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Declarant initials:

I, Peter J. Morin, have personal knowledge of the facts stated in this declaration and, if called as a witness, I could and would testify competently thereto under oath. I declare as follows:

- 1. I am a neutrologist certified by the American Board of Psychiatry and Neurology. I practice neurology in Massachusetts and in Maine. I serve as Instructor of Neurology at the Boston University School of Medicine and Director of Outpatient Services for the New England GRECC (Geriatric Research Education and Clinical Center) at the Bedford VAMC. In that capacity, I supervise a 50-bed hospice ward for individuals in the advanced stages of dementia. I perform molecular research in neurology and neuroscience. I am also principal investigator for several neurological clinical research studies and serve as a neurologist for the Framingham Heart Study. I am providing this declaration as a neurologist, and not as a representative of Boston University or of the Department of Veterans Affairs.
- 2. My knowledge of Terri Schiavo's situation is indirect. I have not been provided an opportunity to review the medical record, nor to examine Ms. Schiavo. What knowledge I have about her situation derives from the news media and conversations with individuals who have followed the case carefully.
- 3. Based upon this information, I am concerned about the management of Ms. Schiavo's condition. As a neurologist who cares for the dying, I have a considerable interest in the rational management of such cases, both by attending physicians and by the court.
- 4. My first concern regards the quality of the information used by the court to establish Ms. Schiavo's intentions in the event of serious neurological injury. I have never met a 25-year-old outside of medical school who seriously considered such unusual possibilities regarding her mortality. In my experience, where an individual's wishes are uncertain (that is, where they have not previously been expressed in writing), the medical community defers to the more conservative course of action. This is especially true when there are family members who are

- 5. Secondly, I am concerned that the medical information under consideration is not contemporaneous with the decision at hand. It is my understanding that Ms. Schiavo has not had a medical, let alone a neurological, evaluation since 2002 or before. Since her neurological status is disputed, a decision should not be made based on outdated information. Given the uncertainty of her neurological status, current technology should be brought to bear on her current situation. It is my understanding that Ms. Schiavo had a neuroimaging test some three or more years ago and that her last electroencephalogram was more than ten years ago. Unless these tests were conclusive (i.e., showed indisputable, severe or extensive brain injury), they can hardly be considered relevant to her current neurological status. A quality and unbiased neurological evaluation and appropriate diagnostic tests need to be obtained at the time of decision-making, even when the decision-making process has been prolonged. Failure to do so in a high-profile case such as this can only lower the threshold for inappropriately withholding lifesaving medical support for other neurologically compromised individuals.
- 6. I have discussed this matter with other neurologists. While all neurologists understand that some cases of come and PVS are logically managed by not continuing life support, there is a general concern regarding management of highly uncertain cases. Even if it can be established that Ms. Schiavo's chances for a meaningful neurological recovery are very low, it cannot be established that her chances are zero. If there is no written indication that she would not wish to remain alive under these circumstances, and if there are family members willing to assume the burden of her care, then good neurological practice would give those family members the opportunity to provide that care. Established ethics societies and committees, some independent

Occlarant initials:

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and some associated with national neurological organizations, should consider this matter and its implications prior to a final decision being rendered.

I declare under the penalty of perjury under the laws of the Commonwealth of Massachusetts that the foregoing is true and correct.

Executed this 8th day of March, 2005, in Needham, Massachusetts

Peter J. Morin, M.D., Ph.D., Declarant

BIOGRAPHICAL SKETCH

Provide the following information for the key personnel in the order listed for Form Page 2 Follow the sample format for each person. DO NOT EXCEED FOUR PAGES.

NAME	POSITION TITU	*** ***	**************************************	
Peter John Morin	Neurologist			
EDUCATION/TRAINING (Bogin with baccaloureate or other initial profe	ssional education, s	such as neasing, and ine	clude postdoctoral training.)	
INSTITUTION AND LOCATION	DEGREE (# applicable)	YEAR(s)	FIELD OF STUDY	
Colby College, Waterville, Maine	B.A.	1984	Physics	
Boston University School of Medicine	Ph.D.	1992	Cellular Neurobiology	
Boston University School of Medicine	M.D.	1992	Internal Medicine	

RESEARCH AND PROFESSIONAL EXPERIENCE: Concluding with present position, list, in chronological order, previous employment, experience, and honors, include present membership on any Federal Gavamment public advisory committee. List, in chronological order, the titles, all authors, and complete references to all publications during the past three years and to representative earlier publications pertinent to this application. If the list of publications in the last three years exceeds two pages, select the most pertinent publications.

