Sixth Edition

Primer of DIAGNOSTIC IMAGING

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To all of the radiologists whose knowledge, research, and wisdom contributed to this book

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Preface

The sixth edition of *Primer of Diagnostic Imaging* will continue to function as a integral learning source for residents and fellows, as well as a refresher text for faculty and practicing physicians. This book serves not only as the core curriculum for our fast-evolving specialty but also as a current reference text for practitioners. We have sought to update and include material to reflect current imaging algorithms, while at the same time retaining older material that is pertinent to a variety of practice patterns. Since the preparation of the last edition, we have incorporated further advances in the various imaging modalities and have to the best of our knowledge corrected any

inaccurate information. Furthermore, by continuing to have our successful graduates evaluate the content, we have ensured that the subject matter covers the information assessed by the current American Board of Radiology examination. We are particularly indebted to our reviewers for their feedback. It is our hope that *Primer of Diagnostic Imaging* will continue to serve the next generation of radiologists, helping them navigate the stream of continuously emerging new information.

Mukesh G. Harisinghani John W. Chen Ralph Weissleder

Abbreviations

2D	two-dimensional	ANCA	antineutrophil cytoplasmic antibody
3D	three-dimensional	AP	anteroposterior
4D	four-dimensional	APKD	adult polycystic kidney disease
5-HIAA	5-hydroxyindoleacetic acid	APS	anterior pararenal space
) IIIIII) nydroxymdoleacette acid	APUD	amine precursor uptake and
AA	aortic arch	мов	decarboxylation
A–a	alveolar-arterial Po ₂ difference	APW	absolute percentage washout
AAA	abdominal aortic aneurysm	AR	autosomal recessive
ABC	aneurysmal bone cyst	ARDS	acute respiratory distress syndrome
ABPA	allergic bronchopulmonary aspergillosis	ARF	anterior renal fascia
ABS	amniotic band syndrome	ARKD	autosomal recessive kidney disease
AC	abdominal circumference;	ARPCKD	autosomal recessive polycystic kidney
	acromioclavicular; alternating current		disease
ACA	anterior cerebral artery	ARVD	arrhythmogenic right ventricular dysplasia
ACC	agenesis of corpus callosum; adenoid	AS	ankylosing spondylitis; aortic stenosis
	cystic carcinoma	ASA	anterior spinal artery
ACL	anterior cruciate ligament	ASD	airspace disease; atrial septal defect
ACLS	advanced cardiac life support	a-Se	amorphous selenium
ACOM	anterior communicating (artery)	ASNR	American Society of Neuroradiology
ACR	American College of Radiology	ATA	American Thyroid Association Guidelines
ACS	anterior cervical space	ATN	acute tubular necrosis
ACT	activated clotting time	ATP	adenosine triphosphate
ACTH	adrenocorticotropic hormone	\mathbf{AV}	arteriovenous; atrioventricular
AD	abdominal diameter; autosomal dominant;	AVF	arteriovenous fistula
	average distance	AVM	arteriovenous malformation
ADC	apparent diffusion coefficient	AVN	avascular necrosis
ADEM	acute disseminated encephalomyelitis	AZV	azygos vein
ADH	antidiuretic hormone	D.	1 .
ADPKD	adult dominant polycystic kidney disease	Ва	barium
AFB	aortofemoral bypass amniotic fluid index	BBOT	2,5-bis(5-ter-butyl-2-benzoxasolyl)
AFI AFL	air-fluid level	BI	thiophene Billroth I
AFP	alpha-fetoprotein	BII	Billroth II
AFV	amniotic fluid volume	BAC	bronchoalveolar carcinoma
Ag	silver	BAI	basion-axial interval
AHA	American Heart Association	BBB	blood-brain barrier
AHD	acquired heart disease	BBBD	blood-brain barrier disruption
AI	aortic insufficiency; aluminum	BCDDP	Breast Cancer Detection Demonstration
AICA	anterior inferior cerebellar artery	20221	Program
AICD	automatic implantable cardioverter-	BCNU	1,3-bis(2-chloroethyl)-1-nitrosourea
	defibrillator	BCP	basic calcium phosphate
AICV	anterior intercostal vein	BCS	Budd-Chiari syndrome
AIDS	acquired immunodeficiency syndrome	BE	barium enema; binding energy
AIP	acute interstitial pneumonia	BEIR	Biological Effects of Ionizing Radiation
AJCC	American Joint Committee on Cancer	BF	Bucky factor
ALD	adrenoleukodystrophy	BFM	bronchopulmonary foregut malformation
ALS	amyotrophic lateral sclerosis	BGO	bismuth germanium oxide
AM	abnormal motility	β-НСG	beta-human chorionic gonadotropin
AMI	acute myocardial infarction	BIP	bronchiolitis obliterans interstitial
AML	angiomyolipoma; anterior mitral leaflet		pneumonitis
amu	atomic mass unit	BI-RADS	Breast Imaging Reporting and Data
ANC	acute necrotic collections		System

BLC	benign lymphoepithelial cyst	CLC	corpus luteum cyst
BOOP	bronchiolitis obliterans and organizing	CMC	carpometacarpal (joint)
	pneumonia	CMD	corticomedullary differentiation
BP	blood pressure	CMS	Centers for Medicare and Medicaid
BPD	biparietal diameter; bronchopulmonary		Services
	dysplasia	CMV	cytomegalovirus
BPE	background parenchymal enhancement	CN	cranial nerve
BPF	bronchopleural fistula	CNS	central nervous system
BPH BPM	benign prostatic hyperplasia	CO	carbon monoxide cobalt
BPOP	beats per minute	CO COP	
БРОР	bizarre parosteal osteochondromatous proliferation	COPD	cryptogenic organizing pneumonia chronic obstructive pulmonary disease
BPP	biophysical profile	CP	cerebellopontine; choroid plexus
Bq	becquerel	CPA	cerebellopontine angle
BRBPR	bright red blood per rectum	CPAM	congenital pulmonary airway
BRTO	balloon-occluded retrograde transvenous		malformation
	obliteration	CPAP	continuous positive airway pressure
BS	buccal space	CPC	choroid plexus cysts
		CPM	central pontine myelinosis
C	caudate; coulomb	cpm	counts per minute
CA	carcinoma	CPPD	calcium pyrophosphate dihydrate
CABG	coronary artery bypass graft	cps	counts per second
CAD	coronary artery disease	CR	computed radiography
CAH CAPD	chronic active hepatitis	Cr	creatine/phosphocreatine
CAPD CaWO ₄	chronic ambulatory peritoneal dialysis calcium tungstate	CREST	calcinosis, Raynaud's, esophageal dysmotility, sclerodactyly, telangiectasia
CBD	common bile duct		(syndrome)
CBF	cerebral blood flow	CRL	crown-rump length
CBV	cerebral blood volume	CRMO	chronic recurrent multifocal osteomyelitis
CC	corneal clouding; craniocaudad;	CRPS	complex regional pain syndrome
	craniocaudal	C-section	cesarean section
CCA	common carotid artery	C-spine	cervical spine
CCAM	congenital cystic adenoid malformation	CS	central sulcus; carotid space
CCF	carotid-cavernous sinus fistula	Cs	cesium
CCK	cholecystokinin	CSF	cerebrospinal fluid
CCU	coronary care unit	CsI	cesium iodide
CD CD4	cystic duct	CSP CT	corrected sinusoidal pressure
CD4 CDH	cluster designation 4 antigen congenital diaphragmatic hernia;	CTA	computed tomography computed tomographic angiography
CDII	congenital dislocation of the hip	СТАР	computed tomographic angiography
CDI	color Doppler imaging	Onn	portography
cd/m ²	candela per square meter	CTC	computed tomography colonography
CEA	carcinoembryonic antigen	CTDI	CT dose index
CECT	contrast-enhanced computed tomography	CTP	computed tomographic perfusion
CF	cystic fibrosis	CTPA	CT pulmonary angiography
CFA	common femoral artery; cryptogenic	CTV	computed tomographic venography
	fibrosing alveolitis	CU	clinical unit
CHA	calcium hydroxyapatite; common hepatic	Cu	copper
	artery	CVA	cerebrovascular accident
CHAOS	congenital high airway obstruction	CVR	CPAM volume ratio
CHD	common hepatic duct; congenital heart disease	CVS	calcium volume score; chorionic villus
CHF	congestive heart failure	CWP	sampling coal workers' pneumoconiosis
Cho	choline	CXR	chest radiograph
CHP	chronic hypersensitivity pneumonitis	0211	effect factograph
CI	cardiothoracic index	D	dilatation
Ci	Curie	DA	double arch
CIDP	chronic inflammatory demyelinating	DAI	diffuse axonal injury
	polyneuropathy	dB	decibel
CIN	contrast-induced nephropathy	DC	direct current
CJD	Creutzfeldt-Jakob disease	DCIS	ductal carcinoma in situ

