The 33rd Annual Carrell-Krusen

NEUROMUSCULAR SYMPOSIUM

February 17-18, 2011

A SEMINAR FOR MUSCULAR DYSTROPHY CLINIC DIRECTORS, CASE MANAGERS, NURSES
AND MEMBERS OF INTERDISCIPLINARY CARE TEAMS

THE 33RD ANNUAL CARRELL-KRUSEN
NEUROMUSCULAR SYMPOSIUM Awardee:

ROBERT T. LEshner, MD

Sponsored by:
The Departments of Pediatrics and Neurology at UT Southwestern
and the Office of Continuing Medical Education
COURSE DIRECTOR
Susan T. Iannaccone, M.D., F.A.A.N.
Professor of Neurology and Pediatrics
Jimmy Elizabeth Westcott Distinguished Chair in Pediatric Neurology
The University of Texas Southwestern Medical Center
Director of Child Neurology
Children's Medical Center
Dallas, Texas

COURSE CO-DIRECTOR
Gil I. Wolfe, M.D., F.A.A.N.
Professor of Neurology
Dr. Bob and Jean Smith Foundation Distinguished Chair in Neuromuscular Disease Research
The University of Texas Southwestern Medical Center
Dallas, Texas

We gratefully acknowledge the following for their financial support of this program:

CHILDREN’S MEDICAL CENTER
HILL-ROM, INC.
THE MUSCULAR DYSTROPHY ASSOCIATION
TALECRIS BIOTHERAPEUTICS

WHO SHOULD ATTEND
The Carrell-Krusen Neuromuscular symposium is designed for muscular dystrophy clinic directors, case managers, nurses and members of interdisciplinary care teams.

OBJECTIVES
At the end of the symposium, participants should be able to:
• Formulate protocols for treatment and management of both common and unusual neuromuscular disorders.
• Identify characteristic findings on muscle biopsies and EMG studies on neuromuscular disorders.
• Describe recent advances in basic and clinical research on neuromuscular disorders.

CONCEPT AND METHOD
The symposium provides an open and energetic interchange to discuss rare and challenging cases, as well as patients with common problems in order to improve their treatment and management. Physical exams will be demonstrated on patients present at the symposium with ample time for discussion of each case. Sessions will include slide presentations of unusual muscle and nerve biopsies and EMG findings presented by several symposium registrants in an informal manner for open discussion. The scientific session will include formal presentations reporting recent research results. The Carrell-Krusen Neuromuscular Lecture will be a formal one-hour presentation by visiting professor, Robert T. Leshner, MD.

ACCREDITATION
This program has been planned and implemented in accordance with the Essential Areas and Policies of the Accreditation Council for Continuing Education through the joint sponsorship of the University of Texas Southwestern Medical Center, Children's Medical Center and Texas Scottish Rite Hospital for Children. The University of Texas Southwestern Medical Center is accredited by the ACCME to provide continuing medical education for physicians.

CREDIT DESIGNATION-PHYSICIANS
The University of Texas Southwestern Medical Center at Dallas designates this live activity for a maximum of 12.75 AMA PRA Category 1 Credit(s)™. Physicians should claim only those credits commensurate with the extent of their participation in the activity.

CREDIT DESIGNATION
UT Southwestern University Hospitals is an approved provider of continuing nursing education by the Texas Nurses Association, an accredited approver by the American Nurses Credentialing Center’s Commission on Accreditation.

This activity provides 7.5 contact hours Day 1 (2/17/11).
This activity provides 5.33 contact hours Day 2 (2/18/11).

Respiratory Therapists: An application has been submitted to the American Association for Respiratory Care for approval.
CONFLICT OF INTEREST
It is UT Southwestern’s policy that participants in CME activities should be made aware of any affiliation or financial interest that may affect the speaker’s presentation(s). Each speaker has completed and signed a conflict of interest statement. The faculty members’ relationships will be disclosed in the course syllabus.

