

### A GUIDE TO CLINICAL DECISION-MAKING: THE PSAP ALGORITHMS

### Online Handbook

FOURTH EDITION

American College of Clinical Pharmacy



For order information or questions, write or call: *A Guide to Clinical Decision-Making: The PSAP Algorithms, Fourth Edition* American College of Clinical Pharmacy 13000 W. 87th St. Parkway Lenexa, KS 66215-4530 *accp@accp.com* 

Copyright © 2008 by the American College of Clinical Pharmacy. All rights reserved. This book is protected by copyright. No part of this publication may be reproduced, stored in a retrieval system, or transmitted, in any form or by any means, electronic or mechanical, including photocopy, without prior written permission of the American College of Clinical Pharmacy.

Printed in the United States of America

Library of Congress Control Number: 2007929131 ISBN-13: 978-1932658293 ISBN-10: 1-932658-29-7

## **Introduction**

Clinicians continue to find the information in the *Pharmacotherapy Self-Assessment Program* (PSAP) modules not only a superb educational resource, but also a valuable patient care resource, replete with up-to-date information. The algorithms particularly can be used by clinicians as a quick reference to help guide patient care decisions. Although the format of PSAP is almost ideal as a teaching tool, it may not lend itself as readily to user-friendly practice applications. Thus, we have created this reference, a condensation of the pharmacotherapy algorithms from PSAP-VI, to provide a ready reference for a busy clinician.

The goal of this reference is to provide an affordable collection of clinically useful pharmacotherapy decisionmaking algorithms that are easy to find, understand, and apply. We have attempted to provide algorithms to help guide experienced clinicians in making optimal choices in concert with their patients without the need to consult additional references extensively. Thus, we have included algorithms leading to selection of type of therapy or selection between or among therapeutic options. Diagnostic algorithms have not been included. The material that is included is listed in the same general therapeutic categories as the PSAP modules. Please note that some algorithms may have been modified from their original PSAP version to reflect "current thinking."

We hope you find this collection of algorithms a useful addition to your patient care armamentarium.

Bruce R. Canaday, Pharm.D., BCPS, FASHP, FAPhA, Editor *Clinical Professor and Vice Chair* Department of Pharmacy Practice and Experiential Education University of North Carolina School of Pharmacy and *Director* Department of Pharmacotherapy Coastal Area Health Education Center Wilmington, North Carolina

## A Guide to Clinical Decision-Making: The PSAP Algorithms, Fourth Edition

Pharmacotherapy Self-Assessment Program, Sixth Edition

#### Authors

Thank you to the following PSAP-VI authors for preparing their algorithms for inclusion in this publication.

#### Cardiology I

Joseph J. Saseen, FCCP, BCPS (AQ Cardiology) Associate Professor Departments of Clinical Pharmacy and Family Medicine University of Colorado and Health Science Center Denver, Colorado

> C. Michael White, Pharm.D., FCCP, FCP Associate Professor of Pharmacy Department of Pharmacy Practice University of Connecticut Hartford, Connecticut

#### Cardiology II

Denise H. Rhoney, Pharm.D., FCCP Associate Professor Eugene Applebaum College of Pharmacy & Health Sciences Department of Pharmacy Practice Wayne State University Detroit, Michigan

Sarah A. Spinler, Pharm.D., FCCP, BCPS Professor of Clinical Pharmacy and Residency and Fellowship Program Coordinator

Philadelphia College of Pharmacy University of the Sciences in Philadelphia Philadelphia, Pennsylvania

#### Cardiology III

Cynthia A. Sanoski, Pharm.D. Associate Professor of Clinical Pharmacy Philadelphia College of Pharmacy University of the Sciences in Philadelphia Philadelphia, Pennsylvania

#### Nephrology I

Thomas C. Dowling, Pharm.D., Ph.D. Associate Professor University of Maryland Baltimore, Maryland

Neeta Bahal O'Mara, Pharm.D., BCPS *Clinical Pharmacist* Dialysis Clinic, Inc. North Brunswick, New Jersey

#### Nephrology II

Scott Bolesta, Pharm.D. Assistant Professor Department of Pharmacy Practice Nesbitt College of Pharmacy and Nursing Wilkes University Wilkes-Barre, Pennsylvania

#### Neurology

Michael E. Ernst, Pharm.D., BCPS *Clinical Associate Professor of Medicine* Division of Clinical and Administrative Pharmacy The University of Iowa College of Pharmacy *Clinical Pharmacist* Department of Family Medicine Carver College of Medicine Iowa City, Iowa

#### Psychiatry II

Tami R. Argo, Pharm.D., M.S., BCPP *Clinical Assistant Professor* Division of Pharmacy Practice University of Texas at Austin College of Pharmacy Austin, Texas

> Angela D. Hughes, Pharm.D. *Psychiatric Pharmacy Resident* College of Pharmacy The University of Texas at Austin Austin, Texas

#### Infectious Diseases I

Thomas P. Lodise, Jr., Pharm.D. Assistant Professor Department of Pharmacy Practice Albany College of Pharmacy Albany, New York

Blake Max, Pharm.D. HIV Clinical Pharmacist and Clinical Assistant Professor Ruth M. Rothstein CORE Center Cook County Bureau of Health Services University of Illinois at Chicago College of Pharmacy Chicago, Illinois

Melinda M. Neuhauser, Pharm.D. Clinical Pharmacy Specialist, Infectious Diseases U.S. Department of Veterans Affairs VACO Pharmacy Benefits Management Services Hines, Illinois

#### Infectious Diseases II

Ronald G. Hall II, Pharm.D., BCPS Assistant Professor Department of Pharmacy Practice Texas Tech University Health Sciences Center School of Pharmacy Dallas, Texas NIH Clinical Scholar Department of Clinical Sciences University of Texas Southwestern Medical Center Dallas, Texas Advanced Practice Pharmacist—Infectious Diseases Department of Pharmacy North Texas Veterans Health Care System Dallas, Texas

#### Pulmonary

Hengameh H. Raissy, Pharm.D. Research Assistant Professor of Pediatrics University of New Mexico School of Medicine Albuquerque, New Mexico

#### Critical Care I

Stacy Alan Voils, Pharm.D., BCPS *Clinical Pharmacy Specialist, Critical Care* Department of Pharmacy Services Virginia Commonwealth University Health System Medical College of Virginia Campus Richmond, Virginia

#### **Older Adults**

Justin J. Sherman, M.C.S., Pharm.D. Associate Professor of Pharmacy Practice Department of Clinical and Administrative Sciences University of Louisiana at Monroe College of Pharmacy Monroe, Louisiana

#### Nutrition

Sarah J. Miller, M.S., Pharm.D., BCNSP *Professor* Department of Pharmacy Practice University of Montana Skaggs School of Pharmacy Missoula, Montana

#### Oncology

Sachin Shah, Pharm.D., BCOP Associate Professor Department of Pharmacy Practice Texas Tech University Health Sciences Center – School of Pharmacy Advanced Hematology/Oncology Clinical Pharmacist Pharmacy Department VA North Texas health Care System Dallas/Fort Worth, Texas

#### Chronic Illnesses I

Thomas M. Parker, Pharm.D., CDE Assistant Professor Departments of Pharmacy Practice and Pediatrics Texas Tech University Health Sciences Center School of Pharmacy and School of Medicine Amarillo, Texas

#### Chronic Illnesses II

Susan P. Bruce, Pharm.D., BCPS Associate Professor and Chair Department of Pharmacy Practice Northeastern Ohio Universities Colleges of Medicine and Pharmacy Rootstown, Ohio

#### Chronic Illnesses Ill

Christina L. Aquilante, Pharm.D. Assistant Professor Department of Pharmaceutical Sciences University of Colorado Denver School of Pharmacy Aurora, Colorado

Jayne E. Pawasauskas, Pharm.D., BCPS *Clinical Associate Professor* Department of Pharmacy Practice University of Rhode Island College of Pharmacy Kingston, Rhode Island

### PHARMACOTHERAPY SELF-ASSESSMENT PROGRAM SIXTH EDITION

### **Editorial Board**

Michelle M. Richardson, Pharm.D., FCCP, BCPS (Chair) Special and Scientific Staff William B. Schwartz Division of Nephrology Tufts Medical Center Assistant Professor of Medicine Tufts University School of Medicine Boston, Massachusetts

Clarence Chant, Pharm.D., BCPS *Clinical Pharmacy Specialist* Critical Care/Research, Pharmacy Department St. Michael's Hospital *Assistant Professor* Leslie Dan Faculty of Pharmacy University of Toronto Toronto, Ontario, Canada

Judy W.M. Cheng, Pharm.D., MPH, FCCP, BCPS (AQ Cardiology) Professor of Pharmacy Practice Department of Pharmacy Practice Massachusetts College of Pharmacy and Health Sciences Clinical Pharmacist Department of Pharmacy Brigham and Women's Hospital Boston, Massachusetts

Katherine Hammond Chessman, Pharm.D., FCCP, BCPS, BCNSP Professor Department of Clinical Pharmacy and Outcome Sciences Pediatric Pharmacy Practice Residency Program Director South Carolina College of Pharmacy Medical University of South Carolina Campus Clinical Pharmacy Specialist Pediatrics/Pediatrics Surgery MUSC Children's Hospital Charleston, South Carolina

> Anne L. Hume, Pharm.D., FCCP, BCPS *Professor of Pharmacy* Department of Pharmacy Practice University of Rhode Island Kingston, Rhode Island

Lisa C. Hutchison, Pharm.D., MPH, BCPS Associate Professor Pharmacy Practice University of Arkansas for Medical Sciences Little Rock, Arkansas Adam B. Jackson, Pharm.D., BCPS Clinical Pharmacy Specialist in Infectious Diseases Pharmacy Department Kaiser Permanente—Colorado Region Clinical Assistant Professor University of Colorado Denver School of Pharmacy Denver, Colorado

