly earlier than VTA neurons (Bayer et al., 1995). Neurogenesis of SNc neurons take place between E10-E13 with a peak at E11-E12 and neurogenesis of VTA neurons is ongoing from E10 to at least E14 with the majority being born at E12-13. Within the SNc and VTA there is also an anterior-posterior, lateral-medial gradient such that neurons in posterior parts of the nuclei are born later than neurons in the anterior parts and lateral regions contain more early generated neurons than medial parts (Bayer et al., 1995).

Origin and organization of mesDA neurons

Dopaminergic precursors are generated from the proliferative ventricular zone (VZ) overlying the ventral midline (Fig 3a). They are generated just anterior of the isthmus, as previously mentioned. As they become postmitotic they start to express TH and migrate ventrally along radial glia cells expressing vimentin and tenascin (Kawano et al., 1995; Shults et al., 1990) (Fig 3a). It is thought that the TH-positive cells migrate in this fashion until they reach the ventral pial surface in the mantle zone (MZ) when they instead follow tangentially oriented axons and migrate laterally to form the RRF, the SNc and the VTA (Kawano et al., 1995). However, since AADC, which is expressed 2 days prior to TH in dopaminergic precursors (Teitelman et al., 1983), is found lateral to the mesDA neuron domain it has been suggested that some precursors, that give rise to lateral SNc neurons, are generated from a more lateral progenitor population and migrate perpendicular to the ventricle (Hanaway et al., 1971; Smidt et al., 2004). The beginnings of mesDA nuclei are evident at E17.5 in mice, however the system continues to develop during the weeks after birth. For example, we have noted that the expression pattern of *Girk2* is not fully developed until three weeks after birth.

Genes involved in development of mesDA neurons

Many genes have been shown to be expressed in the mesDA neurons and are important for different aspects of maintenance and development of these neurons. These include *Aldh1*, *Nurr1*, *Pitx3*, *Lmx1b*, *En1* and *En2* (Fig 3b).

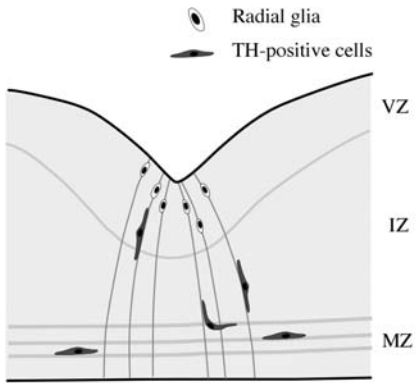
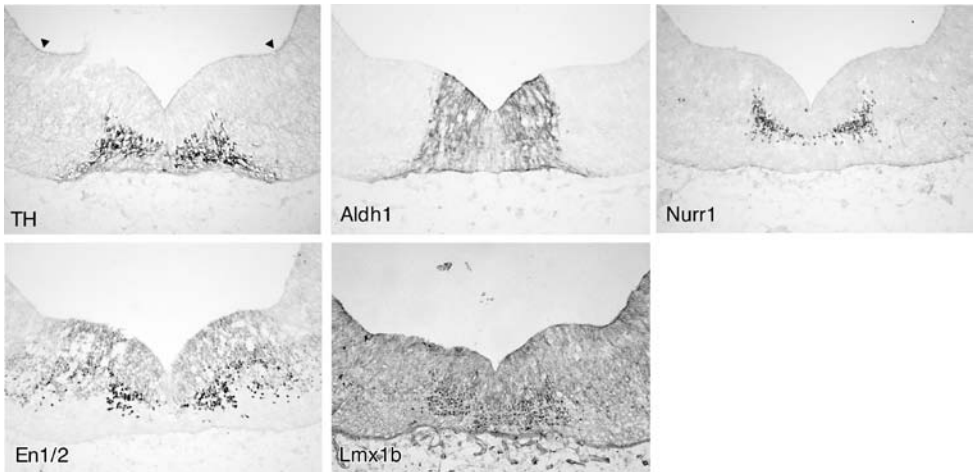
A

Fig. 3A) Schematic drawing of a coronal section of the VM during mesDA neurogenesis. Dividing progenitors of mesDA neurons are located in the ventricular zone (VZ). As they become postmitotic they start migrating ventrally along radial glia through the intermediate zone (IZ). When the mesDA precursors reach the mantle zone (MZ) above the pial surface, they start migrating laterally along horizontal fibers.

B) Coronal sections through the VM of an E11.5 mouse embryo. Developing dopaminergic neurons are identified by the expression of TH. They are first seen at around E11.5 in the MZ. *Aldh1* is expressed in the mesDA progenitors in the VZ and also in more mature mesDA precursors in the IZ and MZ. The postmitotic mesDA precursors also express other markers such as *Nurr1*, *En1/2* and *Lmx1b*. Black arrowheads mark the lateral sulcus which separates the VM from DM and was used as a morphological landmark for VM dissections.

B

Aldh1

Aldehyde dehydrogenase, *Aldh1* (also known as AHD-2) is an enzyme in the retinoic acid pathway, converting retinaldehyde to retinoic acid (Lindhahl and Evces, 1984). It is expressed as early as E9.5 in the midbrain. At this early stage the expression pattern coincides with *SHH* in a narrow wedge encompassing the ventral midline and presumably labels early proliferating dopaminergic precursors (Wallen et al., 1999). Immunostainings at E11.5 show that *Aldh1* at this stage is expressed in both the ventricular zone and budding mantle zone, where it co-localizes with *TH*-expressing neurons. It is thus a marker for both dopaminergic precursors and postmitotic mesDA neurons. At post-natal stages *Aldh1* is mainly confined to the mesDA

neurons of the SNc (McCaffery and Drager, 1994). The function of *Aldh1* in the development and/or maintenance of dopaminergic neurons is as yet unclear as no knockout mice or over-expression data for the gene have been presented.

Nurr1

Nurr1 (Nr4a2) is an orphan nuclear receptor transcription factor (no ligand identified as yet) and widely used as a marker for dopaminergic precursors. It is one of the first genes to be expressed in postmitotic mesDA neuron precursors, the protein expression is evident in midbrain from E10.5 in mice, preceding the expression of TH by about a day (Zetterstrom et al., 1997). *Nurr1* expression remains in the adult mesDA neurons (Backman et al., 1999; Zetterstrom et al., 1996). It is also expressed in other dopaminergic neurons such as the olfactory bulb A16 neurons (Backman et al., 1999). In addition, it is expressed in cells that are not dopaminergic, both during development and in adult tissue (Zetterstrom et al., 1996). *Nurr1* has been shown to bind to the promoter region of the *Th* gene (Iwawaki et al., 2000; Sakurada et al., 1999) and is essential for the neurotransmitter phenotype of dopaminergic neurons, but does not seem to be involved in neurogenesis and other aspects of mesDA neuron specification (Castillo et al., 1998; Saucedo-Cardenas et al., 1998; Smits et al., 2003). In *Nurr1* knockout mice, dopaminergic *Aldh1*-positive precursors are formed and differentiate to express transcription factor *En* and mesDA specific marker *Pitx3* (see below) but do not express TH (Castillo et al., 1998; Saucedo-Cardenas et al., 1998; Wallen et al., 1999; Zetterstrom et al., 1997). *Nurr1* mutants also lack expression of other proteins connected to the neurotransmitter phenotype, DAT and VMAT, and show lower levels of AADC (Castillo et al., 1998; Smits et al., 2003). The role of *Nurr1* is specific for the mesDA neurons as other DA neurons, that also express *Nurr1* during development, were not affected in the *Nurr1* mutants (Castillo et al., 1998; Saucedo-Cardenas et al., 1998; Zetterstrom et al., 1997). The dopaminergic precursors (expressing *Pitx3* and *Aldh1*) that are formed in *Nurr1* mutants, are eventually lost and an increase of apoptotic TUNEL stained cells is seen, demonstrating that *Nurr1* is necessary also for the survival of the dopaminergic neurons (Saucedo-Cardenas et al., 1998; Wallen et al., 1999).

En1/2

Engrailed 1 and 2 (*En1* and *En2*) are homologous homeobox transcription factors that have largely overlapping expression patterns, in particular in the VM (Davis and Joyner, 1988; Simon et al., 2001). *En1* and *2* are expressed early in the mesencephalic region (E8, Davis and Joyner, 1988) and also in the adult mesDA neurons, in the case of *En2* at high levels only by a subset of the neurons (Simon et al., 2001). *En1* and *2* are also expressed by cells in the hindbrain and dorsal mesencephalon but are not expressed by other dopaminergic neurons. *En1* single mutants have severe deletions in the cerebellum and inferior colliculus reflecting the normal expression in the dorsal mesencephalon, whereas *En2* mutants only have minor defects in cerebellar foliation (Joyner et al., 1991; Millen et al., 1994; Wurst et al., 1994). The mesDA neurons appear intact in the single mutants par for a minor alteration in cell density of the VTA in the *En1*^{-/-} mutant (Simon et al., 2001). Thus, in *En1* and *En2* single mutants, the related genes appear to compensate for each other with respect to their function in dopaminergic

neurons. On the contrary, in *En1*^{-/-};*En2*^{-/-} double mutants, TH-positive cells are formed but the expression domain is smaller and by E14 they are completely lost (Simon et al., 2001). The *Engrailed* genes are thus essential for the survival of dopaminergic neurons and have also been implicated in the regulation of α -synuclein (Simon et al., 2001).

Lmx1b

Lmx1b is a member of the LIM homeodomain protein family. Similar to *En1* and *En2*, *Lmx1b* have an early expression pattern in the midbrain. *Lmx1b* is first expressed in neural tissue at E7.5 and the expression is also seen in adult SNc and VTA neurons (Smidt et al., 2000). *Lmx1b* is thus expressed in the mesencephalic region before postmitotic mesDA markers *Nurr1*, *Pitx3* and TH and could be considered to be a marker of both a dopaminergic precursor cell and a mature mesDA neuron. However, it is not a specific precursor marker as it is not restricted to the part of VM that give rise to mesDA neurons. There is also some evidence that the early *Lmx1b* expression is downregulated in the dividing progenitors prior to mesDA neurogenesis and subsequently upregulated in post-mitotic mesDA neurons (unpublished data). In the *Lmx1b* knockout mutant TH-positive cells are formed but are reduced in number. The TH-positive cells that are formed express *Nurr1* but not *Pitx3*, pointing to a link between *Lmx1b* and *Pitx3*. The TH-positive cells in the *Lmx1b* mutant remain up to E16, after which TH-expression is lost (Smidt et al., 2000). This indicates that *Lmx1b* or its downstream targets are necessary for the development of certain mesDA neurons and the survival of the others.

Pitx3

Pitx3 is a paired-like homeobox transcription factor that is expressed exclusively in the mesDA neurons in the CNS. *Pitx3* mRNA is apparent at around E11.5 and overlaps completely with TH at E12.5 (Smidt et al., 1997). The expression continues in the developing mesDA neurons into adulthood. All SNc and VTA neurons in adults show expression of *Pitx3* (Smidt et al., 2004; Smidt et al., 1997; Zhao et al., 2004) although some studies report that only around 50% of the mesDA neurons are *Pitx3*-positive (van den Munckhof et al., 2003).

Several studies on the Aphakia mouse, shown to be a *Pitx3* mutant (Rieger et al., 2001; Semina et al., 2000), have revealed a near complete loss of dopaminergic neurons in the SNc specifically, while VTA neurons were less affected. The innervation of the striatum was also partially lost in the aphakia mice (Hwang et al., 2003; Nunes et al., 2003; Smidt et al., 2004; van den Munckhof et al., 2003). Some studies reported an effect on the morphology of the VTA neurons in the mutants (Hwang et al., 2003; Smidt et al., 2004) but the neurotransmitter phenotype of these cells was intact and they expressed other mesDA related genes such as *En1/2*, *Lmx1b* and *Nurr1*.

The loss of TH-expressing neurons in *Pitx3* mutants is seen as early as E12.5 (Maxwell et al., 2005; Smidt et al., 2004). Maxwell et al used a *Pitx3-GFP* knock-in mouse, where the homozygotes are functional knockouts, instead of the Aphakia mice. Using GFP (green fluorescent protein) to track progenitors in the knockout mutant they reported equal numbers of *Pitx3-GFP* positive cells in heterozygotes and mutants during embryonic stages (E12.5 and onwards), indicating that *Pitx3* is not essential for the generation of mesDA precursors. However,

as there were significantly fewer TH-positive cells among the GFP-expressing precursors in the mutants, Pitx3 is likely to be involved in the regulation of *Th* gene in a subset of dopaminergic neurons. A responsive element for Pitx3 on the TH promoter has also been shown which supports this notion (Cazorla et al., 2000; Lebel et al., 2001). In addition, there were subgroups of mesDA precursors in the heterozygotes that expressed either only *Pitx3*-GFP or TH at E12.5. The *Pitx3*-GFP+/TH- cells were predominately located in a ventrolateral position where the SNc will form. This suggests that mesDA neurons may develop from distinct precursors that express Pitx3 and TH in slightly different sequence and that SNc neurons are derived from precursors that express Pitx3 prior to TH which fail to survive when Pitx3 is missing. The induction of Pitx3-expressing precursors in the *Nurr1* mutants showed that its expression is independent of *Nurr1* (Saucedo-Cardenas et al., 1998; Wallen et al., 1999). The loss of Pitx3 expression but not *Nurr1* expression in *Lmx1b* mutants indicates that Pitx3 form an independent pathway together with *Lmx1b* in mesDA neuron development (Smidt et al., 2000).

Cell replacement therapy

In PD patients there is a progressive loss of dopaminergic neurons and thus lowered transmission of DA in the striatum. Although recent studies show that new neurons can be generated in the adult brain in response to injury (Arvidsson et al., 2002) there is still debate whether or not re-generation of dopaminergic neurons occurs in the adult SNc (Frielingsdorf et al., 2004; Zhao et al., 2003). It does seem clear however that the entire nigrostriatal innervation would be difficult to recreate in adult individuals from potential newly formed dopaminergic neurons. Cell replacement therapy, whereby developing dopaminergic neuroblasts from fetal VM tissue are placed in the striatum at the site of dopamine transmission, have instead emerged as a possible strategy to alleviate motor symptoms (reviewed in Dunnett and Bjorklund, 1999; Winkler et al., 2005). Several studies in rodent models showed that cells in grafted tissue were spontaneously active and established functional synaptic contacts thus restoring dopamine transmission (reviewed in Dunnett et al., 2000). This has prompted research on generating an optimal cell preparation for transplantation purposes. Stem cells, which can be expanded indefinitely and have the potential to differentiate into any cell type would be an ideal source. However, much work remains to identify the optimal stem cell and find ways to expand it without losing its potential.

Stem cells

Stem cells – definitions and concepts

Stem cells have a unique ability for proliferation and subsequent differentiation to specialized cells that make them ideal for use in replacing damaged or lost tissue of the body. The theoretical stem cell is per definition a cell that can self-renew (i.e give rise to more of itself), proliferate indefinitely and differentiate to any given celltype. Stem cells are classified by

their differentiation potential as *totipotent*, being able to give rise to all cell types, *pluripotent*, capable of generating cells derived from all three germ layers or *multipotent*, giving rise to many cell types usually of one particular lineage (Kirschstein and Skirboll, 2001).

Most stem cells are studied by isolating putative stem cell fractions and characterize them *in vitro*. There is an obstacle in reconciling the theoretical stem cells with its *in vitro* counterpart. Self-renewal can be evaluated by *clonal assays*, where the progeny of one single cell is shown to contain cells that can repeatedly give rise to new cells with the same properties as the original cell (Reynolds and Weiss, 1996). Proliferation capacity can be somewhat determined in culture however the longevity of it must by necessity be limited to “long-term”. Many thus prefer to label putative stem cell preparations as *progenitor* cells, which refer to cells with a more restricted potential than true stem cells (McKay, 1997). The term *precursor* is also used in connection with stem and progenitor cells and signifies any cell that is earlier in the developmental pathway than another defined cell (McKay, 1997) without saying anything about the “stemness” of it.

It is also necessary to distinguish between the *fate* of stem cells *in vivo*, within its normal context, and the *potential* of cultured stem cells (Gaiano and Fishell, 1998). The potential of an *in vitro* stem cell can be evaluated by differentiation *in vitro* or by introducing it to its normal environment, usually by transplantation. Assessment of the full differential potential of a stem cell may be difficult if the signals necessary for differentiation to a certain subtype are not known or not provided.

However, it is also possible to manipulate stem cells *in vitro* and expose them to circumstances that they do not normally encounter. In this context it can be valuable to distinguish between the *capacity* of a stem cell, which is an intrinsic differentiation potential and what it will differentiate into under normal circumstances *in vitro* or when transplanted into its original environment, and the *capability* of stem cell, which reflects what it can become under certain exaggerated conditions (Kirschstein and Skirboll, 2001).

Neural stem cells

Cells with at least some of the cardinal features of stem cells have been demonstrated in many different organs in the body, and in recent years also in the brain. However, most of these are restricted in their developmental potential to generate only cells of the organ where they originate (Kirschstein and Skirboll, 2001). *Neural stem cell* is the definition of a cell that can generate all three main neural celltypes: neurons, astrocytes or oligodendrocytes and/or is a cell that is derived from the nervous system (Gage, 2000). Embryonic stem cells (*ES cells*) is one of two stem cell preparation that can definitely be classified as pluripotent (the other being embryonic germ cells). ES cells are derived from the inner cell mass of the blastocyst (the very early embryo) and are subsequently cultured *in vitro* (Kirschstein and Skirboll, 2001). Factors and protocols that direct ES cells towards a neural fate have been reported, both for human and murine ES cells (Bain et al., 1995; Okabe et al., 1996; Perrier et al., 2004; Strubing et al., 1995).

