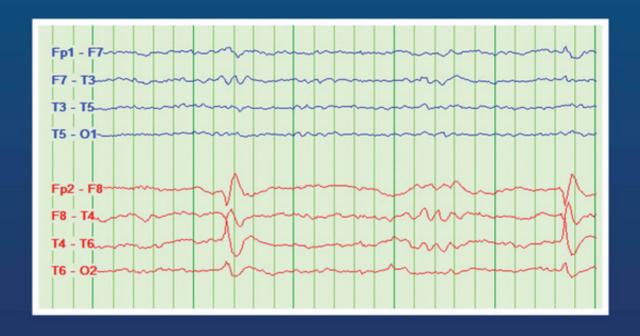
CLINICAL EEG OF ADULTS AND ADOLESCENTS



DAVID P. MOORE, MD

WILEY

Clinical EEG of Adults and Adolescents



Clinical EEG of Adults and Adolescents

David P. Moore University of Louisville Louisville, Kentucky, USA

WILEY

This edition first published 2025 © 2025 John Wiley & Sons Ltd

All rights reserved, including rights for text and data mining and training of artificial intelligence technologies or similar technologies. No part of this publication may be reproduced, stored in a retrieval system, or transmitted, in any form or by any means, electronic, mechanical, photocopying, recording or otherwise, except as permitted by law. Advice on how to obtain permission to reuse material from this title is available at http://www.wiley.com/go/permissions.

The right of David P. Moore to be identified as the author of this work has been asserted in accordance with law.

Registered Offices

John Wiley & Sons, Inc., 111 River Street, Hoboken, NJ 07030, USA John Wiley & Sons Ltd, New Era House, 8 Oldlands Way, Bognor Regis, West Sussex, PO22 9NQ, UK

For details of our global editorial offices, customer services, and more information about Wiley products visit us at www.wiley.com.

The manufacturer's authorized representative according to the EU General Product Safety Regulation is Wiley-VCH GmbH, Boschstr. 12, 69469 Weinheim, Germany, e-mail: Product_Safety@wiley.com.

Wiley also publishes its books in a variety of electronic formats and by print-on-demand. Some content that appears in standard print versions of this book may not be available in other formats.

Trademarks: Wiley and the Wiley logo are trademarks or registered trademarks of John Wiley & Sons, Inc. and/or its affiliates in the United States and other countries and may not be used without written permission. All other trademarks are the property of their respective owners. John Wiley & Sons, Inc. is not associated with any product or vendor mentioned in this book.

Limit of Liability/Disclaimer of Warranty

While the publisher and authors have used their best efforts in preparing this work, they make no representations or warranties with respect to the accuracy or completeness of the contents of this work and specifically disclaim all warranties, including without limitation any implied warranties of merchantability or fitness for a particular purpose. No warranty may be created or extended by sales representatives, written sales materials or promotional statements for this work. This work is sold with the understanding that the publisher is not engaged in rendering professional services. The advice and strategies contained herein may not be suitable for your situation. You should consult with a specialist where appropriate. The fact that an organization, website, or product is referred to in this work as a citation and/or potential source of further information does not mean that the publisher and authors endorse the information or services the organization, website, or product may provide or recommendations it may make. Further, readers should be aware that websites listed in this work may have changed or disappeared between when this work was written and when it is read. Neither the publisher nor authors shall be liable for any loss of profit or any other commercial damages, including but not limited to special, incidental, consequential, or other damages.

Library of Congress Cataloging-in-Publication Data Applied for

Paperback ISBN 9781394315345

Cover Design: Wiley

Cover Image: Courtesy of David P. Moore

Set in 11.5/14pt STIXTwoText by Straive, Pondicherry, India

This book is dedicated to Kerri Remmel, MD, PhD, Professor emiritus and former Chair of the Department of Neurology at the University of Louisville School of Medicine, and to Professors M. Steven Evans, MD, MS and Christopher Shafer, MD.

Brief Contents

Chapter 1:	Arrays and Montages	1
Chapter 2:	The Normal Adult EEG	15
Chapter 3:	Activation Methods	47
Chapter 4:	Patterns of Uncertain Significance	69
Chapter 5:	Artifacts	97
Chapter 6:	The Breach Rhythm	125
Chapter 7:	Slowing	129
Chapter 8:	Rhythmic Delta Activity (RDA)	135
Chapter 9:	Typical Triphasic Waves	147
Chapter 10:	Sporadic Epileptiform Discharges	155
Chapter 11:	Periodic Discharges	173
Chapter 12:	SIRPIDs	191
Chapter 13:	Background Attenuation or Suppression	197
Chapter 14:	Burst-Suppression	201

viii BRIEF CONTENTS

Chapter 15:	Disorders of Cognition and Consciousness	207
Chapter 16:	The Locked-In Syndrome	221
Chapter 17:	Brain Death	225
Chapter 18:	Postanoxic Coma	231
Chapter 19:	Head Trauma	243
Chapter 20:	Stroke	253
Chapter 21:	Epilepsy and Seizures	263
Chapter 22:	Syncope	357
Chapter 23:	Migraine	363
Chapter 24:	EEG Effects of Medications and Intoxicants, Metabolic Disorders, Vitamin Deficiencies, Endocrinologic Disorders, and Heavy Metals	369
Chapter 25:	Infectious Disorders	437
Chapter 26:	Neurodegenerative, Movement, and Related Disorders	459
Chapter 27:	Vascular Dementia	479
Chapter 28:	Developmental, Neurocutaneous, and Other Disorders Typically Presenting in Childhood or Adolescence	483
Chapter 29:	Autoimmune and Demyelinating Diseases	513
Chapter 30:	Vascular Disorders	531
Chapter 31:	Prion Diseases	543
Chapter 32:	Hydrocephalus	557
Chapter 33:	Brain Tumors	559
Chapter 34:	Amnestic Disorders	565
Chapter 35:	Stuporous Catatonia	571
Chapter 36:	Episodic Violence	573
Chapter 37:	Primary Psychiatric Disorders	579
Chapter 38:	Sleep Disorders	597
Chapter 39:	Transcranial Stimulation	609
Index		619