Postdoctoral Training:

7/92-6/93 Internship in Medicine St. Elizabeth's Hospital of Boston, Brighton, MA

7/93-6/96 Longwood Neurology Training Program, Boston Residency Program in Neurology leading to

Board Eligibility House Staff at Brigham and Women's Hospital, Both Israel Hospital, Boston Children's

Hospital, Dana Farber Cancer Institute, New England Deaconess Hospital, and Wost Roxbury and

Brockton Veteran's Administration Hospitals

7/96-7/99 Research and Clinical Fellow, Brigham and Women's Hospital

Molecular Biology of Alzheimer's and Neurodegenerative Disease Center for Neurological Disease at

Brigham and Women's Hospital Behavioral Neurology, Brigham and Women's Hospital, Boston

Recipient of NINDS MSCDA (K08) Award

ENRM VA Hospital, Bedford, MA

Academic Appointments:

1993-1999	Clinical Fellow in Neurology, Harvard Medical School, Boston
1996-1998	Research Fellow in Neurology, Harvard Medical School, Boston
1998-1999	Instructor, Department of Neurology, Harvard Medical School
2002-pres	Department of Neurology, Boston University School of Medicine

Hospital Appointments:

1993-1997	Clinical Fellow in Medicine (Neurology), Brigham and Women's Hospital, Boston
1998-1999	Associate Neurologist, Brigham and Women's Hospital, Boston
1999-pres	Neurologist, St. Joseph Hospital, Bangor, ME
1999-pres	Neurologist, Eastern Maine Medical Center, Bangor, ME
1999-pres	Neurologist, Houlton Regional Hospital, Houlton, ME
2002-pres	Neurologist, Acadia Hospital, Bangor, ME
2002	Neurologist, Boston Medical Center (pending)
2003-pres	Neurologist, Director of Outnationt Services New England GRECC

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Biographical Sketch Format Page

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igator/Program Director (Last, first, middle):

Private Practice:

7/99-6/00 Associate Neurologist, Neurology Associates of Eastern Maine, P.A. Bangor, ME

7/00-11/01 Partner Neurologist, Director, Alzheimer's and Memory Disorders Clinic

Neurology Associates of Eastern Maine, P.A. Bangor, ME.

Professional Societies:

1986	Massachusetts Medical Society
1988	American Medical Society
1995	American Academy of Neurology
1998	American Academy of Science
1998	American Society for Cell Biology
1998	Society for Aging
1999	Maine Medical Society
1999	Penobscot Medical Society
1999	Maine Neurological Association

Awards and Honors:

1992 Alpha Omega Alpha

1992 Ciba-Geigy Prize in Neurological Sciences

Clinical Research Activity

- 1. Treatment of Agitation and Psychosis in Dementia/Parkinsonism (TAP/DAP)
 Principle Investigator
- 2. Pluse II Trial of NS2330 in Alzheimer's Disease Principle Investigator
- 3. Relationship of Pain Threshold and Cognitive Function in AD Principle Investigator
- 4. Cholesterol Lowering Agents to Slow progression of AD (CLASP-AD)
 Principle Investigator
- Effect of Memortine and Food intake in AD Principle Investigator
- 6. Leaprolide for treatment of AD
- Principle Investigator

 7. BUSM Biorepository for Brain Research
- Principle Investigator

 Boston University ADC
- Boston University ADC Staff Investigator

Publications:

- Bielinski DF, Morin PJ, Dickey, BF, Fine RE. Low molecular weight GTP-hinding proteins are associated with neuronal organelles involved in rapid axonal transport. J Biol Chem. 1989; 264: 18363-18367.
- Fishman JM, Cahill M, Morin PJ, McCrory M, Bucher NLR, Ullman D. Specific gangliosides increase rapidly in rat fiver following partial hepateotomy. Biochem Biophys Res Commun. 1991; 174; 638-646.
- Morin PJ. Lui N, Johnson RJ, Leeman SE, Fine RE. Isolation and characterization of rapid transport vesicle subtypes from the rabbit optic nerve. J Neurochem. 1991; 56; 415-427.
- Morin PJ, Johnson RJ, Shachar I, Fine RE, Leeman SE. Characteristics of tachykinin transport vesicles in the optic norve. Ann NY Acad Sci. 1991; 632; 442-443.
- Morin P.I., Johnson R.J., Fine RE. Kinesin is rapidly transported in the optic nerve as π membrane associated protein. Biochem Biophys Acta. 1993; 1146(2): 275-281.
- Morin PJ, Abraham CR, Amaratunga A, Johnson RJ, Huber G, Sandell JH, Fine RE. Amyloid precursor protein is synthesized by retinal gauglion cells, rapidly transported to axolemma, and metabolized. J Neurochem. 1993; 61(2): 464-473.

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Amaratunga A, Morin to, Kosik KS, Fine RE. Inhibition of kinesin synthesis and rapid anterograde axonal 7. transport in vivo by an antisense oligonucleotide. J Biol Chem. 1993; 268(23); 17427-17430.

Morin PJ, Medina M, Brown AMC, Kosik KS. Wnt-1 expression alters APP metabolism. Neurobiology of 8.

Aging, 1998a; 19(4S); \$38

Morin, PJ, Medina, M, Seminov, M, Brown, AMC, and Kosik, K. Wnt-1 induces splicing of exon 15 and 9 expression of L-APP. Neurobiology of Disease. 2004; 16; 59-67.

Dissertation:

Morin PJ. Rapid Transport Vesicles of the Rabbit Optic Nerve. Boston University, 1992.

Patents:

Kosik K, Morin PJ. Methods and compounds for treating Alzheimer's disease. Docket No. 05311/02001.

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