

QUANTIFICATION OF POSTURAL SWAY IN NORMALS AND PATIENTS WITH CEREBELLAR DISEASES¹

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This paper compares data from normals and 5 groups of cerebellar patients collected in an attempt to quantify and characterize their postural sway while standing erect. It extends and supplements earlier work by Dichgans et al. (1976) and Mauritz et al. (1979). The results largely confirm those of the previous authors, describing patterns of sway abnormality that are specific to a given localization of the disease within the cerebellum. New is the differentiation of spinal ataxia in Friedreich's ataxia from anterior lobe atrophies and vestibulo-spinal cerebellar diseases. This study also gives a statistical analysis of the diagnostic importance of single sway parameters.

The 5 patient groups represent the most common localizations of diseases causing cerebellar symptoms: (1) Friedreich's disease, which is prototypical for a disorder of spino-cerebellar afferents, (2) atrophy of the anterior cerebellar lobe, the spino-cerebellum, to which spino-cerebellar afferents mainly project, (3) the vestibulo-cerebellum forming the lower vermis, (4) the cerebellar hemispheres, and (5) diffuse cerebellar atrophies. The pattern of recorded abnormalities has been evaluated as to the significance of every pathological result for localizing the causal disease. As a consequence of this, posturography is now a valuable tool in clinical diagnosis and quantitative documentation of postural ataxia for follow-up studies.

Methods*Apparatus*

Subjects stood on a force measuring platform. With this, strain gauges at the four corners of the platform measured the displacement of the center of foot pressure (CFP) in the antero-posterior and lateral directions. Antero-posterior sway in our experimental condition was computed from the difference between the front and back strain gauges. The lateral sway was computed in an analog manner from the two right and left force transducers. The influence of the subject's weight was electronically compensated.

Sway parameters

The antero-posterior and lateral CFP components were digitalized and, after multiplexing (PCM, $f = 263$ Hz, Kayser K 11 80-20), recorded on tape. On-line computations were done by use of a computer (IN 110). The computer program used a sampling interval of 25 msec and a total sampling duration of 25 sec. Since it was impossible to place different subjects in exactly identical positions on the platform, the first step was a normalization procedure. For this purpose the mean from all antero-posterior and lateral values was calculated and taken as the zero point from which the CFP deviated. The *sway path* (SP) travelled by the CFP on the platform was calculated as in our earlier study (Hufschmidt et al. 1980). The *sway area* covered by the sway path of the CFP within 25 sec was computed as

$$SA = \frac{1}{2} \sum_{i=1}^{n-1} |x_{i+1}y_i - x_iy_{i+1}|$$

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n = number of sample points, x = antero-posterior sway direction and y = lateral sway direction.

The amount of visual stabilization was estimated by the 'Romberg quotient' of the sway path and sway area (SP or SA with eyes closed divided by SP or SA with eyes open (Van Parys and Njiokiktjien 1976).

We also computed a sway *direction histogram* of the CFP. For this purpose the full circle of possible sway directions was divided into 24 intervals of 15° each. The single vectors of the displacement of the CFP within each sampling interval were sorted according to their directions and summed up. Straightforward was defined as zero.

The *amplitude* of body sway in the antero-posterior direction (AP) was calculated as the sum of the 6 direction vectors pointing in the AP direction. The amplitude in the lateral direction (LAT) was calculated in the same manner. The quotient AP:LAT indicates the predominance of antero-posterior body sway. Sway path and sway direction histograms from each subject were depicted on a storage oscilloscope and documented by way of a hard copy unit (Fig. 1).

Simultaneously the displacements of the CFP in the antero-posterior and lateral direction were fed into an on-line Fourier analyser. The total duration of sampling was 40 sec. The span of frequencies analysed ranged from 0.01 to 10 Hz (0.025 Hz steps). The sum of the root mean squares calculated from the power spectrum between 0.01 and 10 Hz served as an additional measure of the amount of postural sway. The peak amplitude within the Fourier spectrum, as identified on the oscilloscope, and its frequency and amplitude were plotted together with the root mean squares on a digital X-Y plotter.

Procedure

Subjects were placed on the platform with their feet parallel 4 cm apart. They were instructed to stand as quietly as possible with their arms folded in front of the chest. Intervals of 1 min each were recorded with eyes open and closed.