DDH	developmental dysplasia of the hip	ESR	erythrocyte sedimentation rate
DDx	differential diagnosis	ESV	end-systolic volume
DE-CMRI	delayed-enhancement cardiac magnetic	ESWL	extracorporeal shock wave lithotripsy
	resonance imaging	ET	endotracheal tube
DECT	dual-energy CT	ETL	echo train length
DES	diethylstilbestrol; diffuse esophageal	eV	electron volt
	spasm	EVLT	endovenous laser treatment
DFTN	diffuse fold thickening with fine	EXP	exponential
	nodularity		
DIC	disseminated intravascular coagulation	FAPS	familial adenomatous polyposis syndrome
DIP	desquamative interstitial pneumonitis;	FBP	filtered back projection
	distal interphalangeal (joint)	FCD	fibrous cortical defect
DIPS	direct intrahepatic portocaval shunt	FD	filling defect
DISH	diffuse idiopathic skeletal hyperostosis	FDA	U.S. Food and Drug Administration
DISI	dorsal intercalated segment instability	FDG	fluorodeoxyglucose
DJD	degenerative joint disease	Fe	iron
DLBCL	diffuse large B-cell lymphoma	FESS	functional endoscopic sinus surgery
DLP	dose length product	FEV	forced expiratory volume
DM	diabetes mellitus	FFDM	full-field digital mammography
DMSA	dimercaptosuccinic acid	FGT	fibroglandular tissue
DNA	deoxyribonucleic acid	FIGO	International Federation of Gynecology
DNET	dysembryoplastic neuroepithelial tumor		and Obstetrics
DORV	double-outlet right ventricle	FL	femur length
DPS	dorsal pleural sinus	FLAIR	fluid-attenuated inversion recovery
DR	digital radiography	FMC	focal myometrial contraction
DRE	digital rectal examination	FMD	fibromuscular dysplasia
DSA	digital subtraction angiography	fMRI	functional magnetic resonance imaging
D-TGA	complete transposition of great arteries	FN	false negative
DTPA	diethylenetriaminepentaacetic acid	FNA	fine-needle aspiration
DU	deep ulcer(s)	FNH	focal nodular hyperplasia
DVT	deep vein thrombosis	FOD	focal spot–object distance
DW	Dandy-Walker	FOV	field of view
DWI	diffusion-weighted imaging	FP	false positive
T.		FRC	functional residual capacity
E	exposure	FS FSE	focal spot; fractional shortening
E _{aver}	average electron energy maximum energy	FSH	fast spin echo follicle-stimulating hormone
E _{max} EA	esophageal atresia	FTA-ABS	fluorescent treponemal antibody
EAC	external auditory canal	TTA-AD3	absorption (test)
EBV	Epstein-Barr virus	FUO	fever of unknown origin
ECA	external carotid artery	FWHM	full width at half maximum
ECD	endocardial cushion defect; ethyl	1 WILWI	ran widan at han maximam
202	cysteinate dimer	g	gram
ECF	extracellular fluid	GA	gestational age
ECG	electrocardiogram	Ga	gallium
ECMO	extracorporeal membrane oxygenation	GB	gallbladder
EDH	epidural hematoma	GBCA	Gd-based contrast agents
EDV	end-diastolic volume	GBM	glioblastoma multiforme
EEG	electroencephalogram	GBPS	gated blood pool study
EF	ejection fraction	G-CSF	granulocyte colony-stimulating factor
EFW	estimated fetal weight	GCT	giant cell tumor
EG	eosinophilic granuloma	Gd	gadolinium
EGA	estimated gestational age	GDA	gastroduodenal artery
EMS	endometrial stripe	GE	gastroesophageal
ENT	ear, nose, throat	GEJ	gastroesophageal junction
EPA	Environmental Protection Agency	GFR	glomerular filtration rate
EPO	erythropoietin	GH	growth hormone
ERCP	endoscopic retrograde	GI	gastrointestinal
	cholangiopancreatography	GIP	giant cell interstitial pneumonia
ERPF	effective renal plasma flow	GIST	gastrointestinal stromal tumor
ERV	expiratory reserve volume	GLAD	glenolabral degenerative joint disease

glut	glucose transporter	ICV	internal cerebral vein
GLUT1	glucose transporter 1	ID	information density; inner diameter
GM	gray matter	IDA	iminodiacetic acid
GnRH	gonadotropin-releasing hormone	IDH	isocitrate dehydrogenase
GRE	gradient-recalled echo	IG	immunoglobulin
GSD	genetically significant dose	IgA	immunoglobulin A
GSO	gadolinium oxyorthosilicate	IgE	immunoglobulin E
GTD	gestational trophoblastic disease	IgG	immunoglobulin G
GU	genitourinary	IHF	immune hydrops fetalis
GVH	graft-versus-host (disease)	IHN	infrahyoid
GWM	gray-white matter	IHSS	idiopathic hypertrophic subaortic
Gy	gray (unit of radiation)	1137	stenosis
Н	hoight	IJV IL-2	internal jugular vein interleukin-2
HA	height hepatic artery	IL-2 ILO	International Labor Organization
HAZV	hemiazygos vein	ILT	inferolateral trunk
Hb	hemoglobin	IM	intramuscular
HbAS	sickle cell trait	IMA	inferior mesenteric artery
HbSS	sickle cell disease	IMV	internal mammary vein
HC	head circumference	In	indium
HCC	hepatocellular carcinoma	INF	inferior
HCG	human chorionic gonadotropin	INH	isoniazid
HD	Hurter and Driffield (curve)	INR	international normalized ratio
HGH	human growth hormone	INSS	International Neuroblastoma Staging
HIDA	hepatic iminodiacetic acid derivative		System
HIP	health insurance plan	IPF	idiopathic pulmonary fibrosis
HIV	human immunodeficiency virus	IPH	idiopathic pulmonary hemorrhage
HLA	human leukocyte antigen	IPKD	infantile polycystic kidney disease
HLHS	hypoplastic left heart syndrome	IPMN	intraductal papillary mucinous neoplasm
HMD	hyaline membrane disease	IPMT	intraductal papillary mucinous tumor
HMDP	hydroxymethylene diphosphonate	IQ	intelligence quotient
HMPAO	hexamethylpropyleneamine oxime	IRV	inspiratory reserve volume
HOCA HPF	high-osmolar contrast agent	ITB IUD	iliotibial band intrauterine device
HPO	high-power field hypertrophic pulmonary osteoarthropathy	IUGR	intrauterine device
HPS	hypertrophic pyloric stenosis	IUP	intrauterine pregnancy
HPT	hyperparathyroidism	IV	intravenous
HPV	human papilloma virus	IVC	inferior vena cava
HQ	high-quality	IVDA	intravenous drug abuse(r)
HR	heart rate	IVP	intravenous pyelogram
hr	hour	IVS	interventricular septum
HRCT	high-resolution computed tomography		•
HS	hepatosplenomegaly; high-speed	JRA	juvenile rheumatoid arthritis
HSA	human serum albumin		
HSG	hysterosalpingogram	keV	kiloelectron volt
HSV	herpes simplex virus	Kr	krypton
HTLV	human T-cell lymphotrophic virus	KS	Kaposi sarcoma
HTN	hypertension	KUB	kidney, urethra, bladder
HU HVA	heat unit; Hounsfield unit homovanillic acid	kV kVo	kilovoltage
HVL	half-value layer	kVp	peak kilovoltage
IIVL	nan-varde layer	L	left; length
IA	intraarterial	LA	left atrium
IAA	interruption of aortic arch	La	lanthanum
IABP	intraaortic balloon pump	LAD	left anterior descending (artery)
IAC	internal auditory canal	LAE	left atrial enlargement
IBD	inflammatory bowel disease	LAM	lymphangioleiomyomatosis
ICA	internal carotid artery	LAO	left anterior oblique
ICRP	International Commission on Radiological	LATS	long-acting thyroid-stimulating (factor)
	Protection	LBBB	left bundle branch block
ICU	intensive care unit	LBWC	limb/body wall complex

LCA	left carotid artery; left coronary artery	MA	meconium aspiration; mesenteric
LCF	lateroconal fascia		adenopathy
LCIS	lobular carcinoma in situ	mA	milliampere
LCL	lateral collateral ligament	MAA	macroaggregated albumin
LCNEC	large cell neuroendocrine carcinoma	MAb	monoclonal antibody
LCP	Legg-Calvé-Perthes (disease)	MAG	methyl-acetyl-gly
LCx	left circumflex (artery)	MAG3	methyl-acetyl-gly-gly
LD	lymphocyte depleted (Hodgkin	MAI	Mycobacterium avium-intracellulare
	lymphoma)	MALT	mucosa-associated lymphoid tissue
LD_{50}	lethal dose, 50%	MAOI	monoamine oxidase inhibitor
LDH	lactate dehydrogenase	MAP	maximum-a-posteriori
LEJV	left external jugular vein	mAs	milliampere second
LES	lower esophageal sphincter	MBq	megabecquerel
LET	linear energy transfer	MC	mixed cellularity (Hodgkin lymphoma)
LFT	liver function test	MCA	middle cerebral artery
LGA	large for gestational age; left gastric	MCD	medullary cystic disease
LON	artery	MCDK	multicystic dysplastic kidney
LH	luteinizing hormone	mCi	millicurie
LHA	left hepatic artery	μCi	microcurie
LHD	left hepatic duct	MCL	medial collateral ligament
	left internal jugular vein	MCP	metacarpophalangeal
LIJV			mixed connective tissue disease
LIMA	left internal mammary artery	MCTD	middle cerebral vein
LIMV	left internal mammary vein	MCV	
LIP	lymphocytic interstitial pneumonia	MD	monochorionic, diamniotic (twins)
LIQ	low intelligence quotient	MDA	metaphyseal-diaphyseal angle
LL	lower lobe	MDCT	multidetector computed tomography
LLI	left lateral inferior	MDP	methylene diphosphonate
LLL	left lower lobe	MELAS	mitochondrial myopathy, encephalopathy,
LLS	left lateral superior		lactic acidosis, stroke like episodes
LM	lateromedial	MEN	(syndrome)
LMB	left mainstem bronchus	MEN	multiple endocrine neoplasia
LMI	left medial inferior	MERRF	myoclonic epilepsy with ragged red fibers
LMP	last menstrual period	3.6-37	(syndrome)
LMS	left medial superior	MeV	megaelectron volt
LN	lymph node	MFH	malignant fibrous histiocytoma
LOCA	low-osmolar contrast agent	MGH	Massachusetts General Hospital
LP	lymphocyte predominant (Hodgkin	MHz	megahertz
TDA	lymphoma)	MI	myocardial infarction
LPA	left pulmonary artery	MIBG	metaiodobenzylguanidine
LPM	anterolateral papillary muscle	MIBI	methoxyisobutyl isonitrile
LPO	left posterior oblique	MIP	maximum-intensity projection
LPV	left portal vein	ML	mediolateral
LR	likelihood ratio	MLCN	multilocular cystic nephroma
L-R shunt LSA	left-to-right shunt left subclavian artery	MLD	maximum transverse diameter to the left from midline
	·	MIEM	
LSCV	left subclavian vein	MLEM	maximum likelihood expectation maximization
LSICV	left superior intercostal vein	MLO	
LSMFT	liposclerosing myxofibrous tumor		mediolateral oblique
LSO	lutetium oxyorthosilicate	MM	monoamniotic, monochorionic (twins)
L-TGA	corrected transposition of great arteries	MNG	multinodular goiter
LTV	lateral thoracic vein	Мо	molybdenum
LUCL	lateral ulnar collateral ligament	mo MOCE	month
LUL	left upper lobe	MOCE	multiple osteocartilaginous exostoses
LUQ	left upper quadrant	MOM	multiples of median
LUS	lower uterine segment	mOsm	milliosmole
LV	left ventricle	MPA	main pulmonary artery
LVA	left vertebral artery	MPD	maximum permissible dose
LVE	left ventricular enlargement	MPM	posteromedial papillary muscle
LVEF	left ventricular ejection fraction	MPV MP	main portal vein
LVH	left ventricular hypertrophy	MR mP	magnetic resonance
LYSO	lutetium ytrium oxyorthosilicate	mR	milliroentgen