Emilie L. Karpiuk, Pharm.D., BCPS, BCOP Oncology Pharmacist Department of Pharmacy Froedtert Hospital Milwaukee, Wisconsin

Linda Gore Martin, Pharm.D., M.B.A., BCPS Associate Professor, Social and Administrative Pharmacy School of Pharmacy University of Wyoming Laramie, Wyoming

Todd P. Semla, Pharm.D., M.S., FCCP, BCPS, AGSF *Clinical Pharmacy Specialist* VACO Pharmacy Benefits Management Service U.S. Department of Veterans Affairs Hines, Illinois *Associate Professor* Departments of Medicine and Psychiatry and Behavioral Sciences The Feinberg School of Medicine Northwestern University Chicago, Illinois

## Table of Contents

### Cardiology

Algorithm for treatment of	
hypertension from JNC 7	
Compelling indications for specific	
pharmacotherapy	2
QTc interval monitoring algorithm	3
Evaluation of the acute coronary	
syndrome patient	4
Risk Stratification for non-ST-segment	
elevation acute coronary syndrome	5
Initial pharmacotherapy for ST-segment	
elevation acute coronary syndromes	6
Initial pharmacotherapy for non-ST-segment	
elevation acute coronary syndrome	7
Acute ischemic stroke	
management algorithm	8
Algorithm for chronic management	
of atrial fibrillation	9
Algorithm for selecting antiarrhythmic drug	
therapy for maintenance of sinus rhythm in	
patients with recurrent paroxysmal or	
recurrent persistent atrial fibrillation	
Algorithm for treatment of pulseless	
ventricular tachycardia/ventricular	
fibrillation	
Algorithm for treatment of asystole	
and pulseless electrical activity	

### Nephrology

Algorithm for assessing kidney function	
Anemia treatment algorithm	14
Causes of ESA hyporesponse	
Options for managing chronic kidney	
disease-mineral and bone disorder for	
patients with elevated iPTH and	
normal/low calcium concentrations	16
Decision algorithm for prevention of CIN	17
Approach to the Risk Assessment of CIN	
after Percutaneous Coronary Intervention	18

### Neurology/Psychiatry

Algorithm for the management	
of Parkinson's disease	19
Algorithm for use in guiding the	
treatment of generalized anxiety	
disorder and panic disorder	20
-	

### **Infectious Diseases**

Management of suspected <i>Staphylococcus</i> <i>aureus</i> skin and soft tissue infections	21
Classification of nosocomial pneumonia	
Management strategy recommended by the	
American Thoracic Society/Infectious	
Diseases Society of America guidelines	23
Treatment algorithm for HIV-associated	
dyslipidemia	

### Pulmonary

Management of acute asthma exacerbations25
--

### **Critical Care I**

Overview of BTF guidelines for the	
management of patients with TBI	

### **Older Adults**

Treatment algorithm for chronic prostatitis/	
chronic pelvic pain syndrome	27

### Nutrition

Decision algorithm for evaluation	
of dietary supplement use	

### Oncology

Proposed algorithm for metastatic renal	
cell cancer therapy based on patient	
presentation	29

### **Chronic Illnesses**

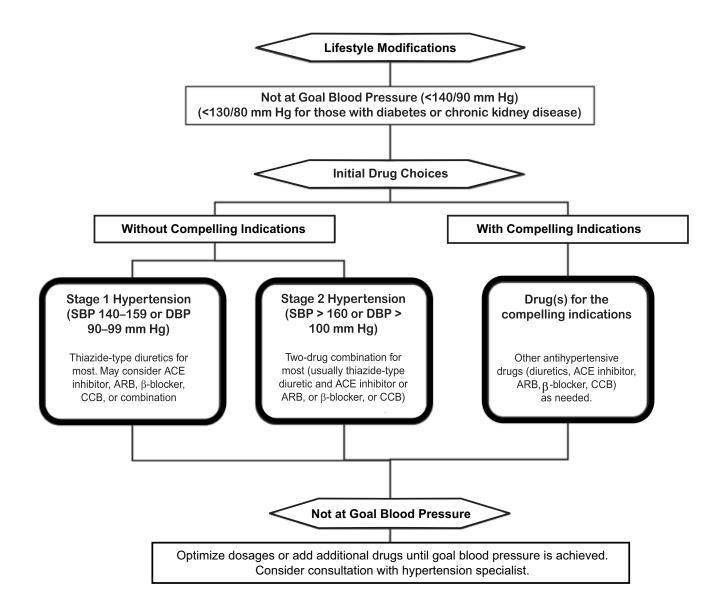
Algorithm for management of a basal/bolus insulin regimen in children and adolescents	30
Treatment algorithm for type 2 diabetes	
mellitus in adolescents	31
Recommendations on indications for the use of	
nonbiologic disease-modifying antirheumatic	
drugs (DMARDs) in rheumatoid arthritis (RA)	
patients who have never received DMARDs	32

Recommendations on indications for the use	
of biologic disease-modifying antirheumatic	
drugs (DMARDs) in patients with rheumatoid	
arthritis (RA)	
Algorithm for the management of	
metabolic syndrome	34
An algorithm for improving drug therapy in	
disabled or frail elderly patients	





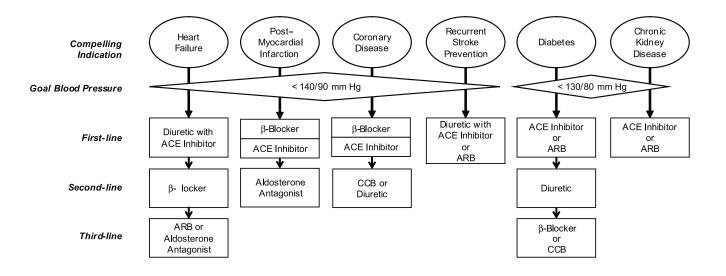
### Algorithm for treatment of hypertension from JNC 7



ACE = angiotensin-converting enzyme; ARB = angiotensin receptor blocker; CCB = calcium channel blocker; DBP = diastolic blood pressure; JNC 7 = Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure. Chobanian AV, Bakris GL, Black HR, Cushman WC, Green LA, Izzo JL Jr, et al; Joint National Committee on Prevention, Detection, evaluation, and

# Cardiology

#### Compelling indications for specific pharmacotherapy



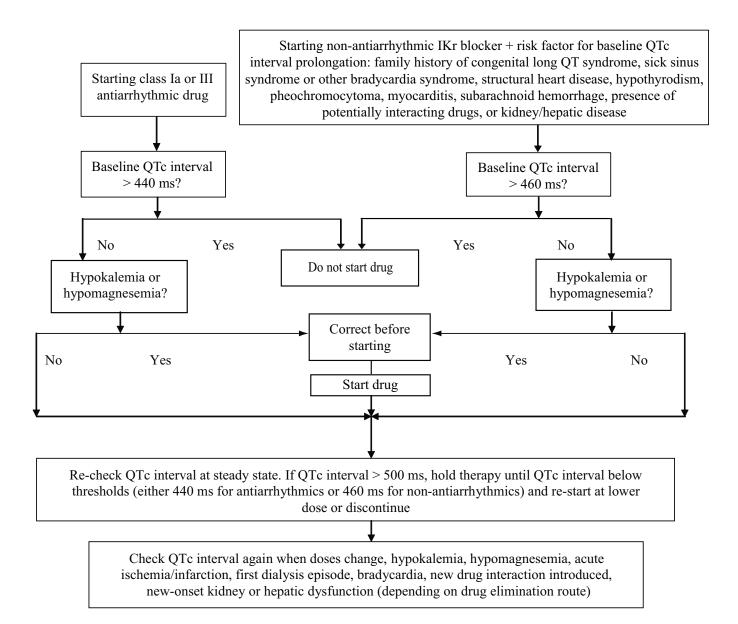
Recommendations are based on evidence demonstrating reduced morbidity and/or mortality related to the compelling indication with recommended pharmacotherapy and adapted from JNC 7 recommendations. Blood pressure should be managed concurrently with the compelling indication using these drugs when possible.

ACE = angiotensin-converting enzyme; ARB = angiotensin receptor blocker; CCB = calcium channel blocker; JNC 7 = Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure.

Chobanian AV, Bakris GL, Black HR, Cushman WC, Green LA, Izzo JL Jr, et al; Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure. National Heart, Lung, and Blood Institute; National High Blood Pressure Education Program Coordinating Committee. Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure. Hypertension 2003;42:1206–52.

# **Cardiology**

### QTc interval monitoring algorithm

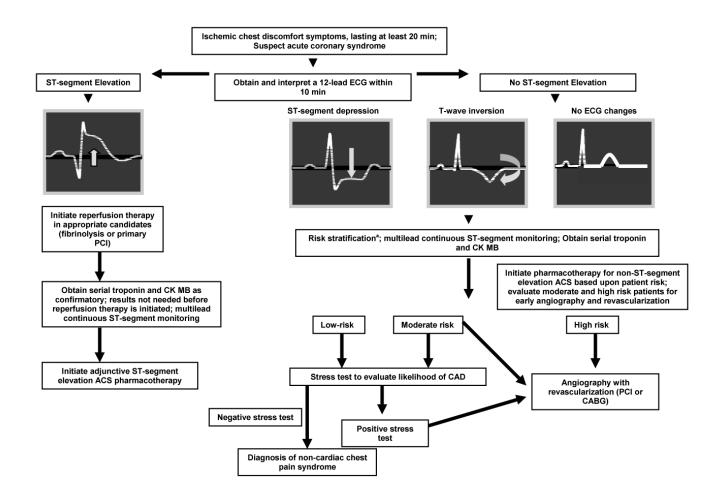


IKr = rapid component of the delayed rectifier potassium channel; ms = millisecond; QTc interval = corrected QT interval.