Neural stem cells can be derived from both fetal and adult tissue and is stimulated to proliferate *in vitro* in response to various growthfactors such as epidermal growth factor (EGF) and basic

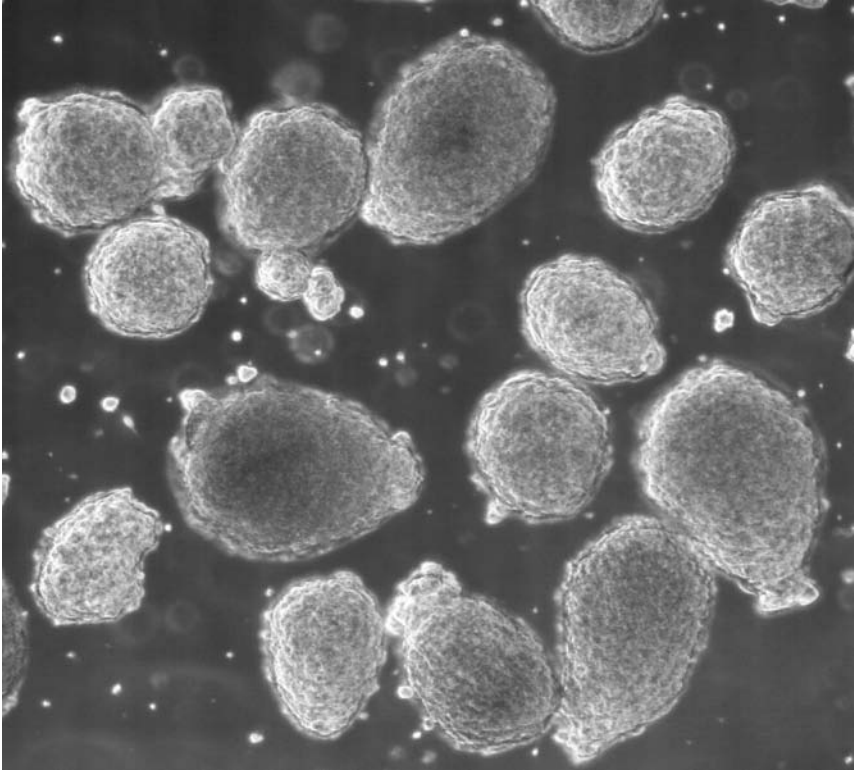


Fig. 4 Neurospheres derived from mouse E11.5 VM tissue

fibroblast growth factor (bFGF) (for review see Gage, 2000; Martinez-Serrano et al., 2001; McKay, 1997). However, there is some debate what this cell is and where it is located at different developmental stages. It has been suggested that neuroepithelial cells in the ventricular zone corresponds to an early *fetal neural stem* cell that is characterized by its responsiveness to bFGF. These early stem cells will in some regions give rise to subventricular zone (SVZ) cells where another, EGF-responsive stem cell appears during late embryogenesis and remain in neurogenic zones in the adult tissue from which it can be isolated as an *adult neural stem* cell (Pevny and Rao, 2003). Alternatively, radial glia cells have been proposed to be the stem cells of the fetal brain and that SVZ cells in adult animals are derived from this population (Alvarez-Buylla et al., 2001). This controversy points to a major obstacle when studying neural stem cells, namely the lack of markers to positively identify the neural stem cells. A few markers such as nestin, *Sox* genes and Musashi have been suggested to label murine neural stem cells (Aubert et al., 2003; Lendahl et al., 1990; Sakakibara et al., 1996). However, although they label cells with the characteristics of neural stem cells they also label more restricted cell types (Barraud et al., 2005).

Neurospheres as a way to expand neural stem cells

Many of the problems in connection with characterizing neural stem cells lies in the difficulty in maintaining them in culture as a pure population. The *neurosphere assay* was initially presented as way to isolate a neural stem cell that self-renewed to form more of itself and whose progeny could give rise to all the celltypes of the CNS, neurons, astrocytes and oligodendrocytes (Reynolds and Weiss, 1992; Reynolds and Weiss, 1996). Neurosphere cultures have since been generated from several subregions of both adult and fetal neural tissue and used to expand neural stem cells. Neural tissue is dissociated into single cells, suspended in defined medium with added mitogens and seeded out in low attachment vessels. After a few days, the cells have divided to form free-floating spheres (Fig 4). However, the neurospheres contain a very heterogenous population of cells of which the self-renewing stem cells only make up a small fraction (Reynolds and Weiss, 1996). It also appears as though other cells with more limited properties than a neural stem cell have sphereforming capacity (Reynolds and Rietze, 2005). It has been shown that culture conditions affect the characteristics of the expanded cells (reviewed in Gage, 2000; Lillien, 1998; Martinez-Serrano et al., 2001). Factors like choice of mitogen, additions to the culture medium or cell density may alter the properties of the cells or select for a specific cell to be expanded. Selection for a specific cell assumes that there are differences between neural stem (or progenitor) cells. Indeed, many studies have shown that fetal neural stem/progenitor cells display region specific characteristics (Hitoshi et al., 2002; Horiguchi et al., 2004; Klein et al., 2005; Parmar et al., 2002; Zappone et al., 2000) indicating that this is the case.

Expansion of midbrain neural stem and progenitor cells

Early studies attempting to expand neural stem cells from mesencephalic tissue showed that EGF was mitogenic for progenitors in embryonic rat VM (Mytilineou et al., 1992) and that mesencephalic tissue from mice as young as E10 could be induced to proliferate with bFGF and serum (Kilpatrick and Bartlett, 1993). More specifically, bFGF was shown to prolong division of dopaminergic precursors and delay the differentiation of TH-positive cells in attached primary cultures from E12 rat embryos (Bouvier and Mytilineou, 1995). Subsequent studies established neurosphere cultures from E14-E16 rat mesencephalon using EGF (Ptak et al., 1995; Svendsen et al., 1995) or EGF/bFGF combined (Caldwell and Svendsen, 1998). The cells could be passaged and remained mitotically active over long time but only a small number differentiated to TH-positive neurons (Caldwell and Svendsen, 1998; Ptak et al., 1995). However, specific inducing protocols showed that expanded neurospheres had the capability to develop into dopaminergic neurons under certain circumstances (Carvey et al., 2001; Ling et al., 1998; Storch et al., 2001; Storch et al., 2003). Attached culture methods proved successful in expanding dopaminergic precursors from rat fetal tissue, however these cultures were not expanded long-term (Studer et al., 1998). Similar protocol has also been used to expand dopaminergic precursors from human fetal mesencephalic tissue (Sanchez-Pernaute, 2001).

AIMS OF THIS THESIS

The overall aim of this thesis work has been to further elucidate mechanisms and events important for the development of the midbrain dopaminergic neurons *in vivo* and explore ways to generate this kind of neurons *in vitro*. These issues have been investigated in the papers included in this thesis in the following way:

- Identify and isolate early dopaminergic precursors – **Paper I**
- Identify new factors involved in the development of dopaminergic neurons – **Paper II**
- Expand neural stem and progenitor cells with a potential to differentiate into mesDA neurons – **Paper III**

Results

RESULTS AND COMMENTS

Identifying early midbrain neural progenitors

One of the aims of this thesis was to identify genes that labelled early precursors in the dopaminergic neuron lineage to gain more insight into how dopaminergic neurons are formed. In order to do this, I evaluated the midbrain expression of several proteins known to be involved in development in the forebrain. The most relevant are pictured in Figure 5. Many of them did not present a pattern that was suggestive of being involved in dopaminergic neuron development but were instead expressed in either lateral or intermediate VM domains (Fig 5). However, pre-B-cell homeodomain protein PBX was found to be located in the same domain as TH at E11.5 and onwards (see Fig 5 and also **paper I**). I also evaluated the embryonic expression of FA-1 or Dlk-1, a protein previously reported to be expressed by SNc and VTA neurons in the adult rat (Jensen et al., 2001). I found that its expression correlated well with TH also during embryonic stages (Fig 5). These results identified two new markers of mesDA neurons.

Sox-genes are differentially expressed in the midbrain

Before the start of this thesis *Aldh1* was the only marker known to be expressed specifically in early mesDA progenitors, however *Aldh1* also labels more mature neurons (McCaffery and Drager, 1994). We wanted to identify genes that labelled mitotic dopaminergic progenitors to be able to isolate cells for expansion in culture and possibly follow the cells in different expansion protocols.

The transcription factor *Sox1* had previously been shown to be one of the earliest markers for neural ectoderm (Pevny et al., 1998; Wood and Episkopou, 1999). It was reported to be expressed by neuroepithelial cells throughout the early neural tube, and associated with dividing neural cells. When I analysed the midbrain expression in a transgenic mouse expressing GFP under the *Sox1* promoter (Aubert et al., 2003), I found that both *Sox1*-GFP and the *Sox1* protein was conspicuously absent from the medial part of the midbrain where progenitors of the mesDA neurons would be located (Fig 5). However, it was expressed profusely in the dorsal midbrain. The dorsolateral expression extended ventrally over the lateral sulcus which separates the VM from the DM, such that a narrow *Sox1* domain was present at the lateral edges of the defined VM, coinciding with expression of e.g *Pax6* and *Meis2* (Fig 5). I subsequently found that *Sox2*, closely related to *Sox1* and expressed in early neuroectoderm as well as in VZ from neurulation (Cheung et al., 2000; Wood and Episkopou, 1999), is expressed in the VZ throughout the midbrain, including the medial region of the ventral midbrain. This suggests that it labels also dopaminergic mitotic progenitors (unpublished results). The *Sox* genes are thus, despite extensive overlap in other parts of the nervous system, differentially expressed in the midbrain during the period of mesDA neuron development.

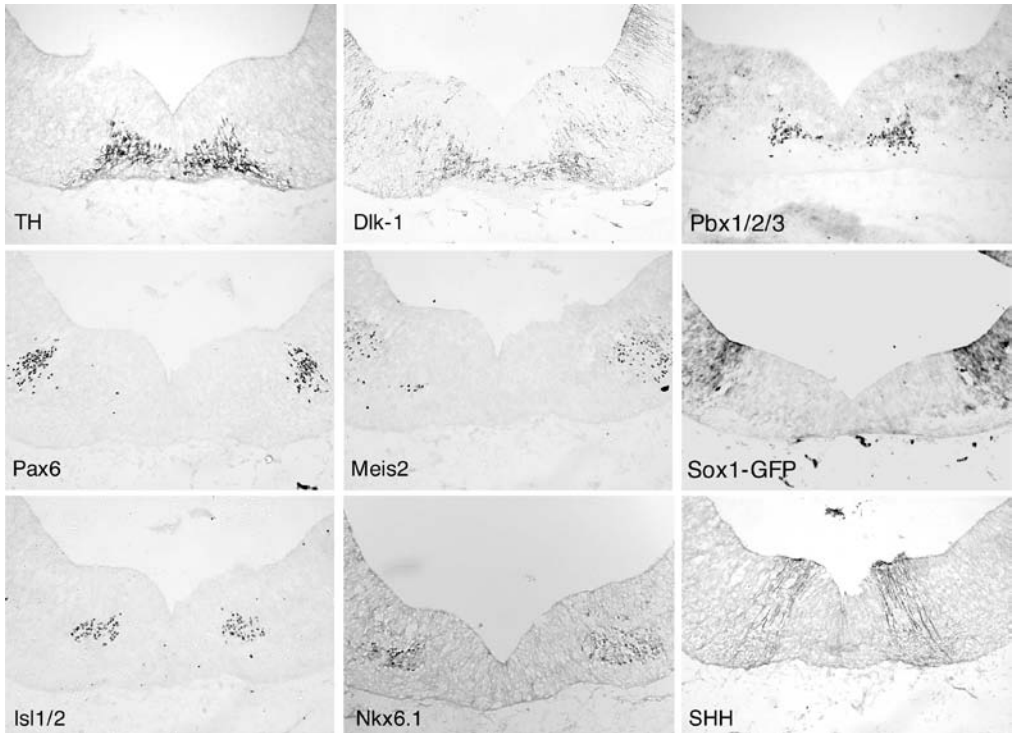


Fig. 5 Coronal sections through the VM of E11.5 mouse. Dlk-1 and Pbx1/2/3 were found to co-localize with TH in developing mesDA neurons. Other markers were found to be expressed in either a lateral domain spanning the lateral sulcus (Pax6, Meis2 and Sox1) or in an intermediate domain (Isl1/2 and Nkx6.1). Curiously the protein expression of SHH appeared to be outside of the medial domain where dopaminergic neurons are generated.

Neurogenin2 labels progenitor cells in the ventral midbrain (Paper I and II)

One group of transcription factors that had not previously been implicated in dopaminergic neuron development was basic helix loop helix (bHLH) factors. bHLH transcription factors are active in determination and differentiation in many tissues including muscle and nerve. In nervous tissue their main function is to select neuronal progenitors and activate genetic programmes for a generic neuronal phenotype (therefore also called proneural genes) however they have also been implicated in subtype specification of certain neurons.

The most common proneural genes are *Mash1* and *Neurogenins*. They are related to two gene families that control neural determination in separate subclasses of neurons in drosophila, the achaete-scute family (Mouse achaete-scute homolog1) and atonal family, respectively. As in drosophila they are expressed in complementary and mostly non-overlapping regions of the peripheral nervous system, PNS and CNS suggesting they define distinct progenitor populations (reviewed in Bertrand et al., 2002).

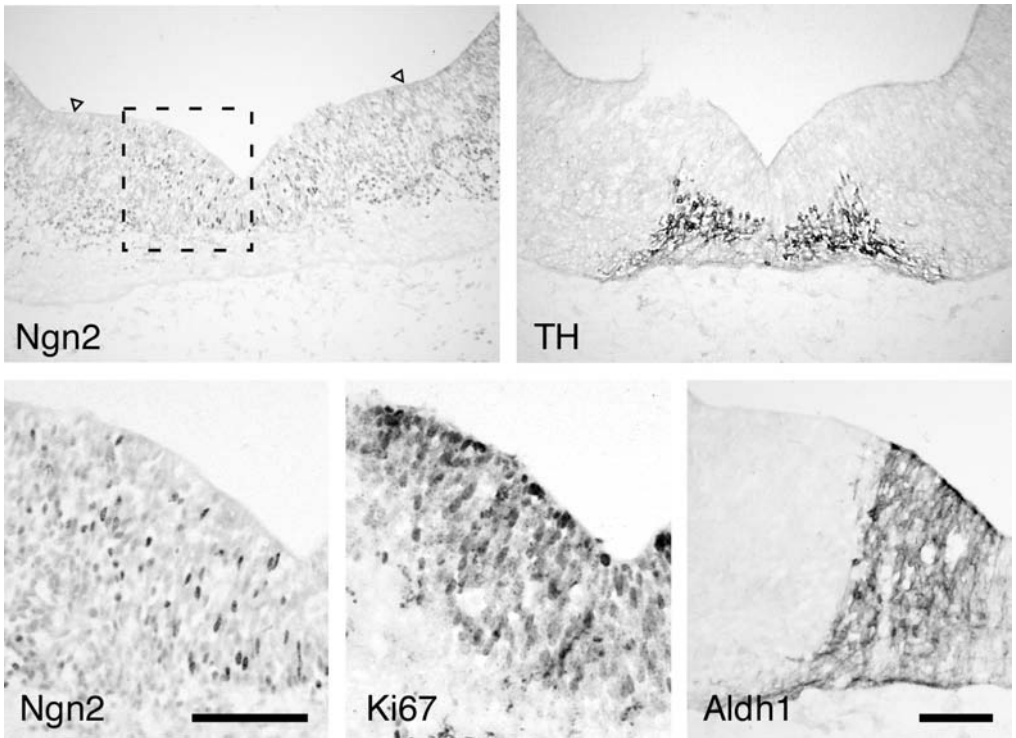


Fig. 6 Coronal sections through the VM at E11.5. Proneural gene Neurogenin2 (Ngn2) is expressed in a restricted pattern in the VM, correlating with the developing TH-positive cells. Ngn2-positive cells are contained within the VZ as determined by staining for Ki67, a marker for dividing cells, in parallel sections. The expression domain of Ngn2 extends lateral to the expression domain of Aldh1 (also in parallel sections), suggesting that it also labels progenitors outside the mesDA domain. The expression of Ngn2 does not overlap with Sox1 expression in the lateral domain of the VM (Sox1 expression limit indicated by open arrowheads). Scale bars: 400 μ m

I screened for expression for proneural genes in the VM and found that Neurogenin2 (Ngn2) was expressed in a pattern that was suggestive of it labeling mitotic dopaminergic neuron precursors. The temporal and spatial expression of Ngn2 and also its expression in comparison to dopaminergic neurons were investigated in detail in **Paper I**. Mitotic nuclei in the VZ undergo a process called interkinetic nuclear migration, i.e they move perpendicular to the ventricular surface depending on where in the cell cycle the cell is. This gives the VZ an impression of being multilayered, termed a pseudo-stratified layer (Jessell and Sanes, 2000a). Ngn2 labeled nuclei that displayed this sort of pattern with strongly labeled and more weakly labeled cells arranged at various distances from the ventricular surface. Parallel sections were stained for the marker Ki67, which labels dividing cells, to determine the extent of the VZ. From this I concluded that the Ngn2 labeled cells were confined to the VZ (Fig 6). Subsequent investigations showed

that although Ngn2 staining appear confined to the VZ, single examples of Nurr1/Ngn2 double stained cells could be detected (**Paper II**). Nurr1 is one of the first proteins to be expressed in postmitotic dopaminergic precursors (see Introduction). If these cells represent postmitotic cells where expression of Ngn2 remain or if they are mitotic cells where Nurr1 is expressed prematurely is at present uncertain, as is the significance of this finding.