Normal subjects

Twenty healthy subjects (15 men, 5 women) aged between 20 and 60 years constituted the

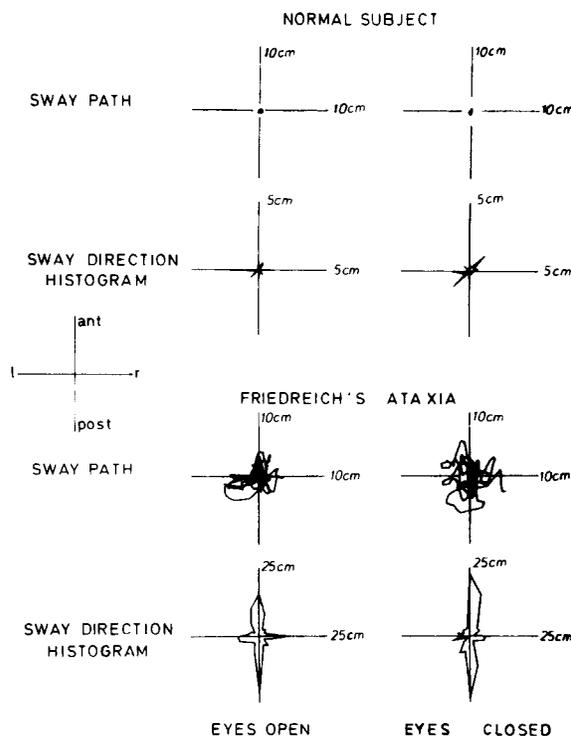


Fig. 1. Sway path and sway area from a normal subject and a patient with Friedreich's ataxia. Note different scalings in the records from the patient.

normal population. The means and standard deviations were calculated from all sway parameters (including root mean squares). The mean + 2 S.D. was defined as the upper limit of normality.

Patients

We examined 41 patients in whom it was possible to define from morphological (CT, surgery) or clinical data the localization of the cerebellar disturbance. The patients were classified into one of the following 5 groups:

(a) Eight patients with *Friedreich's ataxia* (FRIED) were representative of a disorder affecting mainly the afferents to the cerebellum, mediated by the spino-cerebellar tracts. Signs of involvement of the posterior columns (loss of vibration sense and position sense) were observed in all of them. Computerized tomography showed intact cerebellar structures without atrophy in all of these patients.

(b) Six patients had an isolated lesion of the

lower cerebellar vermis, the *vestibulo-cerebellum* (VERMIS). Two of them had a tumor (medulloblastoma, astrocytoma), two a bleed from an angioblastoma. One patient had an aplasia of the vermis. In all of them the vermal lesion was verified by CT and surgery. The anterior lobe and the hemispheres of the cerebellum were not or only little affected. The remaining patient suffered from a Hallgren syndrome (retinitis pigmentosa, deafness, vestibulo-cerebellar ataxia).

(c) Eleven patients suffered from a *late cortical atrophy of the anterior lobe* (ANT. LOBE) a disease which can be observed in chronic alcoholics with a nutritional deficiency (Marie et al. 1922; Victor et al. 1959; Mancall 1975; Adams 1976). Polyneuropathy was either absent or moderate. Computerized tomography of the 10 men and 1 woman showed, to a varying degree, a generalized atrophy of the brain with predominance in the vermal and anterior lobe region of the cerebellum.

(d) Seven patients suffered from diseases which were mainly localized in the cerebellar *hemispheres* (HEMISPHE). Three of them had bilateral ischemic lesions. In the four remaining ones, exhibiting ataxia and dysmetria predominantly of the arms and less of the legs, the cause of the cerebellar disease was unknown.

(e) Nine patients had a cerebellar disease which, according to clinical testing and CT, may have affected *all parts of the cerebellum* (DIFFUSE). Three of them suffered from chronic intoxication (two bromide, one diphenylhydantoin), five suffered from a heredoataxia (Gerstman-Sträussler (1), familial spastic ataxia (2), olivo-cerebellar degeneration of Holmes (2)); in one patient the cause of the cerebellar affection could not be classified.

Results

(a) Normal subjects

The sway path during the sample period of 25 sec increased by about 50% with closed eyes as compared to open eyes (Table I). A similar increase was documented by computing the sway area and the Romberg quotient. The antero-posterior sway was invariably greater than sway in the lateral dimension with and without the visual con-

tribution to postural stabilization. The means of individual antero-posterior/lateral quotients again indicated that the increase of sway path and sway area with eye closure was mainly due to an increase in the antero-posterior sway components.

