

## Preface from your Host

### Dear Colleagues,

The DFG Priority Programme *SPP1356 – Pluripotency and Cellular Reprogramming* ([www.spp1356.de](http://www.spp1356.de)) welcomes you to the International Symposium “Reprogrammed Stem Cells” in Berlin from March 20–22, 2014.

At the end of the funding period of the SPP1356 the symposium is intended to bring together German and international colleagues to discuss the molecular concepts of pluripotency and directed reprogramming. The symposium will represent the current state of the ES cell and reprogramming field and provides a platform for interdisciplinary discussion of new concepts and technologies.

Starting with a public keynote presentation on the opening day we will have two more days of scientific presentation and exchange.

Speaking time is assigned as follows (speaking + discussion time):

Invited talks: 25 + 5 min.  
Contributed talks: 10 + 5 min.

Poster sessions are scheduled during lunch breaks. This symposium is a joint enterprise of the SPP1356, the Berlin-Brandenburg Academy of Sciences and Humanities (BBAW) and the ELSA Research Project (BMBF) „Developmental Totipotency“. The latter covers ethical and legal aspects of totipotency and cellular reprogramming in a concurrent session on Friday, March 21st., 2014.

We wish you an interesting and exiting conference as well as an enjoyable stay in Berlin!


### The Conference Chairs



Albrecht Müller



Hans Schöler



Jörn E. Walter



Martin Zenke

## Congress Venue

### Berlin-Brandenburg Academy of Sciences and Humanities

(BBAW)

Entry: Markgrafenstr. 38 · 10117 Berlin



Bei Anfahr mit dem PKW empfehlen wir die Nutzung der umliegenden Parkhäuser

S-Bahn bis Friedrichstrasse  
U2 bis Hausvogteiplatz oder Stadtmitt  
U6 bis Französische Strasse oder Stadtmitt

### Deadlines

Abstract Submission: January 15th, 2014

Registration: January 15th, 2014

Registration Fee: 100,- € (Academics and Students)

The conference will host an industrial exhibition.

### Organisation and Contact

SPP1356 Office, Kathrin Mantel

ZEMM, Universität Würzburg,

Zinklesweg 10, 97080 Würzburg, Germany

phone (+49) 931-201 45478, fax (+49) 931-201 45148

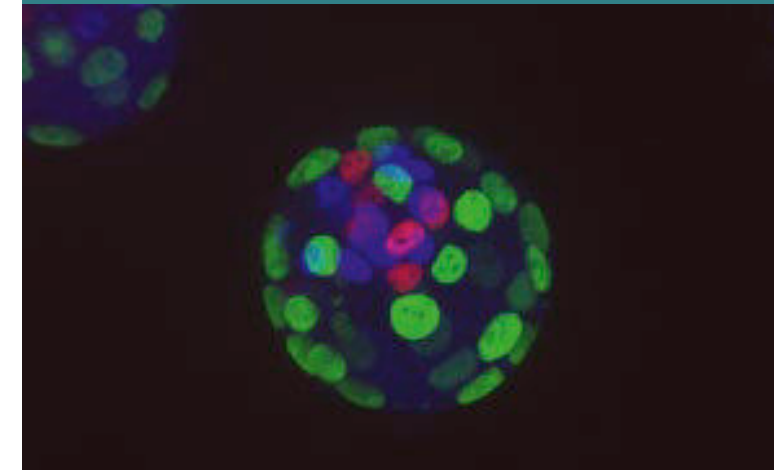
[kathrin.mantel@uni-wuerzburg.de](mailto:kathrin.mantel@uni-wuerzburg.de)

[www.spp1356.de](http://www.spp1356.de)

Gestaltung:  [design@smi.uni-wuerzburg.de](mailto:design@smi.uni-wuerzburg.de)



# International Symposium Reprogrammed Stem Cells at the Berlin-Brandenburg Academy of Sciences and Humanities



March 20–22, 2014  
Berlin



**DFG** Deutsche Forschungsgemeinschaft

in cooperation with the ELSA Research Project (BMBF) “Developmental Totipotency”