The expression of Ngn2 was present in the VM as early as E10.5, before TH-positive cells were detected. At E11.5, when the first TH-positive neurons appeared in the MZ, the Ngn2-positive cells were found to be located in the VZ directly above them. When comparing the expression domain of Ngn2 to that of Aldh1, it extended further lateral of the midline but not as far as the Sox1 expression domain (Fig 6). This quite restricted expression pattern suggested that Ngn2 labeled dopaminergic progenitors and possibly some other progenitor population lateral to them. The caudal limit of Aldh1 coincided with the end of midline expression of Ngn2, however Ngn2 expression domain extended further rostrally into the diencephalon.

At subsequent stages, the expression domain of Ngn2 was restricted more and more towards the midline and included fewer cells, concurrent with fewer cells being committed to a dopaminergic fate (Bayer et al., 1995). Additionally, expression of Ngn2 was first abolished in the rostral part of the mesDA domain, in accordance with the rostrocaudal shift of dopaminergic neurogenesis previously reported (Bayer et al., 1995). Altogether this suggested that Ngn2 labeled a mesDA progenitor cell.

Fate of Ngn2 expression cells in the midbrain

Fate mapping of the Ngn2-positive cells reveal that they are mesDA precursors (Paper I)

The correlation between the spatial and temporal dynamics of Ngn2 staining and dopaminergic neurogenesis was compelling, however it did not prove that Ngn2 was labeling a precursor of mesDA neurons. In order to further elucidate the fate of the ventral midbrain Ngn2-positive cell we made use of a transgenic mouse that had GFP knocked into the *Ngn2* locus (*Ngn2-GFP-KI*, Seibt et al., 2003). We found that GFP expressed from this locus remained in the cells after expression of Ngn2 was downregulated. Thus we could trace the cells that had expressed Ngn2 by their continued expression of GFP. As would be expected, fatemapping of the cells was restricted in time according to the turnover of GFP protein and the detection limit of the antibody for GFP.

At E12.5, Nurr1-positive postmitotic mesDA precursors in the intermediate zone (IZ) co-expressed high levels of GFP and further differentiated, TH-positive cells in the MZ expressed a low level of GFP. This showed that cells expressing Ngn2 will differentiate into mesDA neurons *in vivo*. The high level of GFP in the IZ was an interesting finding. It was clear from GFP IHC that cells in the IZ expressed GFP at a higher level than cells in the VZ (ie cells that also expressed Ngn2). This could be interpreted in many ways, however the most likely explanation is that expression of GFP is slightly lagging and that it reflected very recent events in the dynamics of Ngn2 expression. It is feasible that while the expression of Ngn2 is tightly regulated and the protein is broken down quickly, GFP has a longer half-life and remain. This

high level of GFP in the IZ in cells that have just exited the cell cycle, suggested that Ngn2 is highly upregulated just before or as cells were leaving the cell cycle.

Ngn2-expressing precursors also give rise to other types of neurons (Paper I and II)

We also saw GFP-positive cells lateral to the Nurr1- and TH-positive cells indicating that Ngn2-expressing precursors will give rise to other cell types. The lateral extent of the GFP expression at E12.5 correlated well with the lateral border of Ngn2 expression seen at E11.5 although expression of the Ngn2 itself at this timepoint was more restricted. This suggests that a precursor population expressing Ngn2 and giving rise to a celltype lateral to the dopaminergic neurons was proliferating at E11.5 but no longer at E12.5. Isl-positive neurons of the oculomotor nuclei are located in this lateral region but did not express GFP at E12.5 (**Paper I**). However, when analysed at E11.5 they showed weak GFP expression (**Paper II**). Ngn2-expressing precursors are thus also likley to contribute to oculomotor neurons.

The role of Ngn2 in mesDA neuron development

Absence of Ngn2 results in a dramatic loss of mesDA neurons (Paper II)

Since Ngn2 was expressed by early progenitors in the midbrain, and among them dopaminergic neuron progenitors, we wanted to investigate its role in the development of the dopaminergic neurons. The homozygous *Ngn2-GFP-KI* mice are functional knockout mutants as the coding regions of both alleles are replaced by GFP.

Analysis at the embryonic stages when the dopaminergic neurons are normally generated (E11.5-E13.5), revealed that only a minor fraction of the TH-positive cells were generated in the homozygous knockout mutants. These TH-positive cells were consistently located at the lateral edges of the expected dopaminergic domain but appeared by all accounts to be normal (Fig 7). In postnatal (P0 and P18) mutants the remaining dopaminergic cells were distributed equally between the SN and VTA and shown to project to the correct areas in the forebrain. They also displayed the expected expression of subtype specific markers. The difference between wildtype and mutant mice regarding the number of dopaminergic neurons was less at these postnatal stages however there was still 60-70% reduction of mesDA neurons in the mutant (Fig 7). Interestingly only the dopaminergic neurons, and not the Isl-positive cholinergic neurons, were affected.

Ngn2 is important for the medial mesDA neuron precursors specifically (Paper II)

The loss of dopaminergic neurons in the mesencephalic nuclei appeared to result from an inability of precursors in a medial section of the mesDA domain to develop properly. By the GFP expression and DAPI staining of cell nuclei we could see that cells were generated also in this sector, however they remained close to the ventricle and did not migrate out into the MZ. They did not express NeuroD (a neural differentiation factor downstream of Ngn2), Nurr1 or Pitx3. BrdU and Ki67 stainings showed on the other hand that they were not proliferating. We concluded that Ngn2 is not essential for cell-cycle exit but that in its absence the precursor

cells are arrested from migration and expression of more mature neuronal markers as well as all dopaminergic specific markers are absent. However, they did not express astrocytic markers which indicate that they did not acquire an alternative fate.

Over-expression of Ngn2 does not induce dopaminergic differentiation (Paper II)

In order to determine if Ngn2 by itself promoted a dopaminergic fate we overexpressed Ngn2 in dorsal mesencephalic primary cells and also in primary VM cells. This did not increase the number of TH-positive cells in either culture however it increased the number of β -III-tubulin-positive cells. Thus, Ngn2 is important for the generation of a subset of mesDA neurons but is not on its own sufficient to specify cells to become dopaminergic.

Potential of the Ngn2-GFP positive cells

***In vitro* differentiation (Paper I)**

Since GFP was expressed under the *Ngn2* promoter in the *Ngn2-GFP-KI* mice it was possible to isolate Ngn2-positive cells and their immediate descendants by fluorescence activated cell sorting (FACS) and determine their developmental potential *ex vivo* (outside the brain). As mentioned before we had noticed varying intensities of GFP expression among the cells generated from Ngn2-positive cells. Cells in the VZ, where Ngn2-positive cells are located, exhibited a relatively low GFP expression while the direct descendants, the Nurr1-positive cells of the IZ, showed high level of GFP expression. In order to determine their differentiation potential we separated GFP-positive cells from E12.5 VM into GFP^{high} and GFP^{low}, which corresponded approximately to IZ cells and VZ+MZ cells respectively. The cells were differentiated *in vitro* with minimal serum content. The results confirmed that *Ngn2-GFP* positive cells are precursors to mesDA neurons as only the GFP-positive fractions contained TH-positive cells. However, it also showed that most TH-positive cells that developed *in vitro* were postmitotic at the time of plating, as they rarely incorporated BrdU. This suggested that Ngn2-positive cells from the VZ will not develop into TH-expressing cells *in vitro*, when removed from the surrounding cells and possibly that they are not yet intrinsically specified.

***In vivo* differentiation (Paper I)**

We also wanted to determine the potential of the *Ngn2-GFP*-positive cells to survive and differentiate in a putatively more suitable environment. We therefore transplanted GFP-positive and GFP-negative cells separately to the striatum of newborn rats. The yield of *Ngn2-GFP* cells was too low to further separate the GFP-positive cells into GFP^{high} and GFP^{low} for transplantation. Four weeks after transplantation the grafts were assessed by IHC. By transplanting the mouse-derived cells to neonatal rats it was possible to distinguish them by murine specific antigens. The results showed that, as *in vitro*, mouse derived TH-positive cells were all found in animals which had received grafts from the GFP-positive fraction. These grafts were also enriched in TH-positive cells compared to grafts derived from non-sorted control cells transplanted at the same density. By contrast, all other neuronal types detected; serotonergic, GABAergic and

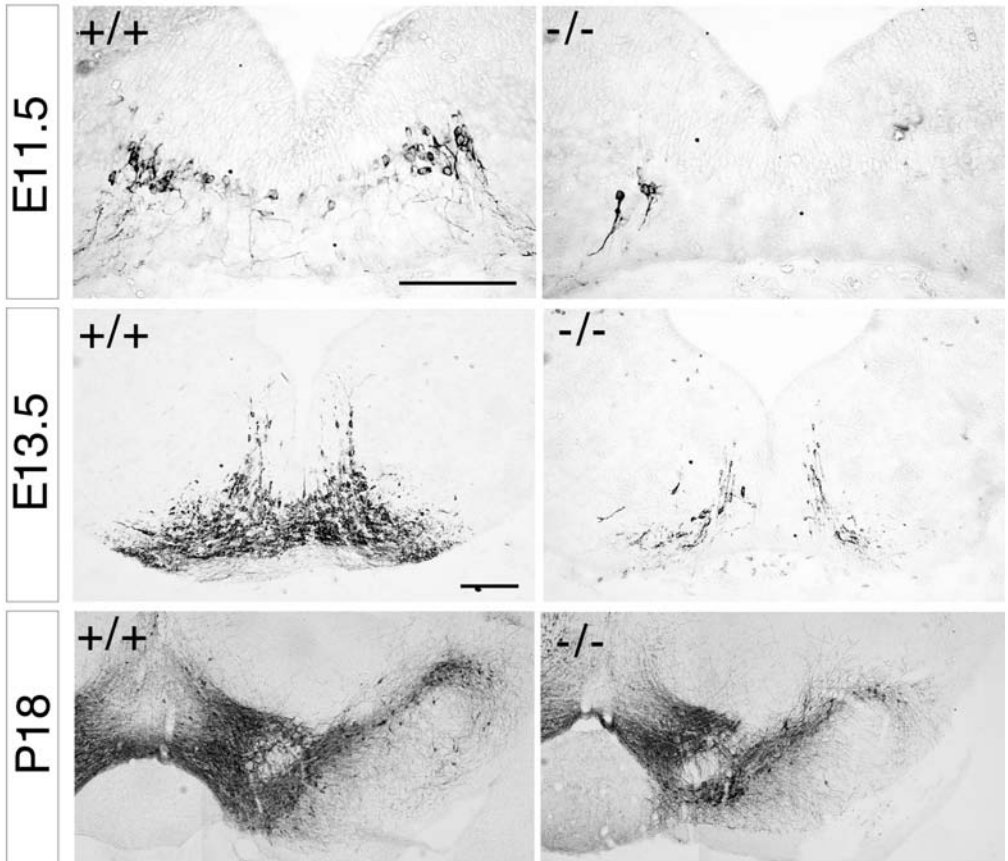


Fig. 7 Coronal sections through midbrain of embryonic and young mice, wt (+/+) and *Ngn2*-KO (-/-) respectively. The majority of TH expressing cells is lost in *Ngn2*-KO mice. At embryonic stages (E11.5-E13.5) when the mesDA neurons are normally generated, only a few cells are seen, located at the lateral edge of the mesDA neuron domain. The number of TH-positive cells in the knockout increase during later embryonic development, however there are still around 60-70% fewer TH-positive cells at postnatal stages (P18).

cholinergic were almost exclusively found in grafts derived from the GFP-negative fraction. Interestingly, although both types of grafts contained many neurons, virtually no glia was found in the GFP-positive fraction. They were however abundant in the GFP-negative fraction and the unsorted control tissue. This suggests that *Ngn2*-GFP positive cells isolated from the VM at E12.5 are marked for a neuronal fate and some of the cells more specifically for a dopaminergic neuronal fate. Notably, GFP^{low} cells (which were also present among the GFP-positive cells), although not specified enough to develop into dopaminergic neurons *in vitro* nevertheless appear specified to become neurons, assuming they survive in the grafts.

Expanding neural stem and progenitor cells from the ventral midbrain

Neurosphere cultures (Paper III)

One of the aims of this thesis was to expand neural stem and progenitor cells that had a potential to differentiate into dopaminergic neurons. Previous studies had indicated that neural stem and progenitor cells are regionally specified and retain some of this specification after expansion *in vitro* (Hitoshi et al., 2002; Horiguchi et al., 2004; Klein et al., 2005; Parmar et al., 2002; Zappone et al., 2000). To increase the possibility of generating mesDA neurons I therefore isolated tissue from E11.5 mouse VM, the time and region where these neurons are generated. Other studies have used mesencephalic tissue of later stages with limited success in generating mesDA neurons (Caldwell and Svendsen, 1998). Dopaminergic neuron progenitors are present and dividing also at earlier stages than E11.5, as shown in [³H] thymidine labelling studies (Bayer et al., 1995). However, I found that E10.5 tissue was difficult to dissect free of contaminating meninges and the yield of cells was limited, and for these reasons avoided as starting material.

The cells were plated in neurosphere medium with growthfactors EGF and bFGF and rapidly formed neurospheres. The neurospheres could be passaged and yielded new spheres even after 10 weeks in culture. The spheres needed to be passaged once a week, and this interval between passages did not change over the 10 weeks suggesting that the division rate remained the same over time. The number of plated cells expanded on average 8-fold once the cultures were established. Although the cells formed spheres with the same approximate size and displayed similar growth characteristics as forebrain cultures there were tangible differences. Midbrain derived spheres were denser and required more force to separate into single cells at passage. Moreover, they attached more easily to plastic surfaces, both during the expansion and proliferation phase. This may be attributed to denser extra-cellular matrix however this was not assessed.

Differentiation potential of progenitors from ventral midbrain expanded as neurospheres (Paper III)

When neurospheres were differentiated by removing growth factors, adding a low concentration of serum to the medium and plated on a permissive surface, the expanded cells gave rise to neurons, oligodendrocytes and astrocytes. I did not grow cells at clonal densities and therefore can not be certain that single cells within the neurospheres were multipotent, however the cultures as whole gave rise to neurons and oligodendrocytes with a frequency of approximately 5% and 1% respectively, even after several passages. The majority of the remaining cells were astrocytes judging by their morphology and immunoreactivity for GFAP. The cultures were neurogenic for several passages, however at passage 10 or later, the cultures generated only GFAP-positive cells when differentiated. This suggests that the potential of the expanded cells changed over time or that a glia restricted progenitor was preferentially expanded.

Neurospheres from E11.5 ventral midbrain are regionally specified (Paper III)

As mentioned above, neural stem and progenitors maintain some of their region character *in vitro* and the progeny matches the subtypes generated in their region of origin. However, all neural phenotypes generated in a certain region do not arise spontaneously from corresponding stem cell preparation, but can require additional factors (Hitoshi et al., 2002; Jensen et al., 2004; Klein et al., 2005). I found that neural stem or progenitor cells in the neurosphere cultures maintained a VM molecular identity, at least on the mRNA level, but that neurons with a dopaminergic phenotype were not generated by standard *in vitro* differentiation. Positional information provided by surrounding cells and other extrinsic cues necessary for certain cellfates are largely lost in neurosphere culture (Jensen et al., 2004). Our attempts to provide some of the extrinsic cues present in the developing VM were not successful which indicate that either we did not provide the correct factors or the concentrations and/or combinations were inadequate. An alternative is that the progenitor cells that were expanded with our protocol could not respond to these cues.

However, over-expression of Nurr1 generated TH-positive cells that also expressed other mesDA neuron markers. The frequency with which this occurred and the phenotype of the TH-positive cells generated was specific for VM progenitors and demonstrated that VM neurosphere cultures are particularly equipped to respond to this stimulus compared to forebrain derived neurospheres. Although not verified by other stainings it is also possible that the Nurr1-induced TH-positive cells generated from forebrain derived neural progenitors were olfactory glomerular dopaminergic neurons as these also express Nurr1 during development (Backman et al., 1999).

Over-expression of Ngn2 and Nurr1 together generate a more complete mesDA phenotype (Paper III)

When Ngn2 was over-expressed together with Nurr1 in the neurosphere expanded cells, this generated morphologically mature TH-positive cells. Some of these TH-positive cells also expressed other dopaminergic markers such as VMAT2, Dlk-1, Pitx3 and En1/2. Over-expression of Ngn2 alone did not induce the expression of either of these markers. Thus, although Ngn2 on its own did not appear to affect the differentiation of cells towards a dopaminergic phenotype, it induced a change possibly to make the cells more receptive to either Nurr1 over-expression or to changes induced by Nurr1. Nurr1 and Pitx3/En1/2 are thought belong to separate developmental pathways as En1/2 and Pitx3-positive cells are found in Nurr1-knockout mice at early stages. It is worth noticing that both En1/2 and Pitx3 appear to have been initiated by Nurr1 and Ngn2 co-transduction but not by transduction of either factor independently.

