American Heart Association Guidelines

Evidence Based Guidelines for Cardiovascular Disease Prevention in Women

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Classification of Evidence

Classification

Class la Intervention is useful and effective

Class IIa Weight of evidence is in favor of efficacy

efficacy

Class IIb Efficacy is less well established

Class III Intervention is not useful and may be harmful

Level of Evidence

- Level
 - A Sufficient evidence from multiple randomized trials
 - B Limited evidence from a single randomized trial
 - C Based on expert opinion, case studies or standard of care

Generalizability Index

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- Very likely that results generalize to women
- 2 Somewhat likely that results generalize to women
- Unlikely that results generalize to women
- Unable to project whether results project to women

CVD Statistics

- CVD remains the leading cause of death in women
- > 500,000 women a year die
 - One death every minute
- Coronary Heart Disease accounts for the majority of CVD deaths
- 2/3 of women who die suddenly have no warning symptoms

CVD Prevention Strategies for Clinical Practice

- Assess and stratify women into high, intermediate, lower or optimal risk categories

CVD Prevention Strategies for Clinical Practice

◆ Other CVD risk-reducing interventions should be prioritized on the basis of the strength of recommendation (Level I >IIa >IIb) and within each class on the basis of the evidence, with the exception of lifestyle which is a top priority for all women (A>B>C)

CVD Prevention Strategies for Clinical Practice

- Highest priority for risk intervention in clinical practice is based on risk stratification with
 - High > intermediate > Low > optimal
- Avoid interventions designated as Class III

Spectrum of CVD Risk in Women

Framingham Global Risk > 20%

- Established CHD
- Cerebrovascular disease
- Peripheral vascular disease
- Abdominal aortic aneurysm
- Diabetes mellitus
- Chronic kidney disease
- → Subclinical CVD with > 20% Framingham risk

Spectrum of CVD Risk in Women

Framingham Global Risk > 20%

- Cerebrovascular disease may not confer CHD risk if the disease is above the carotids
- As chronic kidney disease deteriorates, CHD risk rises substantially
- Most women with a single, severe risk factor have a CVD risk < 10% per decade

Spectrum of CVD Risk in Women

Framingham Global Risk > 10 - 20%

- Subclinical CVD (+ coronary calcification)
- Metabolic Syndrome
- Multiple risk factors
- Markedly elevated levels of a single risk factor
- → First degree relatives with early onset atherosclerosis (< 55 in men and 65 in women)</p>

Spectrum of CVD Risk in Women

Framingham Global Risk < 10 %

May include women with multiple risk factors, metabolic syndrome or one or no risk factors

Framingham Global Risk < 10 %

 Optimal levels of risk factors and heart healthy lifestyle

Clinical Recommendations Lifestyle Interventions

- Cigarette Smoking (Class I, Level B, GI 1)
 - Encourage cessation
- Physical Activity (Class I, Level B, GI 1)
 - Minimum of 30 minutes moderate intensity activity seven days a week
- Cardiac Rehabilitation (Class I, Level B, GI 2)
 - Comprehensive risk reduction setting for women after ACS, coronary intervention or new onset or chronic angina)

Clinical Recommendations Lifestyle Interventions

- Heart Healthy Diet (Class I, Level B, GI 1)
 - Encourage fruits, vegetables, whole grains, low-fat or non-fat dairy, fish, legumes and sources of protein low in saturated fat
 - Limit fat intake to less than 10% of total calories
 - Limit cholesterol intake to < 300 mg/day
 - Limit intake of trans-fats
- Weight maintenance or reduction (Class I, Level B, GI 1)
 - Maintain BMI between 18.5 & 25 Waist < 35"

Clinical Recommendations Lifestyle Interventions

- Psychosocial Factors (Class IIa, Level B, GI 2)
 - Women with CVD should be evaluated for depression and treated or referred if present
- Omega-3 Fatty Acids (Class IIb, Level B, GI 2)
 - As an adjunct to diet, omega-3 FA may be considered in high-risk women
- Folic Acid (Class IIb, Level B, GI2)
 - As an adjunct to diet folic acid supplementation may be considered in high risk women (except after revascularization) if a higher than normal level of homocysteine has been detected

