

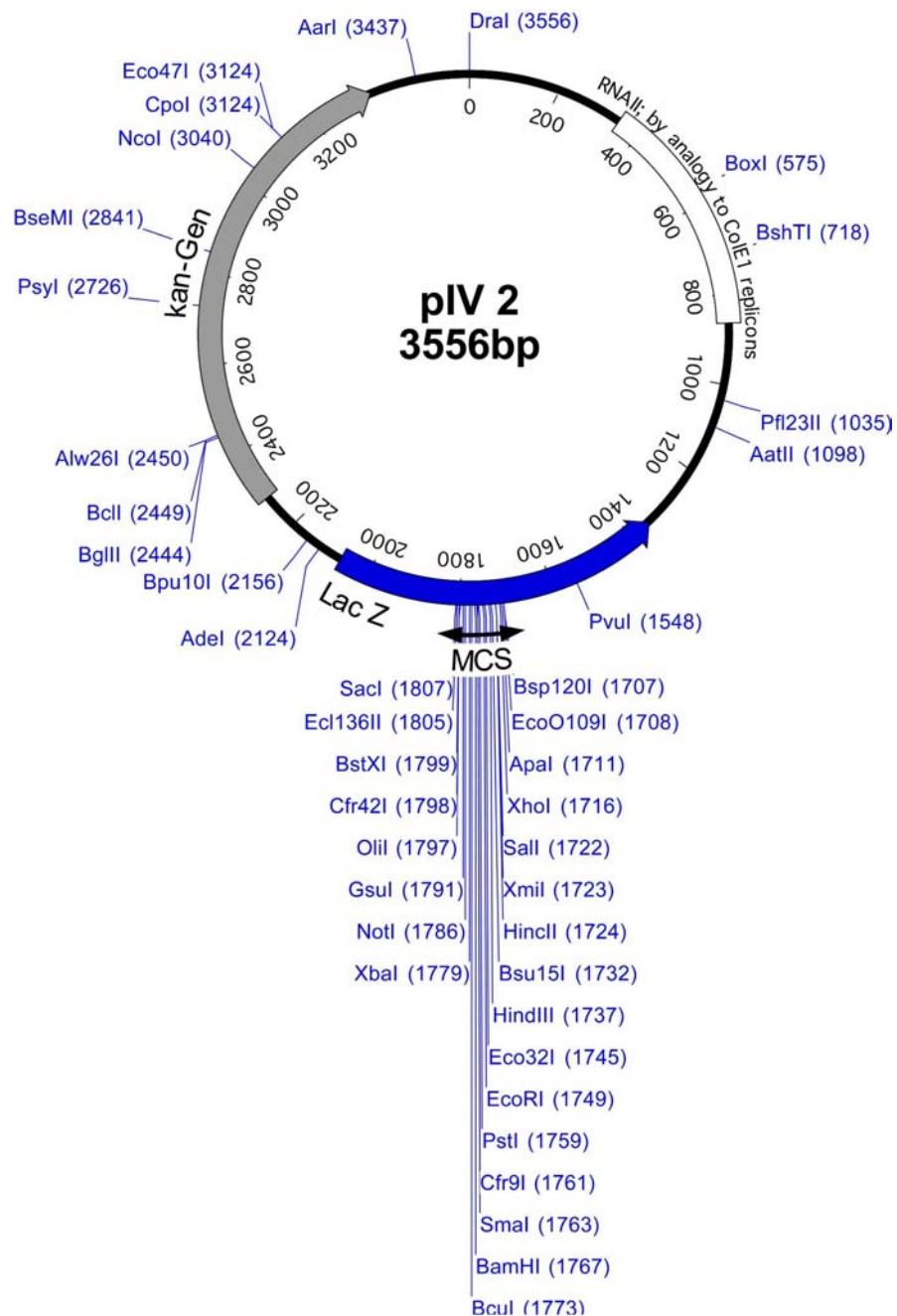
## 8 Appendix

### 8.1 Abbreviations

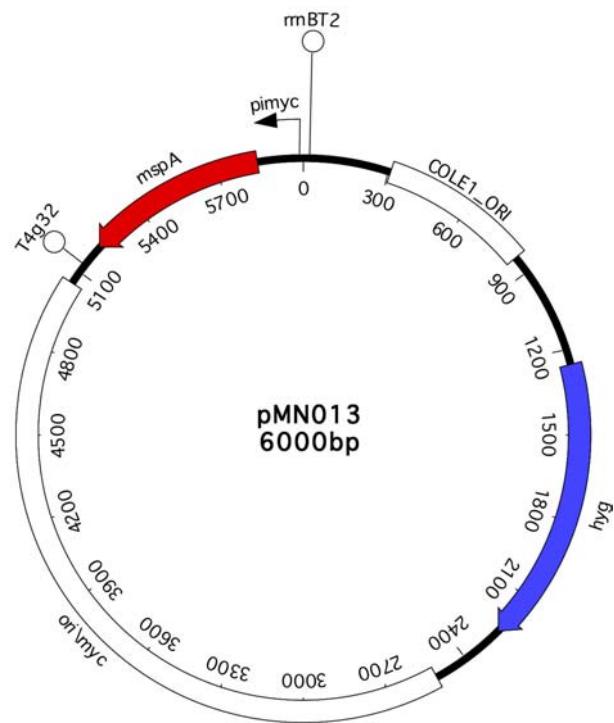
**Table 7:** List of abbreviations.

AG	Arabinogalactan	MOI	Multiplicity of infection
<i>aph</i>	Gene for aminotransferase	<i>mspA</i>	<i>Mycobacterium smegmatis</i> porin A
ATP	Adenosintriphosphate	MW	Molecular weight
att	Attachment site	NCBI	National Center for Biotechnology Information
BCG	Bacille Calmette-Guerin	NL	Non linear
bp	Basepare	nOPOE	n-octylpolyoxyethylene
CFU	Colony forming unit	nS	Nanosiemens
CTAB	Hexa-decyltrimethylammonium bromide	OD	Optical density
dH <sub>2</sub> O	Distilled H <sub>2</sub> O	OM	Outer embrane
DNA	Desoxiribonucleic acid	<i>ompATb</i>	Outer membrane protein A <i>Tuberculosis</i>
dNTP	Desoxiribonucleoside triphosphate	ORF	Open reading frame
ELISA	Enzyme linked immunosorbant assay	PBS	Phosphate buffered saline
EMBL	European Molecular Biology Laboratory	PCR	Polymerase chain reaction
FAM	6-Carboxyl-Flouresein	PG	Peptidoglycan
g	Acceleration of gravity	<i>porM1</i>	Porin <i>Mycobacterium</i> 1
h	Hour	RGM	Rapidly growing mycobacteria
<i>hsp60</i>	Heat shock protein 60	RLU	Relative light unit
HYG	Hygromycin	rMspA	Recombinant MspA
Ig	Immunoglobulin	RNA	Ribonucleic acid
IM	Inner membrane	ROX	6-Carboxy-X-rhodamine
<i>int</i>	Gene for integrase	rpm	Rounds per minute
Ip	Isoelectric point	rRNA	Ribosomal RNA
kDa	Kilodalton	RT	Reverse transcriptase
KM	Kanamycin	s	Second
kV	Kilovolt	SD	Standard deviation
LAM	Lipoarabinomannan	SDS	Sodium dodecyl sulfate
LB	Luria Bertani medium	TAMRA	Tetramethylrhodamine
MA	Mycolic acid	TEM	Transmission electron microscopy
Mb	Mega basepare	U	Unit
min	Minute	WHO	World Health Organization