MRCP magnetic resonance anjography mRCP magnetic resonance cholanglopancreatography magnetic resonance choragophy maximum transverse diameter to the right from midline magnetic resonance enterography magnetic resonance spectroscopy on all cavity on all cavity magnetic resonance spectroscopy on the detector distance on phase control distance of the property				
MRD maximum transverse diameter to the right from midline magnetic resonance enterography MRI magnetic resonance imaging OCH Oriental cholangiohepatitis oral contraceptive; optical colonoscopy; oral cavity optical density; outer diameter object detector distance osmotic dempetable with magnetic resonance imaging OCH Oriental cholangiohepatitis once daily; optical density; outer diameter object detector distance osmotic dempetial optical density; outer diameter object detector distance osmotic dempetial object detector distance osmotic dempetial optical density; outer diameter object detector distance osmotic dempetial object detector distance osmotic distance osmotic distance on object detector distance object detector distance on object detector distance o	MRA	magnetic resonance angiography	NTD	neural tube defect
MRI magnetic resonance marging OCH oriental cholangiohepatitis on cal contraceptive; optical colonoscopy; oral cavity oral cav	MRCP		NTMB	nontuberculous mycobacteria
MRE magnetic resonance enterography MRI magnetic resonance inaging MRS magnetic resonance imaging MRS magnetic resonance spectroscopy MRSA methicillin-resistant Staphylococcus auruus MRV magnetic resonance venography MS methicillin-resistant Staphylococcus auruus MRV magnetic resonance venography MSA methicillin-resistant Staphylococcus auruus MRV magnetic resonance venography MSA multipse sclerosis MSA multip				
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MRSA magnetic resonance spectroscopy MRSA methicillin-resistant Staphylosoccus aureus MRV magnetic resonance venography MS multiple selerosis MSA multisystem atrophy MSAPP maternal serum alpha-fetoprotein MSD mean sac diameter MSD socionation NSD posteroanterior, pulmonary artery perincural arachnolosi pulmonary artery posterior creebral artery posterior communicating (artery) MSD posterior creek sale posterior cor				•
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MSDP mean sac diameter MSD mean sac diameter			OEIS	
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NT nuchal translucency PICA posterior inferior cerebellar artery				
	NT	nuchal translucency	PICA	posterior interior cerebellar artery

PICV	posterior intercostal vein	PUD	peptic ulcer disease
PID	pelvic inflammatory disease	PUL	percutaneous ureterolithotomy
PIE	pulmonary infiltrates with eosinophilia;	PUV	posterior urethral valve
	pulmonary interstitial emphysema	PV	portal vein
PIOPED	prospective investigation of pulmonary	PVA	polyvinyl alcohol
	embolus detection	PVC	premature ventricular contraction;
PIP	postinflammatory polyp; proximal		polyvinyl chloride
	interphalangeal (joint)	PVH	pulmonary venous hypertension
PJP	Pneumocystis jiroveci pneumonia	PVNS	pigmented villonodular synovitis
PKU	phenylketonuria	PVOD	pulmonary venoocclusive disease
PLN	projected length of needle	PVP	portal venous phase
PLPN	projected length to pull back needle	PVS	perivertebral space
PM	photomultiplier (tube)	PWI	perfusion-weighted imaging
PMC	pseudomembranous colitis	PWMA	periventricular white matter abnormality
PMF	progressive massive fibrosis	PZT	lead zirconium titanate
PMHR PML	predicted maximum heart rate posterior mitral leaflet; progressive	04	quality accurance
FINIL	multifocal leukoencephalopathy	QA qid	quality assurance four times daily
PMMA	polymethylmethacrylate	qia	four times daily
PMS	pharyngeal mucosal space	R	range; right; roentgen
PMT	photomultiplier tube	Ra	radium
PNET	primitive neuroectodermal tumor	RA	right atrium; rheumatoid arthritis
PO	orally (per os)	RAI	right anterior inferior
Po_2	partial pressure of oxygen	RAIU	radioactive iodine uptake
post.	posterior	RAO	right anterior oblique
PP	parietal peritoneum	RAS	renal artery stenosis; right anterior
PPF	pterygopalatine fossa		superior
PPHN	persistent pulmonary hypertension of the newborn	RB-ILD	respiratory bronchiolitis–associated interstitial lung disease
ppm	parts per million	RBBB	right bundle branch block
PPO	2,5-ciphenyloxazole	RBC	red blood cell(s) (count)
PPS	posterior pararenal space; parapharyngeal	RBE	relative biologic effectiveness
	space	RCA	right carotid artery; right coronary artery
PPV	positive predictive value	RCC	renal cell carcinoma
PRES	posterior reversible encephalopathy	RCV	red cell volume
PRF	syndrome pulse repetition frequency; posterior renal	RCVS	reversible cerebral vasoconstriction syndrome
	fascia	RDS	respiratory distress syndrome
PRL	prolactin	REJV	right external jugular vein
PROM	premature rupture of membranes	RES	reticuloendothelial system
PRS	perinephric space	RF	radiofrequency; rheumatoid factor
PS	parotid space	RGA	right gastric artery
PSA	prostate-specific antigen	Rh	rhesus (factor)
PSE	partial splenic embolization	RHA	right hepatic artery
PSMA	prostate-specific membrane antigen	RHD	right hepatic duct
PSP	progressive supranuclear palsy	RI	resistive index
PSPMT	pulse spray pharmacomechanical	RIJV	right internal jugular vein right internal mammary artery
PSS	thrombolysis progressive systemic sclerosis	RIMA RIMV	right internal mammary vein
PT	prothrombin time	RIND	reversible ischemic neurologic deficit
PTA	percutaneous transluminal angioplasty	R-L shunt	right-to-left shunt
PTCA	percutaneous transluminal coronary	RLL	right lower lobe
	angioplasty	RLQ	right lower quadrant
PTD	posttransplantation lymphoproliferative	RMB	right mainstem bronchus
	disorder	RML	right middle lobe
PTFE	polytetrafluoroethylene	RMS	retromesenteric anterior interfascial space
PTH	parathormone	Rn	radon
PTLD	posttransplantation lymphoproliferative	RNA	ribonucleic acid
	disorder	ROI	range of interest
PTT	partial thromboplastin time	rPA	ratio of pulmonary artery diameter to
PTU	propylthiouracil		aortic diameter

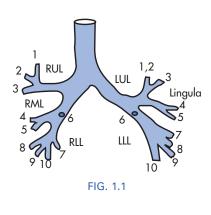
RPI	right posterior inferior	SPIO	superparamagnetic iron oxide
RPN	renal papillary necrosis	SSFP	steady-state free precession
RPO	right posterior oblique	SSFSE	single-shot fast spin echo
RPS	right posterior superior; retropharyngeal	ST	ST complex on ECG
	space	STIR	short tau inversion recovery
RPV	right portal vein	STT	scaphotrapeziotrapezoid
RPW	relative percentage washout	SU	superficial ulcer(s)
RRS	retrorenal posterior interfascial space	sup.	superior
RSA	right subclavian artery	SUV	standardized uptake value
RSCV	right subclavian vein	SV	seminal vesicle
RSV	respiratory syncytial virus	SVC	superior vena cava
RT	radiotherapy	SWI	susceptibility-weighted imaging
RTA	renal tubular acidosis	_	
r-tPA	recombinant tissue plasminogen activator	T	tesla; thalamus; time
RTV	right thoracic vein	T1W	T1-weighted (images)
RUG	retrograde urethrogram	T2W	T2-weighted (images)
RUL	right upper lobe	T3	triiodothyronine
RUQ	right upper quadrant	T4	thyroxine
RV	reserve volume; right ventricle	T18	trisomy 18
RVA	right vertebral artery	T21	trisomy 21
RVEF	right ventricular ejection fraction	TA	truncus arteriosus
RVH DVT	right ventricular hypertrophy	TAPVC	total anomalous pulmonary venous
RVT	renal vein thrombosis	TAPVR	connection
6	second	TAR	total anomalous pulmonary venous return
s S/P		IAK	thrombocytopenia-absent radius (syndrome)
SA	status post sinoatrial; subclavian artery; specific	TAS	transabdominal ultrasound
<i>51</i> 1	activity	TB	tuberculosis
SAH	subarachnoid hemorrhage	TBI	traumatic brain injury
SB	small bowel	TC	thyroid cartilage
SBFT	small bowel follow-through	Tc	technetium
SBO	small bowel obstruction	TCC	transitional cell cancer
SC	subcutaneous	TCD	transcranial Doppler
SCA	superior cerebellar artery	TD	tolerance dose
SCC	squamous cell carcinoma	TDL	true depth of lesion
SCFE	slipped capital femoral epiphysis	TDLU	terminal duct lobular unit
SCLS	small cell lung cancer	TE	echo time
SD	standard deviation	TEE	transesophageal echocardiography
SDAT	senile dementia, Alzheimer type	TEF	tracheoesophageal fistula
SDH	subdural hematoma	TF	thickened folds; transversalis fascia
SE	spin echo	TFA	tibiofemoral angle
seg.	segment	TFCC	triangular fibrocartilage complex
SFA	superficial femoral artery	TFN	thickened folds with nodularity
SGA	small for gestational age	TGA	transposition of great arteries
SGOT	serum glutamic-oxaloacetic transaminase	TGC	time-gain compensator
SHN	suprahyoid	THR	total hip replacement
SI	sacroiliac; signal intensity	THY	thyroid gland
SIN	salpingitis isthmica nodosa	TI	terminal ileum; thallium; time of inversion
SiO ₂	silicone dioxide	TIA	transient ischemic attack
SK	streptokinase	TiO ₂	titanium dioxide
SL SLAC	sublingual	TIPS	transjugular intrahepatic portosystemic shunt
	scapholunate advanced collapse	TIDADE	
SLE SMA	systemic lupus erythematosus	TIRADS TKR	Thyroid Image Reporting and Data System total knee replacement
SMV	superior mesenteric vein	TLA	totai knee repiacement translumbar approach
Sm	superior mesenteric vein tin	TLC	total lung capacity
SNR	signal-to-noise ratio	TLN	true length of needle
SPECT	single photon emission computed	TLPN	true length to pull back needle
JILOI	tomography	TM	tympanic membrane; time motion
SPEN	solid pseudopapillary epithelial neoplasm	TMC	toxic megacolon
SPGR	spoiled gradient-echo	TMJ	temporomandibular joint
	T. Comments		7