Discussion

DISCUSSION

Due to the findings in this thesis one more gene, *Neurogenin2* (*Ngn2*), can be added to the list of markers expressed in the midbrain dopaminergic neuron lineage. Importantly, this is the first gene so far to be expressed in the proliferative progenitors but not in the more differentiated cells and which is selectively, although not exclusively expressed by mesDA neuron precursors in the ventral midbrain.

Ngn2 is not just a marker for mesDA neuron progenitors, although it may be used as such, but also plays a role in the development of dopaminergic neurons. Its precise role in the sequence of events leading to generation of dopaminergic neurons has not been fully elucidated yet.

Drawing from the results of the respective studies I will attempt to assemble a more complete picture of progenitor cells in the midbrain and present some theories of the role of *Ngn2* in the neurogenesis of mesDA neurons

Phenotype of the mitotic progenitor in the ventral midbrain

Members of the *Sox* gene family are detected in mitotic cells in VZ of most the CNS (Bylund et al., 2003; Cheung et al., 2000, Pevny et al., 1998) and have been shown to label cells in the telencephalon that display functional properties of stem cells (Barraud et al., 2005; Zappone et al., 2000). As mentioned in the results section, we found that *Sox1* was not expressed in the medial part of the VM from E11.5 and onwards, however *Sox2* expression is present in the VZ at the time when dopaminergic neurons are generated (unpublished results). GLAST (glutamate transporter EAAT1), which labels radial glia cells in the forebrain (Shibata et al., 1997), was also found to be expressed in the VM at this time (**Paper II** and unpublished results). Radial glia are shown to give rise to the vast majority of neurons in the CNS and are suggested to act as fetal progenitor cells (Anthony et al., 2004; Malatesta et al., 2003; Noctor et al., 2001). One can thus hypothesize that a neural progenitor in the medial VM, which will give rise to dopaminergic neurons, is likely to express GLAST, *Sox2* or both.

Dynamic expression of *Ngn2* within the VZ

We found that proneural gene *Ngn2* have an expression pattern during embryogenesis consistent with it being expressed in a mitotic progenitor in the VM. *Ngn2* is expressed in scattered nuclei in the VZ, and low level of GFP is also seen in the VZ of *Ngn2-GFP-KI* mice (**Paper I and Paper II**). We could not perform double labeling of *Ngn2* and *Sox2* for technical reasons, however *Ngn2* is most likely expressed by the neural progenitors of the VM that also express *Sox2*. Moreover, we saw co-expression of nestin and *Ngn2* (**Paper I**). In other studies *Ngn2* has consistently been reported to be expressed in VZ and SVZ and also to label for BrdU showing that it is expressed in cells that are dividing (Gradwohl et al., 1996; Sommer et al., 1996).

Close inspection of *Ngn2 in situ* hybridisation suggested that *Ngn2* mRNA has its highest expression at the ventral edge of the VZ, close to the postmitotic IZ. Although this is not as clear from the protein expression, which shows *Ngn2*-positive cells throughout the thickness

of VZ, *Ngn2* may be expressed at low levels (too low for detection by IHC) by the majority of VZ cells and be upregulated during last cell cycle. In *Ngn2-GFP-KI* mice the GFP expression, which is slightly lagging compared to *Ngn2* protein expression (presumably due to longer half-life of the protein), show the highest expression in the IZ cells, which also suggest a surge in *Ngn2* expression prior to cell cycle exit (**Paper I and II**).

According to the established model for neurogenesis, mitotic progenitors initially express proneural genes at low levels while they are not yet committed to differentiation (Bertrand et al., 2002; Bylund et al., 2003; Kintner, 2002) Positive feedback loops result in high levels of proneural genes in a subpopulation of progenitor cells, concurrent with the cells being irreversibly committed to differentiation. The initial upregulation of proneural gene expression is the result of lateral inhibition where Notch signalling repress proneural expression and reduce levels of notch ligand in some cells, thereby enabling higher levels of proneural gene expression in the progenitors selected for neuronal fate. However, other positive feedback mechanisms are needed to increase and/or maintain the elevated levels of proneural gene expression (Bertrand et al., 2002; Kintner, 2002). Interestingly, in the *Ngn2* knockout mutants we detected cells with high expression of GFP that maintained contact with the ventricle, similar to radial glia, and which were expressing GLAST (**Paper II**). This indicates both that the neural progenitor of the VZ normally experiences an upregulation of *Ngn2* and that this neural progenitor is a GLAST-positive radial glia-like cell. It is notable that the GFP in the *Ngn2* locus was still upregulated despite the lack of *Ngn2* suggesting that *Ngn2* protein activity is not necessary for the regulatory feedback loops.

Possible interaction of *Ngn2* with *Sox2* in neurogenesis of ventral midbrain cells

Proneural genes function on several levels of neurogenesis. Progenitors are selected for a neuronal fate by an upregulation of proneural genes, which promotes cell cycle exit and start developmental programmes for neuronal differentiation (Bertrand et al., 2002; Kintner, 2002). Bylund et al (2003) showed that activity of Sox proteins in neural progenitors in chick spinal cord maintains them in an undifferentiated state and that high levels of *Ngn2* protein most likely mediate neurogenesis by downregulating *Sox1-3* expression.

We saw that a lack of *Ngn2* caused an accumulation of cells arrested in differentiation in the medial part of the mesDA neuron domain in the *Ngn2* knockout mutants (**Paper II**). Although we did not ascertain the expression of *Sox2* in the mutants it is likely that the lack of *Ngn2* enabled continued *Sox2* expression in these progenitors. Maintenance of some progenitor character such as contact to the ventricle and expression of GLAST may thus be a result of this prolonged *Sox2* expression.

Interestingly, the lack of *Ngn2* did not greatly affect cell cycle exit of the progenitor cells in the VM, as we did not see an increase in proliferating cells (**Paper II**). Several studies connect over-expression of *Ngn2* with premature cell cycle exit (Lo et al., 2002; Mizuguchi et al., 2001). What drives the cells out of cell cycle in the VM remains unclear.

Ngn2 involved in acquiring full neuronal phenotype of mesDA neuron precursors

Proneural genes regulate the expression of general neuronal character at neurogenesis and over-expression of these genes often leads to increased neuronal differentiation both *in vivo* and in cultured cells (Farah et al., 2000; Mizuguchi et al., 2001; Sun et al., 2001). We also found that over-expression of Ngn2 increased the number of β -III-tubulin positive neurons in both primary cells from the dorsal mesencephalon (**Paper II**) and in neural progenitor cells expanded as neurospheres (**Paper III**).

Specific members of the bHLH transcription factor family act as neuronal differentiation genes downstream of earlier expressed proneural genes (Bertrand et al., 2002). The Ngns have previously been shown to co-localize with, and in many areas precede, the expression of NeuroD, a neuronal differentiation gene (Sommer et al., 1996). In the VM we found that NeuroD was expressed by cells bordering the IZ, suggesting that NeuroD is a downstream effector of Ngn2. The lack of NeuroD coupled with the loss of all neuronal markers in the accumulated cells confirmed that NeuroD act as neuronal differentiation factor regulated directly or indirectly by Ngn2 and the absence of Ngn2 (and thus NeuroD) prevent further neuronal differentiation of the newly postmitotic cells. It is worth noticing that NeuroD expression was seen in the dorsal midbrain and regions lateral to the mesDA neuron domain also in the Ngn2 mutants, suggesting that other proneural genes control expression of NeuroD in other regions of the midbrain. We also saw β -III-tubulin positive cells that did not express TH in the MZ of the Ngn2 mutants, suggesting that neurogenesis is maintained for other types of neurons that are thus not dependent on either Ngn2 or NeuroD (**Paper II**).

Does Ngn2 have a role in both differentiation and specification of mesDA neurons?

Although the major role of proneural genes such as *Ngn2* is to regulate genes for a generic neuronal phenotype, they have also been implicated in specification of neuronal subtype. Proneural gene *Mash1* specifies noradrenergic phenotype both in the PNS (Hirsch et al., 1998; Lo et al., 1998) and in the CNS (Pattyn et al., 2000), GABAergic interneurons in the ventral forebrain (Fode et al., 2000) and serotonergic neurons in the hindbrain (Pattyn et al., 2004). Ngn2 on the other hand is involved in the specification of motor neurons in the spinal cord (Scardigli et al., 2001) and glutamatergic neurons in the dorsal forebrain (Fode et al., 2000).

Loss of Ngn2 both affects the neuronal differentiation of the medial VM progenitors and abolishes the expression of genes signifying a dopaminergic phenotype (**Paper II**). However, whether Ngn2 is involved in specifying the dopaminergic phenotype in VM precursors remains to be determined. The cells were arrested in an undifferentiated state, which could leave them unable to respond to differentiation cues without Ngn2 being involved in the phenotypic specification per se. A similar block in neuronal differentiation is seen for precursors of cranial ganglia sensory neurons in the Ngn2 mutants (Fode et al., 1998). However, expression of subtype specific homeodomain proteins that regulates the neurotransmitter phenotype of these cells is unaffected, in contrast to our results.

Substituting the proneural function of Ngn2 by another proneural gene may reveal if the role of Ngn2 for the generation of dopaminergic neurons goes beyond its proneural function and if so, how specific this role is for Ngn2. In a separate study to ours, Kele et al (submitted) have used

Ngn2^{Mash1/Mash1} mice, where the coding sequence for Ngn2 is replaced by the coding sequence for Mash1 in both alleles, to study the contribution of Ngn2 to neuronal differentiation and subtype specification in the VM. They found that Mash1 in the Ngn2 locus increase the number of postmitotic Nurr1-positive precursors but that it could not completely rescue the loss of dopaminergic neurons (Kele et al, submitted). This suggests that Ngn2 do indeed have a specific role in the specification of mesDA neurons.

Possible mechanisms for Ngn2 fate specification

It is clear that over-expression of Ngn2 does not induce a dopaminergic phenotype in expanded neural progenitors (**Paper III**) or primary mesencephalic cultures (**Paper II** and unpublished results). However, this finding is not surprising and Ngn2 may nevertheless have a role in specification of dopaminergic neurons. Many gain of function studies have shown that while Mash1 have the capacity to re-specify progenitors, *Neurogenin* genes, when involved in specification, generally function as permissive factors rather than instructive (Parras et al., 2002). Both Ngn2 and Mash1 are expressed in several areas of the brain that give rise to many different neuronal subtypes. To generate specific subtypes, local co-factors are thought to act together with the proneural genes. For example, Ngn2 interacts with Olig2, another bHLH family member, to specify motor neurons in the spinal cord (Mizuguchi et al., 2001). Mash1 cannot replace Ngn2 to generate these neurons, which shows that Ngn2 is specific in this role (Parras et al., 2002). *Neurogenin* genes are thought to be more dependent on cell context than Mash1 and it is hypothesized that the co-factors needed are very localized, thus explaining that Ngns are rarely instructive when over-expressed. As an example, Ngn1 can induce sensory neuron markers when misexpressed in chick but only if the local concentration of extrinsic factor BMP2 is adequate (Lo et al., 2002).

If Ngn2 is involved in the specification of dopaminergic neurons it is likely there is a co-factor acting together with Ngn2 also in the midbrain. A plausible contender is extrinsic factor SHH, with similar dorso-ventral patterning function as BMPs and known to be essential for dopaminergic neuron development in the midbrain (Jessell, 2000; see also introduction). SHH mRNA expression is lost in neural progenitor cultures (**Paper III**) and presumably not present in high enough concentration in the dorsal midbrain, which could explain why over-expression of Ngn2 does not induce a dopaminergic phenotype in these cultures (**Paper II**). However, a recent study has showed that SHH responsive cells only contribute to dopaminergic neurons before E9.0 (Zervas et al., 2004). This makes SHH unlikely as a co-factor in the Ngn2-positive cells seen at E11.5 and onwards.

Interaction of Ngn2 and homeodomain proteins

Another option for co-factor(s) possibly acting together with Ngn2 to specify a dopaminergic fate is homeobox transcription factors. Mash1 has been shown to interact with homeodomain (HD) protein Phox2b to induce other, lineage specific homeobox transcription factors and determine noradrenergic phenotype (Hirsch et al., 1998; Lo et al., 1998). Ngn2 is part of specifying the phenotypic identity of ventral spinal cord motorneurons, but is unable to promote motor neuron development in chick neural tube without the presence of HD proteins Isl1 and Lhx3 (Lee and Pfaff, 2003).

Homeobox transcription factors play several roles in the specification of neuronal subtypes. In the spinal cord early HD proteins are expressed in response to the opposing patterning action of SHH ventrally and BMPs dorsally, to set up discrete progenitor domains (for review see Jessell, 2000; Lee and Jessell, 1999). Later expressed HD proteins function to specify the identity of newborn neurons from the different domains (Tanabe et al., 1998). Evidence suggests that activation of proneural genes such as *Ngn2* is coordinated with the activation of these subtype specifying homeobox transcription factors, not least since proneural genes also in some cases display specification functions. In the spinal cord motoneurons, a direct transcriptional interaction between *Ngn2* and HD proteins *Isl1* and *Lhx3* provide the mechanism for integrating these developmental programmes (Lee and Pfaff, 2003).

A homeobox transcription factor that could function as a co-factor for *Ngn2* would need to be expressed in the progenitor cells in the VZ together with *Ngn2*. The homeobox transcription factors known to be involved in mesDA neuron development are mainly expressed in the postmitotic neurons. *Lmx1b* is expressed early in the neural tube, but is not present at high levels in the VZ at E11.5 (Fig 3). *En1/2* antibody staining shows a weak expression in the VZ (**Paper I**) but is mainly confined to postmitotic cells (**Paper I** and Simon et al., 2001). However, the possibility exist that there is an as yet unidentified homeobox transcription factor expressed by the progenitors in the VM that could act together with *Ngn2* to specify dopaminergic neurons. A very recent study has revealed the presence of two homeobox genes in the VZ at the time of dopaminergic neurogenesis, *Lmx1a* and *Msx1*, which were hitherto unknown to be expressed in the ventral midbrain. The study also shows that they are essential for the generation of dopaminergic neurons. *Lmx1a* induce the expression of *Msx1* while, interestingly, *Msx1* is shown to induce expression of *Ngn2* at the ventral midline (Andersson et al, *in press*). Thus *Ngn2* acts in the presence of these HD proteins to determine neuronal fate and possibly specify some aspect of the dopaminergic phenotype. However, any direct interaction such as in spinal motoneurons remains to be determined.

Induction of mesDA phenotype *in vitro* by over-expression of *Ngn2* and *Nurr1*

An interesting finding which gives more clues about the role of *Ngn2* in dopaminergic neuron development is that we managed to get TH-positive neurons that also expressed other dopaminergic markers from expanded fetal tissue, but only when over-expressing *Nurr1* and *Ngn2* together in the neural progenitors (**Paper III**). TH-positive cells seen when over-expressing *Nurr1* alone were immature and did not express *En1/2* or *Pitx3*. This indicates again that *Ngn2* is in fact involved in the specification of mesDA neurons. However, if *Ngn2*, alone or together with a co-factor, induce the expression of specific dopaminergic transcription factors such as *En1/2* and *Pitx3* one would expect expression of these markers also in cultures where only *Ngn2* is overexpressed. Interaction with *Nurr1* or its downstream targets seems plausible, yet loss of *Nurr1* has been shown to not affect initial expression of *Pitx3* or *En1/2* (Wallen et al., 1999).

What about the lateral population?

A curious finding in the Ngn2 mutant is that neurogenesis of some mesDA neurons is unaffected. A lateral population of mesDA neurons is formed on time, and the cells express NeuroD, Nurr1, Pitx3 and TH in the expected manner and sequence (**Paper II**). This would suggest that Ngn2 is not involved in the neurogenesis of a subtype of the dopamine neurons or that another gene can compensate for its function in these particular progenitors. Assessment of presence and/or localization of other proneural genes in medial versus lateral mesDA progenitors may explain why this lateral population is spared in the Ngn2 mutants.

Neurogenin1 (Ngn1) is reported to have a weak expression in the rostral midbrain (Sommer et al., 1996) and could possibly assume the function of Ngn2 in the lateral population. Ngn1 and Ngn2 have overlapping expression in the spinal cord (Sommer et al., 1996) and presence of Ngn1 enables expression of general neuronal markers in ventral spinal cord interneuron and motorneuron precursors in the Ngn2 mutants. However, Ngn1 cannot compensate for the role of Ngn2 in expression of the correct subtype specific HD proteins (Scardigli et al., 2001). If Ngn1 compensates for the lack of Ngn2 in the lateral population of dopaminergic neurons in the VM it must assume both the differentiation and specification functions of Ngn2 in these precursors. Alternatively, Ngn2 has only a differentiation function at least in these lateral progenitors.

Another option for a compensating proneural gene is Mash1, which is expressed broadly in the ventral midbrain, including both the lateral and medial mesDA neuron domains (Vernay et al., 2005; **Paper I** and unpublished observation). Mash1 is known to specify GABAergic neurons in the forebrain (Fode et al., 2000), a cell type that is abundant in the adult VM. Even if the normal function of Mash1 does not involve specifying the dopaminergic phenotype it is nevertheless possible that it acquires this function when Ngn2 is absent. For example, Mash1 is upregulated in dorsal telencephalic progenitors in the absence of Ngn2 (Fode et al., 2000). However, this ectopic expression of Mash1 re-specifies the neurons produced from the dorsal telencephalon (Fode et al., 2000), which is apparently not the case in the VM (**Paper II**).

It is also interesting to note that while Ngn2-positive progenitors give rise to Isl-positive motorneurons lateral to the mesDA domain these neurons are unaffected by the lack of Ngn2 (**Paper II**). If there is a rescue mechanism for these neurons or if Ngn2 is redundant for their formation remains to be determined.