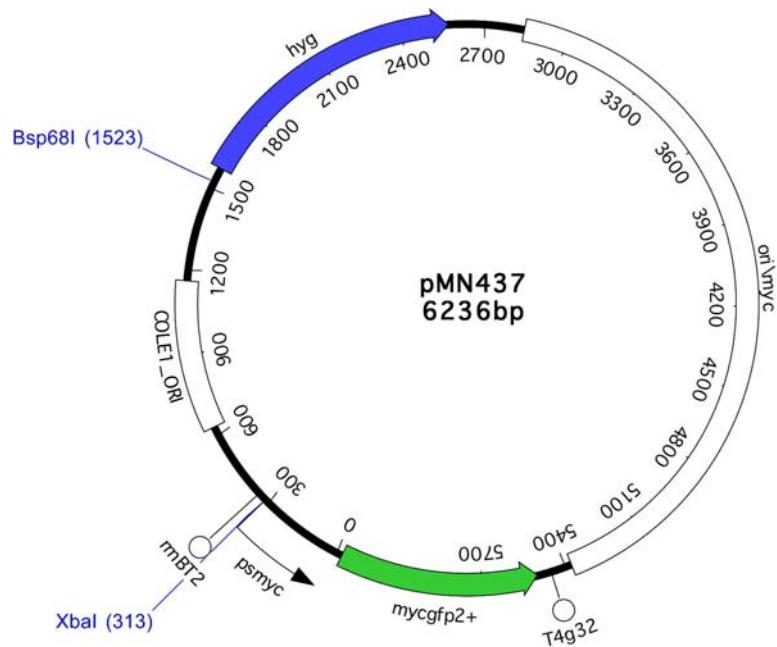
## 8.2 Maps of plasmids



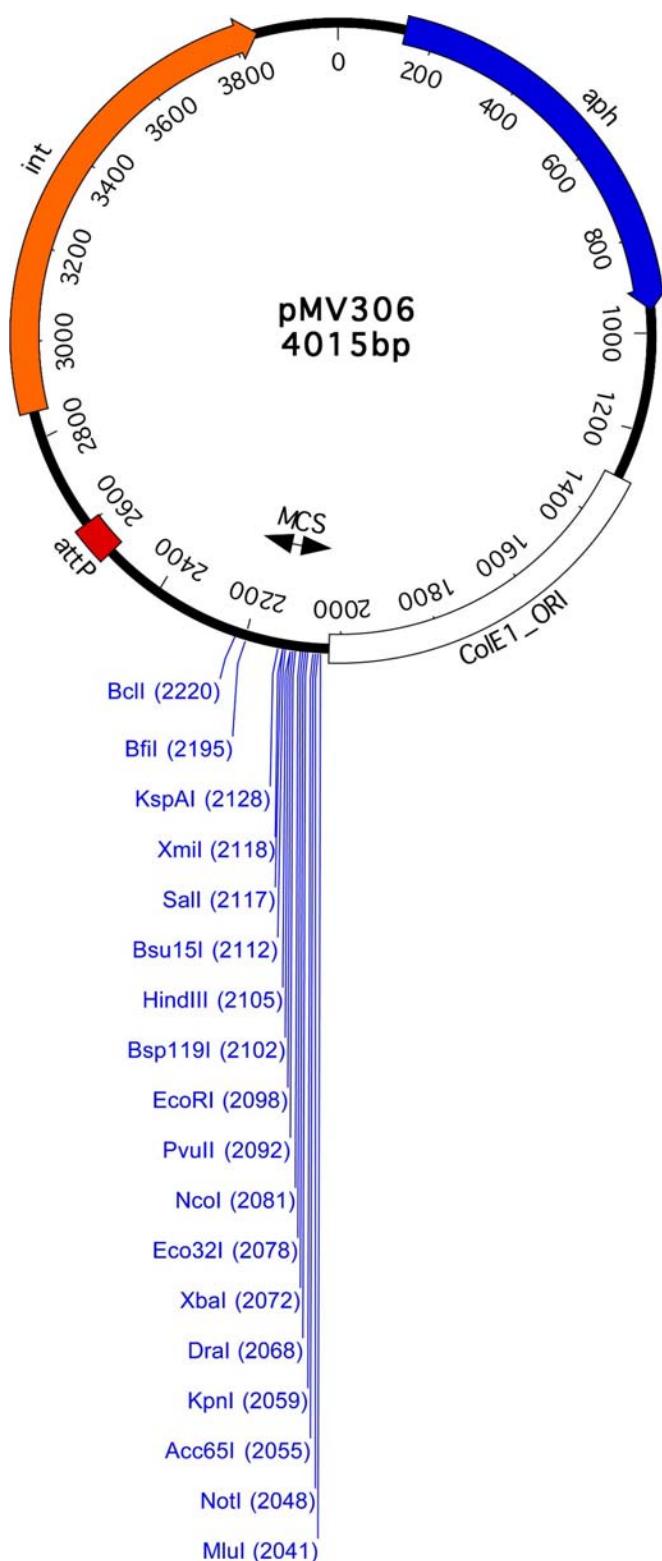
**Figure 21:** Cloning plasmid pIV2 (Strauch et al., 2000).