PSAP

#### Evaluation of the acute coronary syndrome patient



<sup>a</sup>As described in table on p. 5.

ACS = acute coronary syndrome; CABG = coronary artery bypass graft surgery; CAD = coronary artery disease; CK, MB = creatine kinase, myocardial bound; ECG = electrocardiogram; PCI = percutaneous coronary intervention.

Adapted with permission from McGraw-Hill. Spinler SA, de Denus S. Acute coronary syndromes. In: DiPiro JT, Yee GC, Matzke GR, Wells BG, Posey LM, eds. Pharmacotherapy: The Pathophysiologic Approach, 6th ed. New York: McGraw-Hill, 2005:291–320.

## **Cardiology**



#### Risk stratification for non-ST-segment elevation acute coronary syndrome

Using the TIMI Risk Score		
Past Medical History	Clinical Presentation	
<ul> <li>✓ Age = 65 years</li> <li>✓ ≥ = 3 Risk Factors for CAD Hypercholesterolemia HTN DM Smoking Family history of premature CHD<sup>a</sup></li> </ul>	<ul> <li>✓ ST-segment depression (≥ = 0.5 mm)</li> <li>✓ ≥ = 2 episodes of chest discomfort within the past 24 hours</li> <li>✓ Positive biochemical marker for infarction<sup>b</sup></li> </ul>	
✓ Known CAD (= 50% stenosis of coronary artery)		
<ul> <li>Use of aspirin within the past 7 days</li> </ul>		
One point is assigned for each of the seven medical history and clinical presentation findings. The score (point) total is calculated and the patient is assigned a risk for experiencing the composite and point of death muccardial information or urgent need for rayassularization as		

One point is assigned for each of the seven medical history and clinical presentation findings. The score (point) total is calculated and the patient is assigned a risk for experiencing the composite end point of death, myocardial infarction or urgent need for revascularization as follows:

High-Risk	Medium Risk	Low-Risk	
TIMI Risk Score 5–7 poin	ts TIMI Risk Score 3–4 points	TIMI Risk Score 0–2 points	
Other Ways to Identify High-Risk Patients:			
Other findings which alone, or in combinatio	n, may identify a patient at high risk of d	eath or MI:	
ST-segment depression			
Positive biochemical marker for infarction <sup>b</sup>			
Deep symmetric T-wave inversions ( $\geq 2 \text{ mm}$ )			
Acute heart failure			
Diabetes mellitus			
Chronic kidney disease			
Recent myocardial infarction (within the past	2 weeks)		

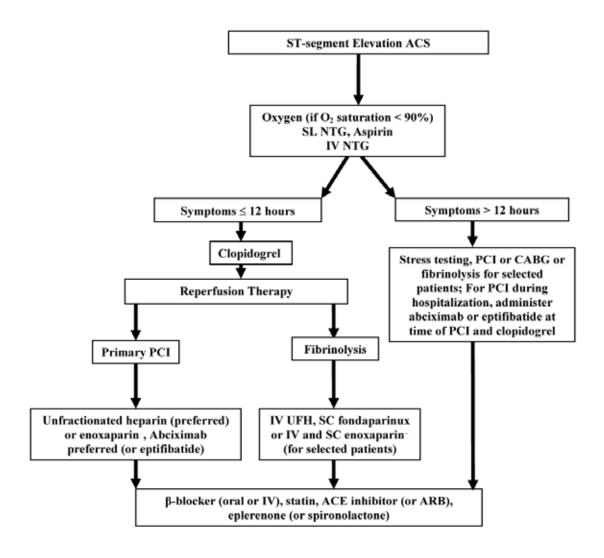
<sup>a</sup>As defined by the National Cholesterol Education Program Adult Treatment Panel III Report (2001): the presence of coronary heart disease in a first degree male relative younger than age 55 or a first-degree female relative younger than age 65.

<sup>b</sup>A positive biochemical marker for infarction is a value of troponin I, troponin T or creatinine kinase MB of greater than the myocardial infarction detection limit. CAD = coronary artery disease; CHD = coronary heart disease; HTN = hypertension; TIMI = Thrombolysis in Myocardial Infarction.

Adapted with permission from McGraw-Hill. Spinler SA, de Denus S. Acute coronary syndromes. In: DiPiro JT, Yee GC, Matzke GR, Wells BG, Posey LM, eds. Pharmacotherapy: The Pathophysiologic Approach, 6th ed. New York, NY: McGraw-Hill, 2005:291–320.



## Initial pharmacotherapy for ST-segment elevation acute coronary syndromes

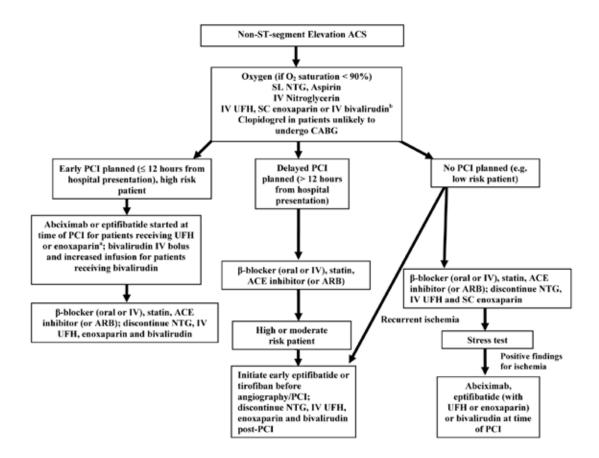


ACE = angiotensin enzyme; ACS = acute coronary syndrome; ARB = angiotensin receptor blocker; CABG = coronary artery bypass graft surgery; IV = intravenous; NTG = nitroglycerin;  $O_2$  = oxygen; PCI = percutaneous coronary intervention; SC = subcutaneously; SL = sublingual; UFH = unfractionated heparin.





## Initial pharmacotherapy for non-ST-segment elevation acute coronary syndrome



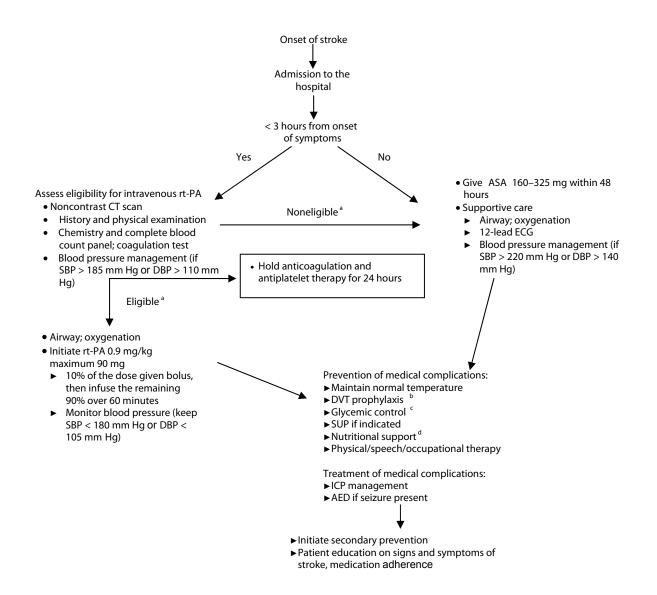
<sup>a</sup>May require supplemental IV dose of enoxaparin.

<sup>b</sup>Fondaparinux may be used as anticoagulant if no PCI planned.

ACE = angiotensin-converting enzyme; ARB = angiotensin receptor blocker; ACS = acute coronary syndrome; CABG = coronary artery bypass graft surgery; IV = intravenous; NTG = nitroglycerin;  $O_2$  = oxygen; PCI = percutaneous coronary intervention; SC = subcutaneous; SL = sublingual; UFH = unfractionated heparin.

# Cardiology

#### Acute ischemic stroke management algorithm



<sup>a</sup>Eligibility for intravenous rt-PA: onset within 3 hours; negative for hemorrhagic stroke or history; blood pressure < 185/110 mm Hg; no head trauma, prior stroke, or myocardial infarction in the past 3 months; no history or gastrointestinal or urinary tract hemorrhage in the past 21 days; no major surgery in the previous 14 days; not taking oral anticoagulant or INR  $\leq$  1.7; platelet count  $\geq$  100,000/mm<sup>3</sup>; blood glucose  $\geq$  50 mg/dL; no seizure with postictal residual neurologic impairments at onset of stroke.

<sup>b</sup>All pharmacologic prophylaxis should be held for 24 hours post-thrombolytic therapy.

 $^{\circ}$ Hyperglycemic control should be managed judiciously. The consensus is to treat when glucose > 300 mg/dL. The role of tight glucose control is currently unknown in patients with stroke.

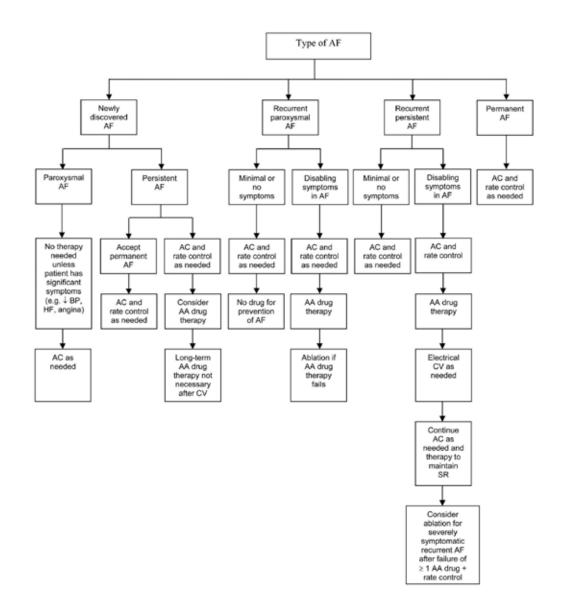
<sup>d</sup>All patients should receive prompt swallow evaluation. If patients fail swallow evaluation, enteral or parenteral nutrition should be initiated.

