

A Multi-Center Study of Follow-Up Intervals in Patients with Cushing's Disease

Geer EB¹, Shafiq I², Gordon MB³, Ayala A⁴, Bonert V⁵, Surampudi V⁶, Katznelson L⁷, Carmichael JD⁵, Manuylova E², Pulaski-Liebert KJ⁸, Lalazar Y¹, Neary MP⁹, Ludlam WH⁹, Romanus D¹⁰, Cherepanov D¹⁰, Eagan M¹⁰, Lee J¹⁰, Broder MS¹⁰, Biller BMK⁸

¹Icahn School of Medicine at Mount Sinai, New York, NY, USA; ²University of Rochester School of Medicine and Dentistry, Rochester, NY, USA; ³Allegheny Neuroendocrinology Center, Allegheny General Hospital, Pittsburgh, PA, USA; ⁴University of Miami and Jackson Memorial Hospital, Miami, FL, USA; ⁵Cedars-Sinai Medical Center, Los Angeles, CA, USA; ⁶Harbor-UCLA Medical Center, Los Angeles, CA, USA; ⁷Stanford University School of Medicine, Stanford, CA, USA; ⁸Massachusetts General Hospital, Boston, MA, USA; ⁹Novartis Pharmaceuticals Corporation, East Hanover, NJ, USA; ¹⁰Partnership for Health Analytic Research, LLC, Beverly Hills, CA, USA

BACKGROUND

- Cushing's disease (CD) has an annual incidence of 1.2-2.4 per million in Europe and up to 8 per million in the U.S.¹⁻³
- Uncontrolled CD has significant morbidity and mortality.⁴
- Long-term follow-up of CD is essential to mitigate risks of increased morbidity/mortality.⁵
- No recommendations exist to guide follow up frequency, in part because the actual frequency with which CD patients are followed in clinical practice is unknown.

OBJECTIVE

- To determine the pattern and frequency of follow-up for CD patients in the US.

METHODS

Study Design and Data Source

- Retrospective data collected from medical records at 8 US pituitary/endocrine centers, selected based on volume of CD patients, location, and patient diversity.
- The study was approved by institutional IRBs.

Patient Selection

- Inclusion criteria:
 - Diagnosed with CD or CD recurrence within past 20 years; AND
 - ≥18 years old at diagnosis.

Data Collection

- Data collected from onset of CS symptoms through 2014
 - Demographics (age, sex, race/ethnicity),
 - Disease characteristics: onset of CS symptoms, date of diagnosis of CD, CD recurrence, biochemical status,
 - CD treatments delivered at study centers and local practices,
 - Comorbidities
 - Prevalence was based only on comorbidities reported at the study centers,
 - Final disposition: date of last visit, evidence of transfer of care, insurance status at last visit.

- Data quality measures included rigorous abstractor training, data quality checks, and follow-up abstraction for inconsistencies and missing entries.

Statistical Analysis

- Results stratified by length of time since last visit.
- Descriptive statistics, including mean, median, standard deviation, and percentage, were reported.
- Data transformations and analyses performed with SAS® version 9.4 (SAS Institute, Cary, NC).

RESULTS

Patient and Follow-Up Characteristics

- By 1/15/15, data regarding 163 patients had been entered in the database, and formed the basis of this analysis.
- 90 patients (55%) were last seen within 1 year, 32 (20%) within 1-2 years, and 41 (25%) >2 years prior (Table 1).
- Recurrent or residual disease was more commonly observed for those seen 1-2 years or >2 years prior (Table 1).
- Evidence of transfer of care, with presumed follow up elsewhere, was noted in 4%, 13% and 37% with last visit ≤1, 1-2, and >2 years prior, respectively (Table 1).