(b) Patients with cerebellar diseases

Table I shows the frequency of pathological results ($> \text{mean} + 2 \text{ S.D.}$) in the different groups of cerebellar diseases for the 12 parameters. The mean percentage of pathological results amounted to 73.9% for the patients with atrophy of the anterior lobe. Patients with diffuse cerebellar lesions (63.9%) and Friedreich's ataxia (58.3%) followed. Patients with lower vermal lesions (48.6%) were more unstable than those with hemispherical cerebellar lesions (32.2%). The most effective parameters to detect postural instability were sway path, sway area, and the antero-posterior component of sway (AP), when the stabilizing effect of vision was suppressed by eye closure. Using these 3 parameters 77–86% of all patients showed pathological results (Fig. 2). The least effective parameters were the registration of the lateral components of sway with open eyes and the quotient of antero-posterior and lateral sway. The particular importance of visual stabilization of posture in patients with late atrophy of the anterior lobe and those with Friedreich's ataxia was illustrated by the frequency of a pathological Romberg quotient

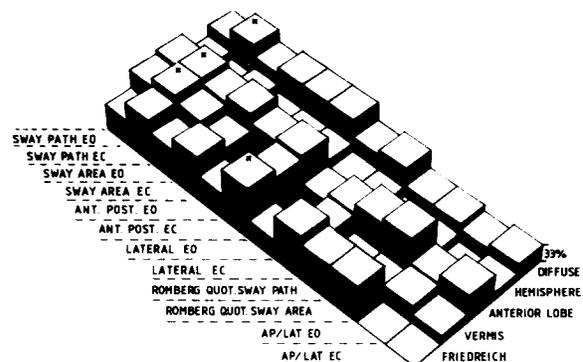


Fig. 2. Frequency of pathological results with different sway parameters (from left to right) across the 5 patient groups (from the foreground to the background). Columns with a cross on the top indicate 100% pathology.

TABLE I
Frequency (%) of pathological results (upper limit = AM + 2 S.D. from normals).

N	Sway path		Sway area		AP		LAT		Romberg quotient		Quotient AP/LAT		Mean (%) Mean ^a S.D.
	EO	EC	EO	EC	EO	EC	EO	EC	Sway path	Sway area	EO	EC	
8	50	86	62	86	50	100	38	86	71	71	0	0	58.3
6	83	100	83	67	33	83	0	50	17	17	33	17	48.6
11	90	100	90	100	82	90	36	54	73	82	36	54	73.9
7	57	43	57	43	43	43	29	29	14	14	0	14	32.2
9	78	100	78	89	89	89	56	67	33	33	22	33	63.9
Mean (%)	71.6	85.8	74	77	59.4	81	31.8	57.2	41.0	43.4	18.2	23.6	

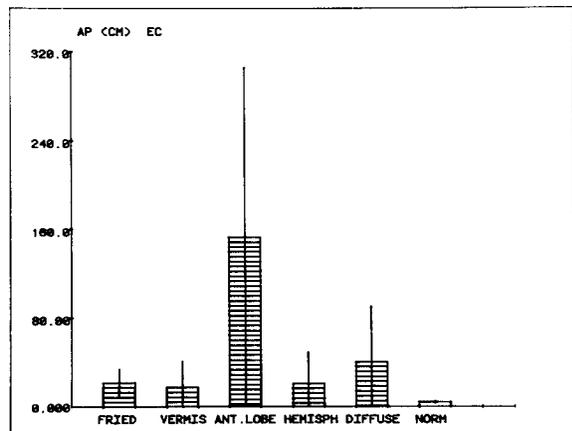
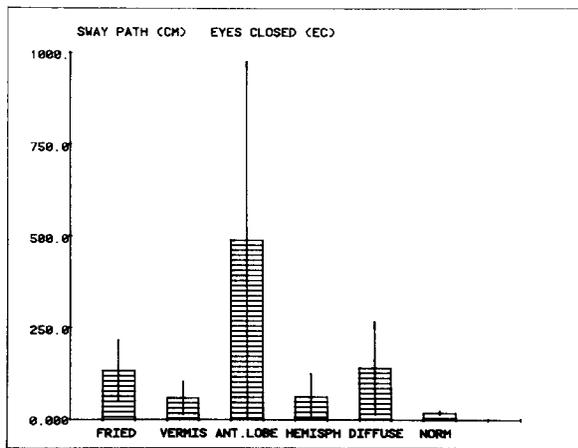
AP = antero-posterior sway; LAT = lateral sway; EO = eyes open; EC = eyes closed.

^a Mean and standard deviation from normals.

TABLE II

Analysis of variance for the results comparing the 5 groups of patients and normals.

