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Review Article

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Review/perspective on hysterical paralysis: A diagnosis of exclusion for spinal surgeons

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ABSTRACT

Background: Hysterical paralysis (HP) and/or conversion disorders (CD) are diagnoses of exclusion for spine surgeons. Before assigning this diagnosis to a patient, they must first undergo a full neurodiagnostic evaluation (i.e., X-rays, MR, CT/Myelo-CT) to rule out organic spinal pathology. Here, we reviewed select articles highlighting how to differentiate HP/CD patients from those with spinal disease.

Methods: Several case studies and small series of patients with HP/CD were included in our analysis. Notably, prior to being assigned the diagnoses of HP/CD, patients had to first undergo X-ray, MR, CT, and/or Myelo-CT evaluations to rule out spinal disorders; typically, their neurodiagnostic studies were normal.

Results: Patients with HP/CD often presented with varying clinical complaints of motor paralysis despite intact reflexes, normal sensory examinations, and lack of sphincter disturbance (i.e. intact rectal tone). Further, go and nogo functional MRI (fMRI) examinations demonstrated inconsistencies in areas of brain activation for patients with HP/CD complaints.

Conclusions: HP/CD are diagnoses of exclusion, and patients should first undergo a full panel of neurodiagnostic studies to rule out organic spinal disease. While those with HP/CD should not have unnecessary operations, those with real "surgical pathology" should have appropriate spine surgery performed in a timely fashion.

Keywords: Computed Tomography (CT), Conversion Disorder (CD), Diagnosis of Exclusion, Functional MRI (fMRI), Hysterical Paralysis, Magentic Resonance Imaging (MR/MRI)

INTRODUCTION

Hysterical paralysis (HP) or conversion disorders (CF) are diagnoses of exclusion [Table 1].^[1-7] Therefore, before assigning patients the diagnosis of HP/CD, spine surgeons must first complete full evaluations (i.e., X-rays, MR, CT, Myelo-CT studies) to rule out organic spinal disease that may warrant surgical intervention. Further, if and when fMRI is available, these studies may further supplement and differentiate between organic spinal pathology and HP/CD.^[4,10,16]

MULTIPLE FACTORS ASSOCIATED WITH HYSTERICAL PARALYSIS/ **CONVERSION DISORDERS**

Maxion et al. (1989) identified multiple factors in patients with "classical conversion syndromes" and/or psychogenic seizures/paralysis [Table 1].^[11] Out of 172 patients, 55% had