AED = antiepileptic drug; ASA = aspirin; CT = computed tomography; DBP = diastolic blood pressure; DVT = deep vein thrombosis; ECG = electrocardiogram; ICP = intracranial pressure; INR = international normalized ratio; rt-PA = recombinant tissue plasminogen activator; SBP = systolic blood pressure; SUP = stress ulcer prophylaxis.





#### Algorithm for chronic management of atrial fibrillation

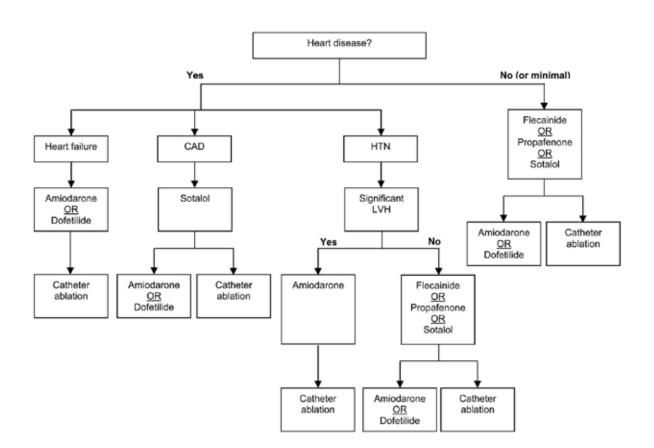


AA = antiarrhythmic; AC = anticoagulation; AF = atrial fibrillation; BP = blood pressure; CV = cardioversion; HF = heart failure; SR = sinus rhythm. Adapted from the American College of Cardiology and American Heart Association. Fuster V, Rydén LE, Cannom DS, Crijns HJ, Curtis AB, Ellenbogen KA, et al. ACC/AHA/ESC 2006 guidelines for the management of patients with atrial fibrillation: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines and the European Society of Cardiology Committee for Practice Guidelines (Writing Committee to Revise the 2001 Guidelines for the Management of Patients with Atrial Fibrillation). J Am Coll Cardiol 2006;48:e149–246.

# **Cardiology**



Algorithm for selecting antiarrhythmic drug therapy for maintenance of sinus rhythm in patients with recurrent paroxysmal or recurrent persistent atrial fibrillation<sup>a</sup>



<sup>a</sup>Within each of the boxes, the drugs are listed alphabetically, not in order of suggested use. However, the sequence of the boxes does imply the order of suggested use.

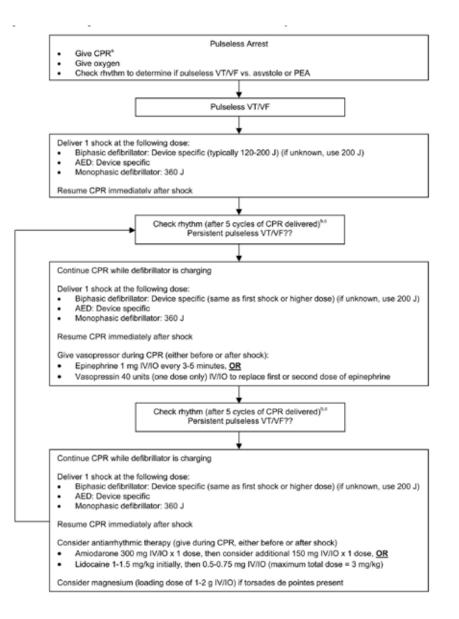
CAD = coronary artery disease; HTN = hypertension; LVH = left ventricular hypertrophy.

Adapted from the American College of Cardiology and American Heart Association. Fuster V, Rydén LE, Cannom DS, Crijns HJ, Curtis AB, Ellenbogen KA, et al. ACC/AHA/ESC 2006 guidelines for the management of patients with atrial fibrillation: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines and the European Society of Cardiology Committee for Practice Guidelines (Writing Committee to Revise the 2001 Guidelines for the Management of Patients with Atrial Fibrillation). J Am Coll Cardiol 2006;48:e149–246.





## Algorithm for treatment of pulseless ventricular tachycardia/ventricular fibrillation



<sup>a</sup>If arrest is witnessed and defibrillator is readily available, 2 rescue breaths can be given before defibrillation. If arrest is unwitnessed, 5 cycles of CPR should be administered before defibrillation.

<sup>b</sup>One cycle of CPR = 30 chest compressions, then 2 breaths; 5 cycles about 2 minutes.

<sup>c</sup>After advanced airway established, cycles of CPR no longer need to be given. Instead, continuous chest compressions should be given without pauses for breaths. Give 8–10 breaths/minute.

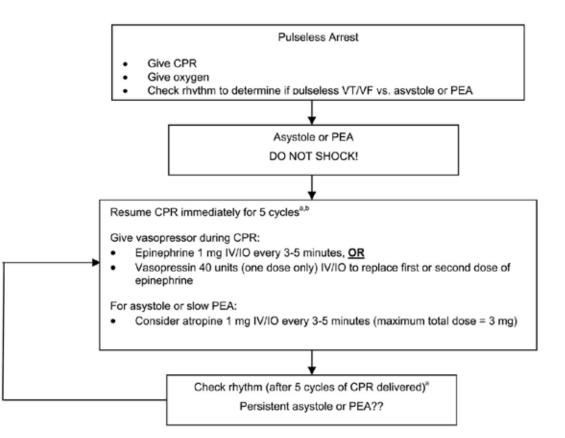
AED = automated external defibrillator; CPR = cardiopulmonary resuscitation; IO = intraosseous; IV = intravenous; J = joules; PEA = pulseless electrical activity; VF = ventricular fibrillation; VT = ventricular tachycardia.

Adapted from the American Heart Association. 2005 American Heart Association Guidelines for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care. Circulation 2005;112(Suppl 1):IV-58–IV-66.

# Cardiology



#### Algorithm for treatment of asystole and pulseless electrical activity



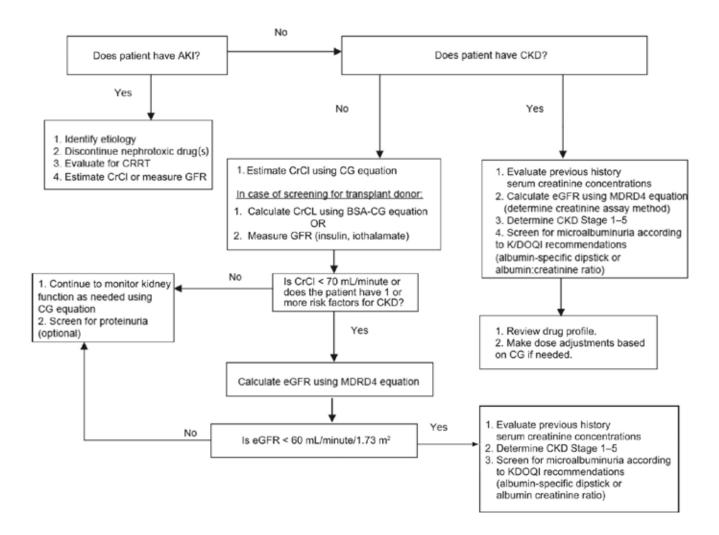
<sup>a</sup>One cycle of CPR = 30 chest compressions; then, 2 breaths; 5 cycles about 2 minutes.

Adapted from the American Heart Association. 2005 American Heart Association Guidelines for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care. Circulation 2005;112(suppl 1):IV-58–IV-66.

<sup>&</sup>lt;sup>b</sup>After advanced airway established, cycles of CPR no longer need to be given. Instead, continuous chest compressions should be given without pauses for breaths. Give 8–10 breaths/minute.

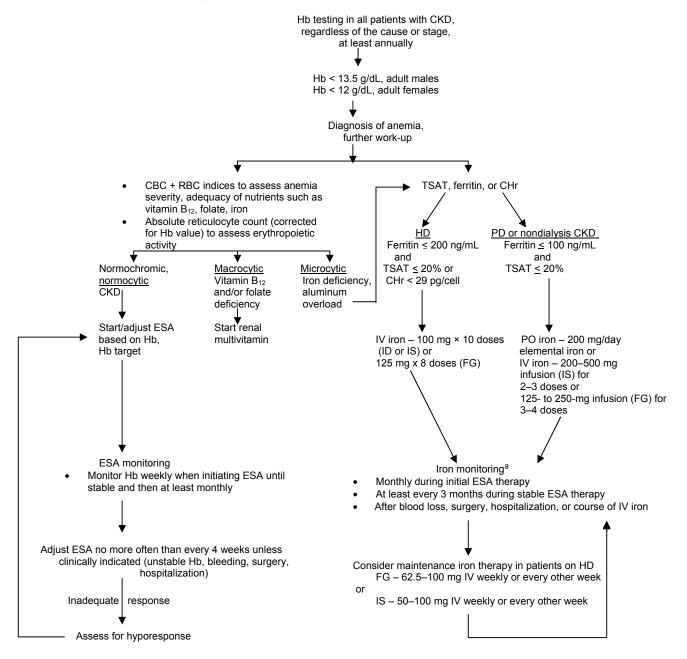
CPR = cardiopulmonary resuscitation; IO = intraosseous; IV = intravenous; PEA = pulseless electrical activity; VF = ventricular fibrillation; VT = ventricular tachycardia.

### Algorithm for assessing kidney function



AKI = acute kidney injury; BSA = body surface area; CG = Cockcroft-Gault; CKD = chronic kidney disease; CrCl = creatinine clearance; CRRT = continuous renal replacement therapy; eGFR = estimated glomerular filtration rate; GFR = glomerular filtration rate; KDOQI = Kidney Disease Outcomes Quality Initiative; MDRD4 = 4-variable Modification of Diet in Renal Disease.