DISCUSSION OF OFF LABEL USES
Because this course is meant to educate physicians with what is currently in use and what may be available in the future, there may be “off-label” uses discussed in the presentations. Speakers have been requested to inform the audience when off-label use is discussed.

REGISTRATION
The early registration fee of $175 (by February 10, 2011), and late registration fee of $200 (after February 10, 2011), entitles the participants to admission to the conference, all course materials, continental breakfasts, refreshment breaks and Thursday and Friday lunch.

Registration is confirmed upon receipt of registration fee.

LOCATION
Texas Scottish Rite Hospital For Children
2222 Welborn Street
Dallas, Texas 75219

ACCOMMODATIONS
A block of rooms is being held at the The Stoneleigh Hotel & Spa until January 31, 2011. After January 31, 2011, reservations will be accepted on a rooms and rate available basis. Identify the group as UTSMC/Public Education.

The Stoneleigh Hotel & Spa
2927 Maple Ave
Dallas, TX 75201
800.921.8498
214.871.7111
Rate: $139/Night

Please make reservations directly with the hotel. A block of rooms have been reserved for the Carrell-Krusen Neuromuscular Symposium. The daily rate is $139 Single/Double. Valet parking for overnight guests is $20 and is charged to the guest’s room. The special rate cannot be guaranteed past Monday, January 31, 2011.

Registration forms must be received no later than Thursday, February 10, 2011. Space is limited! Walk-in registration may not be available. All symposium attendees (including faculty and presenters) must complete the attached registration form.

CANCELLATION POLICY
The Office of Continuing Education reserves the right to limit registration and cancel courses no less than one week prior to the course, should circumstances make this necessary.

REFUND POLICY
A $50 handling fee will be deducted from cancellation refunds. Refund requests must be received by mail or fax before February 7, 2010. No refunds will be made thereafter.
INFORMATION
For additional information, please call the Office of Continuing Education, 214-648-3138, 1-800-688-8678, fax 214-648-2317, or email cmeregistrations@utsouthwestern.edu

To obtain information on other UT Southwestern programs, go to http://www.utsouthwestern.edu/cme for a free email subscription of the Office of Continuing Education’s monthly calendar, send your name, medical specialty and current email address to: cmecalendar@utsouthwestern.edu

PROGRAM ACCESSIBILITY
We accommodate people with disabilities. Please call 214-648-3138 for more information, or mark the space indicated on the registration form. To ensure accommodation, please register as soon as possible.

The University of Texas Southwestern Medical Center at Dallas is committed to providing programs and activities to all persons regardless of race, color, national origin, religion, sex, age, veteran status, or disability.

UT Southwestern is an equal opportunity institution.

THE 33rd ANNUAL CARRELL-KRUSEN NEUROMUSCULAR SYMPOSIUM AWARDEE:

ROBERT T. LESHER, MD
Professor of Neurology and Pediatrics
Children’s National Medical Center
Washington, DC

PAST LECTURERS
1978 - K. Engel and Valerie Askanas
1979 - Michael Brooke
1980 - Daniel B. Drachman
1981 - Stanley Appel
1982 - Lewis P. Rowland
1983 - Walter G. Bradley
1985 - George Karpati
1986 - Salvatore DiMauro
1987 - Peter Dyck
1988 - Theodore L. Munsat
1989 - Louis M. Kunkel
1990 - P.K. Thomas and Anita Harding
1991 - Victor Dubowitz
1992 - Robert C. Griggs
1993 - Andrew Engel
1994 - Harvey B. Sarnat
1995 - Jerry M. Mendel
1996 - Robert A. Ouvrier
1997 - Richard T. Moxley, III
1998 - Lewis P. Rowland
1999 - Robert G. Miller
2000 - Arthur K. Asbury
2001 - Robert B. Layzer
2002 - Alan Pestronk
2003 - Valerie Askanas, W. King Engel and Francesco Muntoni
2004 - John Sladky
2005 - David Pleasure
2006 - Kenneth Fischbeck
2007 - Austin J. Sumner
2008 - Angela Vincent
2009 - John T. Kissel
2010 - Yadollah Harati
# DAY 1 PROGRAM AGENDA – THURSDAY, FEBRUARY 17, 2011