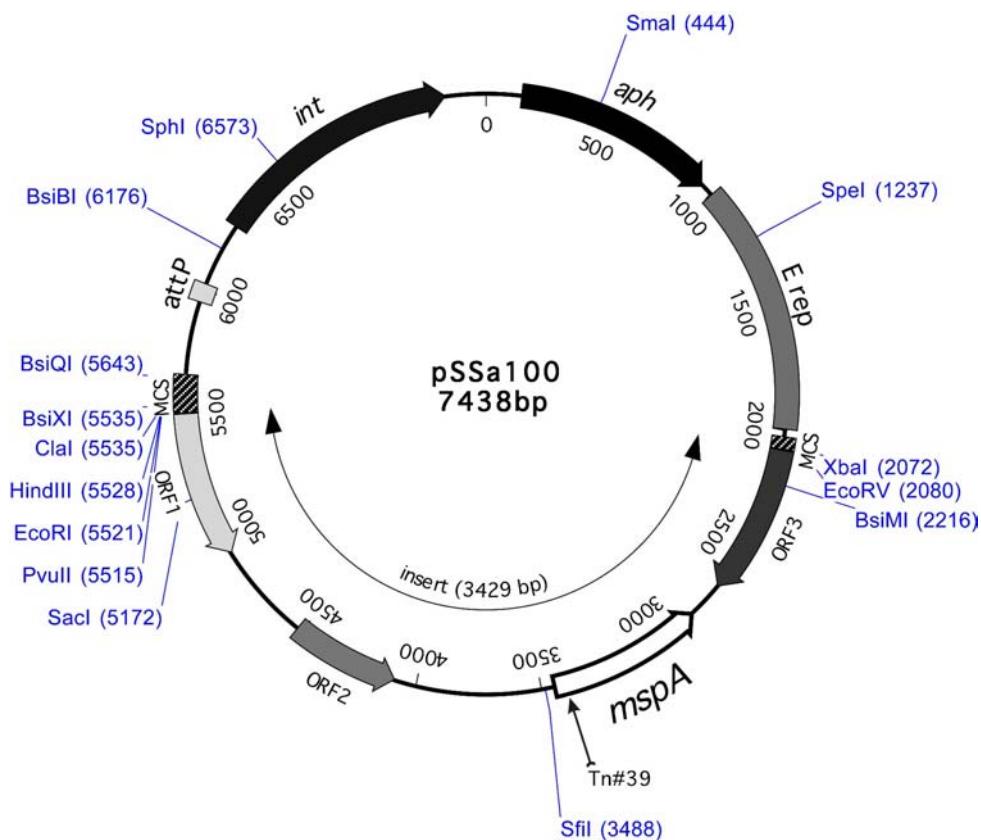
**Figure 22:** Plasmid pMN013 (Mailaender et al., 2004).