Contents

Cha	pter 1:	Arrays and Montages	1
1.1	Arrays	· ·	
	1.1.1	International 10–20 System	1 1
	1.1.2	Anatomical Position of Electrodes of the International 10–20 System	3
	1.1.3	Modified Combinatorial 10–10 System	6
	1.1.4	Supplemental Noninvasive Electrodes	8
1.2	Monta	qes	8
	1.2.1	Bipolar Montages	8
		1.2.1.1 Longitudinal Bipolar Montages	9
		1.2.1.2 Transverse Bipolar Montages	9
	1.2.2	Ipsilateral Ear Referential Montages	11
	1.2.3	Choosing Among Montages	12
	Refere	nces	12
Cha	pter 2:	The Normal Adult EEG	15
2.1	Alpha Rhythm		15
	2.1.1	General Characteristics	16
	2.1.2	Reactivity	17
	2.1.3	Alpha Variants	19
	2.1.4	Abnormalities of the Alpha Rhythm	19
		2.1.4.1 Attenuation and Slowing	19
		2.1.4.2 The Bancaud Phenomenon	21

X CONTENTS

2.2	Beta A	Activity			
2.3	The D	Pesynchronized EEG	23		
2.4	The E	ffect of Aging	24		
	2.4.1	Aging and the Alpha Rhythm	24		
	2.4.2	Aging and Focal Slowing	25		
2.5	Mu Rl	hythm	25		
2.6		da Waves	28		
2.7	Sleep		30		
	2.7.1	The Stages of Sleep	30		
		2.7.1.1 Stage N1	30		
		2.7.1.2 Stage N2	31		
		2.7.1.3 Stage N3	32		
		2.7.1.4 Stage R	32		
	2.7.2	Specific Electroencephalographic Features of Sleep	33		
		2.7.2.1 V Waves	33		
		2.7.2.2 Sleep Spindles	35		
		2.7.2.3 K Complexes	36		
		2.7.2.4 Sawtooth Waves	38		
		2.7.2.5 POSTS	39		
		2.7.2.6 Mittens	40		
	2.7.3	Sleep Deprivation	41		
	Refere	ences	42		
Cha	pter 3:	Activation Methods	47		
3.1	-	and Sleep Deprivation	47		
	3.1.1	Sleep	47		
	3.1.2	Sleep Deprivation	48		
3.2	Intern	nittent Photic Stimulation (IPS)	50		
	3.2.1	Methodology	50		
	3.2.2	The Photoparoxysmal Response (PPR)	52		
	3.2.3	Photomyoclonic Response	57		
	3.2.4	Photic Driving Response	58		
3.3	Нурег	rventilation	61		
	3.3.1	Methodology	61		
	3.3.2	The Buildup	62		
	3.3.3	Activation	63		
	3.3.4	Contraindications and Safety	64		
	Refere		65		
Cha	pter 4:	Patterns of Uncertain Significance	69		
4.1	-	Sharp Spikes (Benign Sporadic Sleep Spikes, Benign Epileptiform			
		ients of Sleep)	69		
4.2	Wicke	•	72		
4.3	6 Hz Spike and Wave Bursts ("Phantom" Spike and Wave)				

Contents xi

4.4	14- and 6-Hz Positive Bursts (14- and 6-Hz Positive Spike				
	Discharges, "Cteno	ids")	78		
4.5	Rhythmic Tempore	al Theta Burst of Drowsiness (Rhythmic Mid-temporal			
	Discharges; Psycho		80		
4.6	SREDA (Subclinica	al <u>R</u> hythmic <u>E</u> lectrographic <u>D</u> ischarge of <u>A</u> dults)	81		
	4.6.1 Typical SR	EDA	83		
	4.6.2 Atypical S	REDA	86		
	4.6.3 Concludin	g Remarks	86		
4.7	Rhythmic Midline	Theta	87		
	4.7.1 The Cigan	ek Rhythm	88		
	4.7.2 Frontal Mi	idline Theta	89		
	4.7.3 Texting Rl	ıythm	91		
	References		92		
Cha	oter 5: Artifacts		97		
5.1	Eye Movement		98		
		je Movements	98		
	_	e Movements	100		
		o-oculogram	100		
		Eyes and Retinal Detachment	102		
5.2	Muscle Artifact		102		
5.3	Glossokinetic Artif	act	104		
5.4	Sweat Artifact		104		
5.5	EKG Artifact		105		
5.6	Ballistocardiograpl	nic Artifact	108		
5.7	Pulse Artifact		108		
5.8	Tremor Artifact		109		
5.9	Shiver Artifact		111		
5.10	Patient Movement		112		
5.11	Alternating Currer	nt	116		
5.12	Electrode Pop		116		
5.13	Dental Amalgam A	Artifact	118		
5.14	Drip Artifact		118		
5.15	-	act and Electroretinogram	119		
5.16	Salt Bridge		121		
5.17	Ventilator Artifact		122		
5.18	Snoring Artifact		123		
	References		124		
Cha	oter 6: The Breac	h Rhythm	125		
	References	-	128		
Chaj	oter 7: Slowing		129		
7.1	Generalized Slowin	ng	129		
7.2	Lateralized Slowing 1				

