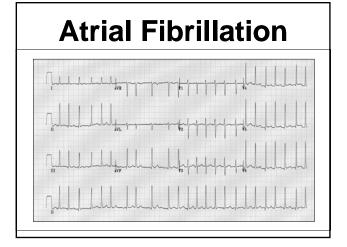
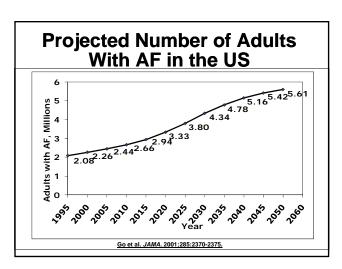
Atrial Fibrillation

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Division of Cardiovascular Medicine,
Electrophysiology
Ohio State University Medical Center

Learning Objectives

- Review the growing incidence and importance of AF in the population
- Discuss the use of anticoagulation in AF for stroke prevention
- Summarize pharmacologic and nonpharmacologic options for AF management





Costs to the Health Care System

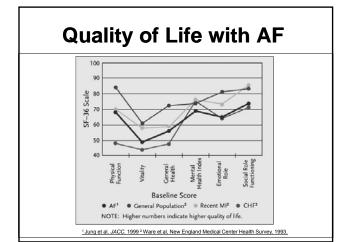
Estimated US cost burden 15.7 billion annually

- · 35% of arrhythmia hospitalizations
- Average hospital stay = 5 days
- Mean cost of hospitalization = \$18,800
- · Does not include:

Costs of outpatient cardioversions

Costs of drugs/side effects/monitoring

Costs of AF-induced strokes



Diagnostic Evaluation

Minimum Evaluation

- History and physical Sx with AF, CV disease
- Electrocardiogram LVH, MI, BBB, WPW
- Echocardiogram LVH, LAE, LVEF, Valves
- · Labs TSH, Renal fxn
- · Sleep history

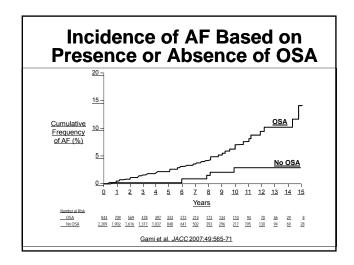
AHA / ACC / ECS Guidelines 2006

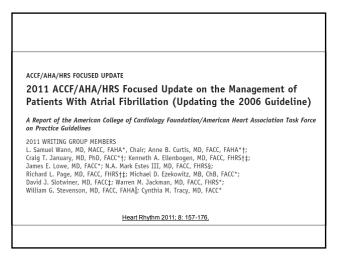
Diagnostic Evaluation

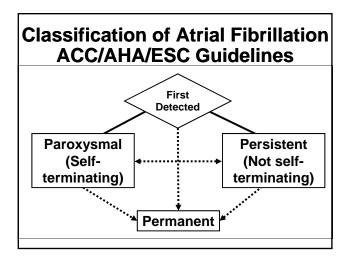
Additional Testing

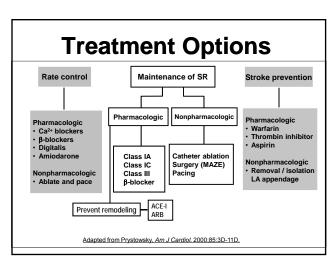
- ETT CAD, Exercise induced SVT / AF
- Holter / Event Monitor Confirm AF and Sxs
- TEE LA clot
- EPS SVT triggered AF
- Sleep Study

AHA / ACC / ECS Guidelines 2006









Atrial Fibrillation and Stroke

- 5 fold increased risk of CVA
- · AF accounts for 1 out of every 6 CVAs
- · Paroxysmal same risk as persistent
- Thromboemboli originating from LAA



Stroke Risk Assessment in AF: CHADS₂ Score

Clinical Parameter	Points
CHF	1
Hypertension	1
Age > 75yo	1
Diabetes	1
Stroke	2

CHADS ₂ Score	Annual Stroke Risk %	NNT
0	1.9	417
1	2.8	125
2	4.0	81
3	5.9	33
4	8.5	27
5 or 6	12-18	44

Gage et al, JAMA 2001; 285:2864.