Thursday, March 20th, 2014

17:00–17:45 **Registration**

17:45–18:00 **Welcome and Opening**

18:00–19:30 **Public Keynote Lecture**  
**Ernst-Ludwig Winnacker,**  
**Human Frontier Science Program,**  
**Strasbourg, F**

*Title:* Stratifizierte Medizin über Stammzellen und andere moderne Technologien

19:30 **Social Hour** (lite bites and drinks)

Friday, March 21st, 2014

09:00–10:30 **The Biology of ES Cells and Reprogramming I**  
 Amanda G. Fisher, Imperial College, London, UK  
 Tobias Cantz, Hannover Medical School, Germany, ELSA Research Project (BMBF)  
 „Developmental Totipotency“  
 SPP1356-presentations

11:00–12:30 **Concurrent Session I:**  
**ELSA Research Project (BMBF) „Developmental Totipotency: Defining a Normative Criterion in Ethics and Law in the Light of Novel Results in Developmental Biology“**

**Concurrent Session II:**  
**The Biology of ES Cells and Reprogramming II**  
 Ana Pombo, MDC, Berlin, Germany  
 Austin Cooney, Baylor College of Medicine, Houston, TX, USA  
 SPP1356-presentations

12:30–14:00 **Lunch & Posters & Exhibition**

14:00–15:30 **Pluripotency and Chromatin I**  
 Takashi Tada, Kyoto University, JP  
 Niall Dillon, Imperial College London, UK  
 Asifa Akhtar, MPI of Immunobiology and Epigenetics, Freiburg, Germany  
 SPP1356-presentations

16:00–17:30 **Pluripotency *in silico***  
 Thomas Dandekar, University of Würzburg, Germany  
 SPP1356-presentations

17:30–18:30 **Totipotency and Bioethics**  
**Joined Session of ELSA Research Project (BMBF) and SPP 1356**

20:00 **Evening Event**  
 (Location: Saarländische Landesvertretung)

Saturday, March 22nd, 2014

09:00–10:30 **Pluripotency and Chromatin II**  
 Wendy Bickmore, MRC, Human Genetics Unit, University of Edinburgh, UK  
 Ian Chambers, University of Edinburgh, UK  
 SPP1356-presentations

11:00–12:30 **Molecular Regulators of ES Cells and Development I**  
 Rainer Renkawitz, University of Giessen, Germany  
 SPP1356-presentations

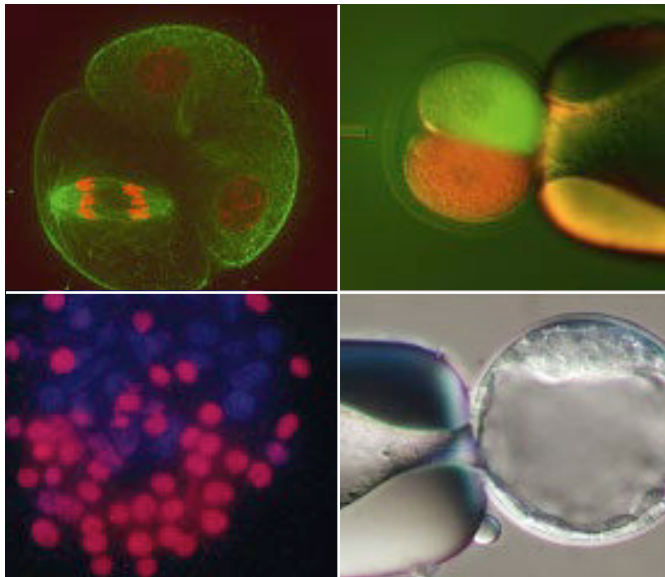
12:30–13:30 **Lunch & Posters & Exhibition**

13:30–15:00 **Molecular Regulators of ES Cells and Development II**  
 SPP1356-presentations