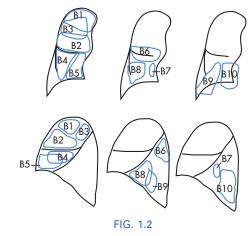
TN	true negative	VC	vital capacity
TNM	tumor-node-metastases	VCUG	voiding cystourethrogram
TOA	tubooyarian abscess	VD	vas deferens
TOF	time of flight	VDRL	Venereal Disease Research Laboratory
TORCH	toxoplasmosis, rubella, cytomegalovirus,	VHL	von Hippel-Lindau (disease)
TORON	herpes simplex virus (syndrome)	VISI	volar intercalated segment instability
TP	true positive	VMA	vanillylmandelic acid
tPA	tissue plasminogen activator	VP	ventriculoperitoneal
TPN	total parenteral nutrition	ŸŹ	ventilation/perfusion
TPO	tracheopathia osteoplastica	VR VR	Virchow-Robin (space)
TR	repetition interval	VRE	vancomycin-resistant enterococcus
TRAM	transverse rectus abdominis	VS	visceral space
1100101	musculocutaneous	VSD	ventricular septal defect
TRAPS	twin reversal arterial perfusion sequence	VUR	vesicoureteral reflux
TRUS	transrectal ultrasound	V Z	varicella zoster
TSH	thyroid-stimulating hormone	V Z	varicena zoster
TSI	thyroid stimulating immunoglobulin	W	width; tungsten
TTN	transient tachypnea of the newborn	WBC	white blood cell(s) (count)
TURP	transurethral resection of prostate	WES	wall-echo-shadow (triad)
TV	tidal volume; transvaginal	WHO	World Health Organization
TVS	transvaginal sonography	WM	white matter
140	transvaginar somograpmy	WPW	Wolf-Parkinson-White (syndrome)
U	uranium	w/w	weight (of solute) per weight (of total
UA	umbilical artery	**/ **	solvent)
UBC	unicameral bone cyst		sorvene)
UC	ulcerative colitis	XCCL	exaggerated craniocaudal
UCD	uremic cystic disease	Xe	xenon
UCL	ulnar collateral ligament	XGP	xanthogranulomatous pyelonephritis
UGI	upper gastrointestinal		
UIP	usual interstitial pneumonia	y r	year
UK	urokinase	•	•
UL	upper lobe	Z	atomic number
UPJ	ureteropelvic junction		
US	ultrasound	Symbols	
U.S.	United States	<	less (common) than
USPIO	ultrasmall superparamagnetic iron oxide	«	much less (common) than
UTI	urinary tract infection	≤	less than or equal to
UV	ultraviolet; umbilical vein	>	more (common) than
UVJ	ureterovesical junction	>>	much more (common) than
			greater than or equal to
VA	vertebral artery	≥ ⇒ Ø ↑	leads to
VACTERL	vertebral body, anal, cardiovascular,	Ø	normal, unchanged
	tracheoesophageal, renal, limb	↑	increased
	anomalies (association)	\downarrow	decreased
VATS	video-assisted thorascopic surgery		

Chest Imaging

CHAPTER OUTLINE Imaging Anatomy, 1 **Chronic Lung Disease, 27** Pleura, 53 Gross Lung Anatomy, 1 Idiopathic Diseases, 27 General, 53 Fluid Collections, 54 Parenchymal Anatomy, 4 Lymphoproliferative Disorders, 31 Pulmonary Function, 5 Collagen Vascular Diseases, 32 Pleural Tumors, 56 Mediastinum, 5 Vasculitis and Other, 57 Imaging Protocols, 5 Granulomatoses, 33 Mediastinum, 57 Other Chronic Disorders, 34 Infection, 6 General, 57 Inhalational Lung Disease, 35 General, 6 Anterior Mediastinal Tumors, 57 Bacterial Infections, 8 Middle Mediastinal Tumors, 60 Pneumoconiosis, 35 Posterior Mediastinal Viral Pneumonia, 12 Antigen-Antibody-Mediated Tumors. 61 Fungal Infections, 14 Lung Disease, 38 Toxin-Induced Interstitial Other Mediastinal Disorders, 62 **Acquired Immunodeficiency** Pneumonitis/Fibrosis, 39 Syndrome, 18 **Differential Diagnosis, 62** Airway Disease, 39 General, 18 General, 62 Chest, 18 Chronic Bronchial Disease, 41 Atelectasis, 63 Pneumocystis Jiroveci Consolidation, 64 Lung Injury, 44 Pneumonia, 19 Pulmonary Masses, 66 Postoperative Chest. 47 Cystic and Cavitary Lesions, 68 Neoplasm, 20 Pulmonary Vasculature, 49 Interstitial Lung Disease, 70 General, 20 Pulmonary Arterial Abnormal Density, 72 Bronchogenic Carcinoma, 21 Hypertension, 49 Tracheobronchial Lesions, 73 Tumor Staging, 22 Pulmonary Edema, 50 Pleural Disease, 74 Specific Lung Tumors, 25 Pulmonary Embolism, 51 Mediastinum, 74 Lung Metastases From Other Vasculitis, 52 Primary Lesions, 27 Venous Abnormalities, 52

Imaging Anatomy		Lower lobe	Superior	B6	
GROSS LUNG ANATOMY			Medial basal Anterior basal Lateral basal	B7 B8 B9	
SEGMENTAL ANATOMY (Figs. 1.1–1.2)			Posterior basal	B10	
Right Lung			Left Lung		
Upper lobe	Apical	B1	Upper lobe		
• •	Anterior	B2	Upper	Apicoposterior	B1, B3
	Posterior	В3		Anterior	B2
Middle lobe	Lateral	B4	Lingula	Superior	B4
	Medial	В5		Inferior	B5





Lower lobe	Superior	В6
	Medial basal	В7
	Anterior basal	В8
	Lateral basal	В9
	Posterior basal	B10

SEGMENTAL COMPUTED TOMOGRAPHY (CT) ANATOMY (Fig. 1.3)

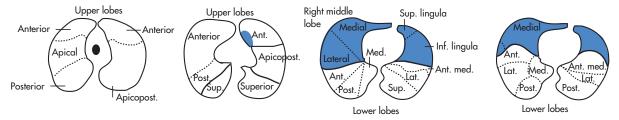


FIG. 1.3

BRONCHIAL CT ANATOMY (Fig. 1.4)

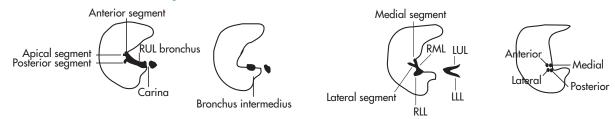


FIG. 1.4

PLAIN RADIOGRAPH ANATOMIC LANDMARKS (Figs. 1.5–1.9)

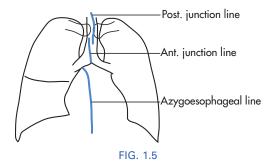
Thoracic Inlet

The thoracic inlet represents the junction between structures at the base of the neck and those of the thorax. It parallels the first rib and is higher posteriorly than anteriorly.

Lines

• Anterior junction line: 2-mm linear line that projects over the trachea. Represents the

- approximation of the visceral and parietal pleura of the right and left lungs anterior to the mediastinum (composed of four layers of pleura)
- Posterior junction line (four layers of pleura): extends above clavicles and can often be seen on a frontal radiograph as a vertical line traversing the tracheal air column
- Posterior tracheal stripe (normally measures <4 mm in diameter): thickening or presence of a focal opacity in the region of the posterior tracheal stripe should raise the possibility of esophageal carcinoma



Normal Hemorrhage, adenopathy

FIG. 1.6

Azygos

Sup.

FIG. 1.7

Major

Inf. accessory

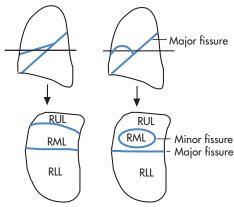
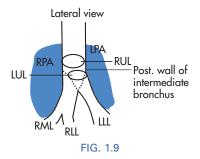


FIG. 1.8



- Azygoesophageal line: interface between RLL air and mediastinum
- Left paraspinal line: extends from aortic arch to diaphragm
- Right paraspinal line

Paratracheal Stripe

- Abnormal if >4 mm
- Never extends below right bronchus

Fissures

The normal major fissures are seldom seen on a posteroanterior radiograph.

The top of the left lower lobe (LLL) is usually higher than the top of the right lower lobe (RLL).

- Minor (horizontal) fissure
- Major (oblique) fissure
- Azygos fissure
- Azygos fissuresOther fissures

Superior accessory fissure Inferior accessory fissure Left minor fissure

Pulmonary Ligament

- Consists of a double layer of pleura that connects the medial aspect of the lower lobe (LL) to the adjacent mediastinum and diaphragm
- Not seen on posteroanterior or lateral chest radiographs (CXRs)
- Determines the shape of the collapsed LL in patients with atelectasis and the shape of the collapsed lung in patients with pneumothorax

Trachea

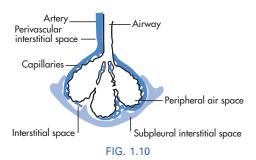
- The trachea is a midline structure
- The aorta commonly causes a smooth indentation on the left side
- The trachea measures 10-12 cm in length
- 16–20 U-shaped cartilage rings on its lateral and anterior aspects
- Calcification of the cartilage rings is a common normal finding in patients older than 40 years, particularly women, but it is seldom evident on radiographs

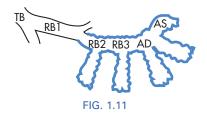
 Divides into the left and right main bronchi at the carina (approximately at the level of the fifth thoracic vertebra)

Upper Lobe (Ul) Bronchi (Figs. 1.10–1.11)

The right main bronchus divides into the RUL bronchus and the bronchus intermedius.

- RUL bronchus is always higher than LUL bronchus on lateral view
- Posterior wall of bronchus intermedius (right) is normally less than 2 mm thick and bifurcates into middle and LLs bronchi
- Tracheal bronchus (bronchus suis): 0.1% of population, arises from right wall of trachea (left much less common), supplies apical segment or occasionally entire RUL
- The left main bronchus is approximately 5 cm in length and divides into the LUL and LLL bronchi
- Accessory cardiac bronchus: 0.1% of population, extends inferomedially from medial wall of bronchus intermedius or RLL bronchus toward mediastinum; may be blind ending





PARENCHYMAL ANATOMY

ACINUS

- Includes all structures distal to one terminal bronchiole. The terminal bronchiole is the last purely air-conducting structure.
- Acinus measures 7 mm
- Acinus contains about 400 alveoli

SECONDARY PULMONARY LOBULE

- Smallest anatomic unit of the lung visible on high-resolution CT (HRCT)
- Polygonal structure bounded by interlobular septa, 1.5–2 cm in diameter
- Three to five acini per secondary lobule
- Supplied by several terminal bronchioles

EPITHELIUM

The alveolar epithelium is made up of two cell types:

- Type 1 pneumocytes
- Type 2 pneumocytes: produce surfactant, have phagocytic ability, and regenerate

HIGH-RESOLUTION COMPUTED TOMOGRAPHY (HRCT) (Fig. 1.12)

Technique

- 1–1.5-mm thin collimation
- · High spatial frequency reconstruction

This helps to improve spatial resolution, thereby improving the ability to detect subtle abnormalities—thick interlobular septa, cyst walls, small nodules, ground glass opacities and bronchiectasis.