Measured parameters	Eyes	$F_{P < 0.05} = 2.38$ $df (5/55)$	Multiple comparisons $P < 0.05$
Sway path	Open	3.74	Ant. lobe > norm
Sway path	Closed	7.29	Ant. lobe > all others
Sway area	Open	2.69	-
Sway area	Closed	3.10	Ant. lobe > norm
AP sway	Open	4.04	Ant. lobe > norm
AP sway	Closed	7.22	Ant. lobe > all others
LAT sway	Open	1.85 n.s.	-
LAT sway	Closed	5.71	Ant. lobe, Fried > norm
AP/LAT quotient	Open	2.62	-
AP/LAT quotient	Closed	2.26	-
Romberg quotient sway path		6.32	Ant. lobe > vermis. Hemisph. norm
Romberg quotient sway area		2.90	-



in these two groups. With lesions of the lower vermis and the cerebellar hemispheres the visual stabilization played a minor role.

Table II shows the results of an analysis of variance and of multiple comparisons. Despite the fact that F values showed a significant difference between the patient groups (except for the lateral sway component with open eyes) only the patients with atrophy of the anterior lobe could be clearly differentiated from the others. This effect was most clearly visible in the sway path and the antero-posterior sway component with closed eyes (Fig. 3). Patients with Friedreich's ataxia who showed also an increased sway path and sway area could be differentiated from those with anterior lobe atrophy by the fact that their AP/LAT quotient was lower than that of normals, indicating a stronger lateral sway component.

Fig. 3. Means and standard deviations of the sway path (cm, upper part) and the antero-posterior sway components (AP, lower part) recorded with the eyes closed from the 5 groups of patients. Patients with Friedreich's ataxia (FRIED), lesions of the cerebellar vermis (VERMIS), the anterior lobe (ANT. LOBE), the cerebellar hemispheres (HEMISP), diffuse cerebellar atrophies (DIFFUSE) compared to normals (NORM). Both parameters clearly differentiate patients with the anterior lobe disease.

TABLE III

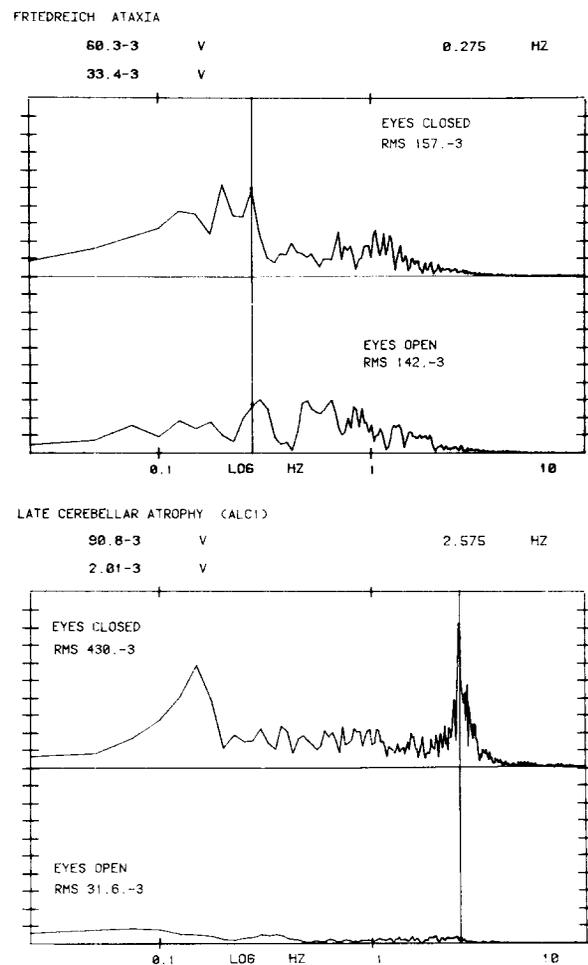
Analysis of variance of root mean squares. Five patient groups and normals.

Parameter	$F_{P < 0.05} = 2.38$ $df = 5/55$	Multiple comparisons
CFP AP EO	3.54	-
CFP AP EC	6.96	Ant. lobe > vermis, Hemisp, norm
CFP LAT EO	3.26	-
CFP LAT EC	7.92	Fried > verm, hemisp, Norm/ant. lobe > norm

CFP = center of foot pressure; EO = eyes open; EC = eyes closed.