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Gould <i>et al.</i> ^[6] 1986	Validity Hysteria	7 Features Hysteria	7 Features Hysteria	Hysteria Findings with	Movement
Nerv Ment Dis	30 Consec	Hypo-chondria	Cont	Acute Brain Lesions	Paralysis Often
	Neurology	Secondary Gain	Changing	Invalidates 7 Features as	Mislabeled as
	Admits Acute Brain	BI, Nonanat Sensory Loss,	Hypalgesia Give way Weak	Typical for Hysteria	Hysteria Caution Diagnose
	Lesions	Pain/Vib Split	30 had 1 Feature		Hysteria-Often
		Midline	Most 3–4		Incorrect
Maxion <i>et al.</i> ^[11] 1989	HP Syndromes	172	Symptoms Ages	80% Severe Conflicts	Long-Lasting
Psychother	Not Rare	55%-Fits	15-25 or 45-55	Men-Professional	Syndrome
Psychosom Med	Younger Patients	Psychogenic	50% Unskilled	Problems	More in Females
Psychol		45% HP 2/3 F	Workers 12% Perinatal Brain	Females - Relationship Problems	PP
		275 F 25% GI Surgery	Damage	Problems	
Apple ^[1] 1989	HP-Spinal	Variety Patterns	15 of 17 Normal	16 of 17 Improved After	HP Spine
Paraplegia	17 Patients	Motor-	DTR	Avg 3.8 Day Hospital Say	Normal DTR
		Sensory Loss	All Sphincter	13 Fully Recovered	Normal Sph
			Control Intact		Routine X-rays
Marshall <i>et al.</i> ^[10]	Functional	UD Foignad?	Casar Famala Laft	fMRI	Not Organic Concluded
1997	Anatomy HP/CD	HP -Feigned? - Failure to Find	Case: Female Left Leg Paralysis +	Left Leg Paralyzed Go-	These 2 Areas
Cognition	Use fMRI Test	Organic Cause	Sensory Loss	Nogo Test Not Activate	Inhibit Prefrontal
0	Motor Paralysis	0	No Organic Lesion-	Right Primary Motor	Effects on Right
	Case		Psychological	Cortex-Right	Primary Motor
			Trauma	Activated Orbito-Frontal-	Cortex
Letonoff <i>et al.</i> ^[9] 2002	HP 3 F LE	Diagnosis Evolusion	Normal: Labs,	Anterior Cingulate Cortex Inconsistent Exams	Turnically
Spine	Paralysis	Diagnosis Exclusion. Not Organic	Electrical,	Loss Motor Sensory	Typically Walked Out of
opine	Prevalence	2 Trauma	Imaging Studies	Function-Normal DTR-	Hospital
	5-22 per 100,000	1 Prior HystS	Spontaneous Return	Claim Incontinence But	Told Exams
	Typically Female No Education		Function 6 mos	Normal Rectal Tone	Normal
Vuilleumier ^[16] 2005 Prog Brain Res	Hysterial CD-No	Psychogenic	fMRI See	fMRI <activity <="" frontal="" td=""><td>Better Know</td></activity>	Better Know
	Organic Lesion	Stress-	Increased Activation	Subcortical Motor Control with HP-<	NeuroPsych Bases
	Use fMRI	Emotional-Conflicts Use fMRI, EEG, PET	of cingulate or Orbitofrontal	Somatosensory Cortex	Hysterical CD Improve Clinical
		or SPECT	Regions During CD	HP	Management
			Symptoms	< Visual Cortex	
				Hysterical Blindness	
Okun and Koehler ^[13]	Astasia and	Despite Normal Leg	Blocq 19 th Century	Paralysis	CD Differentiated
2007	Abasia=	Function in Bed	Neurologists	Jump fits	from Hysteria
Mov Disord	Inability to stand and walk CD			Tremor Bizarre behavior=	
Cojan <i>et al.</i> ^[4] 2009	Motor Inhibition	Study Mechanism	Failure to Move	Identification	Conversion
Neuroimage	HP Go-NoGo	Inhibition Motor	Due to Activation	Inhibitory Mechanisms	Circuits Not
	fMRI Conversion	Brain Pathways by	Percuneus	in Simulation and	Usual Inhibition
	CD	Emotional Status	Ventrolateral Frontal	Conversion Paralysis	Circuits—
			Gyrus Cinculate Contex		Activation Midlin
			Cingulate Cortex, VMPFc		Brain-Emotional Regulation
Hsieh <i>et al</i> . ^[7] 2010	CD Paral After	Normal Till 1 hr	Motor Fx Recovered	Reexplore Negative	Normal SEP
Spine	LDH	Postop-LLE Weak	Postop Until	Normal SEP/	and MEP Help
	37 yo F Left Lam	Surgical Reexplore	Recurrent Paralysis	MEP+Imaging	Confirm CD with
	L5S1-LDH	No Clot	10 hrs later		Normal
		No Lesion			Motor/Sensory

(Contd...)