Induction of Ngn2 in ventral midbrain

A study by Vernay et al (2005) recently revealed a role for late expressed Otx2 in regulating Ngn2 and Mash1 in the midbrain (as opposed to early expressed Otx2 which function to establish the position of the MHO; Broccoli et al., 1999). In Otx2 conditional knock-out mice (CKO), where the expression of Otx2 is downregulated in nestin-expressing cells from the age of E10.5 and completely abolished by the age of E12.5, the expression of Ngn2 and Mash1 is reduced in the ventral midbrain already at E11.5. The Otx2 CKO display a phenotype similar to what we see in the Ngn2 mutant mice, with dopaminergic neurons lost predominantly around the ventral midline, although the reduction of TH-positive neurons in these Otx2 CKO is not as severe as what we have found in the Ngn2 mutants (**Paper II**). In the ventral spinal cord, early expressed HD protein Pax6 have been shown to directly regulate Ngn2 (Scardigli et al., 2003)

and both Ngn2 and fellow bHLH transcription factor Olig2 are lost in motorneuron progenitors in the Pax6 mutant (Mizuguchi et al., 2001). Whether Otx2 act in a role similar to Pax6 in the spinal cord to directly regulate Ngn2 is not yet clear. Msx1, which is also shown to induce expression of Ngn2 (Andersson et al, *in press*) is likely to act in the context of Otx2.

Why is it so hard to get a dopaminergic neuron *in vitro*?

Many attempts have been made to generate mesDA neurons from expandable neural stem and progenitors and few successful approaches have been reported. The question thus arise whether the difficulty lies in providing the correct induction for a dopaminergic phenotype in an otherwise responsive progenitor population or if there is a specific dopaminergic progenitor population that is hard to expand.

Previous studies have identified several key factors in the dopaminergic neuron development, such as SHH, FGF8, Lmx1b, Nurr1, En1/2 and Pitx3 (see Introduction). Our results showed that Ngn2 is required for proper generation of a subset of dopaminergic neurons *in vivo* and over-expression of Ngn2 together with Nurr1 in expanded neural progenitors can give rise to mesDA neurons with some of the distinct features of mesDA neurons (**Paper III**). The recent results regarding new HD proteins in the developing dopaminergic progenitors reveal that there are other cell-intrinsic factors previously unknown that are both sufficient and required to generate dopaminergic neurons (Andersson et al, *in press*). It is also clear that the nature of the progenitor is essential. To generate a dopaminergic phenotype, only progenitors with a ventral midbrain character are relevant (**Paper III** and Andersson et al, *in press*). Judging from our results and others, it seems clear that the emergence of a mesDA phenotype requires a complex interaction of extrinsic signals at precise concentrations and cell-intrinsic factors.

Nonetheless, certain culture conditions may be more suited to preserve the expression of cell-intrinsic factors and perhaps also to expand specialized cells that provide the extrinsic signals. We tried to optimize the culture conditions by adding B27 in the medium, which is reported to be beneficial for dopaminergic neuron development of cultured cells (Kim et al., 2003; Svendsen et al., 1995) and removing HEPES that inhibits differentiation of TH-positive cells in ES cell cultures (Lee et al., 2000). We also explored several different growth factors but none of the options appeared superior in generating dopaminergic neurons and only the use of EGF/bFGF combined resulted in long-term expansion (**Paper III**). In separate experiments attached cultures were established using EGF/bFGF but these cultures were also unable to robustly and spontaneously give rise to clearly dopaminergic neurons (unpublished results).

An aspect of culture variables we did not explore was using hypoxic conditions that are more similar to the *in vivo* situation. Lowered oxygen (<5% O₂) during incubation is reported to alter the proliferation and differentiation of mesencephalic precursor cells *in vitro*. Both neurogenic potential and dopaminergic yield are improved over cultures grown in standard oxygen levels and the effect is specific for midbrain precursors (Milosevic et al., 2005; Storch et al., 2001; Studer et al., 2000). Lowered oxygen is hypothesized to preserve an immature, multipotent phenotype of the mesencephalic progenitor. High oxygen levels also affect the survival of progenitors and/or more mature cells possibly by increasing formation of free radicals to which midbrain dopaminergic neurons appear particularly sensitive (Milosevic et al., 2005).

Are we expanding the correct progenitor?

Although we did see cells capable of responding to Nurr1 and Ngn2 and forming dopaminergic neuron in our expanded cultures the frequency of such cells was low.

This raised a question if we were expanding the correct progenitor in neurosphere cultures. Using heterozygous *Ngn2*-GFP embryos as starting material for neurospheres we attempted to follow the progenitor cells in culture by assessing the expression of *Ngn2*-GFP. However, we and others (F. Guillemot, personal communication), have experienced that *Ngn2*-GFP expression is lost after dissociation and plating, and in our experiments the expression of *Ngn2*-GFP was not significantly upregulated even when cells were differentiated (unpublished results). Both upregulation and maintenance of proneural genes is the result of positive feedback loops, initially dependent on cell to cell signalling (Bertrand et al., 2002; Kintner, 2002). Lack of cell contact at plating contact could possibly hinder the upregulation of *Ngn2*-GFP.

We also explored the possibility that another subpopulation of neural progenitor cells were preferentially expanded under the conditions used. A study in this lab found that *Sox1*-GFP-positive cells derived from the forebrain are enriched in neurosphere forming cells and that *Sox1*-GFP negative cells did not form neurospheres when plated (Barraud et al., 2005). We had found, as previously mentioned, that Sox1 expression was absent from medial VM (Fig 5). However, the subdissected VM pieces used to start neurosphere cultures included the lateral *Sox1*-GFP-positive region at the edges (see Material and methods). This raised the question whether this fraction was preferentially expanded in neurosphere cultures. Sub-dissected VM from *Sox1*-GFP mice were sorted by FACS, and the GFP^{pos} and GFP^{neg} fractions were plated. Interestingly, only the GFP-positive cells formed neurospheres (unpublished data). However, an explanation for this could be that *Sox1*-GFP positive cells are more resistant to the FACS procedure than medial progenitors. Furthermore, when VM tissuepieces were dissected into medial and lateral fractions, and plated (without sorting), both subregions were found to form neurospheres with similar expansion characteristics (unpublished data). Gene expression data also indicated that cells expanded as neurospheres had a ventral midbrain character, and mRNA expression of both Ngn2 and Aldh1 was also seen (**Paper III**). Taken together this shows that neural stem and progenitors from the VM can be expanded in neurosphere cultures however the proportion of different midbrain progenitors within the spheres are unknown and the proportion of mesDA progenitors is likely to be low.

Material and methods

MATERIAL AND METHODS

Experimental animals

In this thesis wild type NMRI mice were used as a standard for histological analysis (Paper I) and establishment of tissue cultures (Paper III).

Featured in Paper I and II is a transgenic mouse strain where the first exon of the proneural gene *Ngn2* is replaced with the coding sequence for green fluorescent protein (Ngn2-KI-GFP). The mouse was generated by homologous recombination in ES cells, on a129Sv/CD1 background (Seibt et al., 2003) and maintained on the same background in house. Heterozygous mice (Ngn2+/-) were used as reporter mice to detect and isolate the cells expressing Ngn2 and their progeny (paper I and II). Homozygous *Ngn2*-GFP mice are functional knockouts (Ngn2-/-) and used to study the effect of deleting Ngn2 (Paper II). For generation of heterozygote embryos used for cell sorting (Paper I) Ngn2+/- males were mated with wild-type NMRI females. The embryos were genotyped by GFP-fluorescence and the expected 50/50 ratio was generally found. For Paper II wildtype, heterozygote and homozygote littermates were collected from Ngn2+/- x Ngn2+/- matings and genotyped by PCR (see below).

In Paper III a transgenic mouse strain expressing GFP under the β -actin promoter was used (Okabe et al., 1997). We also utilized a transgenic mouse expressing GFP under the *Sox1* promoter (Aubert et al., 2003) in some unpublished studies.

For generation of embryos, adult females (>6 weeks) were mated with adult males overnight and checked for vaginal plug before noon the following day. This day was considered embryonic day (E) 0.5. At the day of collection of embryos, the females were euthanized by CO₂ exposure before noon to ensure that repeated experiments generated embryos of similar developmental stage. For paper II we also used neonatal and P18 pups littermates. The size of the litters were reduced to <5 pups on P0 to increase survival of Ngn2-/-.

PCR genotyping

Ngn2 wildtype allele was detected by primers Ngn2KI5 and Ngn2KI3 generating a 813 bp product. For genotyping of the mutant allele primers Ngn2KI5 and Ngn2KImutant3 generated a 440 bp product. The primer sequences were

Ngn2KI5 5'-GGA CAT TCC CGG ACA CAC AC-3'

Ngn2KI3 5'-AGA TGT AAT TGT GGG CGA AG-3'

Ngn2KImutant3 5'-GCA TCA CCT TCA CCC TCT CC-3'

Histological analysis

Preparation of tissue

Embryos were immersion fixed in 4% PFA overnight at 8° and rinsed in PBS. P0 pups were decapitated, the brain removed and placed in 4% PFA overnight. P18 mice and adult rats received lethal doses of pentobarbitone and were perfused trans-cardially with 0.9% saline

followed by 4% PFA. The brains were then dissected out and post-fixed for 2 hours. All tissue was cryo-protected in sucrose before being sectioned, young embryos (E10.5-E13.5) in 30% sucrose and older embryos (E15.5-E17.5), P0 and adult brain in 25% sucrose. Embryos and P0 brains were mounted in OCT compound and sectioned on a cryostat. Coronal and saggittal sections, 12 μm (E10.5-E13.5), 14 μm (E15.5-E17.5) and 16 μm (P0) thick were mounted on Super FrostPlus glass-slides in series of 10. The slides were stored in -20° . P18 mice and adult brains were cut on a sliding microtome at a thickness of 30 μm and 40 μm respectively, in series of 5. Sections were stored in freezing media in -20° .

Immunohistochemistry

Sections mounted on slides were air-dried to ensure that the sections remained on the glass. Slides and free-floating sections were rinsed in KPBS 3x10 min to remove OCT compound and freezing medium respectively. Depending on type of immunostaining and antibody one or more of the following pre-treatments were performed on the tissue:

Quenching – Sections were incubated with 3% H_2O_2 and 10% methanol in KPBS for 10 min to exhaust endogenous peroxidase in the tissue. This was performed if the peroxidase-based ABC-kit (Vectastain) was used for antibody detection.

Antigen retrieval – Sections were boiled in 10 mM citrate buffer, pH 6 for 20 min. Alternatively, DNA was denatured in 1M HCl at 65°C for 30 min.

Pre-incubation – As a standard, sections were incubated with a blocking solution made up of 2-5% normal serum (from the species the secondary antibody was raised in) and 0.25% triton-X in KPBS. Some antibodies (see table 1) required special blocking solution consisting of 1% dry-milk/10% normal serum/1mg/ml BSA in KPBS. Sections were pre-incubated for an hour. Following these pre-treatments, primary antibodies, diluted in blocking solution, were incubated with the sections overnight at room temperature (for exceptions see table 2). Slide-mounted sections were kept in humidified chamber. The following day the sections were rinsed with KPBS 3x10 min and incubated with secondary antibody for two hours, in the dark in cases where fluorescent secondary antibodies were used. Biotinylated secondary antibodies were followed by incubation with streptavidin-horseradish peroxidase complex (ABC elite kit, Vectastain) for 1-1.5 hours. Addition of Di-amino-benzidine, DAB (0.5 mg/ml, Sigma) and H_2O_2 to the sections visualized the antibody binding.

To visualize nuclei in fluorescent immunostained tissue, DAPI (1:1000, Sigma) were added to the final rinse in KPBS. Slides were subsequently coverslipped in PVA-DAPCO, an anti-fading mounting media. DAB-stained slides were dehydrated in increasing ethanol and xylen baths and coverslipped with DBX.

In situ hybridization

A subcloned fragment of Ngn2 cDNA (F. Guillemot) was used as a template to generate an antisense digoxigenin (DIG)-labelled cRNA probe. 1 μg of DIG-labeled probe and 50 μg yeast tRNA was added per ml hybridization solution. After proteinase K treatment, postfixation and dehydration in ethanol the slides were incubated overnight at 55°C in a sealed humidified

chamber. Following hybridization the slides were washed in 2X SSC/50% formamide at 65°C for 30 min and RNase treated (20 mg/ml) at 37°C for 30 min. The slides were washed twice with 2X SSC/50% formamide, at 65°C for 20 min each, then 2X SSC and 0.1X SSC at 37°C for 15 min each and finally PBT (PBS + 0.1% Tween-20) for 15 min. Colour reaction was performed using BM purple (Boehringer Mannheim).

Tissue culture

Dissection

Embryos were kept in ice-cold PBS. Sub-dissection of the VM from the embryos were performed in L15 medium (Gibco). The VM was restricted rostrally by the diencephalon, caudally by the mid-hindbrain constriction and contained within the sulcus separating alar and basal plates (see Fig 3). All meninges were carefully removed from the neural tube. Tissue pieces were kept in L15 on ice before being processed further.

Neurosphere cultures

Tissue pieces were incubated with DMEM (Gibco) containing 0.05% DNase/0.1% trypsin for 15 min at 37°C. The trypsin solution was inactivated with DMEM containing 10% fetal bovine serum (FBS) and removed. The tissue pieces were dissociated mechanically to a single cell suspension in neurosphere culture medium. Cell number was estimated by diluting cell suspension 1:10 in Trypan Blue (Sigma). Viable, non-stained cells were counted in a Bürker chamber.

The cells were plated in tissue culture flasks in neurosphere medium with 10 ng/ml bFGF (R&D systems), 20 ng/ml EGF (R&D systems) and 40 U heparin/ml (Sigma) at a concentration of 100 000 - 500 000 cells/ml.

Neurospheres were passaged once a week and replated in proliferation medium at 100 000 cells/ml. Alternatively the cells were cryo-preserved in freezing medium and stored at -150°C. When re-established the cells were plated at 2X normal cell density.

Differentiation of neurospheres

Neurospheres were differentiated on plastic 4-well slides that had been coated by PLL (4 µg/cm², slides incubated for an hour RT, then rinsed and let to dry) and directly prior to plating, were coated with laminin (50 µg/ml, incubated on slides 2 hours in 37°C). Whole spheres were re-suspended in basic neurosphere medium without growthfactors but with added 1% FBS. 0.625 ml of neurospheres were re-suspended in 1 ml differentiation medium and added to each 4-well chamber (≈625 000 cells/cm²).

Primary cultures and co-cultures

Subdissected VM were mechanically dissociated and plated at a density of 120 000 cells/cm² on PLL coated plastic chamberslides (Paper I) or 150 000 cell/cm² on matrigel (BD Biosciences) coated plastic chamberslides (Paper II). The cells were processed further for BrdU labeling and viral transduction, see separate sections.

For co-cultures, primary cells from E11.5 and E12.5 VM were plated in differentiation medium on PLL coated plastic slides (150 000 cells/cm²). Expanded neurosphere cells were added to the wells 24 hours later (\approx 400 000 cells/cm²) and the slides were maintained under differentiating conditions for 5-7 days.

Immunocytochemistry

Cells were fixed in 4% PFA or 4% PFA/0.4% Glutaraldehyd (for GABA staining) for 15 min and rinsed in PBS or KPBS for 3x10 min. A pre-incubation for 1 hour with 10% normal serum and 0.1% triton-X in KPBS was always performed before applying primary antibodies. Cells were incubated with primary antibody overnight at room temperature, followed by gentle rinses with KPBS, 3x10 min. Appropriate Cy2-, Cy3- and Cy5-conjugated secondary antibodies (1:200, Jackson lab) were applied and left on for 2 hours, room temperature. Double and triple stainings were performed simultaneously. For BrdU staining the cells were permeabilized with -20°C methanol for 20 min then treated with 1M HCl for 20 min at 37°C . The acid was neutralized with 0.1 borate buffer. DAPI nuclear stain (1:1000, Sigma) was included to visualize all cells for subsequent cell counts. Slides were coverslipped using PVA-DAPCO mounting medium and kept covered at 8°C .

RT-PCR

mRNA was isolated from freshly dissected tissue as well as proliferating neurospheres using the kit RNAqueousTM-4PCR from Ambion. In brief, proliferating spheres and freshly dissected tissue were dissociated in the supplied Lysis/Binding solution. The samples were immediately transferred to -80°C if they were not processed further directly after collection.

The supplied protocol was followed using a centrifuge to draw solutions through the filter cartridge. Suggested maximum volumes, centrifugation times and speed were used throughout. The RNA was eluted with $40\ \mu\text{l} + 20\ \mu\text{l}$ Elution solution.

The isolated RNA was DNase treated twice to remove remaining DNA. RNase free DNase1 (Ambion) and DNase buffer were incubated with each sample at 37° , 1 hr. The DNase was inactivated with DNase inactivation reagent, supplied with the kit and the sample transferred to a fresh tube. Heat treatment, 95° for 4 min, was followed by another round of DNase treatment and inactivation.

The samples were analyzed on agarose gel to ensure that the integrity of RNA was intact. PCR using primers for the housekeeping gene *Gpdh* established that the samples were free of genomic DNA.

cDNA was synthesized from the samples using the SUPERSRIPT II for RT-PCR protocol (Invitrogen). Approximately $1\ \mu\text{g}$ of total RNA was used per $20\ \mu\text{l}$ reaction.