Stroke Risk Assessment in AF: CHADS₂ Score

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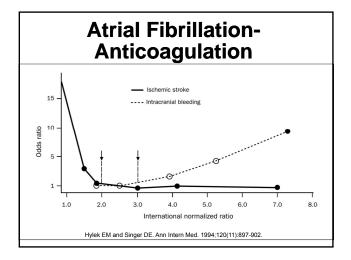
CHADS ₂ Score	Treatment	
0	ASA	
1	ASA or Warfarin (INR 2-3)	
2+	Warfarin (INR 2-3)	

Gage et al, JAMA 2001; 285:2864.

Anticoagulation

- Overall
 - √ 62% reduction with warfarin
 - ✓ 19% with ASA
- AFFIRM
 - √ 80% of CVAs occurred after coumadin was stopped or was subtherapeutic

CHADS2 Score	Events per years	NNT	
	Warfarin	No Warfarin	
0	0.25	0.49	417
1	0.72	1.52	125
2	1.27	2.50	81
3	2.20	5.27	33
4	2.35	6.02	27
5 or 6	4.60	6.88	44



Warfarin Limitations

- · Slow onset/offset
- Unpredictable dosing
- · Drug/diet interactions
- · Warfarin resistance (genetic)
- Narrow therapeutic index
- · Routine monitoring
- Patient dissatisfaction ("rat poison")
- Prescriber dissatisfaction

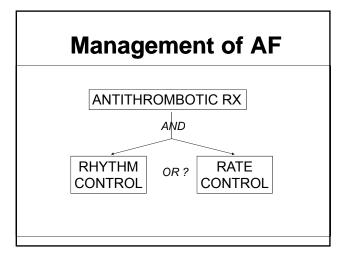


Dabigatran

- · Direct thrombin inhibitor
 - Reversible binding
 - Free & clot-bound thrombin
- Inhibits platelet aggregation
- Inhibits tissue factor-induced thrombin generation
- · Renally cleared
- No antidote

FDA-Approved Labeling

- · Who it's for:
 - Non-valvular AF patients for stroke prevention
- Who it's NOT for:
 - Mechanical heart valves
 - PE
 - DVT
 - Prophylaxis for knee/hip replacements
 - HIT



Rate Control

Atrial Fibrillation

Rate control - Drug Therapy

Digoxin – controls resting rate, OK in CHF patients.

Beta, Calcium channel blockers – controls resting and exercise rates.

Best therapy – combination of beta blocker and digoxin.

Primary Goal – Avoid Tachycardia Induced Cardiomyopathy

What is optimum rate control?

- AFFIRM trial
 - Resting heart rate less than 80 bpm
 - Peak heart rate less than 110 bpm
- RACE II

The NEW ENGLAND
JOURNAL of MEDICINE

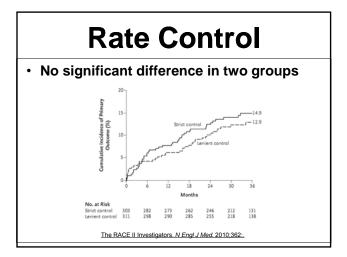
Lenient versus Strict Rate Control in Patients with Atrial Fibrillation

Isabelle C, Van Gelder, M.D., Hessel F, Groenveld, M.D., Hanry J, G.M. Crijns, M.D., Ype S. Tuininga, M.D., Jan GP, Tijssen, Ph.D., A. Marco Alings, M.D., Hans L. Hillige, M.D., Johanna A. Bergsma-Kadjik, M.S. at H. Cornell, M.D., Otto Kamp, M.D., Ragmooff Tuikle, M.D., Hans A. Booker, M.D., Drijk, Yan Velchiusen, M.T.

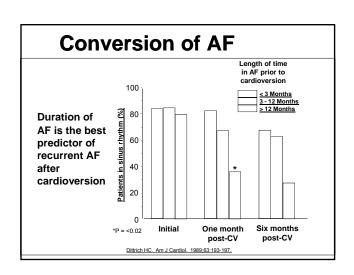
RACE II

- 614 patients
- Lenient Control (<110 bpm) versus strict control (<80 at rest, <110 at peak).
- Mean follow up 2 years.
- Primary Outcomes of death, CHF, stroke embolism, life threatening arrhythmias

The RACE II Investigators. N Engl J Med. 2010;362: 1363-1373.