xii CONTENTS

7.3	Focal Slowing	131
7.4	Zeta Waves	132
	References	133
Char	oter 8: Rhythmic Delta Activity (RDA)	135
8.1	Intermittent Rhythmic Delta Activity (IRDA)	136
	8.1.1 Frontal Intermittent Rhythmic Delta Activity (FIRDA)	136
	8.1.2 Occipital Intermittent Rhythmic Delta Activity (OIRDA)	137
	8.1.3 Temporal Intermittent Rhythmic Delta Activity (TIRDA)	140
8.2	Generalized Rhythmic Delta Activity (GRDA) and Lateralized	
	Rhythmic Delta Activity (LRDA)	143
	8.2.1 GRDA	143
	8.2.2 LRDA	144
8.3	Rhythmic Delta Activity in the Critically Ill	144
	References	145
Char	oter 9: Typical Triphasic Waves	147
ona	References	151
01	40 C 1: F :1 :C F: 1	
-	oter 10: Sporadic Epileptiform Discharges	155
10.1	Typical Sporadic Epileptiform Discharges	155
10.2	Secondary Bilateral Synchrony and the Possible Development of a	160
10.2	Mirror Focus	160
10.3	Capturing a Sporadic Epileptiform Discharge After a Seizure	165
10.4	Sporadic Epileptiform Discharges as Predictors of Relapse During	166
10.5	Anti-seizure Medication Withdrawal	166
10.5	Sporadic Epileptiform Discharges Occurring in Normal Individuals	1//
10.6	and in Patients Without a History of Unprovoked Seizures Transitory Cognitive Impairment Caused by a Sporadic	166
10.0	Epileptiform Discharge	167
10.7	Music Hath Charms: The Mozart Effect	168
10.7	References	168
	references	100
Chap	oter 11: Periodic Discharges	173
11.1	Generalized Periodic Discharges (GPDs)	174
	11.1.1 Description of GPDs	174
	11.1.2 Etiology of GPDs	174
	11.1.3 GPDs and Seizures	174
11.2	Lateralized Periodic Discharges (LPDs)	176
	11.2.1 Description of LPDs	177
	11.2.2 Etiology of LPDs	177
	11.2.3 LPDs and Seizures	179
	11.2.4 Lateralized Periodic Discharges Plus Fast (LPDs+F) Activity	183
11.3	Bilateral Independent Periodic Discharges (BIPDs)	184
11.4	Unilateral Independent Periodic Discharges (UIPDs)	185
11.5	Multifocal Periodic Discharges (MfPDs)	186
	References	188

Contents xiii

Cha	pter 12: SIRPIDs	191
12.1	-	191
12.2	1	192
12.3	Association of SIRPIDs with Spontaneously Occurring Seizures	194
	References	194
Cha	pter 13: Background Attenuation or Suppression	197
13.1	Symmetric Background Attenuation or Suppression	197
13.2	Asymmetric Background Attenuation	198
	Reference	199
Cha	pter 14: Burst-Suppression	201
14.1	Description of Burst-Suppression	201
14.2	Etiology of Burst-Suppression	205
	References	205
Cha	pter 15: Disorders of Cognition and Consciousness	207
15.1	Dementia	207
15.2	Delirium	208
15.3	Coma	209
	15.3.1 Typical Background Seen in Coma	209
	15.3.2 Specific Electroencephalographic Patterns Seen in Coma	210
	15.3.2.1 Alpha Coma	210
	15.3.2.2 Spindle Coma	212
	15.3.2.3 Beta Coma	214
	15.3.3 Reactivity and the Prognostic Value of the EEG in Coma	214
	15.3.4 Nonconvulsive Seizures in Coma	215
15.4	Persistent Vegetative State and Minimally Conscious State	216
	15.4.1 Persistent Vegetative State	216
	15.4.2 Minimally Conscious State	217
15.5	Akinetic Mutism	218
	References	218
	pter 16: The Locked-In Syndrome	221
16.1	The EEG in Classic Locked-In Syndrome	221
16.2	The EEG in Complete Locked-In Syndrome	222
	References	222
	pter 17: Brain Death	225
17.1	Clinical Criteria for the Diagnosis of Brain Death	226
17.2	Minimum Technical Standards for EEG Recording in Suspected	
	Cerebral Death	227
17.3	Normothermia and an Absence of Drug Intoxication	228
17.4	Sensitivity and Specificity of Electrocerebral Inactivity for Death	229
	References	229