Table 1: (Continued).								
Stone <i>et al.</i> ^[15] 2012 J Neurol Neurosurg Psychiatry Zhu <i>et al.</i> ^[17] 2012	Autobio-graphy Functional Paralysis Book "A Leg To Stand On" HP After Spine	Recovery Fall Norway 1982 Leg Injury Surgery Reattach Quad Nonanatomic	Emotional Period Leg Not Part of Body-Hard to Regain Walking Ortho Surgeon	First Rejected Hysterical Paralysis Later Reconsidered Aware Nonantomic	Recognize Functional Psychogenic Paralysis Post Injury Rapid Recovery			
Rheumatol int	Surgery 2 Cases 2 M	Deficits	Should Learn to Recognize HP Avoid Unneeded Surgery	Functional Deficits	Expected from HI			
Nguyen <i>et al.</i> ^[12] 2013 Int J Obstet Anesth	Recurrent PP in 29 yo F After Dural Puncture Elective Delivery	Symptoms LE Weakness and Sensory Loss ASx Chiari I M	Spinal Anes Followed by GA (Convert to Open C Section)	Extubated -Bilateral LE Weakness 4-6 h later-MR Normal Dx HP/CD Prior Episode After Dural Puncture (i.e. at 27 Weeks Gestation similar event)	HP After Dural Puncture Rate			
De Bustos <i>et al.</i> ^[5] 2014 Font Neurol Neurosci	Clinical HP/CD Hotly Debated 2000 years	Typically F Sexual Disorders Other Diagnoses Neuroses. Parkinson's, Seizures	3 Classes HP- Acute Attacks- Paroxysms Functional Syndromes, Visceral events.	Minor HP-Syncope- Tetany, Twilight States, Paroxysmal Amnesia, Cataleptic Attacks.	Minor HP- Focal Hysterias, Paralyses, Contractures - Spasms, Anes Sensory Abnormality			
Kanchiku <i>et al.</i> ^[8] 2017 Clin Spine Surg	HP in 11 Adolescent	3 M, 8 F Avg Age 16.5 (13–19 yo) Followed Avg 4.67 years (1–10.25)	All Nonorganic Signs	Used MEP Assess Primary Muscle Weakness (Exclude Organic Disease)	Psychiatric Diagnosis 9 Neurosis 2 Psychosomatic Dis-Conservative Rx-Excellent 7 Good 4 Outcomes			
Blashfield ^[2] 2019 J Nerv Ment Dis	16 Class Systems Before WWI	<15% Correlation Over 7 Different American Classes	16 Classes: Mania Acute Mania Chronic Maina Melancolia Paralysis Senile Dementia	16 Classes: General Epileptic Insanity Hysterical Insanity Moral Insanity Idiocy Cretinism	11 Names 19 th Century Views Mental Disorders			
Osman <i>et al.</i> ^[14] 2020 Epilepsy Behav	FND Sudan 40 of 1000 Neuro- psych CD Clinic	Dx Exclusion Neuro Eval Depression BDI, HAD	60% Young F Unemp 82.5% HystS	47.5% Speech Abnl 35% week/Paral 97.5% MD	95% Marked Clinical Response Combined Anti-Dep PsychoTh			

HP: Hysterical Paralysis, Lit: Literature, Rev: Review, Conversion Disorder: CD, Org: Organic, Fx: Function, Ppt: Precipitated, TR: Trauma, Prev: Prevalence, F: Females, M: Males, LE : Lower Extremity, HystS: Hysterical Seizure, DTR: Deep Tendon Reflexes, Sph: Sphincter Function, Inc: Incontinence Bowel/ Bladder, Spont: Spontaneous, Pt: Patient, Educ: Education, Lim: Limited, SocBack: Socioeconomic Background, mos: Months, FND: Functional Neurological Disorder, Neuropsych: Neuropsychological, Neuro: Neurological, Eval: Evaluation, Dis: Disease, BDI: Beck Depression Inventory, HAD: Hospital Anxiety and Depression Scales, Abnl: Abnormalities, Weak: Weakness, Paral: Paralysis, Anti-Dep: Anti-Depressants, PsychoTh: Psychotherapy, Dx: Diagnosis, MD: Mood Disorder, Unemp: Unemployed, LDH: Lumbar Disc Herniation, Surg: Surgery, Lam: Laminotomy, Postop: Postoperatively, hrs: Hours, Electro: Electrodiagnostic. MEP: Motor Evoked Potentials, SEP: Somatosensory Evoked Potentials, CNS: Central Nervous System, SS: Signs/Symptoms, BI: Belle Indifference Vib: Vibration, Nonanat: Nonanatomical, Consec: Consecutive, Avg: Average, yo-Years Old ,Psycho: Psychogenic , Rx: Therapy, Ortho: Orthopedic, PP: Psychogenic Parlaysis, ASx: Asymptomatic, Chiari I M: Chiari I Malformation, Anes: Anesthesia, GA: General Anesthesia, GI Surgery: Gastric/Duodenal Ulcers, CNS: Central Nervous System, MRI: Magnetic Resonance Images, fMRI: Functional MRI, PET: Positron Emission Tomography Scan, SPECT: Single-Photon Emission Computerized Tomography, EEG: Electroencephalography, vmPFC: Ventromedial Prefrontal Cortex, Quad: Quadriceps, Cont: Continued psychogenic fits, and 45% had hysterical paralysis. The age at presentation was bimodal; ages 15-25 vs. 45-55 years of age. Two-thirds of the patients were females who mostly had "relationship problems", while the 1/3 who were males mostly complained of "professional problems". Eighty percent of both children and adults had severe social problems. Fifty percent were unskilled workers. Just 12% of younger patients had a history of perinatal brain damage. Of interest, an additional 25% had undergone prior gastric/ duodenal ulcer surgery.