**Keynote Lecture**  
**Rudolf Jaenisch, Whitehead Institute, Cambridge, MA, USA**

15:30–17:00 **Chromatin and Nuclear Organisation**  
 Danny Reinberg, Howard Hughes Medical Institute, New York, USA  
 Bas van Steensel, NCI, Amsterdam, NL  
 SPP1356-presentations

17:00–17:30 **Concluding Remarks and End of Meeting**



Source: Michele Boiani lab, Münster

## Liebe Kolleginnen und Kollegen

Der deutsche Gesetzgeber bestimmt den menschlichen Embryo sowohl im Embryonenschutzgesetz (1990) als auch im Stammzellgesetz (2002, 2008) mittels des biologisch funktionalen Begriffs der Totipotenz. Geht man hypothetisch davon aus, dass menschliche Körperzellen während einer Reprogrammierung zu iPS-Zellen eine transiente totipotente Phase durchlaufen, stellt sich die Frage, ob auch diesen Zellen der normative Status eines Embryos zukommt. Kann der Rekurs auf die entwicklungsbiologische Totipotenz angesichts neuer Techniken und Erkenntnisse noch eine tragfähige Basis für die normative Statusbestimmung des menschlichen Embryos liefern? Dieser Frage widmet sich unser interdisziplinäres BMBF-gefördertes ELSA-Forschungsverbundprojekt „Entwicklungsbiologische Totipotenz. Bestimmung als normatives Kriterium in Ethik und Recht unter Berücksichtigung neuer entwicklungsbiologischer Erkenntnisse“ und lädt zur Präsentation und Diskussion der Projektergebnisse ein. Das Symposium ist eine gemeinsame Veranstaltung unseres Forschungsprojekts und des DFG-Schwerpunktprogramms SPP 1356 „Pluripotency and Cellular Reprogramming“ in Kooperation mit der Interdisziplinären Arbeitsgruppe Gentechnologiebericht der Berlin-Brandenburgischen Akademie der Wissenschaften (BBAW). Wir freuen uns auf Ihr Kommen und auf anregende Diskussionen.

*Thomas Heinemann Hans-Georg Dederer Tobias Cantz*

T. Heinemann H.-G. Dederer T. Cantz

### Teilprojekt Philosophie, Koordination

Prof. Dr. med. Dr. phil. Thomas Heinemann, Lehrstuhl Ethik, Theorie und Geschichte der Medizin, Philosophisch-Theologische Hochschule Vallendar

Dr. Heike Baranzke, Barbara Advena-Regnery, Kathrin Rottländer



### Teilprojekt Rechtswissenschaft

Prof. Dr. iur. Hans-Georg Dederer, Lehrstuhl für Staats- und Verwaltungsrecht, Universität Passau  
Lena Laimböck



### Teilprojekt Entwicklungsbiologie

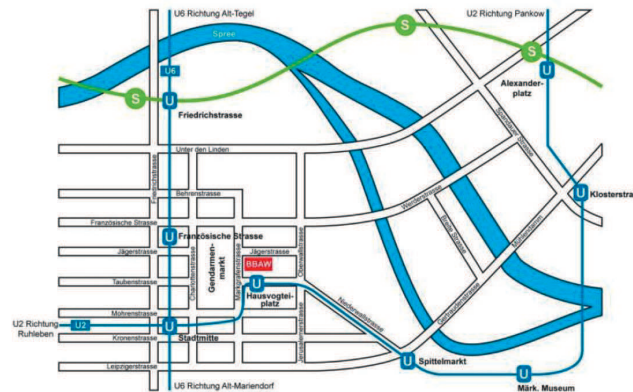
PD Dr. med. Tobias Cantz, Medizinische Hochschule Hannover  
Susan Sgodda, Abbas Peh-Pajoo



## Veranstaltungsort

Berlin-Brandenburgische Akademie  
der Wissenschaften (BBAW)  
Jägerstr. 22/23, 10117 Berlin

Eingang: Markgrafenstr. 38



## Kontakt und Anmeldung

Philosophisch-Theologische Hochschule Vallendar (PTHV)  
Lehrstuhl Ethik, Theorie und Geschichte der Medizin  
Pallottistr. 3, 56179 Vallendar