Treatment for Cushing's Disease

- 96% underwent surgical removal of pituitary adenoma as first-line therapy.
- Among 64 patients with residual/recurrent CD, 36 (56%) received pharmacotherapy, 20 (31%) underwent radiation therapy, and 9 (14%) were treated with adrenalectomy.
 - Fewer patients with last visit >2 years prior had radiotherapy (8%) compared to those seen within 1 year (42%) (Table 2).
- Median elapsed time from start of most recent type of therapy to last visit was shorter among patients last seen >2 years prior compared to those seen within 1 year, 4.7 vs. 11.6 months, respectively.

Table 2. Treatment for Recurrent/Residual CD^a

Patients with recurrent/residual CD	Time since last visit		
	≤1 year	1-2 years	>2 years
No. of patients	N=43	N=9	N=12
Pharmacotherapy	27 (63) ^b	2 (22)	7 (58)
Radiotherapy	18 (42)	1 (11)	1 (8)
Adrenalectomy	7 (16)	0 (0)	2 (17)

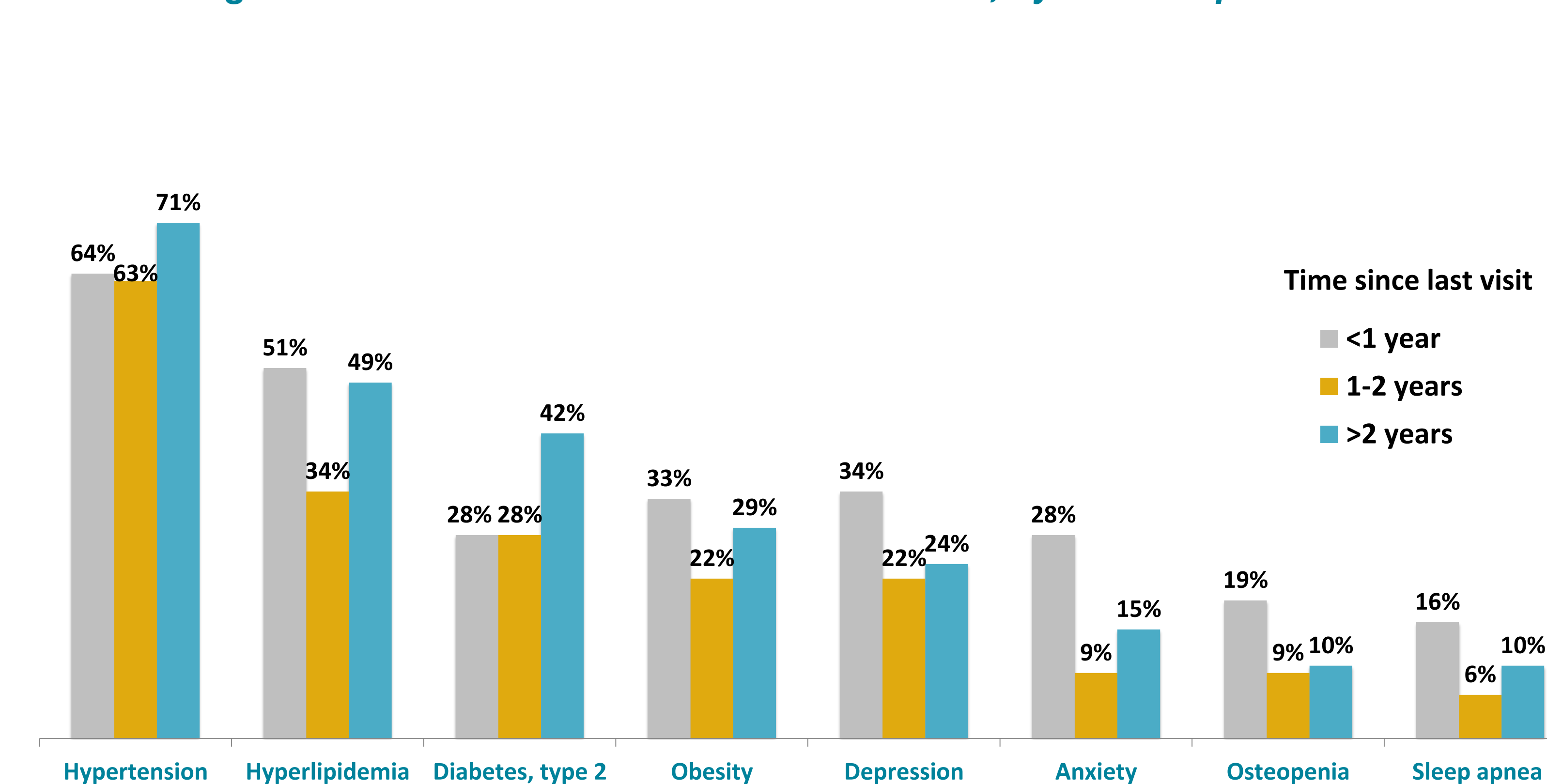
^a any therapies from first documented date of residual/recurrent CD to last visit at study site
^b n (%)