CLASSIFICATION SYSTEMS FOR PSYCHOLOGICAL/MENTAL DISORDERS

In 2019, Blashfield analyzed 16 different classification systems using 22 common terms to describe psychological/ mental disorders that had been used since the 19th century [Table 1].^[2] Three of these descriptions included mania, general paralysis, and senile dementia [Table 1].^[2]

HISTORY OF HYSTERICAL PARALYSIS (HP)/CONVERSION DISORDERS (CD)

Several studies highlighted how inconsistencies in clinical evaluations and ultimately normal neurodiagnostic studies helped differentiate between HP/CD vs. organic spinal disorders [Table 1].^[1,5,13] In 1989, Apple et al. evaluated 17 patients presenting with different complaints/patterns of motor paralysis, sensory loss, and sphincter dysfunction; 15 of 17 had normal reflexes despite motor paralysis, and all 17 had intact sphincter control (i.e. despite some complaints of incontinence) [Table1].^[1] Sixteen of 17 spontaneously improved over an average 3.8 day hospital stay, with 13 recovering full normal function over this period. They recommended obtaining X-ray/other studies early in the work-up of patients with potential HP/CD diagnoses, and that inconsistencies in the neurological examinations were extremely helpful in differentiating between HP/CD and organic spinal disorders. Okun and Koehler (2007) cited Paul BLocq's (1860-1896) definition of astasia-abasia; "... the inability to maintain an upright posture, despite normal function of the legs in...bed", while further differentiating this syndrome from typical hysteria [Table 1].^[13] DeBustos (2014) et al. reassessed the inconsistent clinical presentation of patients with HP/CD, and broke hysteria down into several main groups [Table 1].^[5] These included; "...paroxysms, attacks, acute manifestations, long-lasting functional syndromes, and visceral events." The first group included minor/major hysterical attacks, while the second group included "...focal hysterical symptoms, paralyses, contractures and spasms, anesthesia, and sensory disorders."