(c) Results of Fourier analysis

The root mean squares of the Fourier spectra of the displacement of the center of foot pressure give a frequency independent global measure of postural instability (Kapteyn and Bles 1976). Table III shows the results of an analysis of variance for the 5 patient groups and the normals. It reveals significant *F* values for all measured parameters. The multiple comparisons indicate that with open eyes no patient group produced parameters that exceeded those of normals. With closed eyes, patients with Friedreich's ataxia and anterior lobe atrophy could be differentiated from those with lower vermal lesions and those with diseases which affected the cerebellar hemispheres. The last two frequently displayed root mean squares which were within the range of normality. Again, patients with Friedreich's ataxia had the highest CFP root mean squares when the lateral component was computed. The AP root mean squares were highest in patients with anterior lobe atrophy. The most prominent peak of the frequency spectrum of CFP movement was always within the range of 0.1-1 Hz for patients with Friedreich's ataxia and between 2 and 4 Hz (mainly around 3 Hz) for the patients with anterior lobe lesions (Fig. 4).



Discussion

A parametric comparison of sway path and sway area in normals with the data from other authors is impossible since the recording techniques and methods of data processing differ too much. Relative measurements, such as the differences between closed and open eyes or the ratio

Fig. 4. Comparison of Fourier power spectra from a patient with Friedreich's ataxia (upper part) and a patient with atrophy of the anterior lobe. The main frequency components in the patient with Friedreich's ataxia are between 0.1 and 1 Hz. The second patient shows the pathognomonic 2.6 Hz peak with closed eyes but not with open eyes. The root mean squares of sway (upper limit = 38.00) are also normal with open eyes. The scaling on the ordinate indicates 100 mV/steps. The vertical line indicates the measured frequency (0.275 and 2.575 Hz). The numbers at the top of the figures indicate the power at this frequency (mV).

possible by means of posturography, dynamic tests such as tilting the platform sinusoidally around the ankle joint, thereby injecting a low frequency disturbance, or suddenly displacing the platform and measuring the reflex responses in leg muscles supply valuable additional information (Dichgans et al. 1983).

Summary

Posturography was performed in 41 patients with cerebellar diseases by means of a force measuring platform using an on-line computer program which calculated sway path, sway area, antero-posterior and lateral sway components and the amount of visual stabilization. Postural ataxia was quantitatively studied in 8 patients with spinal ataxia (Friedreich's), 6 patients with vestibulo-cerebellar lesions, 11 patients with anterior lobe atrophy, 7 patients with hemispherical cerebellar lesions, and 9 patients with a disease affecting all parts of the cerebellum. Patients with lesions of the cerebellar hemispheres could not be separated from normals by means of posturography. Lesions of the spino-cerebellar afferents (Friedreich's ataxia) caused an omnidirectional low frequency sway with preserved visual stabilization. Patients with anterior lobe atrophy showed a predominant antero-posterior sway, often with a spontaneous high frequency body tremor around 3 Hz. Vestibulo-cerebellar lesions exhibited omnidirectional low frequency sway poorly stabilized by vision. Quantitative posturography helps to localize cerebellar lesions and allows for quantitative follow-up studies of cerebellar diseases.

Résumé

Quantification des oscillations posturales chez des sujets normaux et des patients cérébelleux

On a soumis à l'analyse posturographique 41 patients atteints de maladies du cervelet, ceci à l'aide d'une plateforme mesurant les forces, avec programme de calcul en ligne de la direction de l'oscillation, de sa surface, de ses composantes

antéro-postérieure et latérale, et du degré de stabilisation visuelle.

L'ataxie posturale a été étudiée quantitativement chez 8 patients avec ataxie spinale (de Friedreich), 6 patients avec lésion vestibulo-cérébelleuse, 11 avec une atrophie du lobe antérieur, 7 avec lésion des hémisphères cérébelleux et 9 avec une affection touchant toutes les parties du cervelet. Les sujets atteints aux hémisphères cérébelleux n'ont pas, avec la méthode posturographique, pu être séparés des normaux. L'ataxie de Friedreich était accompagnée d'oscillations à fréquence lente, dans toutes les directions, mais avec maintien de la stabilisation visuelle. Les patients avec atrophie du lobe antérieur montraient des oscillations avec prédominance antéro-postérieure, souvent accompagnées de tremblements corporels à fréquence élevée (autour de 3 Hz). Les lésions vestibulo-cérébelleuses ont entraîné des oscillations omnidirectionnelles à basse fréquence, très mal stabilisées par la vision.

La posturographie quantitative aide ainsi à localiser les lésions du cervelet et permet de suivre quantitativement l'évolution des affections cérébelleuses.

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