## TIME
7:00 am  
7:00 am-7:45 am  
7:45 am-8:00 am  
7:00 am registration  
Continental Breakfast  
Introduction & Announcements  

## SESSION / TITLE

<table>
<thead>
<tr>
<th>TIME</th>
<th>ABSTRACT #</th>
<th>TITLE</th>
<th>PRESENTER(S)</th>
<th>LOCATION</th>
</tr>
</thead>
<tbody>
<tr>
<td>7:00 am-7:45 am</td>
<td>L-1</td>
<td>Motor neuron disease*</td>
<td>L. Sharp</td>
<td>Auditorium</td>
</tr>
<tr>
<td>7:45 am-8:00 am</td>
<td>L-2</td>
<td>A curious case of generalized denervation with markedly elevated CK</td>
<td>R. Coates</td>
<td>Auditorium</td>
</tr>
<tr>
<td>8:00 am-8:30 am</td>
<td>L-3</td>
<td>Experience with cardiac transplantation in muscular dystrophy patients</td>
<td>K. Ding</td>
<td>Auditorium</td>
</tr>
<tr>
<td>8:00 am-8:30 am</td>
<td>L-4</td>
<td>Variable phenotypes of Ullrich congenital muscular dystrophy</td>
<td>H. Gilbreath</td>
<td>Auditorium</td>
</tr>
<tr>
<td>8:30 am-9:00 am</td>
<td>L-5</td>
<td>Progressive weakness, wasting and white matter disease</td>
<td>M. Wicklund</td>
<td>Auditorium</td>
</tr>
<tr>
<td>9:00 am-9:30 am</td>
<td>L-6</td>
<td>CMT and Tremor – Is it really CMT?</td>
<td>E. Patton</td>
<td>Auditorium</td>
</tr>
<tr>
<td>9:00 am-9:30 am</td>
<td>L-7</td>
<td>An unusual case of muscle stiffness</td>
<td>M. Brock</td>
<td>Auditorium</td>
</tr>
</tbody>
</table>

**Session I Moderators:** Jaya Trivedi and Carlyne E. Jackson

**Abstract:** Title abbreviated from abstract.

## SESSION II

<table>
<thead>
<tr>
<th>TIME</th>
<th>ABSTRACT #</th>
<th>TITLE</th>
<th>PRESENTER(S)</th>
<th>LOCATION</th>
</tr>
</thead>
<tbody>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-1</td>
<td>Variability of muscle biopsy in nemaline rod myopathy</td>
<td>S. Kim</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-2</td>
<td>Nemaline myopathy: A case having partial trisomy of 17p12 and 16q23</td>
<td>B. Rabin</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-3</td>
<td>Acute onset neuromuscular junction dysfunction associated with statin</td>
<td>A. Mahmood</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-4</td>
<td>Demyelinating polyneuropathy in Waardenburg syndrome</td>
<td>T. Scarff</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-5</td>
<td>Novel heterozygous FIG4 mutation causing CMT1 phenotype</td>
<td>S. Baker</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-6</td>
<td>Abnormal spontaneous activity in Lambert-Eaton myasthenic syndrome</td>
<td>G. Roy</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-7</td>
<td>Limb-girdle myasthenia gravis</td>
<td>I. Laccheo</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-8</td>
<td>Oculomotor muscle weakness and generalized fatigue in a child with inflammatory myopathy</td>
<td>E. Gaitour</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-9</td>
<td>Statin-induced motor neuron disease</td>
<td>A. Swenson</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>P-10</td>
<td>Chronic distal weakness</td>
<td>D. Jabari</td>
<td>Woodlawn A&amp;B</td>
</tr>
</tbody>
</table>

**Session II Moderators:** Mazen Dimachkie and Steven Vernino

*Title abbreviated from abstract.*
# Day 1 Program Agenda – Thursday, February 17, 2011