XİV CONTENTS

Chap	oter 18: Postanoxic Coma	231
18.1	The Background	232
	18.1.1 Slowing	232
	18.1.2 Suppressed Background	232
	18.1.3 Background Reactivity	232
	18.1.4 Alpha Coma Pattern	233
	18.1.5 Spindle Coma Pattern	233
	18.1.6 Electrocerebral Inactivity	233
18.2	Discharges and Burst-Suppression	234
	18.2.1 Discharges	234
	18.2.1.1 Sporadic Epileptiform Discharges	234
	18.2.1.2 Periodic Discharges	234
	18.2.2 Burst-Suppression	234
18.3	Seizures	235
	18.3.1 Myoclonic Seizures	235
	18.3.2 Burst-Suppression with Eye-Opening	236
	18.3.3 Nonconvulsive Seizures	237
18.4	Stimulus-Induced Rhythmic, Periodic, or Ictal Discharges (SIRPIDs)	238
18.5	The EEG in Prognosis	238
	References	239
Char	oter 19: Head Trauma	243
19.1	Moderate to Severe Traumatic Brain Injury	243
	19.1.1 The Background	244
	19.1.2 Discharges and Rhythmic Delta Activity	244
	19.1.3 Seizures	245
	19.1.3.1 Early Seizures	245
	19.1.3.2 Late Seizures, Including Post-traumatic Epilepsy	245
19.2	Mild Traumatic Brain Injury	246
	19.2.1 Concussion	247
	19.2.2 Post-concussion Syndrome	247
19.3	Blast and Whiplash Injuries	248
19.4	Chronic Traumatic Encephalopathy	248
19.5	Subdural Hematoma	249
	References	250
Char	oter 20: Stroke	253
20.1	Infarction	253
-	20.1.1 Supratentorial Infarction	253
	20.1.1.1 Large Vessel Occlusions	253
	20.1.1.2 Lacunes	254
	20.1.2 Brainstem Infarction	255

Contents xv

20.2	Intrace	erebral He	morrhage	256
	20.2.1	Suprater	ntorial, Nontraumatic, Non-aneurysmal	
		Intracere	ebral Hemorrhage	256
	20.2.2	Brainster	m Hemorrhage	257
20.3	Aneur	ysmal Subo	arachnoid Hemorrhage	257
	Refere	nces		258
Chap	oter 21:	Epilepsy	and Seizures	263
21.1	Inherit	ed or Idio _l	pathic Epilepsies	263
	21.1.1	Idiopath	ic Generalized Epilepsies	264
		21.1.1.1	Childhood Absence Epilepsy	264
		21.1.1.2	Juvenile Absence Epilepsy	271
		21.1.1.3	Juvenile Myoclonic Epilepsy	272
		21.1.1.4	Epilepsy with Generalized Tonic–Clonic Seizures Alone	277
	21.1.2	Epilepsy	with Eyelid Myoclonia (Jeavons Syndrome)	277
	21.1.3	Typical A	Absence Status Occurring in Inherited Epilepsies	279
		21.1.3.1	Childhood and Juvenile Absence Epilepsies	280
		21.1.3.2	Juvenile Myoclonic Epilepsy	281
		21.1.3.3	Jeavons Syndrome	281
	21.1.4	Epilepsy with Myoclonic Absences		
	21.1.5	Self-Lim	ited Focal Epilepsies of Childhood	283
		21.1.5.1	Self-Limited Epilepsy with Centrotemporal Spikes	284
		21.1.5.2	Self-Limited Epilepsy with Autonomic Seizures	285
		21.1.5.3	Childhood Occipital Visual Epilepsy	286
	21.1.6	Focal Ep	ilepsies with Autosomal Dominant Inheritance	287
		21.1.6.1	Sleep-Related Hyperkinetic Epilepsy (SHE)	288
		21.1.6.2	Familial Mesial Temporal Lobe Epilepsy	288
		21.1.6.3	Epilepsy with Auditory Features	289
		21.1.6.4	Familial Focal Epilepsy with Variable Foci	289
	21.1.7	Progressi	ive Myoclonus Epilepsies	290
		21.1.7.1	Unverricht–Lundborg Disease	290
		21.1.7.2	Lafora Disease	290
		21.1.7.3	Neuronal Ceroid Lipofuscinoses	291
		21.1.7.4	Myoclonus Epilepsy with Ragged Red Fibers (MERRF)	292
21.2	Other	Epilepsies	Typically of Childhood Onset	293
	21.2.1	Lennox-	Gastaut Syndrome	294
		21.2.1.1	The Electroencephalographic Background	294
		21.2.1.2	Generalized Paroxysmal Fast Activity and Tonic Seizures	294
		21.2.1.3	Bursts of Atypical Generalized Spike–Wave Complexes	
			and Atypical Absence Seizures	296
		21.2.1.4	Atypical Absence Status Epilepticus	298
	21.2.2	Landau-	Kleffner Syndrome	298
	21.2.3	Rasmuss	en's Syndrome	300

