Cervical Dystonia Subtypes

Baseline Results From the <u>Cervical Dystonia Patient Registry for Observation of</u> **OnabotulinumtoxinA** <u>Efficacy</u> (CD PROBE)

P. David Charles,¹ Mark Stacy,² Joseph Jankovic,³ Marc Schwartz,⁴ Mitchell Brin,^{5,6} Spyridon Papapetropoulos^{5,7*}

(on behalf of the CD PROBE Study Group)

¹Vanderbilt University, Nashville, TN; ²Duke University, Durham, NC; ³Baylor College of Medicine, Houston, TX; ⁴MedNet Solutions, Minnetonka, MN; ⁵Allergan, Inc., Irvine, CA; ⁶University of California, Irvine, CA; ⁷University of Miami, Miller School of Medicine, Miami, FL

OBJECTIVE

To describe patient, disease, and treatment characteristics associated with subtypes of predominant head and neck postures in subjects with cervical dystonia (CD).

BACKGROUND

CD presents with variable head and neck postures as well as movements (dystonic tremor). Most clinical studies preferentially include subjects with presentations of predominant torticollis or laterocollis. In order to better understand why subjects with predominant retrocollis and anterocollis are often excluded, we have characterized the various CD subtypes with respect to demographic characteristics, severity and disability, and treatment approaches.

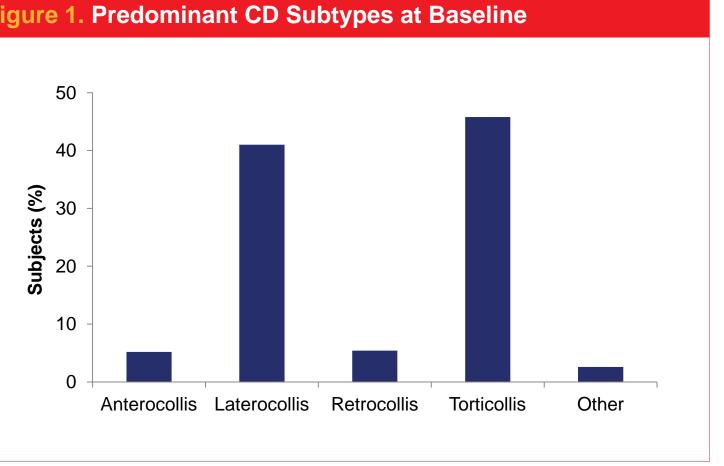
METHODS

CD PROBE (NCT00836017) is an ongoing longitudinal registry enrolling subjects with CD treated with onabotulinumtoxinA (Botox[®]). Subjects with CD and medically appropriate for botulinum toxin treatment that were either naïve to toxin or new to the physician and ≥16 weeks since the last injection are eligible for enrollment.¹ The predominant subtypes of CD at baseline are examined by the Toronto Western Spasmodic Torticollis Rating Scale (TWSTRS), Physician Global Assessment of Severity, Cervical Dystonia Impact Questionnaire (CDIP-58), and dose at initial treatment.

RESULTS

- As of September 13, 2011, 786 subjects had enrolled in CD PROBE and were analyzed for this report. Baseline characteristics are presented in Table 1.
- Predominant presentation subtypes are presented in Figure 1. Subjects presenting with predominant anterocollis or retrocollis experienced onset at a later age when compared with laterocollis or torticollis (p<0.05; **Table 2**). The time from diagnosis to treatment was shortest for retrocollis when compared with torticollis (p<0.0001; **Table 2**).
- Physicians more often rated subjects with anterocollis as severe when compared with other subtypes (p<0.02; Figure 2), and anterocollis subjects scored higher on TWSTRS and its disability subscale (p<0.05; Figure 3).
- Subjects with anterocollis or retrocollis scored higher on the CDIP-58 (Figure 4), and subjects with retrocollis were least likely to be employed (p<0.001; Table 2).
- Retrocollis was treated with the highest median dose (190U) when compared with the other subtypes (p<0.05) and anterocollis the least (115U; Figure 5).