## Session II (continued)

<table>
<thead>
<tr>
<th>Time</th>
<th>Session</th>
<th>Title</th>
<th>Presenter(s)</th>
<th>Location</th>
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<tbody>
<tr>
<td>1:00 pm-2:30 pm</td>
<td>ABSTRACT #</td>
<td>TITLE: Respiratory failure triggered by a car accident</td>
<td>V. Hoyos</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td></td>
<td>P-11</td>
<td>Myasthenia gravis presenting as dropped head syndrome</td>
<td>R. Paduga</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td></td>
<td>P-12</td>
<td>Leg weakness, isolated quadriiceps atrophy and elevated CPK</td>
<td>C. Phan</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td></td>
<td>P-13</td>
<td>A case of rapidly progressive quadraparesis with a twist</td>
<td>P. Lavian</td>
<td>Woodlawn A&amp;B</td>
</tr>
<tr>
<td></td>
<td>P-14</td>
<td>Transient Neonatal Autoimmune Autonomic Ganglionopathy</td>
<td>S. Baker</td>
<td>Woodlawn A&amp;B</td>
</tr>
</tbody>
</table>

## Session III

<table>
<thead>
<tr>
<th>Time</th>
<th>Session</th>
<th>Title</th>
<th>Presenter(s)</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>2:30 pm-2:40 pm</td>
<td>ABSTRACT #</td>
<td>TITLE: Actin filament aggregation myopathy – its nosological profile from the microscope to the molecule</td>
<td>H. Goebel</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-1</td>
<td>Heat or exercise-induced rhabdomyolysis*</td>
<td>W. Tang</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-2</td>
<td>The unusual case of recurrent tea-colored urine</td>
<td>C. Konersman</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-3</td>
<td>A 65 year old female with subacute onset of limb weakness</td>
<td>C. Phan</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-4</td>
<td>Riboflavin-responsive lipid myopathy</td>
<td>A. Verma</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-5</td>
<td>Chronic proximal and distal weakness</td>
<td>S. Majmudar</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-6</td>
<td>A young patient with amyloid deposits in the muscle</td>
<td>K. Schlesinger</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-7</td>
<td>Unknown case presentation: Muscle pain and weakness in a 69 year old man, 4 years after an autologous bone marrow transplant</td>
<td>J. Fratkin</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td>3:50 pm-4:10 pm</td>
<td>ABSTRACT #</td>
<td>TITLE: Recurrent respiratory failure</td>
<td>Y. Hannawi</td>
<td>Pre-Function-C</td>
</tr>
<tr>
<td></td>
<td>BM-9</td>
<td>A 16 year old boy with exercise-induced rhabdomyolysis</td>
<td>X. Li</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-10</td>
<td>A 10 year old boy with muscle pain and hemolytic anemia</td>
<td>N. Kuntz</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-11</td>
<td>Early onset ocular, facial and limb weakness with high CK</td>
<td>J. Rawson</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-12</td>
<td>Unusual case of myasthenia gravis and high white blood cell count</td>
<td>E. Patton</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-13</td>
<td>Progressive onset diplopia and dysphagia in a post-operative patient</td>
<td>B. Ray</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td>5:00 pm-5:10 pm</td>
<td>ABSTRACT #</td>
<td>TITLE: Familial hoarseness and distal weakness</td>
<td>A. Shaibani</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td>BM-14</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>BM-15</td>
<td></td>
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</tbody>
</table>

*Title abbreviated from abstract
# Day 2 Program Agenda – Friday, February 18, 2011