#### Anemia treatment algorithm

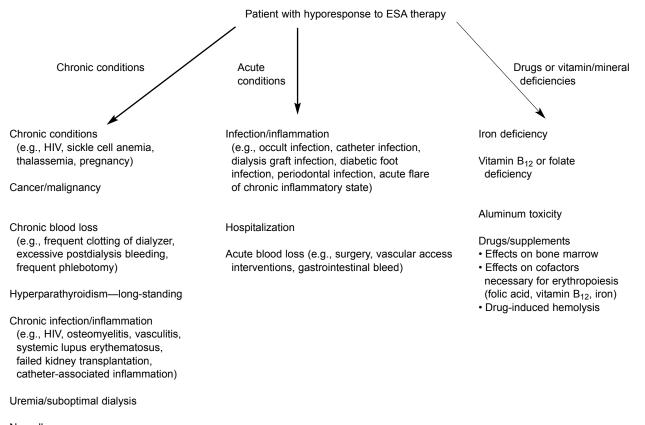


<sup>a</sup>Wait 1-2 weeks to evaluate iron status if intravenous doses greater than 200 mg are administered.

CBC = complete blood cell (count); CHr = reticulocyte hemoglobin; CKD = chronic kidney disease; ESA = erythropoiesis-stimulating agent; FG = ferric gluconate; Hb = hemoglobin; HD = hemodialysis; ID = iron dextran; IS = iron sucrose; IV = intravenous; PD = peritoneal dialysis; PO = by mouth; RBC = red blood cell; TSAT = transferrin saturation.

Adapted from the American College of Clinical Pharmacy. Stamatakis MK. Chronic kidney disease. In: Mueller BA, Bertch KE, Dunsworth TS, Fagan SC, Hayney MS, O'Connell MB, et al, eds. Pharmacotherapy Self-Assessment Program, 4th ed. Nephrology Module. Kansas City; MO: American College of Clinical Pharmacy, 2003:187.

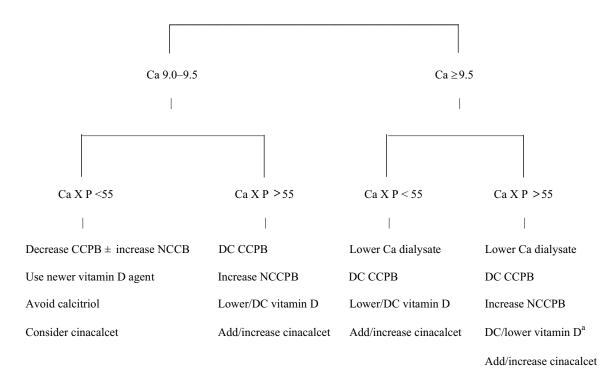
#### **Causes of ESA hyporesponse**



Nonadherence

ESA = erythropoiesis-stimulating agent; HIV = human immunodeficiency virus.

Options for managing chronic kidney disease-mineral and bone disorder for patients with elevated iPTH and normal/low calcium concentrations



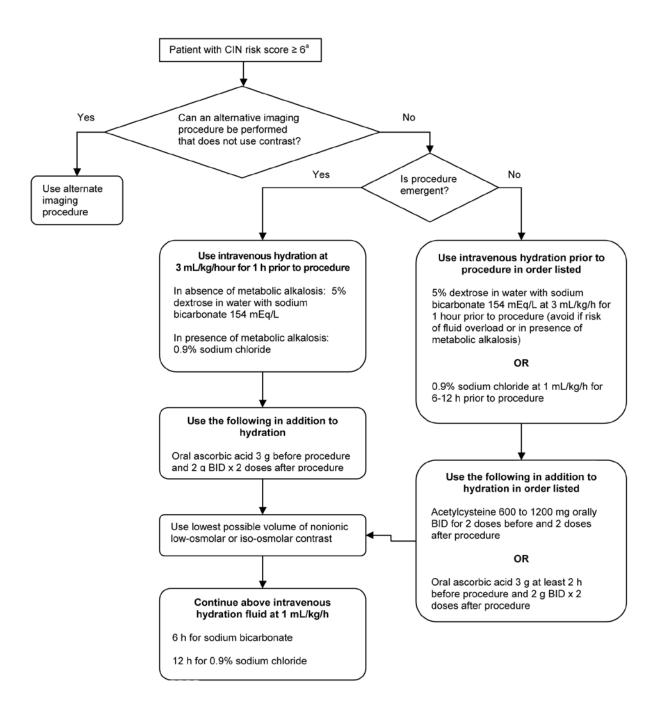
<sup>a</sup>Lower/DC vitamin D = decrease dose or discontinue calcitriol/paricalcitol/doxercalciferol until Ca X P  $\leq$  55 mg<sup>2</sup>/dL<sup>2</sup>.

Ca = calcium mg/dL;  $Ca \times P = calcium/phosphate product mg^2/dL^2$ ; CCPB = calcium-containing phosphate binder; DC = discontinue; NCCPB = noncalcium containing phosphate binder; P = phosphorus in mg/dL.





#### **Decision algorithm for prevention of CIN**



<sup>a</sup>Based on risk score calculated using table on p. 18. BID = 2 times/day; CIN = contrast-induced nephropathy.



#### Approach to the risk assessment of CIN after percutaneous coronary intervention

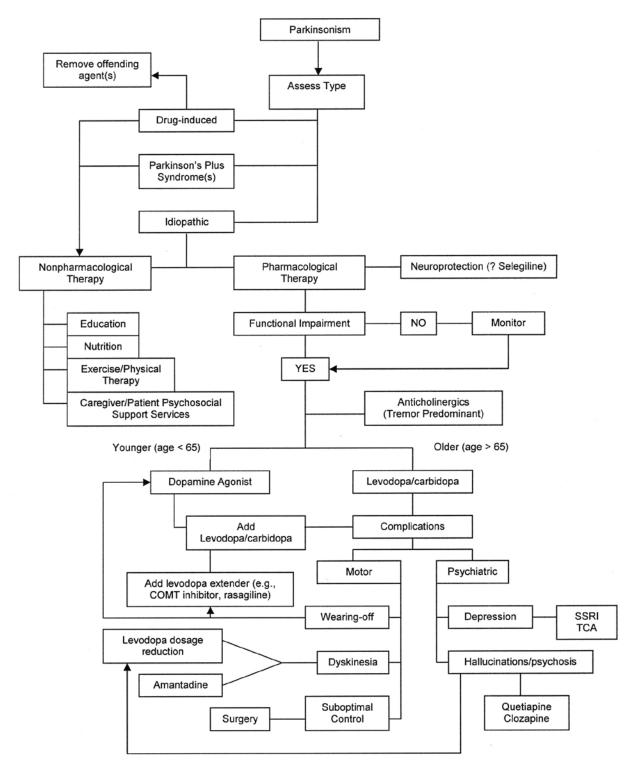
Risk Factor	Definition		Risk Score	
Hypotension	SBP < 80 mm Hg for at leas support or IABP within 24	st 1 hour requiring inotropic hours periprocedural	5	
Use of IABP			5	
CHF	NYHA class III/IV and/or h	istory of pulmonary edema	5	
Elderly	Age > 75 years		4	
Anemia	Het $< 39\%$ for men or Het $< 36\%$ for women		3	
Diabetes mellitus			3	
Contrast volume	For every 100 mL		1	
Kidney disease				
Baseline SCr	SCr >1.5		4	
OR				
Baseline GFR <sup>a</sup>	40-59 mL/minute/1.73 m <sup>2</sup>		2	
	20-39 mL/minute/1.73 m <sup>2</sup>		4	
	< 20 mL/minute/1.73 m <sup>2</sup>		6	
<b>Risk Assessment</b>				
Total Risk Score	Risk Group	Risk of CIN	Risk of Dialysis	
≤ 5	Low	7.5%	0.04%	
6–10	Moderate	14%	0.12%	
11–15	High	26.1%	1.09%	
≥ 16	Very high	57.3%	12.5%	

<sup>a</sup>Estimated using the Modification of Diet in Renal Disease Equation. CIN = contrast-induced nephropathy; CHF = congestive heart failure; GFR = glomerular filtration rate; Hct = hematocrit; IABP = intra-arterial balloon pump; NYHA = New York Heart Association; SBP = systolic blood pressure; SCr = serum creatinine. Reprinted with permission from Elsevier. Mehran R, Aymong ED, Nikolsky E, Lasic Z, Iakovou I, Fahy M, et al. A simple risk score for prediction of

contrast-induced nephropathy after percutaneous coronary intervention. J Am Coll Cardiol 2004;44:1393-9.



### Algorithm for the management of Parkinson's disease

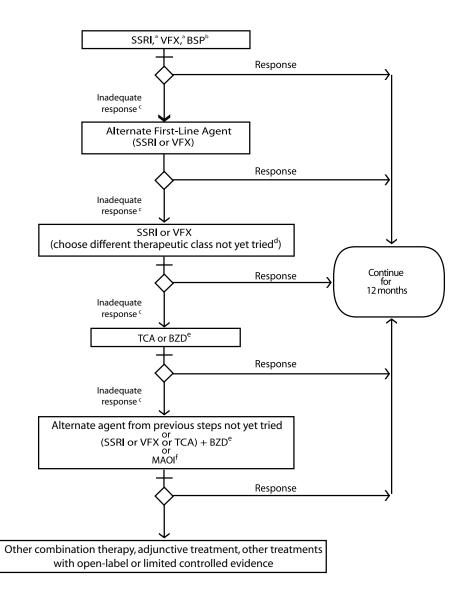


COMT = catechol-O-methyl-transferase; SSRI = selective serotonin reuptake inhibitor; TCA = tricyclic antidepressant.