XVI CONTENTS

21.3	Localiz	zation-Rela	nted Epilepsies	300
	21.3.1	Tempora	l Lobe Epilepsy	301
		21.3.1.1	Medial Temporal Lobe Epilepsy	301
		21.3.1.2	Lateral Temporal Lobe Epilepsy	306
		21.3.1.3	Electroencephalographic Differences Between Mesial	
			Temporal Sclerosis and Lateral Temporal Lobe Epilepsy	307
	21.3.2	Frontal L	obe Epilepsy	309
	21.3.3		Lobe Epilepsy	311
	21.3.4		l Lobe Epilepsy	312
	21.3.5	-	e Epilepsy	313
21.4	Seizure	Types		313
	21.4.1		zed Tonic–Clonic Seizures	314
	21.4.2	Focal Aw	vare (Simple Partial) Seizures	314
	21.4.3		ic Seizures	315
	21.4.4	Absence		316
	21.4.5	Focal Im	paired Awareness (Complex Partial) Seizures	316
21.5	Status	Epilepticus	S	317
	21.5.1	Generaliz	zed Tonic–Clonic Status Epilepticus	317
	21.5.2		ic Status Epilepticus	322
	21.5.3	Focal Aw	vare (Simple Partial) Motor Status Epilepticus	323
	21.5.4	Nonconv	ulsive Status Epilepticus (NCSE)	323
		21.5.4.1	Focal Impaired Awareness (Complex Partial)	
			Status Epilepticus	327
21.6	The Po	stictal EEC	J.	328
21.7	Ictal-I1	nterictal C	ontinuum (IIC) and Brief Potentially Ictal Rhythmic	
	Discha	rges (B[I]R	Ds)	334
	21.7.1	Ictal-Inte	erictal continuum	334
	21.7.2	Brief Pot	entially Ictal Rhythmic Discharges (B[I]RDs)	335
21.8	Psycho	ses of Epile	epsy	337
	21.8.1	Ictal Psyc	chosis	337
	21.8.2	Postictal	Psychosis	338
	21.8.3	Psychosis	s of Forced Normalization	338
	21.8.4	Interictal	l Psychosis	339
	Refere	nces		340
Chap	oter 22:	Syncope		357
22.1	Neural	ly Mediate	ed Syncope	358
	22.1.1	Vasovago	al Syncope	358
	22.1.2	Carotid S	Sinus Hypersensitivity	359
	22.1.3	Micturiti	on Syncope	359
22.2	Cardio	ic Syncope		360
22.3		laneous Sy1	ncopes	360
	22.3.1	Cough Sį	yncope	360
	22.3.2	Fainting	Larks	361
22.4			into a Seizure	361
	Refere	nces		362

Contents xvii

Cha	pter 23:	Migrain		363
23.1				364
23.2	Migrai	Migraine with Brainstem Aura		
23.3	Hemip	miplegic Migraine		
23.4	4 Ictal Migrainous Headache			365
	Referer	nces		366
Cha	pter 24:	EEG Effe	ects of Medications and Intoxicants, Metabolic	
		Disorder	s, Vitamin Deficiencies, Endocrinologic Disorders,	
		and Hea	vy Metals	369
24.1	Medica	tions and	Intoxicants	369
	24.1.1	Antidepr	ressants	369
		24.1.1.1	Tricyclics	370
		24.1.1.2	Mirtazapine	371
		24.1.1.3	SSRIs	371
		24.1.1.4	Bupropion	371
	24.1.2	Serotonii	n Syndrome	372
	24.1.3	Lithium		372
	24.1.4	Antipsyc	hotics	373
		24.1.4.1	First-Generation Agents	373
		24.1.4.2	Second-Generation Agents	374
		24.1.4.3	Clozapine	375
	24.1.5	Tardive I	Dyskinesia	376
	24.1.6	Neurolep	rtic Malignant Syndrome	376
	24.1.7	Levodopo	1	376
		Amantac		377
	24.1.9	Antichol	inergics	377
	24.1.10	Antihisto	nmines	378
	24.1.11	Antiseizu	are Medications	378
	24.1.12	Buspiron	e	382
	24.1.13	Benzodio	zepines	382
	24.1.14	Barbitur	ates	383
	24.1.15	Propofol		385
	24.1.16	Etomida	te	385
	24.1.17	Dexmede	rtomidine	386
	24.1.18	Opioids		386
	24.1.19	Alcohol o	and Alcohol-Related Disorders	387
		24.1.19.1	Alcohol Intoxication	387
		24.1.19.2	Blackouts	388
		24.1.19.3	Pathological Intoxication	388
		24.1.19.4	Uncomplicated Alcohol Withdrawal	388
		24.1.19.5	Delirium Tremens	388
		24.1.19.6	Alcohol Withdrawal Seizures	389
		24.1.19.7	Subacute Encephalopathy with Seizures in	
			Alcoholics (SESA)	390
	24.1.20	Disulfira	m	390

xviii CONTENTS

	24.1.21	390	
	24.1.22	Cannabis Sativa	391
	24.1.23	Solvents	391
		24.1.23.1 Acute Intoxication	392
		24.1.23.2 Chronic Cognitive Deficits	392
	24.1.24	Hallucinogens	393
		24.1.24.1 LSD	393
		24.1.24.2 Mescaline	393
	24.1.25	Ketamine and Phencyclcidine	393
		24.1.25.1 Ketamine	394
		24.1.25.2 Phencyclidine	394
	24.1.26	Certain Antibiotics	394
		24.1.26.1 Penicillins	394
		24.1.26.2 Cephalosporins	395
		24.1.26.3 Isoniazid	396
	24.1.27	Acyclovir	396
	24.1.28	Baclofen	396
	24.1.29	Aminophylline	397
		Lidocaine	397
	24.1.31	Diqoxin	397
	24.1.32	Corticosteroids and ACTH	397
24.2	Metabo	olic Disorders	398
	24.2.1	Hyponatremia	398
	24.2.2	Hypernatremia	398
	24.2.3	Hypocalcemia	398
	24.2.4	Hypercalcemia	399
	24.2.5	Hypomagnesemia	399
	24.2.6	Hypermagnesemia	399
	24.2.7	Hypoglycemia	400
	24.2.8	Hyperglycemia	401
	24.2.9	Uremia	401
	24.2.10	Hepatic Encephalopathy	402
	24.2.11	Sepsis	403
	24.2.12	Respiratory Failure	404
	24.2.13	Carbon Monoxide Intoxication	405
	24.2.14	Hepatic Porphyria	405
24.3	Vitami	n Deficiencies	406
	24.3.1	B12 Deficiency	406
	24.3.2	Wernicke's Encephalopathy	406
	24.3.3	Pellagra	407
24.4	Endocr	inologic Disorders	408
	24.4.1	Cushing's Syndrome	408
	24.4.2	Adrenocortical Insufficiency	408
	24.4.3	Hyperthyroidism	409
	24.4.4	Hypothyroidism	409