Characteristic	
Sex Female Male	596 (76.1) 187 (23.9)
Race/Ethnicity Asian Black Hispanic Native American White Other	16 (2.0) 14 (1.8) 25 (3.2) 1 (0.1) 725 (92.6) 2 (0.3)
Age, y	57.7 ± 14.3 (19.4–100.0)
Body mass index	26.5 ± 5.4 (3.6–50.1)
Toxin status Naïve Non-naïve	502 (64.2) 280 (35.8)



CD = cervical dystonia

Data are presented as n (%) or mean ± standard deviation (range)

Table 2. Baseline Patient CD History

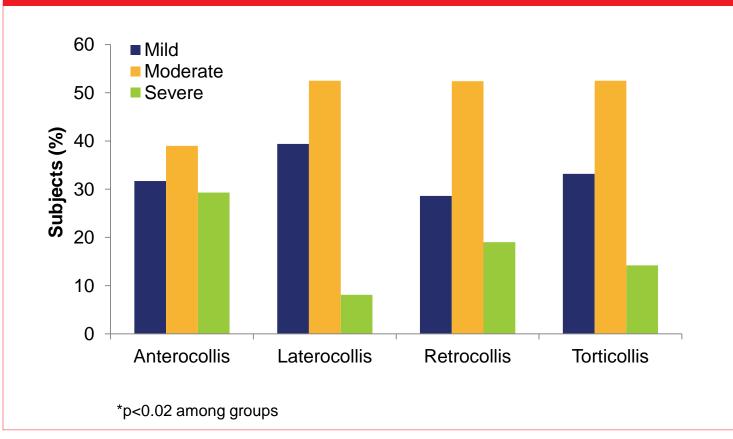
	Anterocollis	Laterocollis	Retrocollis	Torticollis	Total	p value
Age at symptom onset, y	53.0 ± 16.9	48.7 ± 16.8	53.1 ± 17.2	47.2 ± 15.4	48.5 ± 16.2	0.044
Time from CD onset to diagnosis, y	3.5 ± 6.4	5.5 ± 8.0	3.9 ± 7.5	5.4 ± 8.9	5.3 ± 8.4	0.217
Time from CD diagnosis to first treatment, y	0.7 ± 2.5	0.9 ± 3.2	0.1 ± 0.2	1.4 ± 5.3	1.1 ± 4.2	<0.0001
Currently employed	15 (37.5)	142 (46.7)	6 (15.4)	165 (47.8)	328 (45.1)	0.001