American College of Clinical Pharmacy



Algorithm for use in guiding the treatment of generalized anxiety disorder and panic disorder



<sup>a</sup> If acute relief needed or increase in anxiety or insomnia on initiation, may overlap with a short term (2-4 weeks) BZD.

- <sup>c</sup>After trial of 8–12 weeks at adequate dose, may try increasing dose if good tolerability.
- <sup>d</sup> Try VFX if not yet tried, or try and alternate SSRI therapy if patient has not responded to an adequate trial of both an SSRI and VFX.
- <sup>e</sup> BZD not recommended in patients with past or current comorbid substance abuse or dependence.

<sup>&</sup>lt;sup>b</sup>Only for initial, nonrefractory treatment of GAD.

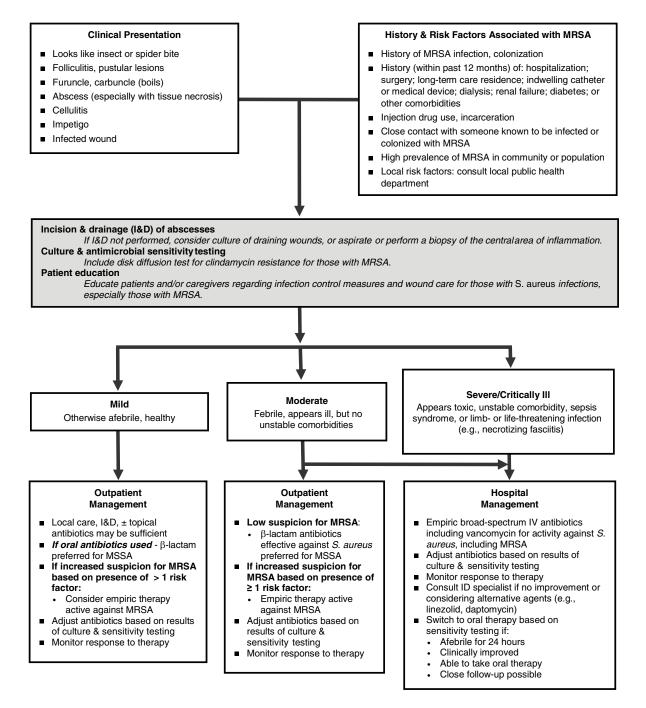
<sup>&</sup>lt;sup>f</sup>Only for treatment of PD.

BSP = buspirone; BZD = benzodiazepine; GAD = generalized anxiety disorder; MAOI = monoamine oxidase inhibitor; PD = panic disorder; SSRI = selective serotonin reuptake inhibitor; TCA = tricyclic antidepressant; VFX = venlafaxine.

# Infectious Diseases



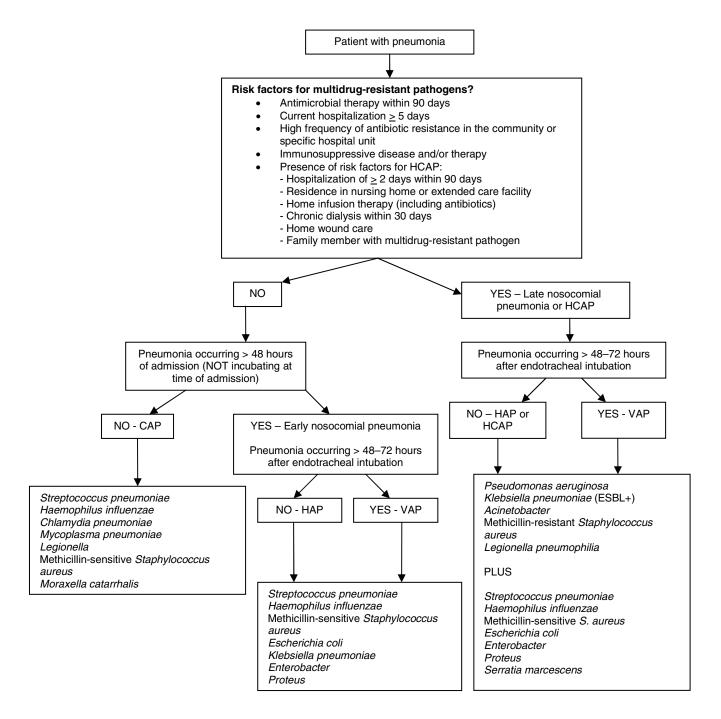
## Management of suspected *Staphylococcus aureus* skin and soft tissue infections



Adapted with permission from Interim guidelines for evaluation & management of community-associated methicillin-resistant *Staphylococcus aureus* skin and soft tissue infections in outpatient settings, which was developed collaboratively by the Infectious Diseases Society of Washington and Public Health—Seattle and King County, Tacoma-Pierce County Department of Health, and Washington State Department of Health. ID = infectious diseases; IV = intravenous; MRSA = methicillin-resistant *S. aureus*; MSSA = methicillin-sensitive *S. aureus*.

# Infectious Diseases

#### Classification of nosocomial pneumonia



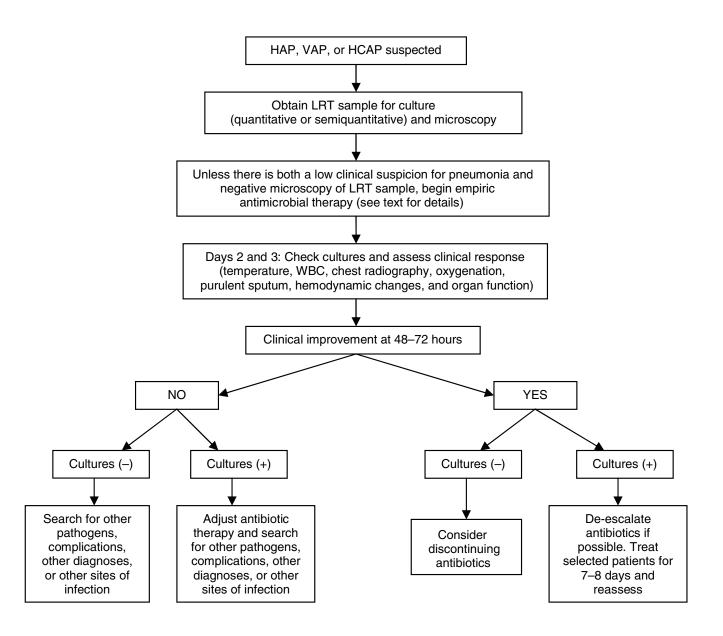
CAP = community-acquired pneumonia; ESBL = extended-spectrum  $\beta$ -lactamase; HAP = hospital-acquired pneumonia; HCAP = health care-associated pneumonia; VAP = ventilator-associated pneumonia.

Adapted with permission from: Guidelines for the management of adults with hospital-acquired, ventilator-associated, and healthcare-associated pneumonia. Am J Respir Crit Care Med 2005;171:388–416.

# Infectious Diseases



#### Management strategy recommended by the American Thoracic Society/ Infectious Diseases Society of America guidelines

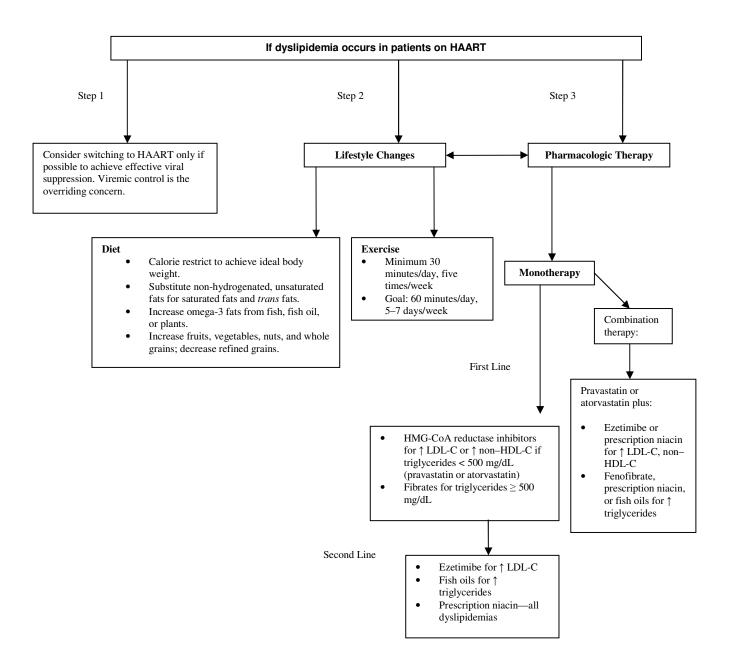


HAP = hospital-acquired pneumonia; HCAP = health care-associated pneumonia; LRT = lower respiratory tract; VAP = ventilator-associated pneumonia; WBC = white blood cell count.

Adapted with permission from: Guidelines for the management of adults with hospital-acquired, ventilator-associated, and healthcare-associated pneumonia. Am J Respir Crit Care Med 2005;171:388–416.



### Treatment algorithm for HIV-associated dyslipidemia



HAART = highly active antiretroviral therapy; HDL-C = high-density lipoprotein cholesterol; HIV = human immunodeficiency virus; HMG-CoA = 3-hydroxy-3-methylglutaryl coenzyme A; LDL-C = low-density lipoprotein cholesterol.