DIFFERENTIATING HYSTERICAL PARALYSIS/ CONVERSION DISORDERS FROM ORGANIC SPINAL PATHOLOGY

The incidence of HP/CD is 5-22/100,000 in the overall population. Here we have summarized how the diagnoses of HP/CD can be differentiated from organic spinal pathology [Table 1].^[6,8,9,12,14,17] Gould et al. (1986) observed that 30 patients with organic brain lesions demonstrated at least one of 7 "pathognomonic" findings of hysteria, while many exhibited 3-4."[6] These criteria included a; "....history of hypochondriasis, secondary gain, la belle indifference, nonanatomic sensory loss, splitting of midline by pain or vibratory stimulation, changing boundaries of hypoalgesia, and giveaway weakness". Their observations invalidated using these findings to differentiate HP/CD from organic disease.^[6] Additionally; "...movement disorders and paralysis are most often mislabeled as hysteria", and, "A diagnosis for hysteria must be made with great caution as it so often proves incorrect".^[6] Further, they showed how major biases impacted who was misdiagnosed with HP/CD; those typically misdiagnosed included women, homosexual men, and those with psychiatric illnesses. Letonoff et al. (2002) evaluated 3 females who presented with complaints of complete paralysis/sensory/sphincter loss in the lower extremities, but had intact reflexes/rectal tone, normal laboratory studies, electrodiagnostic evaluations, and imaging studies [Table 1].^[9] Interestingly, all were from low income groups with little education, 2 patients had histories of trauma, and a third patient had prior "hysterical" seizures. Once they were told that they had no "organic disease", all 3 patients recovered within 6 months during which time they typically required psychiatric help and physical therapy. Zhu et al. (2012) cited 2 males who presented with inconsistent neurological deficits that led to the diagnoses of HP/CD; both patients spontaneously fully recovered. Nguyen et al. (2013) reported a 29 year-old female who, 4-6 hours following a cesarean section performed under general anesthesia once a spinal anesthetic had failed, developed bilateral lower extremity weakness [Table 1].^[12] Following the normal lumbar MR, she was diagnosed with psychogenic paresis (HP/CD). Interestingly, she had experienced a similar episode several months earlier after a lumbar puncture. Kanchiku et al. (2017) used motor evoked potentials (MEP) to rule out organic disease and rule in hysterical motor deficits in 11 teenagers averaging 16.5 years of age [Table 1].^[8] Osman et al. (2020) diagnosed 40 functional neurological disorders (FND) out of 1000 new admission to a neuropsychiatric clinic [Table 1].^[14] Those with FND were typcially females (60%) with; "...psychogenic nonepileptic seizures (82.5%), speech abnormalities (47.5%), limb paralysis or weakness (35%)...". About 95% of these patients improved on antidepressant medication and/or with psychotherapy.

CONVERSION PARALYSIS AFTER SPINE SURGERY

Few papers focused on episodes of HP/CD "paralysis" immediately following spine surgery [Table 1].^[7,17] In 2010, Hsieh *et al.* reported a 37 year old female who, one hour after a left L5S1 laminotomy, newly developed left leg weakness; the immediate reoperation showed no significant hematoma or other pathology.^[7] Ten hours following the second surgery, the patient developed recurrent left leg weakness. This time, however, normal neurodiagnostic studies (i.e. including somatosensory evoked (SEP) and motor evoked potentials MEP)), led to the correct diagnosis of HP/CD.

CONVERSION PARALYSIS AFTER LEG INJURY

In 2012, Stone reported his own experience with transient psychogenic paralysis of a leg following a fall resulting in a torn quadriceps muscle [Table 1].^[15] He developed transient hysterical paralysis postoperatively that required him to regain the ability to walk.

FUNCTIONAL MRI OF PARALYSIS

fMRI, performed in patients with unexplained/inconsistent neurological deficits and normal neurodiagnostic studies, helped identify/differentiate patients with HP/CD vs. organic spinal disease [Table 1].^[4,10,16] In 1997, Marshall et al. had a patient with no organic lesion to explain her intermittent left-leg paralysis/sensory loss [Table 1].^[10] On the fMRI, the request to prepare to move the right leg, and actually move the good right leg, resulted in appropriate activation of the left motor/premotor cortex. However, when asked to move the paralyzed left leg, there was just activation documented in the right orbito-frontal and right anterior cingulate cortices. In 2005, Vuilleumier used fMRI to assess the locations of brain activation during HP/CD-related motor paralysis, sensory disturbances, and "blindness"; rather than showing activation in the motor/premotor, somatosensory, and visual cortices respectively, they typically demonstrated increased activity in the cingulate and orbitofrontal areas [Table 1].^[16] Cojan et al. (2009) used the go-nogo fMRI tests to demonstrate activation in the precuneus, ventral lateral frontal gyrus, and ventromedial prefrontal cortices rather than the motor/premotor cortex in patients presenting with HP/CD paralysis [Table 1].^[4]