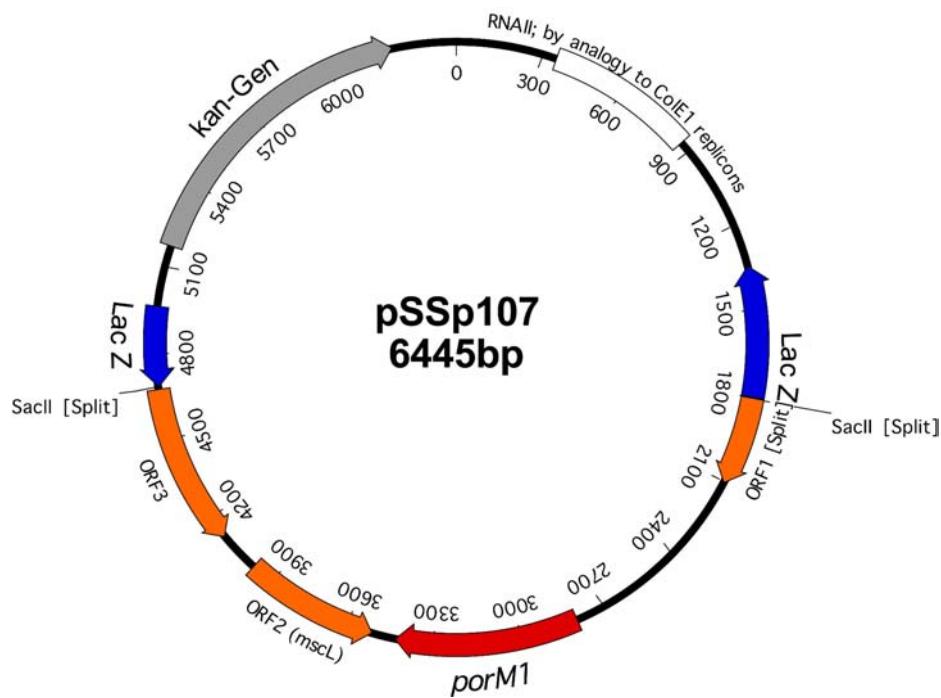
**Figure 23:** Plasmid pMN437 (Kaps et al., 2001).



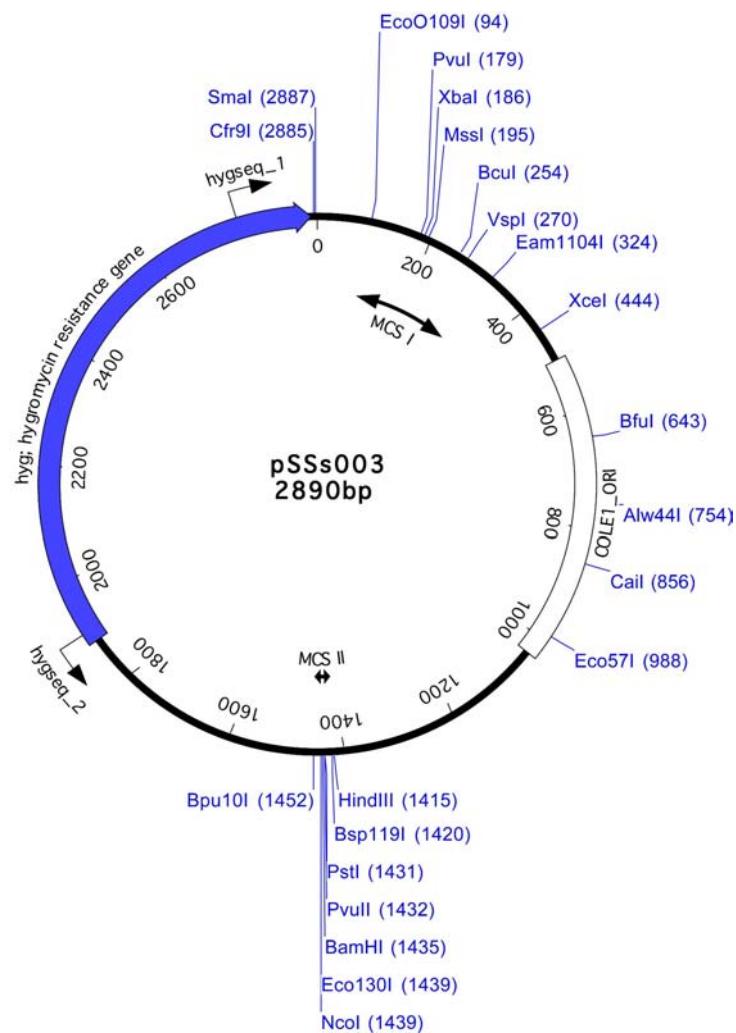
**Figure 24:** Cloning plasmid pMV306 (Stover et al., 1991).