## Session IV

<table>
<thead>
<tr>
<th>Time</th>
<th>Session</th>
<th>Title</th>
<th>Presenter(s)</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>7:00 am-7:45 am</td>
<td>Registration</td>
<td>Continental Breakfast</td>
<td>Gil I. Wolfe</td>
<td>Pre-Function A</td>
</tr>
<tr>
<td>7:45 am-8:00 am</td>
<td>Welcome and Announcements</td>
<td></td>
<td></td>
<td>Pre-Function C</td>
</tr>
<tr>
<td>8:00 am-8:15 am</td>
<td>Abstract S-1</td>
<td>Statin induced immune-mediated necrotizing myopathy</td>
<td>A. Taneja</td>
<td>Auditorium</td>
</tr>
<tr>
<td>8:15 am-8:30 am</td>
<td>Abstract S-2</td>
<td>Necrotizing autoimmune statin associated myopathy</td>
<td>I. Muzyk</td>
<td>Auditorium</td>
</tr>
<tr>
<td>8:30 am-8:45 am</td>
<td>Abstract S-3</td>
<td>Severe weakness in an adolescent girl with anorexia nervosa</td>
<td>P. Mancias</td>
<td>Auditorium</td>
</tr>
<tr>
<td>8:45 am-9:00 am</td>
<td>Abstract S-4</td>
<td>Juvenile myasthenia gravis: A twenty year experience</td>
<td>D. Castro</td>
<td>Auditorium</td>
</tr>
<tr>
<td>9:00 am-9:05 am</td>
<td>Carrell Krusen Lecture</td>
<td>Introduction of Dr. Leshner The Cooperative International Neuromuscular Group (CINRG): Lessons Learned from a 10 year experience</td>
<td>G. I. Wolfe, Robert Leshner</td>
<td>Auditorium</td>
</tr>
<tr>
<td>9:05 am-10:00 am</td>
<td>Coffee Break</td>
<td></td>
<td></td>
<td>Auditorium</td>
</tr>
<tr>
<td>10:00 am-10:30 am</td>
<td>Abstract S-5</td>
<td>Use of dynamic pupillometry as an autonomic testing tool</td>
<td>S. Muppidi</td>
<td>Auditorium</td>
</tr>
<tr>
<td>10:30 am-10:45 am</td>
<td>Abstract S-6</td>
<td>Sporadic inclusion body myositis: A case cohort</td>
<td>B. Estephan</td>
<td>Auditorium</td>
</tr>
<tr>
<td>11:00 am-11:15 am</td>
<td>Abstract S-7</td>
<td>Intra and inter familial variability in clinical phenotype in autosomal dominant centronuclear myopathy from dynamin 2 mutations</td>
<td>L. Davis</td>
<td>Auditorium</td>
</tr>
<tr>
<td>11:15 am-11:30 am</td>
<td>Abstract S-8</td>
<td>Focal neuropathies treated with rimabotulinum toxin B</td>
<td>P. Grogan</td>
<td>Auditorium</td>
</tr>
<tr>
<td>11:45 am-12:00 pm</td>
<td>Abstract S-10</td>
<td>Collagen VI: Recognizing a spectrum of muscle disease</td>
<td>D. Arrington</td>
<td>Auditorium</td>
</tr>
<tr>
<td>12:00 pm-1:30 pm</td>
<td>Lunch</td>
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<td>Reverchon A&amp;B</td>
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## Session V

<table>
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<th>Session</th>
<th>Title</th>
<th>Presenter(s)</th>
<th>Location</th>
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<tbody>
<tr>
<td>1:30 pm-1:40 pm</td>
<td>Abstract BN-1</td>
<td>Late onset Tay Sachs disease*</td>
<td>V. Krishna</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td>1:40 pm-1:50 pm</td>
<td>Abstract BN-2</td>
<td>The atrophic hands</td>
<td>S. Beh</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td>1:50 pm-2:00 pm</td>
<td>Abstract BN-3</td>
<td>Seeing the forest for the trees: A different approach to progressive quadraparesis</td>
<td>K. Kolostyak</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td>2:00 pm-2:10 pm</td>
<td>Abstract BN-4</td>
<td>A young man with motor neuron-like syndrome</td>
<td>B. Estephan</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td>2:10 pm-2:20 pm</td>
<td>Abstract BN-5</td>
<td>A case of chronic sensory, motor and autonomic neuropathy</td>
<td>S. Chen</td>
<td>Reverchon A&amp;B</td>
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*Title abbreviated from abstract