MEDICOLEGAL IMPACT OF MISDIAGNOSING HYSTERICAL PARALYSIS

The diagnoses of HP/CD are diagnoses of exclusion, and should not be established without first obtaining appropriate neurodiagnostic tests and other studies [Table 1].^[1-17] Too frequently females are misdiagnosed with HP/CD, where in fact, they have real spinal pathology. In these cases the

failure to "diagnose and treat", especially in a timely fashion, can lead to irreversible neurological deficits that should have been avoided. Reviewing a case decades ago from a major academic medical center involved a middle aged female who, following a lumbar laminectomy, was "paralyzed." Rather than obtaining a MR, she was dismissed as exhibiting "hysterical paralysis"; the next day, the MR showed a hematoma that was then removed. Nevertheless, by that time, her paralysis was permanent/irreversible. There are likely many similar medicolegal cases out there where spine surgeons/other specialists have failed to rule out organic disease and differentiate patients' real organic complaints (i.e. attributable to spinal-surgical disease) from HP/CD.

CONCLUSION

Patients should not be labeled with HP/CD paralysis until neurodiagnostic/other studies (i.e., variously including MR, CT-Myelo-CT, fMRI, SEP/EMG/ MEP) have ruled out the presence of organic disease.

Declaration of patient consent

Patient's consent not required as there are no patients in this study.

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Conflicts of interest

There are no conflicts of interest.

REFERENCES

- 1. Apple DF Jr. Hysterical spinal paralysis. Paraplegia, 1989;27:428-31.
- 2. Blashfield RK. Pre-kraepelin names for mental disorders. J Nerv Ment Dis 2019;207:726-30.
- Boudissa M, Castelain JE, Boissiere L, Mariey R. Pointillart, Vital JM. Conversion paralysis after cervical spine arthroplasty: A case report and literature review. Orthop Traumatol Surg Res 2015;101:637-41.
- Cojan Y Waber L, Carruzzo A, Vuilleumier P. Motor inhibition in hysterical conversion paralysis. Neuroimage 2009;47:1026-37.
- De Bustos EM, Galli S, Haffen E, Moulin T. Clinical manifestations of hysteria: An epistemological perspective or how historical dynamics illuminate current practice. Front Neurol Neurosci 2014;35:28-42.
- Gould R, Miller BL, Goldberg MA, Benson DF. The validity of hysterical signs and symptoms. J Nerv Ment Dis 1986;174:593-7.
- 7. Hsieh MK, Chang CN, Hsiao MC, Chen WJ, Chen LH. Conversion paralysis after surgery for lumbar disc herniation.

Spine 2010; 35:E308-10.

- 8. Kanchiku T, Suzuki H, Imajo Y, Yoshida Y, Nishida N, Taguchi T. psychogenic low-back pain and hysterical paralysis in adolescence. Clin Spine Surg 2017;30:E1122-5.
- 9. Letonoff EJ, Williams TR, Sidhu KS. Hysterical paralysis: a report of three cases and a review of the literature. Spine 2002;27:E441-5.
- Marshall JC, Halligan, Fink GR, Wade DT, Frackowiak RS. The functional anatomy of a hysterical paralysis. Cognition 1997:64:B1-8.
- Maxion H, Fegers S, Pfluger R, Wiegand J. Risk factors of classical conversion syndrome--psychogenic seizures and paralyses--observations at a neurologic clinic with 172 patients]. Psychother Psychosom Med Psychol 1989;39:121-6.
- 12. Nguyen J, Abola R, Schabel J. Recurrent psychogenic paresis after dural puncture in a parturient. Int J Obstet Anesth 2013;22:160-3.

- 13. Okun MS, Koehler PJ. Paul Blocq and (psychogenic) astasia abasia. Mov Disord 2007;22:1373-8.
- 14. Osman AH, Alsharief SM, Siddig HE. Functional neurological disorder: Characteristics and outcome in a limited-resources country (Sudan). Epilepsy Behav 2020;111:107151.
- 15. Stone J, Perthen J, Carson AJ. "A leg to stand on" by Oliver Sacks: A unique autobiographical account of functional paralysis. J Neurol Neurosurg Psychiatry 2012;83:864-7.
- 16. Vuilleumier P. Hysterical conversion and brain function. Prog Brain Res 2005;150:309-29.
- 17. Zhu L, Ni B, Guo Q. Hysterical paralysis after spinal surgery. Rheumatol Int 2012;32 4077-8.

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