To enable a semi-quantative analysis of the samples, PCR for *Gpdh* was performed on each sample comparing amplified products after increasing number of cycles on an agarose gel. Based on the amount of product after 20, 25, 30 and 35 cycles respectively the cDNA from each sample was diluted to approximately the same concentration and thereafter used for PCR analysis. Full list of PCR primers and programs are found in table 1.

FACS procedure

Sub-dissected VM from E12.5 Ngn2^{+/-} or wild-type embryos was dissociated mechanically in PBS without Ca²⁺ and Mg²⁺ (Gibco) with 1 mM EDTA and 0.5% BSA. The cells were diluted to approximately 3 x 10⁶ cells/ml. To exclude dead cells and cell debris at sorting, 7-aminoactinomycin-D (7AAD, Sigma) was added at 10 μ l/ml. Cells were sorted using DIVA flow cytometer (Becton-Dickson) equipped with 488 nm argon and 633 nm helium-neon lasers. Gates for GFP-negative cells were set using wild-type cells while gates for GFP^{high} and GFP^{low} were chosen arbitrarily.

Transplantation

Unsorted, GFP^{neg} or GFP^{pos} cells were transplanted to neonatal rats under deep hypothermic anaesthesia. 50 000 cells were grafted unilaterally into the striatum of each animal. Injection co-ordinates (mm) were 0.7 anterior and 1.9 lateral to bregma, and 2.9 below the dural surface. The rats were allowed to survive for 4 weeks before processed for IHC (as described above).

BrdU labeling of embryonic tissue and primary cells

Pregnant females were injected i.p with BrdU dissolved in 37°C PBS (8 mg/ml) such that they received 50 mg BrdU/g body weight. BrdU was allowed to incorporate for 45 min before the embryos were collected.

FACSorted GFP^{high}, GFP^{low}, GFP^{neg} primary cells were maintained either under proliferating conditions for two days, with BrdU present in the culture medium (0.2 μ M BrdU, Sigma) and subsequently differentiated during 5 days, or plated in differentiation medium and maintained for 7 days.

Retroviral transduction

Primary cells (Paper II) and neurosphere expanded cells (Paper III) were transduced with VSV-G pseudotyped retrovirus with IRES2-eGFP, Ngn2-IRES2-eGFP and Nurr1-IRES2-eGFP vectors respectively. All viral vectors were derived from Moloney Murine Leukemia virus, murine stem cell based (Clontech). The Ngn2-IRES2-GFP construct was received from Dr. Jonas Frisén and the cDNA for Nurr1 was received from Dr. Thomas Perlmann. The viral vectors were packaged using a packaging cell line (Ory et al., 1996) and had a titre of around 10⁸ TU/ml

Neurosphere expanded cells were passaged and seeded out in proliferation medium (passage 3). 24 hours later the cells were spun down and incubated with retrovirus in a small volume for maximum exposure to the viral particles. Proteamine sulfate was added at 4 mg/ml. The ratio was 1 TU/cell. Cells were resuspended in proliferating medium after four hours and plated out in non-coated 4-well plastic slides (100 000 cells/ml). The cultures were treated as neurosphere cultures and fresh medium (100 μ l/well) and growth factors were added every other day. After

7 days total of expansion the proliferation medium was removed and differentiation medium was added to the cells. They were differentiated for 4 days.

Primary cells were dissociated and plated out in proliferation medium. Fifteen hours later virus was added to the cells (1 TU/cell). Another 12 hours later the medium was switched to differentiation medium and the cells were left to differentiate for 5 days.

List of mediums

Neurosphere medium

<u>To make 100 ml</u>		<u>Final concentration</u>	
10 ml	DMEM/F12 10X	1X	(Gibco no 32500-027)
1 ml	L-Glutamine 200 mM	2 mM	(Sigma no G5763)
1.5 ml	NaHCO ₃ 7.5%	1.125%	(Sigma no S5761)
2 ml	Glucose 30%	0.6%	(Sigma no G7021)
1 ml	Penicillin/Streptomycin	1000 U/ml, 100 mg/ml	(Gibco no 5140-114)
82.5 ml	MilliQ sterile H ₂ O		
2 ml	B27 (sterile)	0.2X	(Gibco no 17504-044)

Differentiation medium

Neurosphere medium and 1% FBS

Freezing medium

Neurosphere medium and 10% BSA, 7.5% DMSO

Table 1. PCR primers and programs

<u>Primer sequence</u>			<u>Program</u>
<i>Aldh1:</i>	sense	5'-ACT CTC AGC AGT GGT ACA CA-3'	A
	antisense	5'-CAG CTG ACT CTG CAG TCA TT-3'	
<i>En1:</i>	sense	5'-TCA AGA CTG ACT CAC AGC AAC CCC-3'	D
	antisense	5'-CTT TGT CCT GAA CCG TGG TGG TAG-3'	
<i>Lmx1b:</i>	sense	5'-AGA CAT TGG CAG CAG AGA CA-3'	D**
	antisense	5'-CTG AGG GAG GTA TCA CTA TC-3'	
<i>Ngn2:</i>	sense	5'-CGT CAA TAC TGA GAC TCT GC-3'	A*
	antisense	5'-ATC TTC GTG AGC TTG GCA TC-3'	
<i>Nurr1:</i>	sense	5'-TAA AAG GCC GGA GAG GTC GTC-3'	A*
	antisense	5'-CTC TCT TGG GTT CCT TGA GCC-3'	
<i>Pbx1:</i>	sense	5'-GTG AGA GAA GTC AGA GCT CA-3'	A
	antisense	5'-ACT GGA AGG AGA TCT ACA GG-3'	
<i>Shh:</i>	sense	5'-GCA CCC CAA AAA GCT GAC CC -3'	E
	antisense	5'-CGT GGT GAT GTC CAC TGC TCG-3'	
<i>Smo:</i>	sense	5'-CTG CCT AGA AGA GCT GTG TA-3'	C
	antisense	5'-TCA CTG CAG TTT TAA GGT GC-3'	
<i>TH:</i>	sense	5'-TTC CGT GTG TTT CAG TGC AC-3'	D
	antisense	5'-TCC AGT ACA TCA ATG GCC AG-3'	
<i>Gpdh:</i>	sense	5'-ACC ACA GTC CAT GCC ATC-3'	B
	antisense	5'-TCC ACC ACC CTG TTG CTG TA-3'	
Program:	A: 94°, 5 min; 94°, 1 min; 55°, 1 min, 72°, 1 min B: 94°, 5 min; 94°, 1 min; 56°, 1 min, 72°, 1 min C: 94°, 5 min; 94°, 1 min; 57°, 1 min; 72°, 1 min D: 94°, 5 min; 94°, 1 min; 60°, 1 min; 72°, 1 min E: 94°, 2 min; 94°, 30 s; 58°, 30 s; 72°, 1 min		

* 0.5 mM MgCl and 5% DMSO in PCR buffer

** 0.5mM MgCl

Table 2. List of antibodies

Antibody	Dilution	Source	Species	Special treatments
AADC	1:1000	Chemicon	rabbit	1
Aldh1	1:500	R. Lindahl	rabbit	
β-III-tubulin	1:333	Sigma	mouse	
BrdU	1:100	Oxford Biotechnology	rat	2a, 2b
Brn3a	1:50	Santa Cruz	mouse	1, 4
Calbindin D28	1:1000	Chemicon	rabbit	1
ChAT	1:1000	Chemicon	rabbit	
CNPase	1:100	Sigma	mouse	
Dlk-1	1:5000	CH Jensen	rabbit	
En1/2	1:500/1:1000	A Joyner	rabbit	
GABA	1:250/1:2000	Sigma	rabbit	(tissue/cells)
GFAP	1:1000	DAKO	rabbit	
GFP	1:5000	Chemicon	chicken	
Girk2	1:80	Alomone Labs	rabbit	1
Isl1	1:100	Hybridoma	mouse	1
Isl1	1:100	Abcam	rabbit	
Isl1/2	1:500	H & T Edlund	rabbit	
Ki67	1:100	Dianova	rabbit	
Ki67	1:50	DAKO	rat	1
M2M6	1:50	Hybridoma	rat	
Nestin	1:10/1:1000	BD Biosciences	rat	adult /embryonic tissue
NeuroD	1:400	Santa Cruz	goat	
Ngn2	1:1000	M Nakafuku	rabbit	
Ngn2	1:20	D. Anderson	mouse	1
Nurr1	1:1000	Santa Cruz	rabbit	
Parvalbumin	1:2000	Sigma	mouse	
Pbx1/2/3	1:400	Santa Cruz	rabbit	
Pitx3	1:400/1:1000	P Burbach	rabbit	1 (tissue/cells)
PSA-NCAM	1:200	Chemicon	mouse	
TH	1:1000	Chemicon	mouse	
TH	1:1000	Pelfreeze	rabbit	
VMAT2	1:1000	Chemicon	rabbit	1
5-HT	1:10000	Incstar	rabbit	

1. Antigen retrieval with 10 mM citrate buffer 20 min
2. Antigen retrieval
 - a) For sections: antigen retrieval with 1M HCl 30 min (65°)
 - b) For cells: MeOH 20 min (-20°), 2M HCl 20 min (37°), 0.1M 2x10 min (RT)
3. Pre-incubation with milk/serum/BSA solution

References

REFERENCES

- Altman, J. and Bayer, S. A.** (1981). Development of the brain stem in the rat. V. Thymidine-radiographic study of the time of origin of neurons in the midbrain tegmentum. *J Comp Neurol* **198**, 677-716.
- Alvarez-Buylla, A., Garcia-Verdugo, J. M. and Tramontin, A. D.** (2001). A unified hypothesis on the lineage of neural stem cells. *Nat Rev Neurosci* **2**, 287-93.
- Andersson, E., Marklund, U., Deng, Q., Friling, S., Alekseenko, Z., Perlmann, T. and Ericsson, J.** (2005). Identification of intrinsic determinants of midbrain dopamine neurons. *Cell, in press*
- Anthony, T. E., Klein, C., Fishell, G. and Heintz, N.** (2004). Radial glia serve as neuronal progenitors in all regions of the central nervous system. *Neuron* **41**, 881-90.
- Arvidsson, A., Collin, T., Kirik, D., Kokaia, Z. and Lindvall, O.** (2002). Neuronal replacement from endogenous precursors in the adult brain after stroke. *Nat Med* **8**, 963-70.
- Aubert, J., Stavridis, M. P., Tweedie, S., O'Reilly, M., Vierlinger, K., Li, M., Ghazal, P., Pratt, T., Mason, J. O., Roy, D. et al.** (2003). Screening for mammalian neural genes via fluorescence-activated cell sorter purification of neural precursors from Sox1-gfp knock-in mice. *Proc Natl Acad Sci U S A* **100 Suppl 1**, 11836-41.
- Backman, C., Perlmann, T., Wallen, A., Hoffer, B. J. and Morales, M.** (1999). A selective group of dopaminergic neurons express Nurr1 in the adult mouse brain. *Brain Res* **851**, 125-32.
- Bain, G., Kitchens, D., Yao, M., Huettner, J. E. and Gottlieb, D. I.** (1995). Embryonic stem cells express neuronal properties in vitro. *Dev Biol* **168**, 342-57.
- Barraud, P., Thompson, L., Kirik, D., Bjorklund, A. and Parmar, M.** (2005). Isolation and characterization of neural precursor cells from the Sox1-GFP reporter mouse. *Eur J Neurosci* **22**, 1555-69.
- Bayer, S. A., Wills, K. V., Triarhou, L. C. and Ghetti, B.** (1995). Time of neuron origin and gradients of neurogenesis in midbrain dopaminergic neurons in the mouse. *Exp Brain Res* **105**, 191-9.
- Bertrand, N., Castro, D. S. and Guillemot, F.** (2002). Proneural genes and the specification of neural cell types. *Nat Rev Neurosci* **3**, 517-30.
- Björklund, A. and Lindvall, O.** (1984). Dopamine-containing systems in the CNS. In *Classical transmitters in the CNS*, vol. 2 (ed. A. Björklund and T. Hökfelt), pp. 55-122. Amsterdam, New York and Oxford: Elsevier.
- Bouvier, M. M. and Mytilineou, C.** (1995). Basic fibroblast growth factor increases division and delays differentiation of dopamine precursors in vitro. *J Neurosci* **15**, 7141-9.
- Broccoli, V., Boncinelli, E. and Wurst, W.** (1999). The caudal limit of Otx2 expression positions the isthmic organizer. *Nature* **401**, 164-8.
- Brodski, C., Weisenhorn, D. M., Signore, M., Sillaber, I., Oesterheld, M., Broccoli, V., Acampora, D., Simeone, A. and Wurst, W.** (2003). Location and size of dopaminergic and serotonergic cell populations are controlled by the position of the midbrain-hindbrain organizer. *J Neurosci* **23**, 4199-207.
- Bylund, M., Andersson, E., Novitsch, B. G. and Muhr, J.** (2003). Vertebrate neurogenesis is counteracted by Sox1-3 activity. *Nat Neurosci* **6**, 1162-8.
- Caldwell, M. A. and Svendsen, C. N.** (1998). Heparin, but not other proteoglycans potentiates the mitogenic effects of FGF-2 on mesencephalic precursor cells. *Exp Neurol* **152**, 1-10.
- Carvey, P. M., Ling, Z. D., Sortwell, C. E., Pitzer, M. R., McGuire, S. O., Storch, A. and Collier, T. J.** (2001). A clonal line of mesencephalic progenitor cells converted to dopamine neurons by hematopoietic cytokines: a source of cells for transplantation in Parkinson's disease. *Exp Neurol* **171**, 98-108.

- Castillo, S. O., Baffi, J. S., Palkovits, M., Goldstein, D. S., Kopin, I. J., Witta, J., Magnuson, M. A. and Nikodem, V. M.** (1998). Dopamine biosynthesis is selectively abolished in substantia nigra/ventral tegmental area but not in hypothalamic neurons in mice with targeted disruption of the *Nurr1* gene. *Mol Cell Neurosci* **11**, 36-46.
- Cazorla, P., Smidt, M. P., O'Malley, K. L. and Burbach, J. P.** (2000). A response element for the homeodomain transcription factor Ptx3 in the tyrosine hydroxylase gene promoter. *J Neurochem* **74**, 1829-37.
- Cheung, M., Abu-Elmagd, M., Clevers, H. and Scotting, P. J.** (2000). Roles of Sox4 in central nervous system development. *Brain Res Mol Brain Res* **79**, 180-91.
- Crossley, P. H., Martinez, S. and Martin, G. R.** (1996). Midbrain development induced by FGF8 in the chick embryo. *Nature* **380**, 66-8.
- Dahlstrom, A. and Fuxe, K.** (1964). Localization of monoamines in the lower brain stem. *Experientia* **20**, 398-9.
- Davis, C. A. and Joyner, A. L.** (1988). Expression patterns of the homeo box-containing genes *En-1* and *En-2* and the proto-oncogene *int-1* diverge during mouse development. *Genes Dev* **2**, 1736-44.
- Di Porzio, U., Zuddas, A., Cosenza-Murphy, D. B. and Barker, J. L.** (1990). Early appearance of tyrosine hydroxylase immunoreactive cells in the mesencephalon of mouse embryos. *Int J Dev Neurosci* **8**, 523-32.
- Dunnett, S. B. and Bjorklund, A.** (1999). Prospects for new restorative and neuroprotective treatments in Parkinson's disease. *Nature* **399**, A32-9.
- Dunnett, S. B., Nathwani, F. and Björklund, A.** (2000). The integration and function of striatal grafts. In *Functional Neural Transplantation II. Novel Cell Therapies for CNS disorders*, vol. 127 (ed. S. B. Dunnett and A. Björklund), pp. 345-380. New York: Elsevier.
- Farah, M. H., Olson, J. M., Sucic, H. B., Hume, R. I., Tapscott, S. J. and Turner, D. L.** (2000). Generation of neurons by transient expression of neural bHLH proteins in mammalian cells. *Development* **127**, 693-702.
- Fode, C., Gradwohl, G., Morin, X., Dierich, A., LeMeur, M., Goridis, C. and Guillemot, F.** (1998). The bHLH protein NEUROGENIN 2 is a determination factor for epibranchial placode-derived sensory neurons. *Neuron* **20**, 483-94.
- Fode, C., Ma, Q., Casarosa, S., Ang, S. L., Anderson, D. J. and Guillemot, F.** (2000). A role for neural determination genes in specifying the dorsoventral identity of telencephalic neurons. *Genes Dev* **14**, 67-80.
- Foster, G. A., Schultzberg, M., Kokfelt, T., Goldstein, M., Hemmings, H. C., Jr., Ouimet, C. C., Walaas, S. I. and Greengard, P.** (1988). Ontogeny of the dopamine and cyclic adenosine-3':5'-monophosphate-regulated phosphoprotein (DARPP-32) in the pre- and postnatal mouse central nervous system. *Int J Dev Neurosci* **6**, 367-86.
- Frielingsdorf, H., Schwarz, K., Brundin, P. and Mohapel, P.** (2004). No evidence for new dopaminergic neurons in the adult mammalian substantia nigra. *Proc Natl Acad Sci U S A* **101**, 10177-82.
- Gage, F. H.** (2000). Mammalian neural stem cells. *Science* **287**, 1433-8.
- Gaiano, N. and Fishell, G.** (1998). Transplantation as a tool to study progenitors within the vertebrate nervous system. *J Neurobiol* **36**, 152-61.
- Gradwohl, G., Fode, C. and Guillemot, F.** (1996). Restricted expression of a novel murine atonal-related bHLH protein in undifferentiated neural precursors. *Dev Biol* **180**, 227-41.
- Hanaway, J., McConnell, J. A. and Netsky, M. G.** (1971). Histogenesis of the substantia nigra, ventral tegmental area of Tsai and interpeduncular nucleus: an autoradiographic study of the mesencephalon in the rat. *J Comp Neurol* **142**, 59-73.