Tel. +49 (0) 261 6402 223

Fax +49 (0) 261 6402 300

lehrstuhl-medizinethik@pthv.de

oder Anmeldeformular unter [www.pthv.de](http://www.pthv.de)

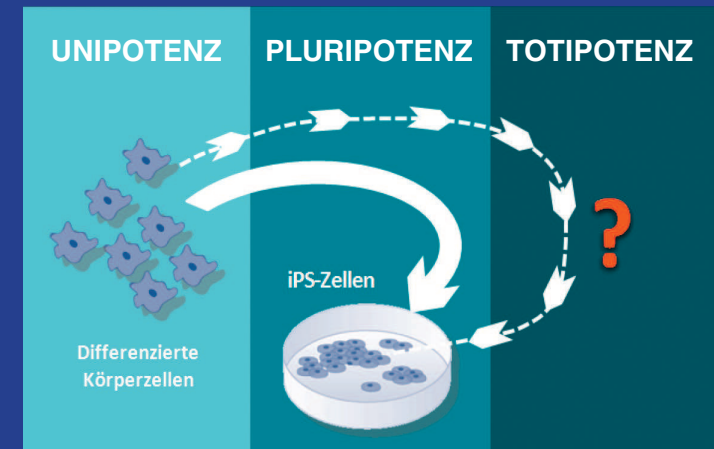
### Anmeldung

ab sofort bis zum 21. Februar 2014

ELSA-Forschungsverbundprojekt „Entwicklungsbiologische Totipotenz: Bestimmung als normatives Kriterium in Ethik und Recht unter Berücksichtigung neuer entwicklungsbiologischer Erkenntnisse“

## Es werde (k)ein Embryo

Zur normativen Bewertung von totipotenten menschlichen Zellen



20. / 21. März 2014  
Berlin



In Kooperation mit dem DFG-Schwerpunktprogramm SPP 1356 „Pluripotency and Cellular Reprogramming“ ([www.spp1356.de](http://www.spp1356.de)) und der Interdisziplinären Arbeitsgruppe Gentechnologiebericht der Berlin-Brandenburgischen Akademie der Wissenschaften ([www.gentechnologiebericht.de](http://www.gentechnologiebericht.de))



# Programm

## Donnerstag, 20. März 2014

- 17:00–17:45 **Registrierung**  
Leibniz-Saal
- 17:45–18:00 **Eröffnung**
- 18:00–19:30 **Öffentliche Keynote Lecture**  
**Ernst-Ludwig Winnacker**  
**Human Frontier Science Program**  
**Straßburg, F**
- Stratifizierte Medizin –  
Über Stammzellen und andere  
moderne Technologien*
- 19:30 **Abendempfang**



©Tobias Cantz

## Freitag, 21. März 2014

- 9:00–10:30 **The Biology of ES Cells and Reprogramming I**  
Amanda G. Fisher  
Imperial College, London, UK  
*Vortragstitel wird angekündigt*
- Tobias Cantz  
Medizinische Hochschule Hannover  
*Reprogramming of Somatic Cells:  
Alternative Approaches Exploiting  
RNA-Interference*  
(ELSA-Forschungsverbundprojekt)
- 10:30–11:00 Kaffeepause
- 11:00–17:30 **Concurrent Session II**  
**The Biology of ES Cells and Reprogramming II**  
SPP 1356 Pluripotency and Cellular Reprogramming (*s.gesondertes Programm*)
- Leibniz-Saal
- Einstein-Saal
- Concurrent Session I**
- Es werde (k)ein Embryo. Zur normativen Bewertung von totipotenten menschlichen Zellen**  
ELSA-Forschungsverbundprojekt  
„Entwicklungsbiologische Totipotenz: Bestimmung als normatives Kriterium in Ethik und Recht unter Berücksichtigung neuer entwicklungsbiologischer Erkenntnisse“
- 11:00–11:20 **Begrüßung und Einführung**
- 11:20–12:30 **Teilprojekt Entwicklungsbiologie**  
(Moderation: Hans-Georg Dederer)
- Susan Sgodda  
Medizinische Hochschule Hannover  
*Der Totipotenzbegriff im Zuge neuer biotechnologischer Verfahren*
- Kommentar: Christopher Baum  
Medizinische Hochschule Hannover
- Diskussion