Contents xix

24.5	Heavy :	Metals	409
	24.5.1	Thallium	410
	24.5.2	Arsenic	410
	24.5.3	Tin	410
	24.5.4	Manganese	410
	24.5.5		411
	24.5.6	Mercury	411
		Aluminum	411
	24.5.8	Bismuth	412
	Referen	nces	412
Chap	oter 25:	Infectious Disorders	437
25.1	Acute I	Bacterial or Viral Meningitis	437
25.2		al Brain Abscess	438
25.3	Subdur	al Empyema	438
25.4		e Viral Infections	439
	25.4.1	Arboviral Encephalitis	439
	25.4.2	Herpes Simplex Virus Encephalitis	440
		SARS-CoV-2	441
	25.4.4	Herpes Zoster	442
	25.4.5	Human Immunodeficiency Virus (HIV) Infection	444
	25.4.6	Progressive Multifocal Leukoencephalopathy (PML)	445
	25.4.7	Cytomegalovirus (CMV) Encephalitis	446
	25.4.8	Human Herpesvirus 6 Infection	446
	25.4.9	Mumps	446
	25.4.10	Infectious Mononucleosis	447
	25.4.11	Rabies	447
	25.4.12	Subacute Sclerosing Panencephalitis (SSPE)	447
25.5		oan Infections	448
	25.5.1	Toxoplasmosis	448
	25.5.2	Malaria	449
25.6	Spiroch	netal Disease	449
	25.6.1	Tertiary Neurosyphilis	449
	25.6.2	Lyme Disease	450
25.7	Tuberci		450
	Referen	nces	450
Chap	oter 26:	Neurodegenerative, Movement, and Related Disorders	459
26.1		ner's Disease	459
	26.1.1	The Alpha Rhythm	459
	26.1.2	Slowing	460
	26.1.3	Sporadic Epileptiform Discharges	460
	26.1.4	Periodic Discharges	461
	26.1.5	Other Abnormalities	461
	26.1.6	The Electroencephalogram as a Differential Diagnostic Test	461
26.2	Pick's I	1 0	462

XX CONTENTS

26.3	Frontotemporal Lobar Degeneration	463			
26.4	Parkinson's Disease				
26.5	Diffuse Lewy Body Disease				
26.6	Progressive Supranuclear Palsy				
26.7	Corticobasal Ganglionic Degeneration	466			
26.8	Huntington's Disease	466			
26.9	Choreoacanthocytosis	467			
26.10	Wilson's Disease	468			
26.11	Spinocerebellar Ataxia	469			
26.12	Tourette's Syndrome	469			
26.13	Myotonic Muscular Dystrophy	470			
26.14	Metachromatic Leukodystrophy	471			
	Adrenoleukodystrophy	471			
	References	472			
Char	oter 27: Vascular Dementia	479			
27.1	Multi-Infarct Dementia	479			
	Lacunar Dementia	480			
27.3	Binswanger's Disease	480			
	References	481			
Char	oter 28: Developmental, Neurocutaneous, and Other Disorders				
Chup	<u> </u>	483			
201	Typically Presenting in Childhood or Adolescence				
28.1	Focal Cortical Dysplasia and Subependymal Heterotopia 28.1.1 Focal Cortical Dysplasia	483 483			
	5 1	486			
28.2	28.1.2 Subependymal Heterotopia Gelastic Seizures	487			
20.2		487			
	28.2.1 Gelastic Seizures and Hypothalamic Hamartomas28.2.2 Gelastic Seizures and Cortical Foci				
28.3	Tuberous Sclerosis	489 489			
28.4	Sturge–Weber Syndrome	489			
28.5		490			
28.6	von Recklinghausen's Disease (Neurofibromatosis Type 1) Down's Syndrome	490			
28.7	Klinefelter's Syndrome	492			
28.8	Fraqile X Syndrome	493			
28.9	Velocardiofacial Syndrome	494			
28.10	· ·	494			
28.11	Prader–Willi Syndrome	495			
	Fetal Alcohol Syndrome Ratr's Syndrome	493			
	Rett's Syndrome	499			
	Pervasive Developmental Disorders	499 501			
	Attention-Deficit/Hyperactivity Disorder	501			
	Developmental Dysphasia				
40.16	Developmental Dyslexia	504 504			
	References	504			