- Hirsch, M. R., Tiveron, M. C., Guillemot, F., Brunet, J. F. and Goridis, C.** (1998). Control of noradrenergic differentiation and Phox2a expression by MASH1 in the central and peripheral nervous system. *Development* **125**, 599-608.
- Hitoshi, S., Tropepe, V., Ekker, M. and van der Kooy, D.** (2002). Neural stem cell lineages are regionally specified, but not committed, within distinct compartments of the developing brain. *Development* **129**, 233-44.
- Horiguchi, S., Takahashi, J., Kishi, Y., Morizane, A., Okamoto, Y., Koyanagi, M., Tsuji, M., Tashiro, K., Honjo, T., Fujii, S. et al.** (2004). Neural precursor cells derived from human embryonic brain retain regional specificity. *J Neurosci Res* **75**, 817-24.
- Hwang, D. Y., Ardayfio, P., Kang, U. J., Semina, E. V. and Kim, K. S.** (2003). Selective loss of dopaminergic neurons in the substantia nigra of Pitx3-deficient aphakia mice. *Brain Res Mol Brain Res* **114**, 123-31.
- Hynes, M., Porter, J. A., Chiang, C., Chang, D., Tessier-Lavigne, M., Beachy, P. A. and Rosenthal, A.** (1995a). Induction of midbrain dopaminergic neurons by Sonic hedgehog. *Neuron* **15**, 35-44.
- Hynes, M., Poulsen, K., Tessier-Lavigne, M. and Rosenthal, A.** (1995b). Control of neuronal diversity by the floor plate: contact-mediated induction of midbrain dopaminergic neurons. *Cell* **80**, 95-101.
- Iwawaki, T., Kohno, K. and Kobayashi, K.** (2000). Identification of a potential nurr1 response element that activates the tyrosine hydroxylase gene promoter in cultured cells. *Biochem Biophys Res Commun* **274**, 590-5.
- Jensen, C. H., Meyer, M., Schroder, H. D., Kliem, A., Zimmer, J. and Teisner, B.** (2001). Neurons in the monoaminergic nuclei of the rat and human central nervous system express FA1/dlk. *Neuroreport* **12**, 3959-63.
- Jensen, J. B., Bjorklund, A. and Parmar, M.** (2004). Striatal neuron differentiation from neurosphere-expanded progenitors depends on Gsh2 expression. *J Neurosci* **24**, 6958-67.
- Jessell, T. M.** (2000). Neuronal specification in the spinal cord: inductive signals and transcriptional codes. *Nat Rev Genet* **1**, 20-9.
- Jessell, T. M. and Sanes, J. R.** (2000a). The generation and survival of nerve cells. In *Principles of neural science*, (ed. E. R. Kandel J. H. Schwartz and T. M. Jessell), pp. 1041-1062. New York: McGraw-Hill.
- Jessell, T. M. and Sanes, J. R.** (2000b). The induction and patterning of the nervous system. In *Principles of neural science*, (ed. E. R. Kandel J. H. Schwartz and T. M. Jessell), pp. 1019-1040. New York: McGraw-Hill.
- Joyner, A. L.** (1996). Engrailed, Wnt and Pax genes regulate midbrain-hindbrain development. *Trends Genet* **12**, 15-20.
- Joyner, A. L., Herrup, K., Auerbach, B. A., Davis, C. A. and Rossant, J.** (1991). Subtle cerebellar phenotype in mice homozygous for a targeted deletion of the En-2 homeobox. *Science* **251**, 1239-43.
- Kawano, H., Ohyama, K., Kawamura, K. and Nagatsu, I.** (1995). Migration of dopaminergic neurons in the embryonic mesencephalon of mice. *Brain Res Dev Brain Res* **86**, 101-13.
- Kele, J., Simplicio, N., Ferri, A.L.M., Mira, H., Guillemot, F., Arenas, E. and Ang, S-L.** Ngn2 is required for the development of ventral midbrain dopaminergic neurons. *submitted*
- Kilpatrick, T. J. and Bartlett, P. F.** (1993). Cloning and growth of multipotential neural precursors: requirements for proliferation and differentiation. *Neuron* **10**, 255-65.
- Kim, J. Y., Koh, H. C., Lee, J. Y., Chang, M. Y., Kim, Y. C., Chung, H. Y., Son, H., Lee, Y. S., Studer, L., McKay, R. et al.** (2003). Dopaminergic neuronal differentiation from rat embryonic neural precursors by Nurr1 overexpression. *J Neurochem* **85**, 1443-54.
- Kintner, C.** (2002). Neurogenesis in embryos and in adult neural stem cells. *J Neurosci* **22**, 639-43.
- Kirschstein, R. and Skirboll, L.** (2001). Stem Cells: Scientific Progress and Future Research Directions. Department of Health and Human Services. June 2001, vol. 2001 (ed).

- Klein, C., Butt, S. J., Machold, R. P., Johnson, J. E. and Fishell, G.** (2005). Cerebellum- and forebrain-derived stem cells possess intrinsic regional character. *Development* **132**, 4497-508.
- Lang, A. E. and Lozano, A. M.** (1998a). Parkinson's disease. First of two parts. *N Engl J Med* **339**, 1044-53.
- Lang, A. E. and Lozano, A. M.** (1998b). Parkinson's disease. Second of two parts. *N Engl J Med* **339**, 1130-43.
- Lebel, M., Gauthier, Y., Moreau, A. and Drouin, J.** (2001). Pitx3 activates mouse tyrosine hydroxylase promoter via a high-affinity binding site. *J Neurochem* **77**, 558-67.
- Lee, K. J. and Jessell, T. M.** (1999). The specification of dorsal cell fates in the vertebrate central nervous system. *Annu Rev Neurosci* **22**, 261-94.
- Lee, S-H., Lumelsky, N., Studer, L., Auerbach, J.M. and McKay, R.D.** (2000). Efficient generation of midbrain and hindbrain neurons from mouse embryonic stem cells. *Nature* **18**, 675-79.
- Lee, S. K. and Pfaff, S. L.** (2003). Synchronization of neurogenesis and motor neuron specification by direct coupling of bHLH and homeodomain transcription factors. *Neuron* **38**, 731-45.
- Lendahl, U., Zimmerman, L. B. and McKay, R. D.** (1990). CNS stem cells express a new class of intermediate filament protein. *Cell* **60**, 585-95.
- Liang, C. L., Sinton, C. M. and German, D. C.** (1996). Midbrain dopaminergic neurons in the mouse: co-localization with Calbindin-D28K and calretinin. *Neuroscience* **75**, 523-33.
- Lillien, L.** (1998). Neural progenitors and stem cells: mechanisms of progenitor heterogeneity. *Curr Opin Neurobiol* **8**, 37-44.
- Lindahl, R. and Evces, S.** (1984). Rat liver aldehyde dehydrogenase. II. Isolation and characterization of four inducible isozymes. *J Biol Chem* **259**, 11991-6.
- Lindvall, O., Brundin, P., Widner, H., Rehncrona, S., Gustavii, B., Frackowiak, R., Leenders, K. L., Sawle, G., Rothwell, J. C., Marsden, C. D. et al.** (1990). Grafts of fetal dopamine neurons survive and improve motor function in Parkinson's disease. *Science* **247**, 574-7.
- Ling, Z. D., Potter, E. D., Lipton, J. W. and Carvey, P. M.** (1998). Differentiation of mesencephalic progenitor cells into dopaminergic neurons by cytokines. *Exp Neurol* **149**, 411-23.
- Liu, A. and Joyner, A. L.** (2001). Early anterior/posterior patterning of the midbrain and cerebellum. *Annu Rev Neurosci* **24**, 869-96.
- Lo, L., Dormand, E., Greenwood, A. and Anderson, D. J.** (2002). Comparison of the generic neuronal differentiation and neuron subtype specification functions of mammalian achaete-scute and atonal homologs in cultured neural progenitor cells. *Development* **129**, 1553-67.
- Lo, L., Tiveron, M. C. and Anderson, D. J.** (1998). MASH1 activates expression of the paired homeodomain transcription factor Phox2a, and couples pan-neuronal and subtype-specific components of autonomic neuronal identity. *Development* **125**, 609-20.
- Lumsden, A. and Krumlauf, R.** (1996). Patterning the vertebrate neuraxis. *Science* **274**, 1109-15.
- Malatesta, P., Hack, M. A., Hartfuss, E., Kettenmann, H., Klinkert, W., Kirchhoff, F. and Gotz, M.** (2003). Neuronal or glial progeny: regional differences in radial glia fate. *Neuron* **37**, 751-64.
- Martinez-Serrano, A., Rubio, F. J., Navarro, B., Bueno, C. and Villa, A.** (2001). Human neural stem and progenitor cells: in vitro and in vivo properties, and potential for gene therapy and cell replacement in the CNS. *Curr Gene Ther* **1**, 279-99.
- Maxwell, S. L., Ho, H. Y., Kuehner, E., Zhao, S. and Li, M.** (2005). Pitx3 regulates tyrosine hydroxylase expression in the substantia nigra and identifies a subgroup of mesencephalic dopaminergic progenitor neurons during mouse development. *Dev Biol* **282**, 467-79.
- McCaffery, P. and Drager, U. C.** (1994). High levels of a retinoic acid-generating dehydrogenase in the meso-telencephalic dopamine system. *Proc Natl Acad Sci U S A* **91**, 7772-6.
- McKay, R.** (1997). Stem cells in the central nervous system. *Science* **276**, 66-71.

- Millen, K. J., Wurst, W., Herrup, K. and Joyner, A. L.** (1994). Abnormal embryonic cerebellar development and patterning of postnatal foliation in two mouse *Engrailed-2* mutants. *Development* **120**, 695-706.
- Milosevic, J., Schwarz, S. C., Krohn, K., Poppe, M., Storch, A. and Schwarz, J.** (2005). Low atmospheric oxygen avoids maturation, senescence and cell death of murine mesencephalic neural precursors. *J Neurochem* **92**, 718-29.
- Mizuguchi, R., Sugimori, M., Takebayashi, H., Kosako, H., Nagao, M., Yoshida, S., Nabeshima, Y., Shimamura, K. and Nakafuku, M.** (2001). Combinatorial roles of *olig2* and *neurogenin2* in the coordinated induction of pan-neuronal and subtype-specific properties of motoneurons. *Neuron* **31**, 757-71.
- Mytilineou, C., Park, T. H. and Shen, J.** (1992). Epidermal growth factor-induced survival and proliferation of neuronal precursor cells from embryonic rat mesencephalon. *Neurosci Lett* **135**, 62-6.
- Noctor, S. C., Flint, A. C., Weissman, T. A., Dammerman, R. S. and Kriegstein, A. R.** (2001). Neurons derived from radial glial cells establish radial units in neocortex. *Nature* **409**, 714-20.
- Nunes, I., Tovmasian, L. T., Silva, R. M., Burke, R. E. and Goff, S. P.** (2003). *Pitx3* is required for development of substantia nigra dopaminergic neurons. *Proc Natl Acad Sci U S A* **100**, 4245-50.
- Okabe, M., Ikawa, M., Kominami, K., Nakanishi, T. and Nishimune, Y.** (1997). 'Green mice' as a source of ubiquitous green cells. *FEBS Lett* **407**, 313-9.
- Okabe, S., Forsberg-Nilsson, K., Spiro, A. C., Segal, M. and McKay, R. D.** (1996). Development of neuronal precursor cells and functional postmitotic neurons from embryonic stem cells in vitro. *Mech Dev* **59**, 89-102.
- Ory, D. S., Neugeboren, B. A. and Mulligan, R. C.** (1996). A stable human-derived packaging cell line for production of high titer retrovirus/vesicular stomatitis virus G pseudotypes. *Proc Natl Acad Sci U S A* **93**, 11400-6.
- Parmar, M., Skogh, C., Bjorklund, A. and Campbell, K.** (2002). Regional specification of neurosphere cultures derived from subregions of the embryonic telencephalon. *Mol Cell Neurosci* **21**, 645-56.
- Parras, C. M., Schuurmans, C., Scardigli, R., Kim, J., Anderson, D. J. and Guillemot, F.** (2002). Divergent functions of the proneural genes *Mash1* and *Ngn2* in the specification of neuronal subtype identity. *Genes Dev* **16**, 324-38.
- Pattyn, A., Goridis, C. and Brunet, J. F.** (2000). Specification of the central noradrenergic phenotype by the homeobox gene *Phox2b*. *Mol Cell Neurosci* **15**, 235-43.
- Pattyn, A., Simplicio, N., van Doorninck, J. H., Goridis, C., Guillemot, F. and Brunet, J. F.** (2004). *Ascl1/Mash1* is required for the development of central serotonergic neurons. *Nat Neurosci* **7**, 589-95.
- Perrier, A. L., Tabar, V., Barberi, T., Rubio, M. E., Bruses, J., Topf, N., Harrison, N. L. and Studer, L.** (2004). Derivation of midbrain dopamine neurons from human embryonic stem cells. *Proc Natl Acad Sci U S A* **101**, 12543-8.
- Pevny, L. and Rao, M. S.** (2003). The stem-cell menagerie. *Trends Neurosci* **26**, 351-9.
- Pevny, L. H., Sockanathan, S., Placzek, M. and Lovell-Badge, R.** (1998). A role for *SOX1* in neural determination. *Development* **125**, 1967-78.
- Ptak, L. R., Hart, K. R., Lin, D. and Carvey, P. M.** (1995). Isolation and manipulation of rostral mesencephalic tegmental progenitor cells from rat. *Cell Transplant* **4**, 335-42.
- Reynolds, B. A. and Rietze, R. L.** (2005). Neural stem cells and neurospheres--re-evaluating the relationship. *Nat Methods* **2**, 333-6.
- Reynolds, B. A., Tetzlaff, W. and Weiss, S.** (1992). A multipotent EGF-responsive striatal embryonic progenitor cell produces neurons and astrocytes. *J Neurosci* **12**, 4565-74.
- Reynolds, B. A. and Weiss, S.** (1992). Generation of neurons and astrocytes from isolated cells of the adult mammalian central nervous system. *Science* **255**, 1707-10.

- Reynolds, B. A. and Weiss, S.** (1996). Clonal and population analyses demonstrate that an EGF-responsive mammalian embryonic CNS precursor is a stem cell. *Dev Biol* **175**, 1-13.
- Rieger, D. K., Reichenberger, E., McLean, W., Sidow, A. and Olsen, B. R.** (2001). A double-deletion mutation in the *Pitx3* gene causes arrested lens development in aphakia mice. *Genomics* **72**, 61-72.
- Sakakibara, S., Imai, T., Hamaguchi, K., Okabe, M., Aruga, J., Nakajima, K., Yasutomi, D., Nagata, T., Kurihara, Y., Uesugi, S. et al.** (1996). Mouse-Musashi-1, a neural RNA-binding protein highly enriched in the mammalian CNS stem cell. *Dev Biol* **176**, 230-42.
- Sakurada, K., Ohshima-Sakurada, M., Palmer, T. D. and Gage, F. H.** (1999). *Nurr1*, an orphan nuclear receptor, is a transcriptional activator of endogenous tyrosine hydroxylase in neural progenitor cells derived from the adult brain. *Development* **126**, 4017-26.
- Saucedo-Cardenas, O., Quintana-Hau, J. D., Le, W. D., Smidt, M. P., Cox, J. J., De Mayo, F., Burbach, J. P. and Conneely, O. M.** (1998). *Nurr1* is essential for the induction of the dopaminergic phenotype and the survival of ventral mesencephalic late dopaminergic precursor neurons. *Proc Natl Acad Sci U S A* **95**, 4013-8.
- Scardigli, R., Baumer, N., Gruss, P., Guillemot, F. and Le Roux, I.** (2003). Direct and concentration-dependent regulation of the proneural gene *Neurogenin2* by *Pax6*. *Development* **130**, 3269-81.
- Scardigli, R., Schuurmans, C., Gradwohl, G. and Guillemot, F.** (2001). Crossregulation between *Neurogenin2* and pathways specifying neuronal identity in the spinal cord. *Neuron* **31**, 203-17.
- Schein, J. C., Hunter, D. D. and Roffler-Tarlov, S.** (1998). *Girk2* expression in the ventral midbrain, cerebellum, and olfactory bulb and its relationship to the murine mutation *weaver*. *Dev Biol* **204**, 432-50.
- Seibt, J., Schuurmans, C., Gradwohl, G., Dehay, C., Vanderhaeghen, P., Guillemot, F. and Polleux, F.** (2003). *Neurogenin2* specifies the connectivity of thalamic neurons by controlling axon responsiveness to intermediate target cues. *Neuron* **39**, 439-52.
- Semina, E. V., Murray, J. C., Reiter, R., Hrstka, R. F. and Graw, J.** (2000). Deletion in the promoter region and altered expression of *Pitx3* homeobox gene in aphakia mice. *Hum Mol Genet* **9**, 1575-85.
- Shibata, T., Yamada, K., Watanabe, M., Ikenaka, K., Wada, K., Tanaka, K. and Inoue, Y.** (1997). Glutamate transporter *GLAST* is expressed in the radial glia-astrocyte lineage of developing mouse spinal cord. *J Neurosci* **17**, 9212-9.
- Shults, C. W., Hashimoto, R., Brady, R. M. and Gage, F. H.** (1990). Dopaminergic cells align along radial glia in the developing mesencephalon of the rat. *Neuroscience* **38**, 427-36.
- Simon, H. H., Saueressig, H., Wurst, W., Goulding, M. D. and O'Leary, D. D.** (2001). Fate of midbrain dopaminergic neurons controlled by the engrailed genes. *J Neurosci* **21**, 3126-34.
- Smidt, M. P., Asbreuk, C. H., Cox, J. J., Chen, H., Johnson, R. L. and Burbach, J. P.** (2000). A second independent pathway for development of mesencephalic dopaminergic neurons requires *Lmx1b*. *Nat Neurosci* **3**, 337-41.
- Smidt, M. P., Smits, S. M., Bouwmeester, H., Hamers, F. P., van der Linden, A. J., Hellemons, A. J., Graw, J. and Burbach, J. P.** (2004). Early developmental failure of substantia nigra dopamine neurons in mice lacking the homeodomain gene *Pitx3*. *Development* **131**, 1145-55.
- Smidt, M. P., van Schaick, H. S., Lanctot, C., Tremblay, J. J., Cox, J. J., van der Kleij, A. A., Wolterink, G., Drouin, J. and Burbach, J. P.** (1997). A homeodomain gene *Ptx3* has highly restricted brain expression in mesencephalic dopaminergic neurons. *Proc Natl Acad Sci U S A* **94**, 13305-10.
- Smits, S. M., Ponnio, T., Conneely, O. M., Burbach, J. P. and Smidt, M. P.** (2003). Involvement of *Nurr1* in specifying the neurotransmitter identity of ventral midbrain dopaminergic neurons. *Eur J Neurosci* **18**, 1731-8.
- Sommer, L., Ma, Q. and Anderson, D. J.** (1996). neurogenins, a novel family of atonal-related bHLH transcription factors, are putative mammalian neuronal determination genes that reveal progenitor cell heterogeneity in the developing CNS and PNS. *Mol Cell Neurosci* **8**, 221-41.