Rhythm Control



Anticoagulation - Cardioversion

- · Atrial stunning
 - Stunning can occur even with one hour of atrial fibrillation
 - If duration < 2 weeks, function may return within 24 hours to one week
 - If duration > 2 weeks, stunning may persist for one month



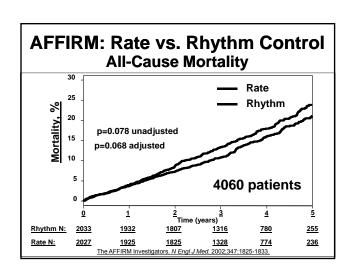
Mattioli, AV. et al. Am J Cardiol 1998; 82:1368.

Cardioversion

- Less than 48 hours duration
 - Cardioversion without TEE
 - Heparin at time of cardioversion
 - Warfarin for a month and reevaluation as outpatient

Cardioversion

- If greater than 48 hours
 - Option 1: Anticoagulate for 4 weeks and then cardiovert
 - Option 2: TEE and if no thrombus, cardiovert
 - If thrombus, 4 weeks warfarin and recheck
 - Anticoagulate for minimum of one month and re-evaluate



Rate vs. Rhythm Control Trials: Implications

- AFFIRM demonstrated that a rate control "strategy" is an acceptable primary therapy in a selected high-risk subgroup of AF patients
- Continuous anticoagulation seems warranted in all patients with risk factors for stroke

Asymptomatic recurrences

 AFFIRM did <u>not</u> define whether it is better to be in NSR.

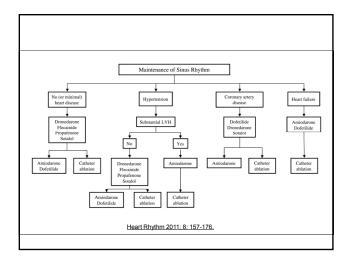
Rhythm Control

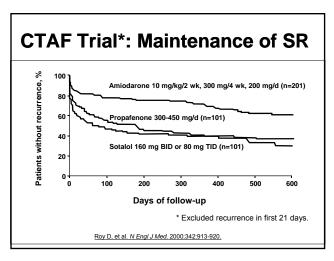
ADVANTAGES

- Avoids electrical and anatomical remodeling
- · Improves hemodynamics
- Enhanced exercise capacity
- · Symptom relief
- · Improves QOL
- Restores atrial transport
- Reduces thromboembolic events?

DISADVANTAGES

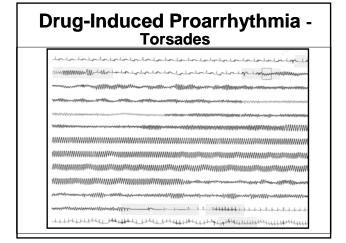
- · Ventricular proarrhythmia
- Increased mortality?
- Drug-induced bradyarrhythmias
- · End-organ toxicity
- Adverse effects
- · Recurrences are likely
- · Asymptomatic (silent) AF





AF Antiarrhythmic Therapy

- · Treatment goals
 - ↓ frequency of recurrences
 - ↓ duration of recurrences
 - ↓ severity of recurrences
 - Not to abolish every episode
- · Safety is primary concern
- Minimize risk of proarrhythmia

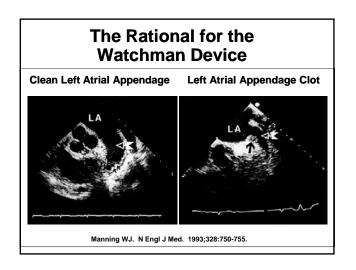


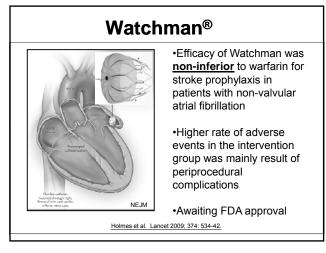
Factors Which Influence Ventricular Proarrhythmia Risk

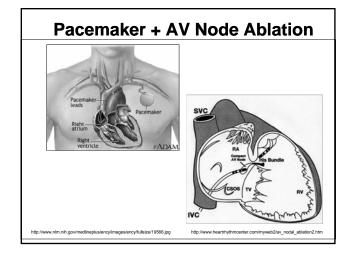
- · Hypokalemia, hypomagnesemia
- Long QT at baseline
- · CHF / Decreased EF
- Ventricular hypertrophy
- Bradycardia
- Female gender
- Reduced drug metabolism or clearance
- Amiodarone has lowest risk