Table 1. Patient Characteristics, by Follow-Up Interval

Characteristic	Time since last visit		
	≤1 year	1-2 years	>2 years
N	90	32	41
Age at last visit (years), mean ± SD	47 ± 13	44 ± 15	44 ± 14
Median [range]	47 [23-83]	43 [21-73]	42 [19-79]
Female , n (%)	65 (72)	25 (78)	39 (95)
Race , Caucasian, n (%) ^a	65 (89)	20 (95)	26 (81)
Black	7 (10)	0 (0)	3 (10)
Other	1 (1)	1 (5)	3 (10)
Ethnicity , Hispanic/Latino, n (%)	13 (14)	4 (13)	7 (17)
No. of comorbidities , mean ± SD ^b	3.9 ± 2.5	2.3 ± 1.6	3.1 ± 2.1
Insurance status			
Insured, n (%)	84 (93)	31 (97)	38 (93)
Uninsured, n (%)	6 (7)	1 (3)	3 (7)
Recurrent/residual disease , n (%)	47 (52)	23 (72)	29 (71)
Time from diagnosis to last visit (years), median [range]	3.8 [0-27.4]	2.8 [0-18.8]	1.6 [0-21.7]
Follow-up duration at study centers^c (years), median [range]	2.3 [1-27.5]	1.2 [1-6.4]	1.1 [1-21.7]
Documented transfer of care , n (%)	4 (4)	4 (13)	15 (37)

^a percent among non-missing observations
^b based on comorbidities reported at study centers
^c from first to last visit

Figure. Prevalence of Individual Comorbidities, by Follow-Up Interval



LIMITATIONS

- Data on care outside the study center was likely underreported in medical records at study sites.
- These results are interim descriptive analyses. At study completion, statistical analyses will be conducted.
- Care and follow-up patterns at the 8 study centers may not be representative of all US centers.

CONCLUSIONS

- Many patients with CD went more than 1 year without a visit and a majority of these patients did not have documented transfer of care, but it is possible that some are being followed elsewhere.
- A higher proportion of patients seen >1 year had evidence of recurrent/residual disease.
- Patients not seen for extended periods may be at risk of undiagnosed recurrence or progression of CD and development of comorbidities, unless they are being seen at other centers.
- Further investigation of factors that increase the risk of becoming lost to follow-up warrant closer examination in order to prevent or mitigate undetected recurrence or progression of CD.

References

- Lindholm J, et al. *J Clin Endocrinol Metab.* 2001;86(1):117-23.
- Extabe J, Vazquez JA. *Clin Endocrinol (Oxf).* 1994;40:479-84.
- Broder MS, et al. *Pituitary.* 2014 May 7. [Epub ahead of print]
- Dekkers OM, et al. *J Clin Endocrinol Metab.* 2007;92:976-981
- Ayala A, et al. *J Neurooncol.* 2014 Jul 1 [Epub ahead of print].



URL: <http://novartis.medicalcongressposters.com/Default.aspx?doc=41c87>
And via Text Message (SMS)
Text : Q41c87 To : 8NOVA (86682) US Only
+18324604729 North, Central and South Americas; Caribbean; China
+447860024038 UK, Europe & Russia +46737494608 Sweden, Europe