- Specht, L. A., Pickel, V. M., Joh, T. H. and Reis, D. J.** (1981). Light-microscopic immunocytochemical localization of tyrosine hydroxylase in prenatal rat brain. I. Early ontogeny. *J Comp Neurol* **199**, 233-53.
- Storch, A., Lester, H. A., Boehm, B. O. and Schwarz, J.** (2003). Functional characterization of dopaminergic neurons derived from rodent mesencephalic progenitor cells. *J Chem Neuroanat* **26**, 133-42.
- Storch, A., Paul, G., Csete, M., Boehm, B. O., Carvey, P. M., Kupsch, A. and Schwarz, J.** (2001). Long-term proliferation and dopaminergic differentiation of human mesencephalic neural precursor cells. *Exp Neurol* **170**, 317-25.
- Strubing, C., Ahnert-Hilger, G., Shan, J., Wiedenmann, B., Hescheler, J. and Wobus, A. M.** (1995). Differentiation of pluripotent embryonic stem cells into the neuronal lineage in vitro gives rise to mature inhibitory and excitatory neurons. *Mech Dev* **53**, 275-87.
- Studer, L., Csete, M., Lee, S. H., Kabbani, N., Walikonis, J., Wold, B. and McKay, R.** (2000). Enhanced proliferation, survival, and dopaminergic differentiation of CNS precursors in lowered oxygen. *J Neurosci* **20**, 7377-83.
- Studer, L., Tabar, V. and McKay, R. D.** (1998). Transplantation of expanded mesencephalic precursors leads to recovery in parkinsonian rats. *Nat Neurosci* **1**, 290-5.
- Sun, Y., Nadal-Vicens, M., Misono, S., Lin, M. Z., Zubiaga, A., Hua, X., Fan, G. and Greenberg, M. E.** (2001). Neurogenin promotes neurogenesis and inhibits glial differentiation by independent mechanisms. *Cell* **104**, 365-76.
- Svensen, C. N., Fawcett, J. W., Bentlage, C. and Dunnett, S. B.** (1995). Increased survival of rat EGF-generated CNS precursor cells using B27 supplemented medium. *Exp Brain Res* **102**, 407-14.
- Tanabe, Y., William, C. and Jessell, T. M.** (1998). Specification of motor neuron identity by the MNR2 homeodomain protein. *Cell* **95**, 67-80.
- Teitelman, G., Jaeger, C. B., Albert, V., Joh, T. H. and Reis, D. J.** (1983). Expression of amino acid decarboxylase in proliferating cells of the neural tube and notochord of developing rat embryo. *J Neurosci* **3**, 1379-88.
- Thompson, L., Barraud, P., Andersson, E., Kirik, D. and Bjorklund, A.** (2005). Identification of dopaminergic neurons of nigral and ventral tegmental area subtypes in grafts of fetal ventral mesencephalon based on cell morphology, protein expression, and efferent projections. *J Neurosci* **25**, 6467-77.
- Ungerstedt, U.** (1971). Stereotaxic mapping of the monoamine pathways in the rat brain. *Acta Physiol Scand Suppl* **367**, 1-48.
- van den Munchhof, P., Luk, K. C., Ste-Marie, L., Montgomery, J., Blanchet, P. J., Sadikot, A. F. and Drouin, J.** (2003). Pitx3 is required for motor activity and for survival of a subset of midbrain dopaminergic neurons. *Development* **130**, 2535-42.
- Vernay, B., Koch, M., Vaccarino, F., Briscoe, J., Simeone, A., Kageyama, R. and Ang, S. L.** (2005). Otx2 regulates subtype specification and neurogenesis in the midbrain. *J Neurosci* **25**, 4856-67.
- Wallen, A., Zetterstrom, R. H., Solomin, L., Arvidsson, M., Olson, L. and Perlmann, T.** (1999). Fate of mesencephalic AHD2-expressing dopamine progenitor cells in NURR1 mutant mice. *Exp Cell Res* **253**, 737-46.
- Wang, M. Z., Jin, P., Bumcrot, D. A., Marigo, V., McMahon, A. P., Wang, E. A., Woolf, T. and Pang, K.** (1995). Induction of dopaminergic neuron phenotype in the midbrain by Sonic hedgehog protein. *Nat Med* **1**, 1184-8.
- Winkler, C., Kirik, D. and Bjorklund, A.** (2005). Cell transplantation in Parkinson's disease: how can we make it work? *Trends Neurosci* **28**, 86-92.
- Wood, H. B. and Episkopou, V.** (1999). Comparative expression of the mouse Sox1, Sox2 and Sox3 genes from pre-gastrulation to early somite stages. *Mech Dev* **86**, 197-201.

- Wurst, W., Auerbach, A. B. and Joyner, A. L.** (1994). Multiple developmental defects in Engrailed-1 mutant mice: an early mid-hindbrain deletion and patterning defects in forelimbs and sternum. *Development* **120**, 2065-75.
- Wurst, W. and Bally-Cuif, L.** (2001). Neural plate patterning: upstream and downstream of the isthmic organizer. *Nat Rev Neurosci* **2**, 99-108.
- Ye, W., Shimamura, K., Rubenstein, J. L., Hynes, M. A. and Rosenthal, A.** (1998). FGF and Shh signals control dopaminergic and serotonergic cell fate in the anterior neural plate. *Cell* **93**, 755-66.
- Zappone, M. V., Galli, R., Catena, R., Meani, N., De Biasi, S., Mattei, E., Tiveron, C., Vescovi, A. L., Lovell-Badge, R., Ottolenghi, S. et al.** (2000). Sox2 regulatory sequences direct expression of a (beta)-geo transgene to telencephalic neural stem cells and precursors of the mouse embryo, revealing regionalization of gene expression in CNS stem cells. *Development* **127**, 2367-82.
- Zervas, M., Millet, S., Ahn, S. and Joyner, A. L.** (2004). Cell behaviors and genetic lineages of the mesencephalon and rhombomere 1. *Neuron* **43**, 345-57.
- Zetterstrom, R. H., Solomin, L., Jansson, L., Hoffer, B. J., Olson, L. and Perlmann, T.** (1997). Dopamine neuron agenesis in Nurr1-deficient mice. *Science* **276**, 248-50.
- Zetterstrom, R. H., Williams, R., Perlmann, T. and Olson, L.** (1996). Cellular expression of the immediate early transcription factors Nurr1 and NGFI-B suggests a gene regulatory role in several brain regions including the nigrostriatal dopamine system. *Brain Res Mol Brain Res* **41**, 111-20.
- Zhao, M., Momma, S., Delfani, K., Carlen, M., Cassidy, R. M., Johansson, C. B., Brismar, H., Shupliakov, O., Frisen, J. and Janson, A. M.** (2003). Evidence for neurogenesis in the adult mammalian substantia nigra. *Proc Natl Acad Sci U S A* **100**, 7925-30.
- Zhao, S., Maxwell, S., Jimenez-Beristain, A., Vives, J., Kuehner, E., Zhao, J., O'Brien, C., de Felipe, C., Semina, E. and Li, M.** (2004). Generation of embryonic stem cells and transgenic mice expressing green fluorescence protein in midbrain dopaminergic neurons. *Eur J Neurosci* **19**, 1133-40.

Acknowledgements

ACKNOWLEDGEMENTS

The acknowledgement, being the most read part of any thesis, should not be written the night before the final, absolutely last (and almost-too-late) submission deadline (Doctors-to-be take note!). But true to form of overestimating my speed and ability to GET THINGS DONE, here I am. However, I hope my appreciation and gratitude towards all you who have made my PhD degree possible and so enjoyable is known and felt even without these final words.

First and foremost I would like to thank the “ladies of the lab” without whom the lab would collapse into a disorganized mess. What you do for the rest of us can never be overestimated! You have all been such a great addition and help during my PhD!

The inimitable Anita Frank who is efficient, smart and one step ahead all the time. There is no problem you can't fix and you do it effortlessly and always with a smile. The great mood of the lab is largely because of you!

Elsy, who worked relentlessly with Ngn2 genotyping, and never lost patience with me even at my most unorganized self. Du är en klippta! This thesis would never have been finished within the time it did without you. (and thanks for lending me the shoes!)

Ulla, you keep us all and now two “snittlabs” in check. Thanks for going that extra mile to make the lab a home away from home with flowers, fruit, decorations, great homebaked cakes and morning coffee that is the envy of everyone around. And it's great to have an ally who understands the unsurpassed enjoyment of a really good flea-market and appreciate a great bargain!

Anna-Karin, thanks for the 1554 billion PLL-coated slides and taking care of my cells at times when I could not be in the lab. I so appreciate your willingness to help out (and your taste in music)!

Anneli, who is one of the coolest grown-ups I know and has a great sense of what is real and worthwhile. Nothing is ever a problem when working with you.

Birgit, thanks for picking up loose ends in the cell lab and helping me out, whenever I asked!

Christina and Eva who just make Monday morning (or Thursday afternoon) coffee so much more fun, with a smile and laugh to share (blev det bättre än Johans, Christina?)

I would like to thank my supervisor Anders Björklund for letting me be a part of his lab and its truly unique atmosphere. You have generously given me the opportunity to meet many great scientist and collaborators, at the lab and at meetings and you have increased my knowledge on so many things. Your time and money have been well spent and I'm grateful for both!

Thanks also to my co-supervisor Malin for taking your role seriously and bestowing such motherly care upon me ;o) ! You've taken time and effort to make my projects, papers and thesis better and I thank you immensely for that.

Thanks also to the people at NS Gene, Bengt, Mette and Lars for support and collaborations!

I've had excellent collaborators who besides being super friendly people have taught me much about the science world:

Josephine, who has wonderful ideas and make sure to follow them through. It's been great having you to bounce ideas off and share music preferences with. You're a fun and smart person and your skills will take you far!

Thanks to Lachlan for taking such care in the presentation of data, you're an absolute wiz at making fabulous images for manuscripts. Also the greatest guy to hang out with outside the lab, always up for a beer (even if I'm not!). The stay at the Spanish Villa was a truly amazing visit. I hope we will be able to see much more of you, Jo and Oliver Downunder.

Dwain and your endless optimism and enthusiasm. You're kool, maaan...! How you manage to stay so cheerful while working more than anyone is a mystery.

Perrine who is a meticulous scientist and the sweetest, most helpful person I know.

Emeli and Jessica for your work!

Thanks a million to Bengt who let me push the deadline more than once and still made the thesis look stunning (don't you think?)

The Björklund lab has been riddled with nice, entertaining, sweet and intelligent people over the years. My PhD would never have been the same without them:

Special salute to Marina, Nathalie and of course Simon for fun, interesting and uninhibited conversations in the microscope room. The truth is in there! Marina, you are so easygoing and normal! Also big up for having such excellent taste in shoes!! (Can I have them?...:o) Simon, you did a great feat in enduring the girls' room but probably learned a few things on the way. I think we have kept each other informed on the ever-so-important boy-girl issues and other things that are soo not lab-related (and some that are....)

Matt for being the nicest guy in the whole place. Don't lose your charming and helpful attitude towards people! Loved your work on the Big L!

My roommates over the years who made it the cosiest and best office in the house: Låtta (I'll always remember the sofa....), Håkan (the desk!), Janne (the screensaver!), Cicci (the "shield"!). Perrine, Josephine and Malin for starting the tradition of little treats after vacations, and newbies Marie and Simon who perpetuated it.

People that came and went, but left an impression hard to erase (and I would want to!) - in particular Bengt J, Lilla-Cilla, Ulrika, Malin A and Kenny.

Our surrounding labs have their share of lovely, LOVELY individuals who all made labparties, meetings and grey day lunches at patienthotellet memories for life. I will miss this! Robban (som är så himla snäll!), Pär (festfixaren och superregissören, I owe you!), Johan H och Malin W (ja ni vet...tycker om er!), Joakim, Therese, Linda, Karin and Anders T.

So many other people have contributed to my PhD by smiling in the corridor, keeping me company at lunch, chatting with me on the bus and helping out scientifically:

Andreas A, Andreas S, Ayse, Biljana, Barbara, Cilla L, Daniel, Deniz, Emma, Gesine, Henrik,

Irene, Jan, Katie, Liselijn, Manolo, Merab, Nina Rog, Sara, Stephan, Tomas B, Tomas C, Zaza and many many more (Sorry if I forgot anyone)!

Sometimes things just fall into place. I've had the best of times, inside and outside the lab, with my Dunk-vänner (Det slank ju ner...). ²Nina – the kitchen wiz who's lovelife was more than complicated but always entertaining, ⁷Johan – the funny, cool cynic with the heart in the right place (somewhere deep below...). Thanks for your company and all the fun things we did when it was just us! Buss 171, this is where it all started! Martin (min Martin!), Jenny – the sassiest baddest babe (kram och tack säger kräftan Andersson!), Hanna and Marie (although you were technically never a dunk) –fun, bubbly buddies. You all brought along a whole new social scene that has made my life during the last couple of years so much fun. Your importance to me cannot be measured!!!!

Skrivandet av den här avhandlingen har gett mig mer kunskap, fler kilon och massor av pek att arkivera. Jag har förlorat sömn och min hälsa men tack och lov inte mina vänner. Fantastiska människor som förstått och stått ut med min frånvaro:

⁴Snäckorna, mina pinglor!! Jenny, Lina, ¹Christel, Marie, Sara, Lotta, Karin och Malin! Vi är en institution nu. Tio år and counting.... Jag är så glad att jag har er, att jämföra doktorandlivet med, men mest att snacka, fixa, festa, äta och mysa ihop med.

Phoenix och Roxanne – divided but still divine....

Mina gamla gamla vänner, ³Lotten, Bodil och Frida, som håller i vått och torrt....

Mina nya nya vänner i bokklubben, snart är jag med igen!

⁸Linda, min nya bästis. Bästa bonusen på doktorandtiden. Snart kan jag vara med och göra roliga saker igen (och försöka låta bli att somna under tiden...)

Sist men mest: Tack till min familj som alltid tror på mig och hejar på mig. Jag vet att jag har varit lyckligt lottad som haft er så nära och för att ni är som ni är.

Största kramar till ⁵mamma och pappa som ställt upp så mycket under alla år. Vad hade jag gjort utan er? Tack för att ni stöttat och hjälpt, hämtat och lämnat, gett mig middag, matlådor och sovplats när hemma och labbet varit för långt ifrån varandra. Till våren blir det promovering tillsammans, pappa!

⁶Jenny och Bugge, för att jag är välkommen till er närhelst jag behöver en paus från vardagen. Hos er är det alltid semester! Tack Sis, för att du frågar vad jag egentligen gör (även om du blir för äcklad för att höra klart....;o) Lill-Sillen, som alltid är så gla' och kan liva upp mig! Nästa år får jag vara mer med dig!

Mormor, farmor och farfar för att ni är såna underbara människor, med perspektiv på livet och med visdom man inte kan forska sig till. I mina tankar, morfar som värdesatte kunskap så högt, Jag tror du hade varit stolt över mig och jag önskar att du kunde ha varit här....

⁹Martin! MSP, BC! Tack för att du lyssnar, för alla peppningar och kramar (telefon- och riktiga). Du var här fast du inte var här! Tack för att du är rolig, klarsynt, förstående, snäll (och snygg...), schysst, hjälpsam och underbar. Du är helt klart högsta vinsten....

Det här var det värsta jag varit med om!!!
– Baloo, Djungelboken

Special thanks to New Tango Orquestra, Svenska Akademien, Timbuktu, Mauro and P3 Svea for inspirational music and lyrics.....