Optional

Increase in kVp or mA (140 kVp, 170 mA) Targeted image reconstruction (one lung rather than both to increase spatial resolution)

HRCT Anatomy

The basic anatomic unit of pulmonary structure and function visible by HRCT is the secondary pulmonary lobule:

- Polyhedral 1.5-cm structure surrounded by connective tissue (interlobular septa) and made up of 5–15 pulmonary acini, which contain the alveoli for gas exchange
- · Central artery and bronchiole
- Peripheral pulmonary veins and lymphatics in septum

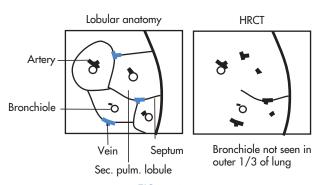


FIG. 1.12

Dominant high-resolution pattern:

- Reticular
- Nodular
- High attenuation (ground glass, consolidation)
- Low attenuation (emphysema, cystic)

Questions:

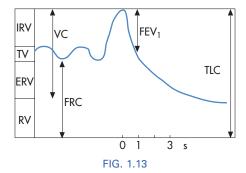
- Location within the secondary lobule
- Upper versus lower zone or a central versus peripheral predominance
- Presence of additional findings (pleural fluid, lymphadenopathy, traction bronchiectasis)

This protocol produces high-definition images of the lung alveoli, airways, interstitium, and pulmonary vasculature. Air trapping is identified on expiratory images.

PULMONARY FUNCTION (Fig. 1.13)

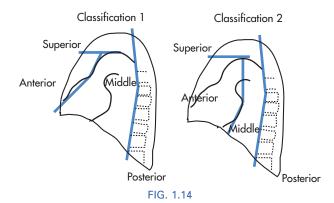
LUNG VOLUMES, CAPACITIES, AND FLOW RATES

- Tidal volume (TV): normal respiratory cycle
- Vital capacity (VC): amount of air that can be expired with force after maximal inspiration
- Functional residual capacity (FRC): volume remaining in lung after quiet expiration
- Total lung capacity (TLC): volume contained in lung at maximum inspiration
- Forced expiratory volume (FEV₁): amount of air expired in 1 second



MEDIASTINUM (Fig. 1.14)

- Superior mediastinum: plane above aortic arch; thoracic inlet structures
- Anterior mediastinum: contains thymus, lymph nodes, mesenchymal tissue; some classifications include the heart and fat
- Middle mediastinum: contains heart, major vessels, trachea and main bronchi, lymph nodes, phrenic nerve, and left recurrent laryngeal nerve



 Posterior mediastinum: starts at anterior margin of vertebral bodies; contains descending thoracic aorta, esophagus, thoracic duct, azygos and hemiazygos veins, lymph nodes, autonomic nerves, paravertebral areas, and fat

IMAGING PROTOCOLS

STANDARD CHEST CT PROTOCOL

Supine position. Scan in suspended inspiration at total lung capacity. Scan setup:

- 5 × 5-mm sections from apex of the lungs to the adrenals
- Six 1.25-mm high-resolution cuts throughout lung at 2.5-cm intervals
- 1-mm reconstructions through pulmonary nodules
- Number of different combinations of pitch and section thickness

In interstitial lung disease the six cuts are repeated with the patient in the prone position. Reconstruction is done with a high-resolution bone algorithm.

Use of IV contrast medium:

- Evaluation of vascular structures, arteriovenous malformation, aortic dissection
- Evaluation of mediastinal tumors, enlarged lymph nodes
- · Hilar masses
- Neck masses

PULMONARY EMBOLISM (PE) CT PROTOCOL

- Patient in supine position
- Scan range: adrenals to lung apex
- Injection of 140 mL of nonionic iodinated contrast at 3 mL/second, with delay of 25–30 seconds. Scanning is performed with suspended respiration.
- Scans are retrospectively reconstructed from the dome of the diaphragm as 2.5-mm-thick slices with 1-mm spacing.

DIAGNOSTIC RADIOLOGY REPORT (AMERICAN COLLEGE OF RADIOLOGY [ACR])

An authenticated written interpretation should be performed on all radiologic procedures. The report should include:

- Name of patient and other identifier (e.g., birth date, Social Security number, or hospital or office identification number)
- 2. Name of the referring physician to provide more accurate routing of the report to one or more locations specified by the referring physician (e.g., hospital, office, clinic)
- 3. History
- 4. Name or type of examination
- 5. Dates of the examination and transcription
- 6. Time of the examination (for ICU/CCU patients) to identify multiple examinations (e.g., chest) that may be performed on a single day
- 7. Body of the report:
 - Procedures and materials

Include in the report a description of the procedures performed and any contrast media (agent, concentration, volume, and reaction, if any), medications, catheters, and devices.

Findings

Use precise anatomic and radiologic terminology to accurately describe findings.

Limitations

Where appropriate, identify factors that can limit the sensitivity and specificity of the examination. Such factors might include technical factors, patient anatomy, limitations of the technique, incomplete bowel preparation, and wrist examination for carpal scaphoid.

Clinical issues

The report should address or answer any pertinent clinical issues raised in the request for the imaging examination. For example, to rule out pneumothorax state: "There is no evidence of pneumothorax." To rule out fracture state: "There is no evidence of fracture." It is not advisable to use such universal disclaimers as "The mammography examination does not exclude the possibility of cancer."

• Comparative data

Comparisons with previous examinations and reports when possible are a part of the radiologic consultation and report and optionally may be part of the "impression" section.

- 8. Impression (conclusion or diagnosis):
 - Each examination should contain an "impression" section.
 - Give a precise diagnosis whenever possible.
 - Give a differential diagnosis when appropriate.
 - Recommend, only when appropriate, followup and additional diagnostic radiologic studies to clarify or confirm the impression.

In normal CXR section the only structures visible in normal lungs are the fissures and the pulmonary vessels.

Lung parenchymal abnormalities are divided into five basic patterns:

- 1. Mass
- 2. Consolidative
- 3. Interstitial
- 4. Vascular
- 5. Airway

Infection

GENERAL

PATHOGENS

Bacterial pneumonia

- Streptococcus pneumoniae (pneumococcus)
- Staphylococcus
- Pseudomonas
- Klebsiella
- Nocardia
- Chlamydia
- Neisseria meningitides
- Haemophilus influenzae
- Anaerobes
- Legionella
- Mycoplasma pneumoniae
- Actinomyces israelii
- Mycobacterium tuberculosis

Viral pneumonia (25% of community-acquired pneumonias)

- Influenza
- Varicella, herpes zoster
- Rubeola
- Cytomegalovirus
- Coxsackievirus, parainfluenza virus, adenovirus, respiratory syncytial virus (RSV)

Fungal pneumonia

- Histoplasmosis
- Coccidioidomycosis
- Blastomycosis
- Aspergillosis
- Cryptococcosis
- Candidiasis
- Zygomycoses

Parasitic pneumonias

- Pneumocystis jiroveci Frenkel 1999 (formerly Pneumocystis carinii)
- Toxoplasma gondii

ACQUISITION OF PNEUMONIA

Community-acquired pneumonia

- S. pneumoniae, Haemophilus
- Mycoplasma

Hospital-acquired pneumonia (incidence 1%, mortality 35%): nosocomial infection

- Gram-negative bacteria: *Pseudomonas, Proteus, Escherichia coli, Enterobacter, Klebsiella*
- Methicillin-resistant *Staphylococcus aureus* (MRSA)
- Vancomycin-resistant enterococcus (VRE)

Pneumonia in immunosuppressed patients

- Bacterial pneumonia (gram negative) still most common
- Tuberculosis
- Fungal
- Pneumocystis pneumonia (PCP)

Endemic pneumonias

- Fungal: histoplasmosis, coccidioidomycosis, blastomycosis
- Viral

Aspiration-associated pneumonia (important)

RISK FACTORS

The radiographic appearance of pulmonary infections is variable depending on the pathogen, underlying lung disease, risk factors, and previous or partial treatment.

COMMUNITY-ACQUIRED INFECTIONS

Risk Factor	Common Pathogens
Alcoholism	Gram-negative bacteria, <i>Streptococcus pneumoniae</i> , <i>Mycobacterium tuberculosis</i> , aspiration (mouth flora)
Old age	S. pneumoniae, Staphylococcus aureus, aspiration
Aspiration	Mouth flora (anaerobes)
Cystic fibrosis	Pseudomonas, S. aureus, Aspergillus
Chronic bronchitis	S. pneumoniae, Haemophilus influenzae

Other risk factors for developing pneumonia:

- Bronchiectasis
- Coma, anesthesia, seizures (aspiration)
- Tracheotomy
- Antibiotic treatment
- Immunosuppression (renal failure, diabetes, cancer, steroids, AIDS)
- Chronic furunculosis (Staphylococcus)

RADIOGRAPHIC SPECTRUM OF PULMONARY INFECTIONS

SUMMARY

Туре	Pathogen	Imaging
Lobar Pneumonia Infection primarily involves alveoli Spread through pores of Kohn and canals of Lambert throughout a segment and ultimately an entire lobe Bronchi are not primarily affected and remain air filled; therefore: Air bronchograms No volume loss because airways are open Nowadays uncommon because of early treatment Round pneumonia (more common in children)	Streptococcus pneumoniae Klebsiella pneumoniae Others Staphylococcus aureus Haemophilus influenzae Fungal	Air bronchogram Consolidation (no volume loss) A B
Bronchopneumonia Primarily affects the bronchi and adjacent alveoli Volume loss may be present as bronchi filled with exudates Bronchial spread results in multifocal patchy opacities	S. aureus Gram-negative bacteria Others H. influenzae Mycoplasma	Patchy consolidation in segmental distribution

SUMMARY—cont'd

Туре	Pathogen	Imaging
Nodules Variable in size Indistinct margins	Fungal Histoplasma Aspergillus Cryptococcus Coccidioides Bacterial Legionella Nocardia Septic emboli S. aureus	
Cavitary Lesions (Infectious) Abscess: necrosis of lung parenchyma ± bronchial communication Fungus ball (air crescent/monad sign) Postprimary TB (favor apical and posterior segments of the upper lobes) Pneumatoceles caused by air leak into pulmonary interstitium	Anaerobic bacteria Aspergillus M. tuberculosis S. aureus	Abscess Pneumatocele
Diffuse Opacities Reticulonodular pattern: interstitial peribronchial areas of inflammation (viral) Alveolar location (PCP) Miliary pattern: hematogenous spread (TB)	Viral <i>Mycoplasma</i> PCP	Reticulonodular Nodular

PCP, Pneumocystis pneumonia; TB, tuberculosis.