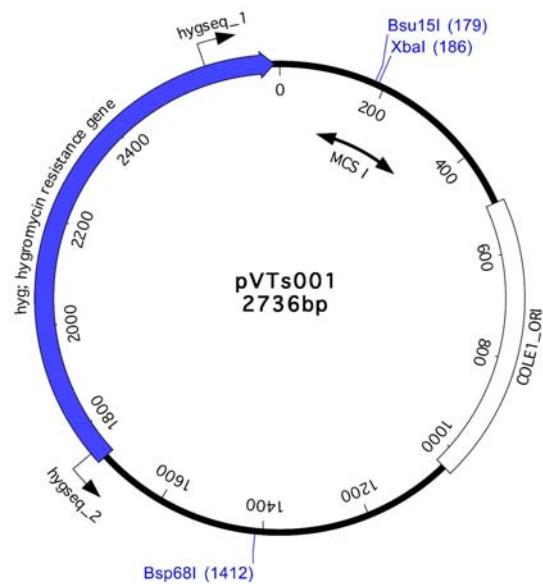
**Figure 25:** Plasmid pSSa100, prepared in this work (Sharbati-Tehrani et al., 2004).



**Figure 26:** Plasmid pSSp107 (identical to pSSp108), prepared in this work.



**Figure 27:** Mycobacterial suicide plasmid pSSs003, constructed in this work.



**Figure 28:** Precursor plasmid pVTs001, constructed in this work.

### 8.3 Nucleotide sequences

*PorM1* sequence from *M. fortuitum* DSM 46621:

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ATGAAGGCATTCACTCGGGTGTGGTCGCGATAGTTGCAGCTATCGCGCGCTGTTACGAG
CACGGCGTATCTCATGCAGGCTGGACAATGAACGTGAGCCTGGTTGATGGTCAGGATCGGA
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TGAGTTCGAGGGCACCTGGAGCTGGCTACCAGATCGGCTTCCCCTGGTCGCTGGTGTGCG
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*PorM1* sequence from *M. fortuitum* 10851/03:

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*PorM1* sequence from *M. fortuitum* 10860/03:

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*PorM1* sequence from *M. peregrinum* 9912/03:

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*PorM1* sequence from *M. peregrinum* 9926/03:

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16S rRNA sequence from *M. fortuitum* DSM 46621:

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**16S rRNA sequence from *M. fortuitum* 10851/03:**

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**16S rRNA sequence from *M. fortuitum* 10860/03:**

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## 8.4 Publications resulted from this study

### 8.4.1 Articles

**Lewin, A., Freytag, B., Meister, B., Sharbati-Tehrani, S., Schafer, H., & Appel, B. (2003).** Use of a quantitative TaqMan-PCR for the fast quantification of mycobacteria in broth culture, eukaryotic cell culture and tissue. *J. Vet. Med. B. Infect. Dis. Vet. Public Health* **50**, 505-509.

**Sharbati-Tehrani, S., Meister, B., Appel, B., & Lewin, A. (2004).** The porin MspA from *Mycobacterium smegmatis* improves growth of *Mycobacterium bovis* BCG. *Int. J. Med. Microbiol.* **294**, 235-245.

**Sharbati-Tehrani, S., Stephan, J., Appel, B., Niederweis, M., and Lewin, A. (2005).** Porins limit the intracellular persistence of *Mycobacterium smegmatis*. (submitted to *Microbiology*).

### 8.4.2 Presentation

**Sharbati-Tehrani, S., Freytag, B., Meister, B., Martinez-Moya, M., Appel, B., and Lewin, A. (2003).** Introduction of a gene from *Mycobacterium smegmatis* encoding a porin into *Mycobacterium bovis* BCG and investigation of its growth physiology. *Infection* 31 (Suppl1): 104. 7. Kongress für Infektionskrankheiten und Tropenmedizin. 27.02.-01.03.2003, Berlin.

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**8.6 Erklärung**

Die dieser Dissertation zugrunde liegenden Arbeiten wurden am Robert Koch-Institut, Berlin, im Zeitraum Mai 2002 bis März 2005 durchgeführt.

Hiermit erkläre ich, dass ich die vorliegende Dissertation selbständig verfasst und keine anderen als die angegebenen Hilfsmittel verwendet habe.

Berlin, 22. März 2005

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Soroush Sharbati-Tehrani