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<thead>
<tr>
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<tr>
<td>2:20 pm-2:30 pm</td>
<td>BN-6</td>
<td><strong>ABSTRACT #</strong></td>
<td>S. Moody</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>TITLE</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>A unique case of atypical acute polyneuropathy</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2:30 pm-2:40 pm</td>
<td>BN-7</td>
<td><strong>TITLE</strong></td>
<td>I. Muzyka</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Unilateral facial numbness and weakness</td>
<td></td>
<td></td>
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<tr>
<td>2:40 pm-2:50 pm</td>
<td>BN-8</td>
<td><strong>TITLE</strong></td>
<td>L. Phillips</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Peripheral neuropathy due to dinitrophenoe used for weight loss</td>
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<td></td>
</tr>
<tr>
<td>2:50 pm-3:00 pm</td>
<td>BN-9</td>
<td><strong>TITLE</strong></td>
<td>T. Nguyen</td>
<td>Reverchon A&amp;B</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Treatment resistant acquired neuropathy presenting with dyspnea</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3:00 pm-3:10 pm</td>
<td>BN-10</td>
<td><strong>TITLE</strong></td>
<td>C. Arif</td>
<td>Reverchon A&amp;B</td>
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<td>Muscle twitching and loss of balance</td>
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<td>3:10 pm-3:20 pm</td>
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<td><strong>TITLE</strong></td>
<td>C. Phan</td>
<td>Reverchon A&amp;B</td>
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<td>A case of the dancer who can no longer dance</td>
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<td>3:20 pm</td>
<td>Adjourn</td>
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</tbody>
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*Title abbreviated from abstract.
Revised 12/13/10
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*Required fields

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Last Name* ___________________________

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Specialty* ___________________________

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INDICATE CREDIT DESIRED: ☐ AMA PRA Category 1 ☐ General
☐ Physical Therapy ☐ Occupational Therapy
☐ Respirational Therapy ☐ Nursing
☐ Certificate of Attendance

REGISTRATION FEES: $175 Early Registration (if received by February 4, 2011)
☐ $200 Late Registration (if received after February 4, 2011)

UT SOUTHWESTERN NEUROLOGY RESIDENTS AND FELLOWS:
☐ $75 Early Registration (if received by February 4, 2011)
☐ $85 Late Registration (if received after February 4, 2011)
☐ Symposium Faculty

THURSDAY DINNER RECEPTION:
☐ Yes I will be attending - $30
☐ Yes I will be attending with a guest - $30 per attendee
☐ Yes I am a UTSW Neurology Resident or Fellow and will be attending - $15

Attendees are also eligible to attend Neurology Grand Rounds at Texas Scottish Rite Hospital for Children at noon on Wednesday, February 16, 2011. There is no cost for this activity.

☐ Please check box if you plan on attending Grand Rounds.

Please indicate which meals you plan on attending:
Thursday, February 17, 2011 ☐ Continental Breakfast ☐ Lunch
Friday, February 18, 2011 ☐ Continental Breakfast ☐ Lunch

METHOD OF PAYMENT: (please select one of the following)
☐ MASTERCARD ☐ VISA Security Code

Card # ___________________________________________ ☐ AMEX Security Code

Amount to be Charged $ ______________

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☐ CHECK

Make checks payable in U.S. Dollars to: CME / UT Southwestern
Include the reference number #RP1102B on the check.
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Fax completed registration form to 214-648-4804.
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Continuing Medical Education / UT Southwestern
523 Harry Hines Blvd., / Dallas, Texas 75390-9059

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A handling fee of $50 will be deducted from cancellation refunds. Refund requests must be received by mail or fax prior to February 7, 2011. No refunds will be made thereafter.

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☐ Please check this box if you require assistance because of a disability to make this program accessible to you. Someone from our office will be in touch with you.

Comments: ___________________________

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