Complications of Pneumonia

• Parapneumonic effusion

Stage 1: exudation: free flowing

Stage 2: fibropurulent: loculated

Stage 3: organization, erosion into lung or chest wall

- Empyema
- Bronchopleural fistula (BPF; fistula between bronchus and pleural space) with eroding pleural-based fluid collections
- Bronchiectasis
- · Pulmonary fibrosis, especially after necrotizing pneumonia or acute respiratory distress syndrome (ARDS)
- Adenopathy

RESOLUTION OF PNEUMONIA

- 80%–90% of cases resolve within 4 weeks.
- 5%-10% resolve within 4-8 weeks (usually in older or diabetic patients). Subsequent radiographs

should always show interval improvement compared with the previous radiographs.

Nonclearance

Antibiotic resistance

Consider other pathogen (e.g., M.

tuberculosis)

Recurrent infection

Obstruction pneumonitis due to tumor

BACTERIAL INFECTIONS

GENERAL

Common Pathogens

- *S. pneumoniae*, 50% (40–60 years)
- Mycoplasma, 30%
- Anaerobes, 10%
- Gram-negative bacteria, 5%
- Staphylococcus, 5%

 Haemophilus, 3% (especially in infants and patients with chronic obstructive pulmonary disease [COPD])

Clinical Findings

Pneumonic syndrome

- Fever
- Cough
- Pleuritic pain
- Sputum

Ancillary findings

- Headache, arthralgia, myalgia
- Diarrhea
- Hemoptysis

STREPTOCOCCAL PNEUMONIA

Radiographic Features

- · Lobar or segmental pneumonia pattern
- Bronchopneumonia pattern
- Round pneumonia (in children)

STAPHYLOCOCCAL PNEUMONIA (Fig. 1.15)

Radiographic Features

- Bronchopneumonia pattern
- Bilateral, >60%
- Abscess cavities, 25%–75%
- Pleural effusion, empyema, 50%
- Pneumatoceles, 50% (check valve obstruction), particularly in children
- Central lines
- · Signs of endocarditis

PSEUDOMONAS PNEUMONIA

Typical Clinical Setting

- Hospital-acquired infection
- Ventilated patient
- · Reduced host resistance
- Patients with cystic fibrosis

Radiographic Features

Three presentations:

- Extensive bilateral parenchymal consolidation (predilection for both LLs)
- Abscess formation

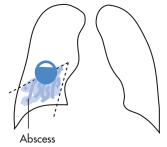


FIG. 1.15

• Diffuse nodular disease (bacteremia with hematogenous spread; rare)

LEGIONNAIRES DISEASE

Severe pulmonary infection caused by *Legionella pneumophila*; 35% of patients require ventilation, 20% mortality. Most infections are community acquired. Patients have hyponatremia. Seroconversion for diagnosis takes 2 weeks.

Radiographic Features

Common features

- Initial presentation of peripheral patchy consolidation
- · Bilateral severe disease
- Rapidly progressive
- Pleural effusions, <50%
- LL predilection

Uncommon features

- Abscess formation
- Lymph node enlargement

HAEMOPHILUS PNEUMONIA

Caused by *H. influenzae*. Occurs most commonly in children, immunocompromised adults, or patients with COPD. Often there is concomitant meningitis, epiglottitis, and bronchitis.

Radiographic Features

- Bronchopneumonia pattern
- LL predilection, often diffuse
- Empyema

MYCOPLASMA PNEUMONIA

Most common nonbacterial pneumonia (atypical pneumonia). Mild course. Age 5–20 years. Positive for cold agglutinins, 60%.

Radiographic Features

- Reticular pattern
- LL predominance, often diffuse
- Consolidation, 50%

Complications

- Autoimmune hemolytic anemia
- Erythema nodosum, erythema multiforme
- Stevens-Johnson syndrome
- Meningoencephalitis

KLEBSIELLA (FRIEDLÄNDER) PNEUMONIA

Gram-negative organism. Often in debilitated patients and/or alcoholics.

Radiographic Features

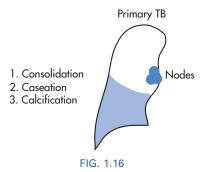
• Consolidation appears similar to that of infection with *S. pneumoniae*

- Lobar expansion
- Cavitation, 30%–50%, typically multiple
- Massive necrosis (pulmonary gangrene)
- Pleural effusion uncommon

TUBERCULOSIS (TB) (Fig. 1.16)

Transmitted by inhalation of infected droplets of *M. tuberculosis* or *M. bovis*. TB acquisition usually requires constant or repeated contact with sputum-positive patients because the tubercle does not easily grow in the immunocompetent human host. Target population includes:

- Patients of low socioeconomic status (homeless)
- Alcoholics



- Immigrants: from Mexico, Philippines, Indochina, Haiti
- Elderly patients
- AIDS patients
- Prisoners

Primary Infection (Fig. 1.17)

Usually heals without complications. Sequence of events includes:

- Pulmonary consolidation (1–7 cm); cavitation is rare: LL (60%) > UL
- Caseous necrosis 2-10 weeks after infection
- Lymphadenopathy (hilar and paratracheal), 95%
- Pleural effusion, 10%
- Spread of a primary focus occurs primarily in children or immunosuppressed patients.

Secondary Infection (Fig. 1.18)

Active disease in adults most commonly represents reactivation of a primary focus. However, primary disease is now also common in adults in developed countries because there is no exposure in childhood. Distribution is as follows:

- Typically limited to apical and posterior segments of ULs or superior segments of LLs (because of high Po_??)
- Rarely in anterior segments of ULs (in contradistinction to histoplasmosis)

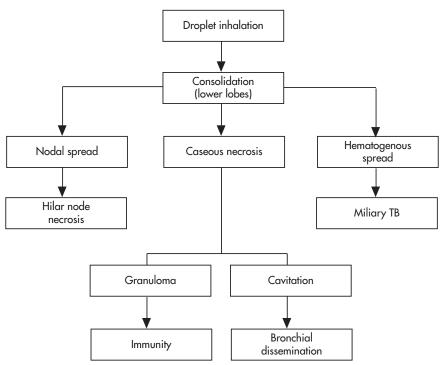


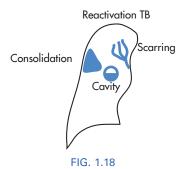
FIG. 1.17

Radiographic Features

- Exudative TB
 - Patchy or confluent air space disease Adenopathy uncommon
- Fibrocalcific TB
 - Sharply circumscribed linear densities radiating to hilum
- Cavitation, 40%

Complications (Fig. 1.19)

- Miliary TB may occur after primary or secondary hematogenous spread.
- Bronchogenic spread occurs after communication of the necrotic area with a bronchus; it produces



an acinar pattern (irregular nodules approximately 5 mm in diameter).

- Tuberculoma (1–7 cm): nodule during primary or secondary TB; may contain calcification
- Effusions are often loculated.
- Bronchopleural fistula
- Pneumothorax

COMPARISON

	Primary TB	Reinfection TB
Location	Usually bases	Upper lobes, superior segment Lower lobes
Appearance	Focal	Patchy
Cavitation	No	Frequent
Adenopathy as only finding	Common	No
Effusion	Common	Uncommon
Miliary pattern	Yes	Yes

TB, Tuberculosis.

NONTUBERCULOUS MYCOBACTERIAL (NTMB) INFECTIONS

The two most common NTMB pathogens are *M. avium-intracellulare* and *M. kansasii* (less common: *M. xenopi, M. chelonei, M. gordonae, M. fortuitum,* "fast grower"). Unlike TB, NTMB infections are not

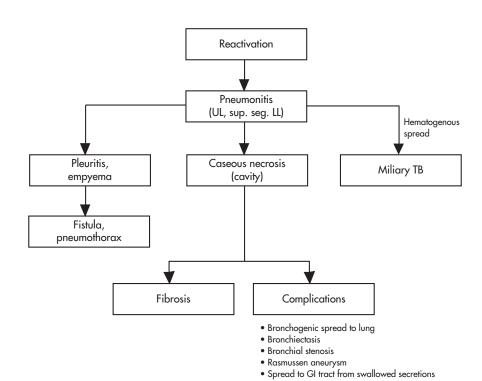


FIG. 1.19

acquired by human-human transmission but are a direct infection from soil or water. There is also no pattern of primary disease or reactivation: the infection is primary, although some infections may become chronic. The infection often occurs in elderly patients with COPD, older women in good health, and AIDS patients.

Radiographic Features

- NTMB infections may be indistinguishable from classic TB.
- Atypical features such as bronchiectasis and bronchial wall thickening are common.
- Nodules are common in older women.

COMPUTED TOMOGRAPHY FINDINGS

Findings	TB (%)	MAI Infection (%)
Nodules <1 cm	80	95
Nodules 1–3 cm	40	30
Mass >3 cm	10	10
Consolidation	50	50
Cavity	30	30
Bronchiectasis	30	95
Bronchial wall thickening	40	95
Septal thickening	50	15
Emphysema	20	20
Calcified granuloma	15	5

MAI, Mycobacterium avium-intracellulare; TB, tuberculosis.

NOCARDIA PNEUMONIA

Caused by *Nocardia asteroides*, worldwide distribution. Common opportunistic invader in:

- Lymphoma
- Steroid therapy; especially transplant patients
- Pulmonary alveolar proteinosis (common)

Radiographic Features

- Focal consolidation (more common)
- Cavitation
- Irregular nodules

ACTINOMYCOSIS

Actinomycosis is caused by *Actinomyces israelii*, a gram-positive normal saprophyte in the oral cavity. Pulmonary disease develops from aspiration of the organism (poor dentition) or from direct penetration into the thorax.

Radiographic Features

- Focal consolidation > cavitating mass
- Lymphadenopathy uncommon
- Extension into the chest wall and pleural thickening is less common today but still occurs and is an important differential feature.

PULMONARY ABSCESS

The spectrum of anaerobic pulmonary infections includes:

- Abscess: single or multiple cavities >2 cm, usually with AFL
- Necrotizing pneumonia: analogous to abscess but more diffuse and cavities <2 cm
- Empyema: suppurative infection of the pleural space, most commonly as a result of pneumonia

Predisposing Conditions

- Aspiration (e.g., alcoholism, neurologic disease, coma)
- Intubation
- Bronchiectasis, bronchial obstruction

Treatment

- · Antibiotics, postural drainage
- Percutaneous drainage of empyema
- Drainage/resection of lung abscess only if medical therapy fails

SICKLE CELL ANEMIA

- Patients with sickle cell disease are at increased risk of pneumonia and infarction. These entities are difficult to differentiate, and hence are called acute chest syndrome.
- Pneumonias were originally due to pneumococci but now are due to viruses or Mycoplasma. Differential diagnosis includes at electasis and infarct.
- Infarcts are more frequent in adults than in children. Rare in children younger than 12 years.
- Consolidation is seen on CXRs; resolves more slowly than in the general population and tends to recur.