Contents xxi

Chap	oter 29: Autoimmune and Demyelinating Diseases	513
29.1	Autoimmune Encephalitis	513
	29.1.1 Anti-Hu	513
	29.1.2 Anti-Ma2	514
	29.1.3 Anti-GABA	514
	29.1.4 Anti-AMPA	514
	29.1.5 Anti-LGI1	514
	29.1.6 Anti-NMDAR	516
29.2	Hashimoto's Encephalopathy	518
29.3	Systemic Lupus Erythematosus	518
29.4	Sjogren's Syndrome	519
29.5	Neurosarcoidosis	520
29.6	Sydenham's Chorea	520
29.7	Delayed Post-hypoxic/Ischemic Leukoencephalopathy	521
29.8	Acute Disseminated Encephalomyelitis	521
29.9	Multiple Sclerosis	522
29.10	Osmotic Demyelination Syndrome	523
	References	524
Chap	oter 30: Vascular Disorders	531
30.1	Granulomatous Angiitis of the Central Nervous System	531
30.2	Polyarteritis Nodosa	531
30.3	Susac's Syndrome	532
30.4	Behçet's Syndrome	532
30.5	Antiphospholipid Syndrome	533
30.6	Thrombotic Thrombocytopenic Purpura	534
30.7	Cerebral Fat Embolism	534
30.8	Radionecrosis	535
30.9	Posterior Reversible Encephalopathy Syndrome	536
30.10	CADASIL	536
30.11	MELAS	537
	References	538
Chap	oter 31: Prion Diseases	543
31.1	Creutzfeldt–Jakob Disease	543
31.2	New-Variant Creutzfeldt–Jakob Disease	550
31.3	Gerstmann–Straussler–Scheinker Disease	550
31.4	Fatal Familial Insomnia	551
	References	551
Char	oter 32: Hydrocephalus	557
	References	558
Char	oter 33: Brain Tumors	559
33.1	Cerebral Lobar Tumors	559
	Diencephalic Tumors	561

XXII CONTENTS

33.3	Tumors of the Posterior Fossa References	561 561
Chap 34.1 34.2 34.3 34.4	Transient Global Amnesia Transient Epileptic Amnesia Alcoholic Blackouts Korsakoff's Syndrome References	565 565 566 567 568 569
Chap	oter 35: Stuporous Catatonia References	571 572
Chap 36.1 36.2 36.3 36.4		573 574 575 576 577 578
Chap 37.1 37.2 37.3 37.4 37.5 37.6 37.7	Schizophrenia Bipolar Disorder Major Depressive Disorder Post-traumatic Stress Disorder Panic Disorder Obsessive—Compulsive Disorder Dissociative Disorders 37.7.1 Depersonalization Disorder 37.7.2 Dissociative Amnesia 37.7.3 Dissociative Fugue 37.7.4 Dissociative Identity Disorder Borderline and Antisocial Personality Disorder 37.8.1 Borderline Personality Disorder 37.8.2 Antisocial Personality Disorder	579 579 580 581 582 583 584 585 586 587 587 588 590 590
37.9 37.10	Pyromania	592 592 592 593 593
Chap 38.1 38.2 38.3 38.4	Somnambulism REM Sleep Behavior Disorder Pavor Nocturnus Nightmares	597 597 599 600 601

Contents	XX111
Comens	XXIII

38.5	Confusional Arousal		
38.6	Narcolepsy		
38.7	Sleep A	apnea .	602
38.8	Pickwi	ckian Syndrome (Obesity Hypoventilation Syndrome)	603
38.9	Kleine–Levin Syndrome		603
38.10	C .		
	Referen	nces	604
Chap	ter 39:	Transcranial Stimulation	609
39.1	Electro	convulsive Therapy	609
	39.1.1	The Ictal EEG	610
		39.1.1.1 The Ictal EEG on a Multiple-Channel Recording	610
		39.1.1.2 The Ictal EEG on a One- or Two-Channel Recording	612
	39.1.2	Relationship of the Duration of the Motor Seizure and of	
		the Electroencephalographic Seizure	613
	39.1.3	Postictal Slowing	613
	39.1.4	Nonconvulsive Status Epilepticus Following an ECT-Induced	
		Tonic-Clonic Seizure	614
39.2	Repetit	tive Transcranial Magnetic Stimulation	614
39.3	Transc	ranial Direct Current Stimulation	615
	Referen	nces	615
Index			619

Chapter 1 Arrays and Montages

1.1 Arrays

As the field of electroencephalography developed, the need for a standardized array of electrodes became increasingly apparent, resulting, in 1958, in the adoption of the International 10–20 System. Subsequently, the need for more closely spaced electrodes to allow for more precise localization led to an expansion of this system into what has come to be known as the Modified Combinatorial 10–10 System.

One, very important, failing of the International 10–20 System has been its inability to capture epileptiform discharges arising from seizure foci in the medial and anterior aspects of the temporal lobe. Of the multiple attempts to remedy this failing, two have found their way into clinical practice: first, utilizing a portion of the 10–10 System, namely, the inferior temporal chain; and, second, using one of the supplemental noninvasive electrodes, namely, the "true" anterior temporal electrode. Both of these are discussed at the end of this chapter.

1.1.1 International 10–20 System

The International 10–20 System was first described by Jasper in 1958 (Jasper 1958) and has become a worldwide standard. In developing this system, Jasper decided that "[p]ositions of electrodes should be determined by measurement from standard landmarks on the skull. . ." and that these "[m]easurements should be proportional to skull size and shape, insofar as

possible." The standard landmarks Jasper chose were the nasion, the inion, and the preauricular points, and to maintain proportionality to the individual skull size, Jasper, rather than using predetermined distances in centimeters, decided on using percentages of the distances between these points on the individual patient's head.

The first line to be drawn lies in the midline and extends from the nasion to the inion. The first point on this line, according to Jasper (1958), is the Fp midline point, and this point "is 10 per cent of the nasion-inion distance above the nasion; the second point (F) is 20 per cent of this distance back from the point Fp, and so on in 20% steps back for the Central, Parietal, and Occipital points (hence the name 10-20 system)."