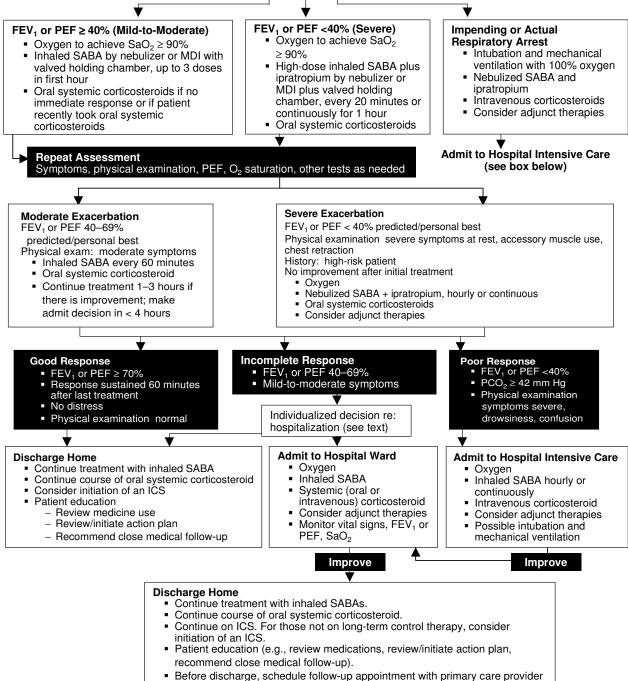
Adapted with permission from Stein JH. Managing cardiovascular risk in patients with HIV infection. J Acquir Immune Defic Syndr 2005;38:115-23.

## Pulmonary



#### Management of acute asthma exacerbations

**Initial Assessment :** Brief history, physical examination (auscultation, use of accessory muscles, heart rate, respiratory rate), PEF or FEV<sub>1</sub>, oxygen saturation, and other tests as indicated



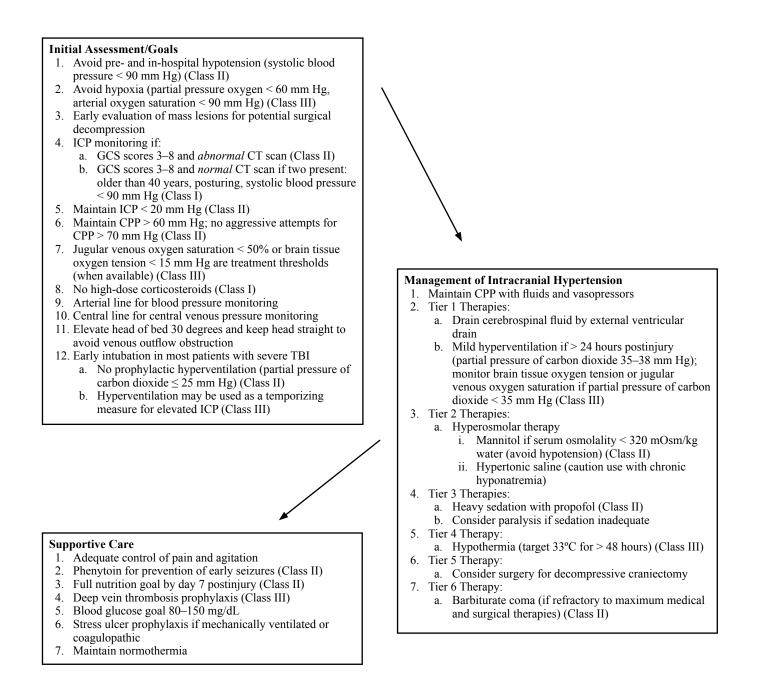
and/or asthma specialist in 1–4 weeks.

Reprinted from NHLBI National Asthma Education and Prevention Program, Expert Panel Report 3. Guidelines for the Diagnosis and Management of Asthma. NIH Publication No. 07-4051. Bethesda, MD: U.S. Department of Health and Human Services, 2007.

# Critical Care



#### Overview of BTF guidelines for the management of patients with TBI

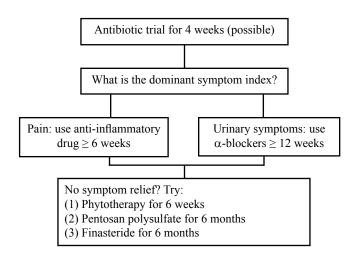


Class I = good-quality randomized controlled trial; Class II = moderate-quality randomized controlled trial, good-quality cohort, or good-quality case control; Class III = poor-quality randomized controlled trial, moderate/poor-quality cohort, moderate/poor-quality case control or database/registry review and case series.

BTF = Brain Trauma Foundation; CPP = cerebral perfusion pressure; CT = computed tomography; GCS = Glasgow Coma Scale; ICP = intracranial pressure; TBI = traumatic brain injury.

# Older Adults

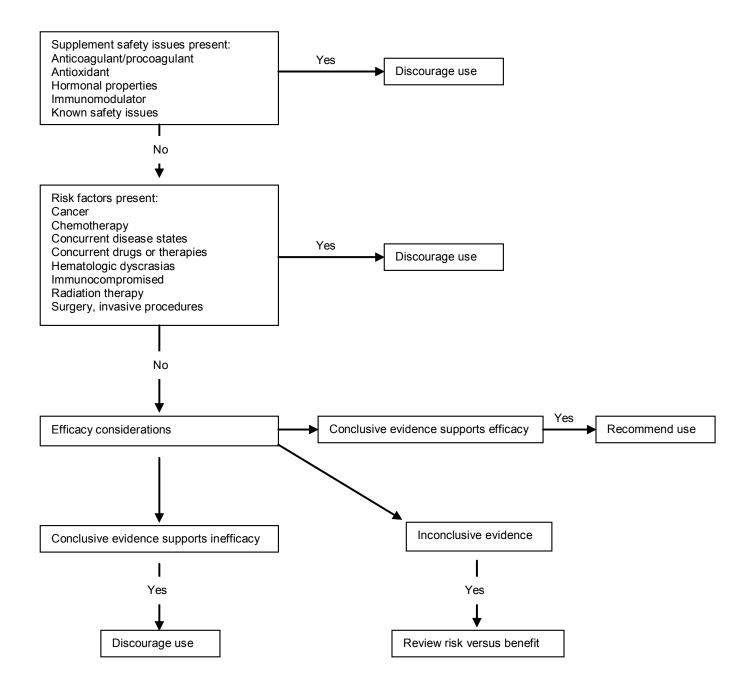
### Treatment algorithm for chronic prostatitis/chronic pelvic pain syndrome



# **Nutrition**



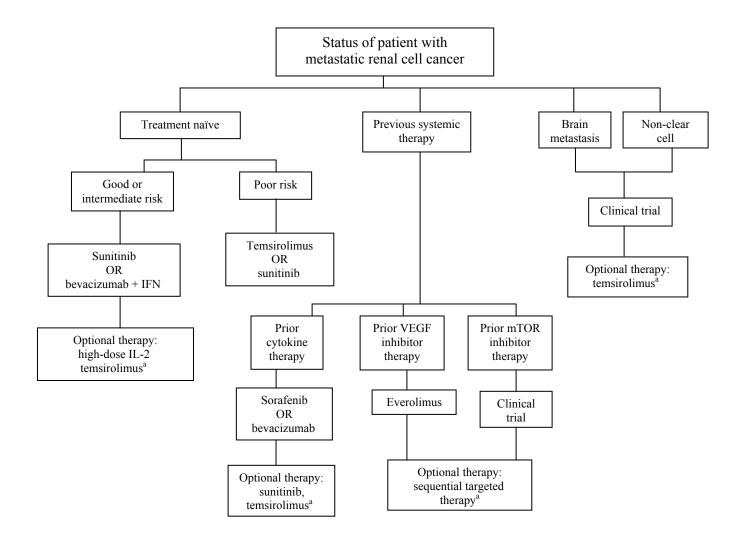
### Decision algorithm for evaluation of dietary supplement use







## Proposed algorithm for metastatic renal cell cancer therapy based on patient presentation

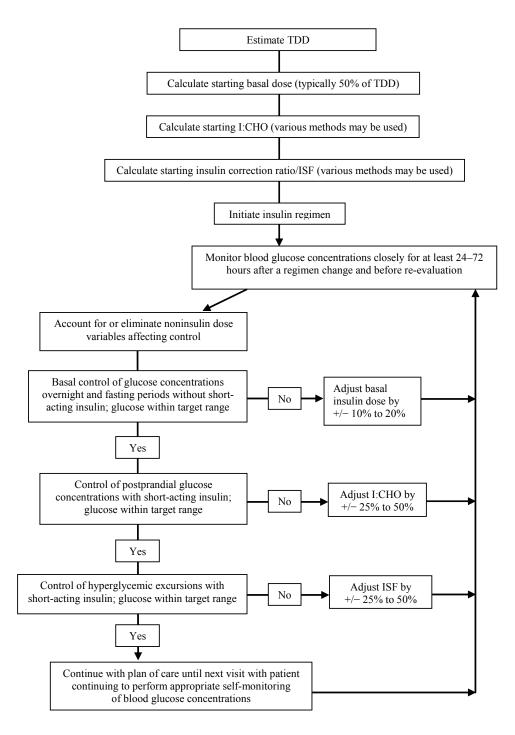


<sup>a</sup>May consider, although limited clinical evidence.

IFN = interferon alfa; IL-2 = interleukin-2 (aldesleukin); mTOR = mammalian target of rapamycin; VEGF = vascular endothelial growth factor.