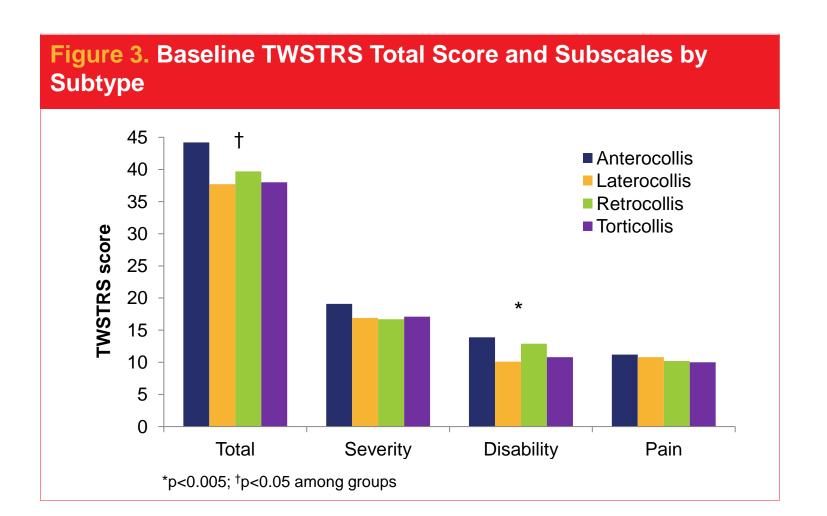
CD = cervical dystonia

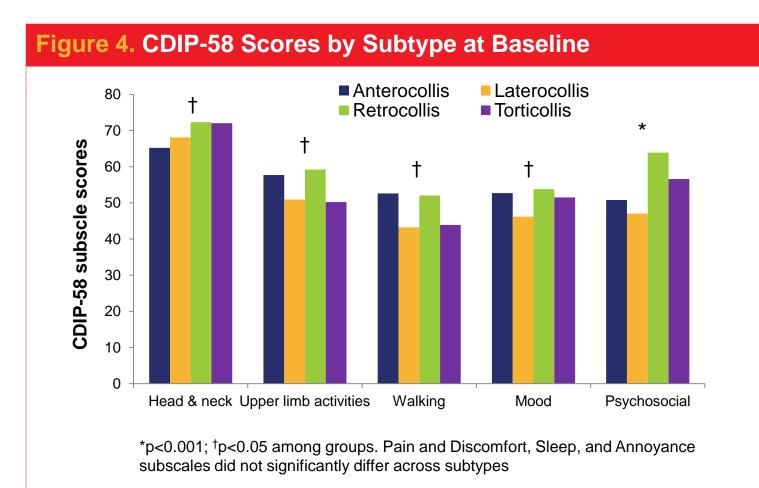
Other included homemaker and student

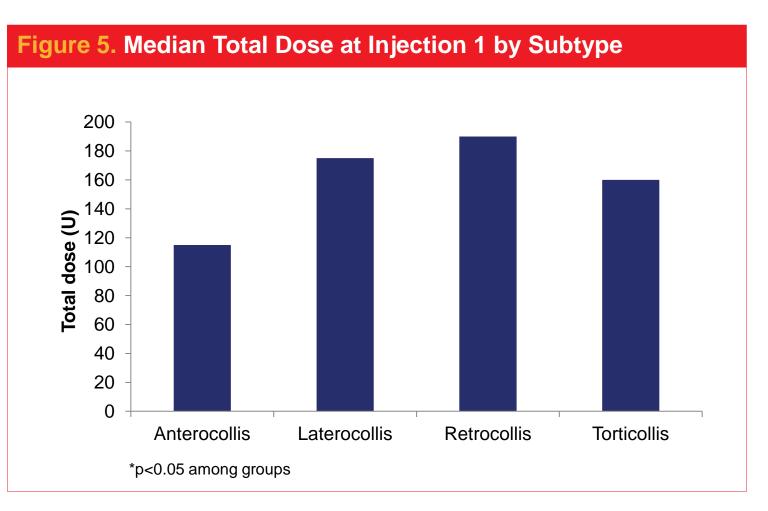
Data are presented as n (%) or mean ± standard deviation (range)











PROBE

CONCLUSIONS

CD presenting with predominant anterocollis or retrocollis is less common but is often associated with increased disease burden, disease severity, and disability; decreased quality of life; and effects on employment status when compared with other subtypes.

REFERENCE

Jankovic J, Adler CH, Charles PD, et al . BMC Neurol. 2011;11:140. 1.