- 12:30–13:30 Mittagspause
- 13:30–15:15 **Teilprojekt Philosophie**  
(Moderation: Tobias Cantz)
- Barbara Advena-Regnery  
Philosophisch-Theologische Hochschule Vallendar  
*Natürlicher Embryo – geeignetes Wertprädikat für die Bioethik?*
- Heike Baranzke, Philosophisch-Theologische Hochschule Vallendar  
*Der menschliche Embryo zwischen Naturzweck und Handlungszweck*
- Kommentar: Geert Keil  
Humboldt-Universität, Berlin
- Diskussion
- 15:15–15:45 Kaffeepause
- 15:45–16:50 **Teilprojekt Rechtswissenschaft**  
(Moderation: Thomas Heinemann)
- Lena Laimböck, Universität Passau  
*Von „Totipotenz“ zu „qualifizierter Entwicklungsfähigkeit“*
- Kommentar: Jens Kersten  
Ludwig-Maximilians-Universität, München
- Diskussion
- 16:50–17:20 **Abschlussdiskussion**  
mit Projektleitern, Projektbearbeiterinnen und Kommentatoren  
(Moderation: Thomas Heinemann)
- 17:30–18:30 **Joint Session of ELSA Research Project (BMBF) and SPP 1356 (DFG)**  
Leibniz-Saal
- Interactive Session: Clinical Transplantation of Reprogrammed Cells and the Need of Normative Frameworks
- 20:00 **Abendveranstaltung in der Saarländischen Landesvertretung**

## Joint Session

ELSA Research Project (BMBF) and SPP 1356 (DFG)

### Clinical Transplantation of Reprogrammed Cells and the Need of Normative Frameworks

#### Session I

##### iPS Cell Research Meets Systems Medicine

**Martin Zenke**

*Technical Note on cDNA Array, mRNA-Sequencing, Array-CGH, Full Genome Sequencing, Epigenetic Profiling*

**Hans-Georg Dederer**

*Availability of Patient-Specific Data to Governmental Institutions*

#### Session II

##### Accidental Findings and Biodata Privacy

**Tobias Cantz**

*Examples of Accidental Findings in iPS-Cell Research*

**Thomas Heinemann**

*Ethical Implications of Accidental Findings in iPS-Cell Research*

#### Session III

##### Transparent Human Beings

**Jörn Walter**

*Big Data Mining: Genetic and Epigenetic High-Throughput Screens*

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## TED-Questions

### Session I

- What is the most important release criterion for the certification of iPS cell lines used in disease modeling / drug screening?
  1. no adverse findings in transcriptome profiling (gene arrays)
  2. no adverse findings in transcriptome profiling (mRNA-sequencing)
  3. no subchromosomal alterations (array-CGH)
  4. no major mutations in full genome sequencing
  5. no major alterations in epigenetic profiling (genome-wide DNA-methylation)
- What is the most important release criterion for the certification of iPS cell derivatives used in cell therapies?
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  5. no major alterations in epigenetic profiling (genome-wide DNA-methylation)

### Session II

- What kind of data would provide accidental findings most problematic for patients?
  1. transcriptome profiling with gene arrays
  2. transcriptome profiling with mRNA-sequencing
  3. analysis of subchromosomal alterations (array-CGH)
  4. genetic analyses using full genome sequencing
  5. epigenetic profiling analyzing genome-wide DNA-methylation

### Session III

- Will we ever be able to predict issues of longevity, personality, sexual orientation in stem cells?
  1. Yes, science will bring us there; no concerns.
  2. No, that is fiction.
  3. This is frightening. It may happen, but we would need a broad consent.

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