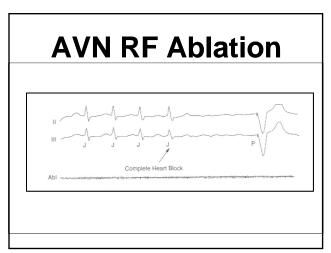
Alternatives to Drug Therapy "Non-Pharmacologic Therapy"

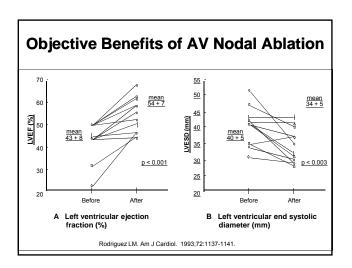
- □Coumadin LAA closure (Watchman)
- □Rate Control AVN RFA + PCMK
- □AARx Adjunctive AFL RFA
- □AARX Curative Afib RFA











AVN Ablation

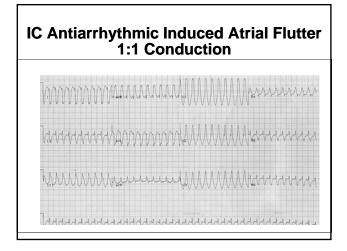
Advantages:

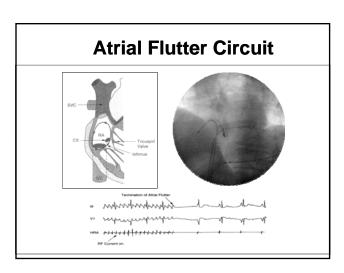
100% efficacy 85% symptomatic improvement Improved EF (LV remodeling) Eliminates need for rate control drugs

Disadvantages:Pacemaker dependant

Good Candidates:

Tachy / Brady Syndrome PPM present – CHF with BiV device Medication refractory / intolerant Elderly





Atrial Flutter Ablation

Approximately 15% of AF patients treated with an AARx will develop AFL

Advantages:

95% efficacy

≈ 80% arrhythmia control if AARx continued As primary Tx RFA more effective than AARx

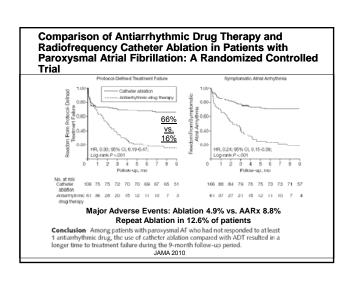
Disadvantages: Invasive

Good Candidates:

Typical AFL (IVC / TV isthmus)
Primary or AARx related Atrial Flutter

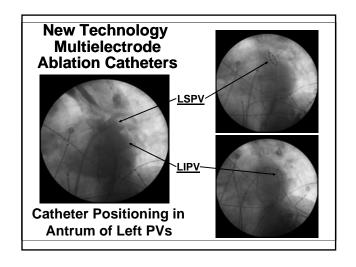
Focal Origin of Atrial Fibrillation Hassaiguerre M, NEJM, 1998 • 94% of AF triggers from Pulmonary Veins • "90-95% of all AF is initiated by PV ectopy"

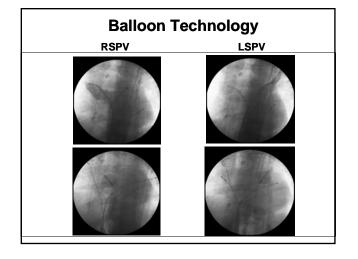
Atrial Fibrillation Ablation Atrial Shell Cardiac MRI

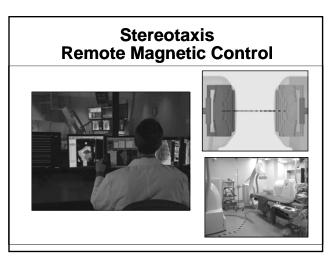


Current State of Curative Catheter-Based RFA Who is a good candidate?

Symptomatic / Frequent AF
Limited Heart Dz
EF > 35%
LA < 5.5cm
No MS / Rheumatic Dz
Younger Patients
No LA thrombus or Hx of CVA
Medically Refractory / Intolerant
(Ablation now second line therapy)







Atrial Fibrillation

New Technology / Studies at Ohio State University

Stereotaxis - Magnetic Catheter Navigation

New Catheter Design / Energy Sources
High Intensity Focused Ultrasound (HIFU)
Ablation Frontiers – Circular Catheters
Cryoablation
Laser Ablation

Cabana trial – Drug vs Ablation (including primary therapy)

Watchman – Left Atrial Appendage Closure Surgical vs Catheter Ablation