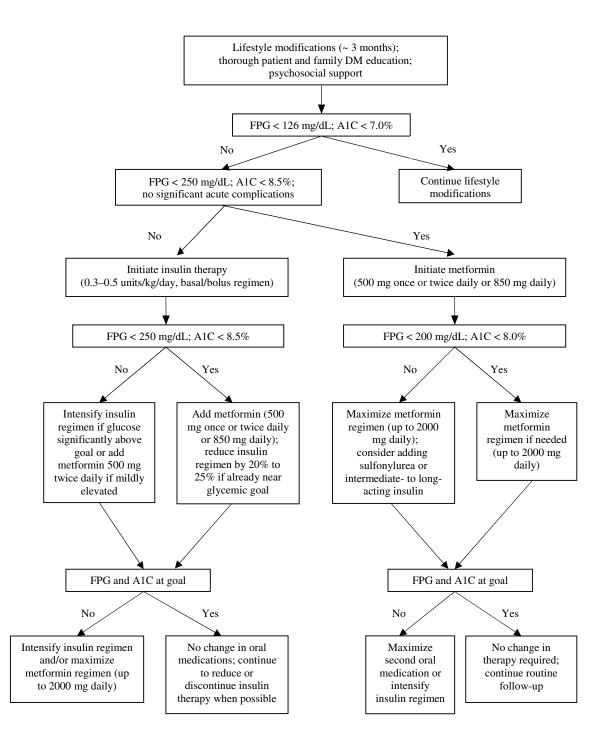
## Algorithm for management of a basal/bolus insulin regimen in children and adolescents



I:CHO = insulin-to-carbohydrate ratio; ISF = insulin sensitivity factor; TDD = total daily insulin dosage or requirement.



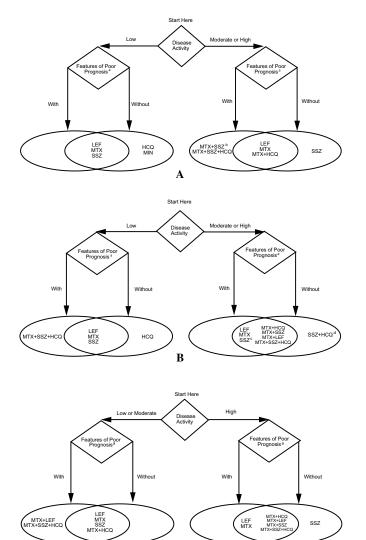
#### Treatment algorithm for type 2 diabetes mellitus in adolescents

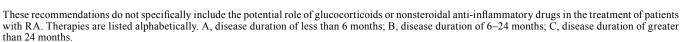


A1C = hemoglobin A1C; DM = diabetes mellitus; FPG = fasting plasma glucose.

# Chronic Illnesses

Recommendations on indications for the use of nonbiologic diseasemodifying antirheumatic drugs (DMARDs) in rheumatoid arthritis (RA) patients who have never received DMARDs





С

aIncludes functional limitation (defined using standard measurement scales such as Health Assessment Questionnaire score or variations of this scale), extra-articular disease (e.g., presence of rheumatoid nodules, secondary Sjögren's syndrome, RA vasculitis, Felty's syndrome, RA lung disease), rheumatoid factor positivity, positive anticyclic citrullinated peptide antibodies, or bony erosions by radiography.

<sup>b</sup>Recommended only for patients with high disease activity with features of poor prognosis.

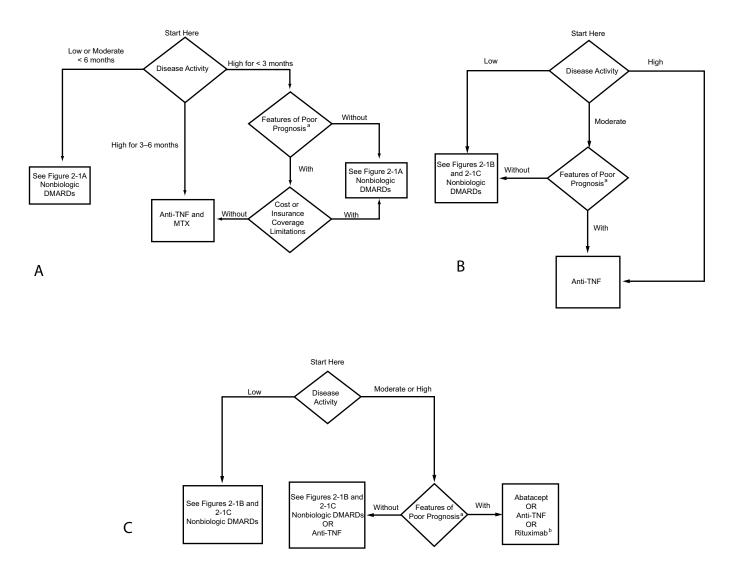
<sup>c</sup> Recommended only for patients with moderate disease activity irrespective of prognostic features and patients with high disease activity without features of poor prognosis.

dOnly recommended for patients with high disease activity without features of poor prognosis.

HCQ = hydroxychloroquine; LEF = leflunomide; MIN = minocycline; MTX = methotrexate; SSZ = sulfasalazine. Reprinted with permission from Saag KG, Teng GG, Patkar NM, Anuntiyo J, Finney C, Curtis JR, et al. American College of Rheumatology 2008 recommendations for the use of nonbiologic and biologic disease-modifying antirheumatic drugs in rheumatoid arthritis. Arthritis Rheum 2008;59:762-84. Available at www.rheumatology.org/publications/guidelines/recommendations.asp?aud=mem. Accessed May 11, 2009.

# Chronic Illnesses

## Recommendations on indications for the use of biologic disease-modifying antirheumatic drugs (DMARDs) in patients with rheumatoid arthritis (RA)



These recommendations do not specifically include the potential role of glucocorticoids or nonsteroidal anti-inflammatory drugs in the treatment of patients with RA. Therapies are listed alphabetically. A, patients with RA less than 6 months; B, patients with RA for 6 months or longer whose prior MTX monotherapy failed; C, patients with RA disease 6 months or longer whose prior MTX combination therapy failed or after sequential administration of other nonbiologic DMARDs.

<sup>a</sup>Includes functional limitation (defined using standard measurement scales such as Health Assessment Questionnaire score or variations of this scale), extraarticular disease (e.g., presence of rheumatoid nodules, secondary Sjögren's syndrome, RA vasculitis, Felty's syndrome, RA lung disease), rheumatoid factor positivity, positive anticyclic citrullinated peptide antibodies, or bony erosions by radiography.

<sup>b</sup>Recommended only for patients with high disease activity with features of poor prognosis.

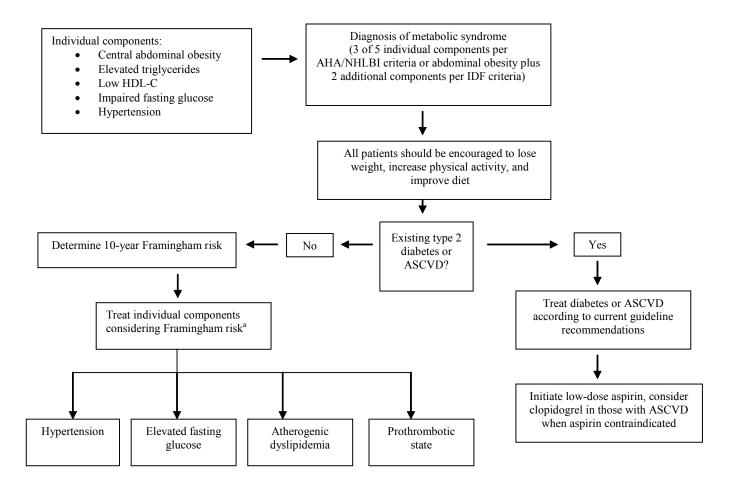
MTX = methotrexate; TNF = tumor necrosis factor.

Reprinted with permission from Saag KG, Teng GG, Patkar NM, Anuntiyo J, Finney C, Curtis JR, et al. American College of Rheumatology 2008 recommendations for the use of nonbiologic and biologic disease-modifying antirheumatic drugs in rheumatoid arthritis. Arthritis Rheum 2008;59:762–84. Available at *www.rheumatology.org/publications/guidelines/recommendations.asp?aud=mem*. Accessed May 11, 2009.

# Chronic Illnesses



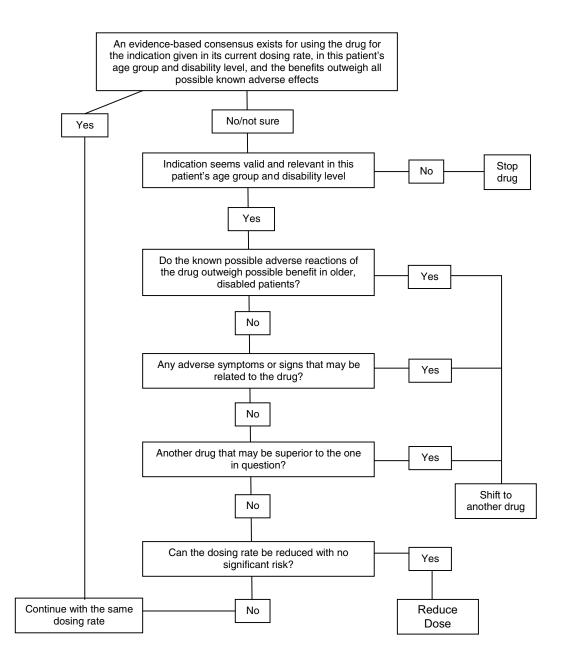
#### Algorithm for the management of metabolic syndrome



<sup>a</sup>Refer to Table 1-2.

AHA/NHLBI = American Heart Association/National Heart, Lung and Blood Institute; ASCVD = atherosclerotic cardiovascular disease; HDL-C = high-density lipoprotein cholesterol; IDF = International Diabetes Federation.

### An algorithm for improving drug therapy in disabled or frail elderly patients



Reproduced with permission from Garfinkel D, Zur-Gil S, Ben-Israel J. The war against polypharmacy: a new cost-effective geriatricpalliative approach for improving drug therapy in disabled elderly people. Isr Med Assoc J 2007;9:430–4. Available at *www.ima.org.il/imaj/ dynamic/web/ArtFromPubmed.asp?year=2007&month=06&page=430*. Accessed May 13, 2009.