VIRAL PNEUMONIA

GENERAL

Classification

DNA viruses

Unenveloped

- Parvoviruses
- Papovaviruses
- Adenoviruses
- Hepatitis viruses (hepatitis B)

Enveloped

- Herpesviruses (herpes simplex virus [HSV], Epstein-Barr virus [EBV], varicella-zoster virus [VZV], cytomegalovirus [CMV])
- Poxviruses (variola virus, molluscum contagiosum virus)

RNA viruses

Unenveloped

- Picornaviruses (hepatitis A virus, coxsackievirus)
- Caliciviruses
- Reoviruses

Enveloped

- Retroviruses (HIV)
- Arenaviruses
- Coronaviruses
- Togaviruses
- Bunyaviruses
- Orthomyxoviruses (influenza virus)
- Paramyxoviruses (mumps virus, measles virus, RSV, parainfluenza virus)

Occurrence

Immunocompetent hosts

Influenza

Hantavirus

EBV

Adenovirus

Immunocompromised hosts

HSV

VZ

CMV

Measles virus

Adenovirus

Spectrum of Disease

- Acute interstitial pneumonia (AIP): diffuse or patchy interstitial pattern, thickening of bronchi, thickened interlobar septa
- Lobular inflammatory reaction: multiple nodular opacities 5–6 mm (varicella; late calcification)
- Hemorrhagic pulmonary edema: mimics bacterial lobar pneumonia
- Pleural effusion: usually absent or small
- Chronic interstitial fibrosis (bronchiolitis obliterans)

INFLUENZA PNEUMONIA

Influenza is very contagious and thus occurs in epidemics. Pneumonia, however, is uncommon.

Involves the upper respiratory tract, including the trachea and major bronchi.

Radiographic Features

- Acute phase: multiple acinar densities
- Coalescence of acinar densities to diffuse patchy airspace disease (ASD; bronchopneumonia type)

VARICELLA-ZOSTER PNEUMONIA

Fifteen percent of infected patients have pneumonias; 90% are older than 20 years.

Radiographic Features

- Acute phase: multiple acinar opacities
- Coalescence of acinar opacities to diffuse patchy ASD
- 1–2-mm calcifications throughout lungs after healing
- HRCT usually shows 1–10-mm well-defined and ill-defined nodules diffusely throughout both lungs.

MEASLES VIRUS PNEUMONIA

Two forms:

- Primary measles virus pneumonia and secondary bacterial pneumonia
- Atypical measles virus pneumonia

Radiographic Features

Primary measles virus:

- Mixed reticular opacities and air space consolidation.
- Lymph node enlargement in the hilum.
- CT findings include ground-glass attenuation, air space consolidation, and small centrilobular nodules.

Virus	Centrilobular Nodules	Lobar Ground Glass	Diffuse Ground Glass	Thickened Interlobular Septa	Consolidation
Influenza virus	+++	+++	+		+
EBV	+	+	+		+
CMV	++	++	++	+	+
VZ	+++	+	+		
HSV	+	+++	+		+++
Measles virus	++	+	+		+
Hantavirus			+++	+	++
Adenovirus	++	+			+++

Atypical measles virus:

- Spherical or segmental consolidation, which clears rapidly
- Hilar lymph node enlargement and pleural effusion are frequently present.

CYTOMEGALOVIRUS (CMV) PNEUMONIA

Occurs most commonly in neonates or immunosuppressed patients.

Radiographic Features

- Predominantly interstitial infection, multiple small nodules (common)
- · Adenopathy may be present

SWINE-ORIGIN INFLUENZA A (H1N1) VIRUS INFECTION

Epidemiologic data to date suggest that the newly emerged H1N1 virus, although transmissible from person to person, is of relatively low virulence. CXRs are normal in more than half of patients. However, the disease can progress to bilateral extensive ASD in severely ill patients. These patients are also at high risk of PE, which should be sought carefully on contrast-enhanced CT scans.

SEVERE ACUTE RESPIRATORY SYNDROME

Emerging highly contagious infection caused by a coronavirus; first outbreak reported in southern China in 2002. The disease has two clinical stages: viral replicative stage and immunopathologic stage. The incubation period for the virus ranges from 2 to 12 days.

Radiographic Features

- During viral replication stage, well-defined areas of ground-glass opacities seen in the LLs and periphery of the lungs. The LLs and peripheral areas of the lungs are most commonly involved.
- During the immunopathologic phase, patients show the appearance of new lesions that are poorly defined and usually localized to the LLs and posterior or dependent regions of the lungs. Following the acute phase, there is a decrease in the extent of ground-glass opacity and consolidation. Some patients may develop spontaneous pneumomediastinum.

FUNGAL INFECTIONS

GENERAL

Two broad categories:

Endemic human mycoses (prevalent only in certain geographic areas):

- Histoplasmosis (Ohio, Mississippi, St. Lawrence river valleys)
- Coccidioidomycosis (San Joaquin Valley)
- Blastomycosis

Opportunistic mycoses (worldwide in distribution) occur primarily in immunocompromised patients (aspergillosis and cryptococcosis may also occur in immunocompetent hosts):

- Aspergillosis
- Candidiasis
- Cryptococcosis
- Mucormycosis

Radiographic Features

- Acute phase: pneumonic type of opacity (may be segmental, nonsegmental, or patchy); miliary (hematogenous) distribution in immunosuppressed patients
- Reparative phase: nodular lesions with or without cavitations and crescent sign
- Chronic phase: calcified lymph nodes or pulmonary focus with fungus (e.g., histoplasmosis)
- Disseminated disease (spread to other organs) occurs primarily in immunocompromised patients

HISTOPLASMOSIS (Fig. 1.20): PULMONARY AND MEDIASTINAL

Histoplasma capsulatum is particularly prevalent in the Ohio, Mississippi, and St. Lawrence river valleys, although the agent has a worldwide distribution. The organism is most prevalent in soil that contains excrement of bats and birds (bat caves, chicken houses, old attics, or buildings).

Clinical Findings

Most patients are asymptomatic or have nonspecific respiratory symptoms, increased complement fixation titer, and *H. capsulatum* antigen positivity.

Radiographic Features

Consolidation (primary histoplasmosis)

- Parenchymal consolidation
- Adenopathy is very common and may calcify heavily later on.



FIG. 1.20

Nodular form (chronic histoplasmosis, reinfection):

- Histoplasmoma: usually solitary, sharply circumscribed nodule, most commonly in LLs
- Fibrocavitary disease in ULs indistinguishable from postprimary TB
- Cavitary nodules

Disseminated form (immunocompromised patients)

- Miliary nodules
- Calcifications in liver and spleen

Mediastinal Histoplasmosis

Mediastinal histoplasmosis may follow pulmonary histoplasmosis. There are two distinct entities (which may not always be separable from each other):

Mediastinal granuloma

- Results from spread of *H. capsulatum* to lymph nodes
- · Granulomas usually calcified
- Displacement of the superior vena cava or esophagus

Mediastinal fibrosis (fibrosing or sclerosing mediastinitis)

- May cause superior vena cava syndrome, airway compression, PA occlusion, pericarditis
- Diffuse infiltration of mediastinum
- Multiple densely calcified nodes (CT is more useful than magnetic resonance imaging [MRI] in making this diagnosis)

COCCIDIOIDOMYCOSIS (Fig. 1.21)

Coccidioides immitis is endemic in the southwest United States (San Joaquin Valley, "valley fever") and in Central America and South America. Infection occurs by inhalation of spores in soil. Human-to-human infection does not occur.

Clinical Findings

Cutaneous manifestations common; 70% are asymptomatic.

Radiographic Features

Consolidation (primary form)

- "Fleeting" parenchymal consolidation, most commonly LLs
- Adenopathy in 20%



FIG. 1.21

Nodular form (chronic form, 5%)

- 15% cavitate
 - 50% have a thin-walled cavity (suggestive of diagnosis)
 - 50% have a thick-walled cavity (i.e., nonspecific)

May present with pneumothorax

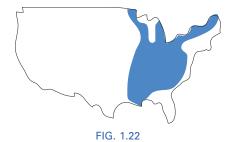
- Nodules rarely calcify
- Hilar or paratracheal adenopathy

Disseminated form (immunocompromised patients; rare: 0.5% of all forms)

- Miliary nodules
- Extrapulmonary spread

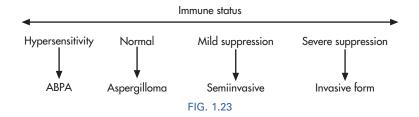
NORTH AMERICAN BLASTOMYCOSIS (Fig. 1.22)

- Caused by *Blastomyces dermatitidis*; uncommon infection. Most infections are self-limited.
- CXR is nonspecific: ASD > nodule (15% cavitate) or solitary mass > miliary spread.
- Focal or diffuse air space consolidation is the most common radiologic finding.
- Focal blastomycosis typically occurs in paramediastinal location and has an air bronchogram, findings that may suggest the diagnosis.
- Satellite nodules around primary focus are common.
- Chronic blastomycosis may mimic lung cancer because it can manifest itself as a focal mass.
 An air bronchogram or presence of satellite nodules is suggestive of correct diagnosis.
- Adenopathy, pleural effusions, and calcifications are very uncommon.
- Bone lesions, 25%
- Skin lesions are common.



ASPERGILLOSIS (Fig. 1.23)

Aspergillus is a ubiquitous fungus that, when inhaled, leads to significant lung damage. The fungus grows in soil, water, decaying vegetation, and hospital air vents. Infection with *A. fumigatus > A. flavus, A. niger*, or *A. glaucus*. There are four unique forms of pulmonary aspergillosis, each associated with a specific immune status.



TYPES OF ASPERGILLOSIS

Туре	Lung Structure	Immune Status	Pathology
Allergic (ABPA)	Normal	Hypersensitivity	Hypersensitivity → bronchiectasis, mucus plugging
Aspergilloma	Preexisting cavity	Normal	Saprophytic growth in preexisting cavity
Invasive	Normal	Severely impaired	Vascular invasion, parenchymal necrosis
Semiinvasive	Normal	Normal or impaired	Chronic local growth, local cavity formation

ABPA, Allergic bronchopulmonary aspergillosis.

ALLERGIC BRONCHOPULMONARY ASPERGILLOSIS (ABPA)

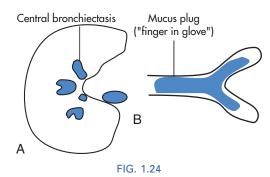
ABPA is a complex type I (IgE-mediated) and type III (IgG-mediated) hypersensitivity reaction to *A. fumigatus*, occurring primarily in individuals with asthma and occasionally in individuals with cystic fibrosis. The hypersensitivity initially causes bronchospasm, mucus production, and bronchial wall edema (IgE mediated); ultimately there is bronchial wall damage due to the type III IgG-mediated response, with resultant cystic bronchiectasis.