CD PROBE Study Group

Pinky Agarwal, WA; Fahd Amjad, DC; Kristin Appleby, DC; Richard Barbano, NY; Peter Barbour, PA; Brandon Barton, IL; Jay Bhatt, IN; Kevin Biglan, NY; David Bowers, TN; David Brown, NY; Michelle Burack, NY; Barbara Changizi, NY; Mahan Chehrenama, VA; Nisha Chhadbria, DC; Cynthia Comella, IL; Francis Conidi, FL; Diane Counce, AL; Paul Cullis, MI; Khashayar Dashtipour, CA; Lisa Davidson, MN; Thomas Davis, TN; J Antonelle De Marcaida, CT; Nancy Diaz, PA; Christina Drafta, NY; Richard Dubinsky, KS; Jeffrey Esper, PA; Virgilio Evidente, AZ; Stanley Fisher, TX; Grace Forde, NY; Karen Frei, CA; Ramon Gil, FL; John Goudreau, MI; Alida Griffith, WA; Laurie Gutmann, WV; Gregory Hanes, FL; Edward Hartmann, GA; Robert Hauser, FL; Vanessa Hinson, SC; Patrick Hogan, WA; Tomas Holmlund, NY; Jyhgong Hou, TX; Christine Hunter, TX; Stuart Isaacson, FL; Bahman Jabbari, CT; Sandra Jacobson, AZ; Joseph Jankovic, TX; Paul Jett, TN; Katie Kompoliti, IL; Daniel Kremens, PA; Rajeev Kumar, CO; Eugene Lai, TX; Julie Leegwater-Kim, MA; Peter LeWitt, MI; Tsao-Wei Liang, PA; Steven Lo, DC; Duarte Machado, CT; Padma Mahant, AZ; Irene Malaty, FL; Bushra Malik, PA; Zoltan Mari, MD; Anthony May, PA; Emilio Melchionna, MA; Eric Molho, NY; Henry Moore, FL; Fatta Nahab, FL; Anthony Nicholas, AL; Suneetha Nuthalapaty, TN; Fernando Pagan, DC; Atul Patel, KS; Mayank Pathak, NY; Gauri Pawar, WV; Diana Pollock, FL; Adolfo Ramirez-Zamore, NY; Ben Renfroe, FL; Diana Richardson, CT; Perry Richardson, DC; Michael Rivner, GA; Ramon Rodriguez, FL; Michael Rossen, MA; Kyle Ruffing, FL; Marwan Sabbagh, AZ; Aliya Sarwar, TX; Cenk Sengun, FL; Kapil Sethi, GA; Scott Sherman, AZ; Holly Shill, AZ; Tanya Simuni, IL; Carlos Singer, FL; Michael Sorrell, MA; Natividad Stover, AL; Thyagarajan Subramanian, PA; David Swope, CA; Martin Taylor, OH; Margaret Tilton, NH; Richard Trosch, MI; Daniel Truong, NY; Winona Tse, NY; Miodrag Velickovic, NY; Aparna Wagle Shukla, FL; Cindy Zadikoff, IL; Lin Zhang, CA; Chong-Hao Zhao, CA

Disclosure

This study and its analysis were sponsored by Allergan, Inc., Irvine, CA. Assistance for poster development was provided by Jennifer Giel, PhD, of Evidence Scientific Solutions, and was funded by Allergan, Inc.

Dr Charles receives income from Allergan, Inc., Ipsen, Medtronic, and the Alliance for Patient Access for education and consulting services; Vanderbilt University received research support from Allergan, Inc., Ipsen, Merz, and Medtronic for research led by Dr Charles. Dr Stacy received compensation from Allergan, Inc., Boehringer-Ingelheim Inc, General Electric, Novartis, Osmotica, Synosia, Schering-Plough, GlaxoSmithKline, Teva, Biogen, and Neurologix for consulting, speaker bureau, protocol steering committee, and/or safety monitoring boards; he received royalties from Informa Press and research support from Ceregene, IMPAX, Michael J. Fox Foundation, Neuraltus Novartis, Parkinson Study Group, and Schering-Plough. Dr Jankovic received compensation from Allergan, Inc., Chelsea Therapeutics, EMD Serono, Merz Pharmaceuticals, Lundbeck Inc., and Teva for consulting services; he received compensation from Medlink: Neurology for serving as an editorial board member and received research support from Allergan, Inc., Allon Therapeutics, Ceregene Inc., Chelsea Therapeutics, Diana Helis Henry Medical Research Foundation, EMD Serono, Huntington's Disease Society of America, Huntington Study Group, Impax Pharmaceuticals, Ipsen Limited, Lundbeck Inc., Michael J. Fox Foundation for Parkinson Research, Medtronic, Merz Pharmaceuticals, National Institutes of Health, National Parkinson Foundation, Neurogen, St. Jude Medical, Teva Pharmaceutical Industries Ltd., University of Rochester, and Parkinson Study Group. Mr Schwartz is an employee of MedNet Solutions. Dr Brin is an employee of Allergan, Inc. and receives salary, stock, and stock options from Allergan, Inc. *Dr Papapetropoulos was an employee of Allergan, Inc. when this study was initiated. He is currently employed at Pfizer Inc in Cambridge, MA. He was previously employed by Biogen Idec, Inc. and received stock options from Biogen Idec, Inc.

Presented at the 16th International Congress of Parkinson's Disease and Movement Disorders (MDS), June 17–21, 2012, Dublin, Ireland