The next line to be drawn is "based upon the Central coronal plane. The distance is first measured from the left to right preauricular points . . ." and Jasper (1958) cautions that we "[b]e sure the tape is passing through the predetermined Central point at the vertex when making this measurement. Ten per cent of this distance is then taken for the Temporal point up from the pre-auricular point on either side. The Central points are then marked 20 per cent of the distance above the temporal points. . ."

The next line to be drawn is an "A-P line of electrodes over the temporal lobe, frontal to occipital, (and this) is determined by measuring the distance between the Fp mid line point . . . through the T position of the Central line, and back to the mid-occipital point. The Fp electrode position is then marked 10 per cent of this distance from the mid-line in front, and the Occipital electrode position 10 per cent of the distance from the mid-line in back. The inferior Frontal and posterior Temporal positions then fall 20 per cent of the distance from the Fp and O electrodes, respectively along this line" (Jasper 1958).

"The remaining mid-Frontal (F3 and F4) and mid-Parietal (P3 and P4) electrodes are then placed along the Frontal and Parietal coronal lines respectively, equidistant between the mid-line and temporal line of electrodes on either side" (Jasper 1958).

In explaining the rationale for the names of the various electrodes, Jasper (1958) commented that "[t]raditional anatomical terms have been employed to designate electrode positions over the various lobes of the brain, with the exception of the Central region which is, strictly speaking, partly frontal and partly parietal."

He goes on to note that "[i]n order to differentiate between homologous portions over left and right hemispheres it was decided to use even numbers as subscripts for the right hemisphere, and odd numbers for the left hemisphere . . . Electrodes at the mid-line in Frontal, Central and Parietal regions were originally designated F_0 , C_0 and P_0 but this led to some confusion since P_0 ,

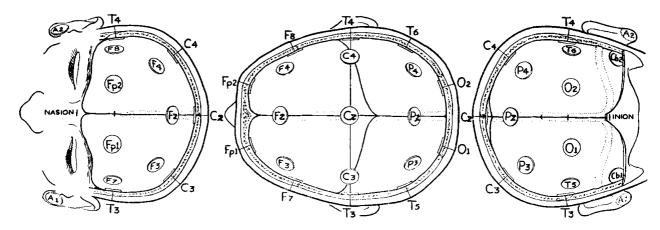


Figure 1.1 The complete International 10–20 System. SOURCE: Jasper (1958)/with permission of Elsevier.

for example, might be interpreted as parieto-occipital. Consequently the midline positions have been changed to F_z , C_z and P_z ($_z$ for zero!)." The complete system of placements is shown in Figure 1.1 and schematically in Figure 1.2.

The full names of each of Jasper's 21 alphanumeric designations are given in Table 1.1. Subsequently, however, two alternative names have gained currency. The first alternative is trivial: "auricle" has been replaced by "ear." The second, however, is far more significant. In the same year that Jasper published his study, Abraham and Ajmone Marsan (1958) referred to F7 and F8 not as "inferior frontal," but as "anterior temporal," and this designation has stuck, despite the fact, as discussed below, that F7 and F8 do not overly the temporal lobe at all.

1.1.2 Anatomical Position of Electrodes of the International 10–20 System

Jasper (1958) stated that "[a]natomical studies should be carried out to determine the cortical areas most likely to be found beneath each of the standard electrode positions," and in his paper he briefly described two such studies.

"Two methods were employed: (1) metal clips placed along the Central and Sylvian fissures at operation were then used to identify these fissures in X-ray studies of the skull after the EEG electrodes had been applied, and (2) electrode positions were carefully marked on the heads of cadavers, drill holes placed through the skull, and the cortex marked with India ink in each position before removing the brain for examination. Brains with gross lesions or focal atrophy were excluded."

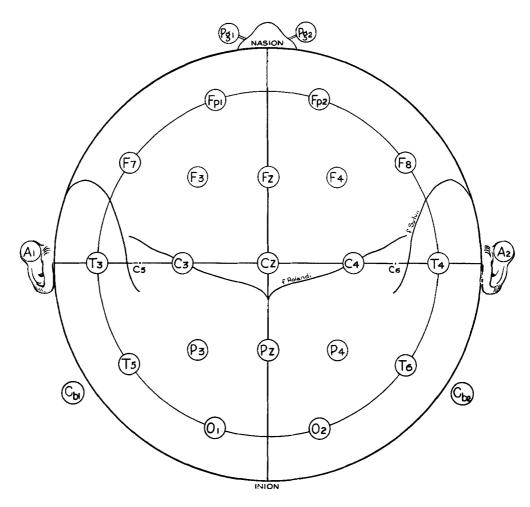


Figure 1.2 The complete International 10–20 System in schematic form. SOURCE: Jasper (1958)/with permission of Elsevier.

 Table 1.1
 Electrode names in the International 10–20 System

		Fp1 left frontopolar		Fp2 right frontopolar		
	F7 Left inferior frontal ^a	F3 Left frontal	F _z Frontal midline	F4 Right frontal	F8 Right inferior frontal ^a	
A1 Left auricle	T3 Left mid-temporal	C3 Left central	C _z Central midline	C4 Right central	T4 Right mid-temporal	A2 Right auricle
	T5 Left posterior temporal	P3 Left parietal	P _z Parietal midline	P4 Right parietal	T6 Right posterior temporal	
	1	O1 Left occipital		O2 Right occipital	1	

 $^{^{\}rm a}$ F7 and F8 are also known as "anterior temporal" electrodes.