Clinical Findings

Elevated levels of specific serum IgE and IgG antibodies to *A. fumigatus*, asthma, peripheral eosinophilia, elevated serum IgE levels (≥1000 IU/mL), positive skin test for *Aspergillus* antigen. Treatment is with oral corticosteroids, antifungal agents, and omalizumab, which is a humanized monoclonal antibody targeted against IgE.

Radiographic Features

- Fleeting pulmonary parenchymal opacities (common manifestation)
- Central, UL saccular bronchiectasis (hallmark) (Fig. 1.24A)



- Mucus plugging ("finger-in-glove" appearance) (Fig. 1.24B) and bronchial wall thickening (common); 25% of patients will demonstrate high-attenuation mucus plugging.
- Tree-in-bud nodularity
- Cavitation, 10%

ASPERGILLOMA (MYCETOMA, FUNGUS BALL)

This is a saprophytic infection that occurs in the setting of structural lung disease (from TB, sarcoid, emphysema). Commonly in ULs, solitary lesions. The fungus grows into the preexisting cavity (e.g., cyst, bulla, bronchiectasis), creating a "fungus ball" consisting of fungus, mucus, and inflammatory cells. Individuals with mycetomas are often asymptomatic but may develop recurrent hemoptysis, which in rare cases can be massive. In these cases, bronchial artery embolization is indicated. The other treatment options include surgical resection, intracavity administration of amphotericin B, and systemic antifungal therapy.

Radiographic Features

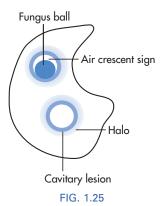
- Focal intracavitary mass (3–6 cm), typically in ULs.
- Air may surround the aspergilloma (Monod sign), mimicking the appearance of cavitation seen with invasive aspergillosis.
- Small area of consolidation around cavity is typical.
- Adjacent pleural thickening is common.
- Fungus ball moves with changing position.

INVASIVE ASPERGILLOSIS

Invasive aspergillosis has a high mortality rate (70%–90%) and occurs mainly in severely immuno-compromised patients (bone marrow transplants, leukemia). The infection starts with endobronchial fungal proliferation and then leads to vascular invasion with thrombosis and infarction of the lung ("angioinvasive infection"). Additional sites of infection (in 30%) are the brain, liver, kidney, and GI tract. Treatment is with systemic and/or intracavitary administration of amphotericin.

Radiographic Features (Fig. 1.25)

- Multiple pulmonary nodules, 40%
- Nodules have a characteristic halo of ground-glass appearance (represents pulmonary hemorrhage)
- Within 2 weeks, 50% of nodules undergo cavitation, which results in the air crescent sign. The appearance of the air crescent sign indicates the recovery phase (increased granulocytic response). The air crescent sign may also be seen in TB, actinomycosis, mucormycosis, septic emboli, and tumors. Do not confuse the air crescent sign with the Monod sign (clinical history helps to differentiate the two).
- Other manifestations:
 Peribronchial opacities
 Focal areas of consolidation



SEMIINVASIVE ASPERGILLOSIS

This form of aspergillosis occurs in mildly immunocompromised patients and has a pathophysiology similar to that of invasive aspergillosis except that the disease progresses more chronically over months (mortality rate 30%). Risk factors include diabetes, alcoholism, pneumoconiosis, malnutrition, and COPD. Treatment is with systemic and/or intracavitary administration of amphotericin.

Radiographic Features

- Appearance similar to that of invasive aspergillosis
- Cavitation occurs at 6 months after infection

CRYPTOCOCCOSIS

Caused by *Cryptococcus neoformans*, which has a worldwide distribution and is ubiquitous in soil and pigeon excreta. Infection occurs through inhalation of contaminated dust.

Clinical Findings

Common in patients with lymphoma, diabetes, or AIDS and in patients receiving steroid therapy.

Radiographic Features

- Most common findings in the lung are pulmonary mass, multiple nodules, or segmental or lobar consolidation.
- Cavitation, adenopathy, and effusions are rare.
- Disseminated form: CNS, other organs

CANDIDIASIS

Caused by *Candida albicans* more frequently than other *Candida* species.

Clinical Findings

Typically in patients with lymphoreticular malignancy; suspect pulmonary disease if associated with oral disease. Often there is disseminated fungemia.

Radiographic Features

- Plain radiograph is nonspecific: opacities (LL) > nodules.
- · Nodular disease in disseminated form
- Pleural effusion, 25%
- Cavitation and adenopathy are rare.

ZYGOMYCOSES

Group of severe opportunistic mycoses caused by fungi of the *Zygomycetes* class:

- Mucormycosis (*Mucor*)
- Rhizopus
- Absidia

Zygomycoses usually have two major clinical manifestations:

- Pulmonary mucormycosis
- Rhinocerebral mucormycosis

Zygomycoses are uncommon infections and occur primarily in immunocompromised patients (leukemia, AIDS, chronic steroid use, diabetes).

Radiographic Features

• Radiographic features are similar to those of invasive aspergillosis because of angioinvasive behavior of fungi.

Acquired Immunodeficiency Syndrome

GENERAL

AIDS is caused by HTLV type III (human T-cell lymphotrophic virus = HIV [human immunodeficiency virus]). HIV-1 and HIV-2 are single-stranded RNA viruses that bind to CD4 present on T lymphocytes (other cells: glial cells, lung monocytes, dendritic cells in lymph nodes). The viral RNA genome is copied into DNA with the help of reverse transcriptase and integrated into the host cellular DNA.

Known routes of HIV transmission:

- Blood and blood products
- Sexual activity
- In utero transmission
- During delivery

CLINICAL FINDINGS

- Lymphadenopathy
- Opportunistic infections
- Tumors: lymphoma—usually B-cell non-Hodgkin's lymphoma (NHL), Kaposi sarcoma (KS)
- Other manifestations:

Associated with lymphocytic interstitial pneumonia (LIP), usually in childhood

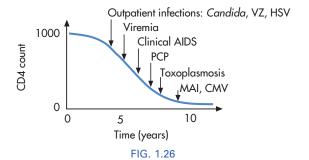
Spontaneous pneumothorax (development of cystic spaces, interstitial fibrosis related to PCP)

Septic emboli

 Clinical findings supportive of AIDS diagnosis (Fig. 1.26):

> CD4 cell count <200/mm³; the dysfunction of the immune system is inversely related to the CD4 cell count; PCP, CD4 cell count <200/mm³; MAI infection, CD4 cell count <50/mm³

> Less than one case of bacterial pneumonia per year



Opportunistic Infections

- P. jiroveci Frenkel 1999, 70%
- Mycobacterial infection, 20%; CD4 cell counts often <50/mm³
- Bacterial infection, 10% (S. pneumoniae, Haemophilus)
- Fungal infection (<5% of AIDS patients)
- Nocardia, <5%: cavitating pneumonia
- CMV pneumonia (common at autopsy)

CHEST

GENERAL

- 50% of all AIDS patients have pulmonary manifestations of infection or tumor.
- A normal CXR does not exclude the diagnosis of PCP.
- CMV infection is common at autopsy but does not cause significant morbidity or death; CMV antibody titers are present in virtually all patients with AIDS.
- Use of chest CT in AIDS patients:

Symptomatic patient with normal CXR; however, patients will commonly first undergo induced sputum or bronchoscopy or be given empirical therapy for PCP.

To clarify confusing CXR findings Workup of focal opacities, adenopathy, nodules

SPECTRUM OF CHEST MANIFESTATIONS

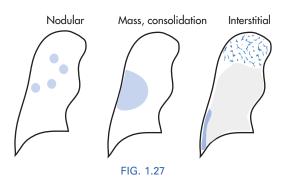
Nodules

(Fig. 1.27)

- KS (usually associated with skin lesions)
- Septic infarcts (rapid size increase)
- Fungal: Cryptococcus, Aspergillus

Large opacity: consolidation, mass

- Hemorrhage
- NHL
- Pneumonia
- Linear or interstitial opacities
- PCI
- · Atypical mycobacteria
- KS



Lymphadenopathy

- Mycobacterial infections
- KS
- Lymphoma
- Reactive hyperplasia, rare in thorax

Pleural effusion

- KS
- Mycobacterial, fungal infection
- Pyogenic empyema

PNEUMOCYSTIS JIROVECI PNEUMONIA

(PJP) (Figs. 1.28–1.30)

General

P. jiroveci is an atypical fungus that can result in pneumonia in immunocompromised individuals. PJP affects individuals with AIDS, transplant recipients, and patients receiving long-term corticosteroid therapy. When PJP occurs in HIV-infected individuals, it is usually only after the CD4 count has decreased to less

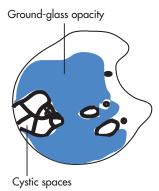


FIG. 1.28

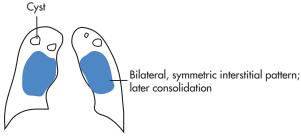


FIG. 1.29





FIG. 1.30

than 200/mm³. The diagnosis of PJP can be confirmed by bronchoalveolar lavage or presence of the organism in a sputum sample.

Radiographic Features

• Interstitial pattern, 80%

CXR: bilateral perihilar ground-glass opacities or thickened interstitium

HRCT: ground-glass opacities, interlobular septal thickening, and in some cases multifocal areas of consolidation

- Up to one-third of patients with PJP will have a normal CXR.
- Multiple cysts of various sizes are seen in onethird of patients with PJP and may cause spontaneous pneumothorax.
- UL *P. jiroveci* involvement is common, because aerosolized pentamidine may not get to ULs; UL disease may mimic TB, but the latter may have pleural effusions or lymphadenopathy, both of which are uncommon in PJP.
- Atypical patterns, 5%

Unilateral disease

Granulomatous inflammation resulting in focal nodules or masses

 PJP as a presenting manifestation of AIDS is decreasing in frequency because of trimethoprimsulfamethoxazole prophylaxis.

MYCOBACTERIAL INFECTION

M. tuberculosis > *M. avium-intracellulare* (MAI) (this pathogen usually causes extrathoracic disease). CD4 cell count usually <50/mm³.

Radiographic Features

• Hilar and mediastinal adenopathy common

Necrotic lymph nodes (TB) have a low attenuation center and only the rim is enhanced with contrast medium.

M. tuberculosis is more commonly associated with necrosis from MAI.

Adenopathy in KS or lymphoma is enhanced uniformly.

- Pleural effusion
- Other findings are similar to those of non-AIDS TB (UL consolidations, cavitations).

FUNGAL INFECTIONS

Fungal infections in AIDS are uncommon (<5% of patients):

- Cryptococcosis (most common); 90% have CNS involvement.
- Histoplasmosis: nodular or miliary pattern most common; 35% have a normal CXR.
- Coccidioidomycosis: diffuse interstitial pattern, thin-walled cavities