- → Blood Pressure: Lifestyle (Class I, Level B, GI 1)
 - Encourage a BP of < 120/80 through lifestyle approaches
- Blood Pressure: Drugs (Class I, Level A, GI 1)
 - Indicated when BP ≥ 140/90 or even lower in evidence of target organ damage or diabetes. Thiazide diuretics should be part of the regimen for most patients unless contraindicated

- Lipids Lipoproteins: (Class I, Level B, GI 1)
 - Optimal levels of lipids and lipoproteins in women:
 - LDL-C < 100 mg/dL
 - HDL-C > 50 mg/dL
 - TG < 150 mg/dL
 - Non HDL-C < 130 mg/dL
 - Should be encouraged through lifestyle approaches

- Lipids: Lifestyle (Class I, Level B, GI 1)
 - In high risk women when LDL-C is elevated, saturated fat intake should be limited to < 7% of calories, cholesterol to < 200 mg/day and trans-fatty acid intake should be reduced

- Lipids: Pharmacotherapy High risk (Class I, Level A, GI 1)
 - Initiate LDL-C lowering therapy (preferably a statin) simultaneously with lifestyle therapy in women with an LDL-C ≥ 100 mg/dL
 - Initiate statin therapy in high risk women with an LDL-C < 100 mg/dL unless contraindicated (Class I, Level B, G1)

- Lipids: Pharmacotherapy High risk (Class I, Level B, GI 1)
 - Initiate niacin or fibrate therapy when HDL-C is low or non HDL-C elevated
 - Dietary supplement niacin must not be used as a substitute for prescription niacin and OTC niacin should only be used if approved & monitored by a physician

- Lipids: Pharmacotherapy -Intermediate risk
 - Initiate LDL-C lowering therapy (preferably a statin) if LDL-C ≥ 130 mg/dL on lifestyle therapy (Class I, Level A, GI 1)
 - Initiate niacin or fibrates when HDL-C is low or non HDL-C is elevated after LDL-C goal is reached (Class I, Level B, GI 1)

- Lipids: Pharmacotherapy Lower Risk (Class IIa, Level B, GI 1)
 - Consider LDL-C lowering therapy in low risk women with 0 or 1 risk factor if LDL-C is > 190 mg/dL or if multiple risk factors are present and LDL-C > 160 mg/dL
 - Consider niacin or fibrate therapy when HDL-C is low or non HDL-C is elevated after LDL-C goal is reached

- ◆ Diabetes (Class I, Level B, GI 1)
 - Lifestyle and pharmacotherapy should be used to achieve near normal HgbA_{1c} (<7%) in women with diabetes

Clinical Recommendations Preventive Drug Interventions

- → Aspirin High Risk (Class I, Level A, GI 1)
 - ASA therapy (81–162 mg) or clopidogrel if the patient is intolerant to ASA should be used unless contraindicated
- Aspirin Intermediate risk (Class IIa, Level B, GI 2)
 - Consider ASA therapy as long as BP is controlled and benefit is likely to outweigh the risk of GI side effects

Clinical Recommendations Preventive Drug Interventions

- → Beta-blockers (Class I, Level A, GI 1)
 - Should be used indefinitely in all high risk women who have had an MI or have chronic ischemic syndromes unless contraindicated
- ◆ ACE Inhibitors (Class I, Level A, GI 1)
 - Should be used in high risk women unless contraindicated
 - Consider ASA therapy as long as BP is controlled and benefit is likely to outweigh the risk of GI side effects
- ARBs (Class I, Level B, GI 1)
 - Should be used in high risk women with clinical evidence of heart failure or an ejection fraction < 40% who are intolerant of ACEI

Clinical Recommendations Class III Interventions

- Hormone Therapy (Class III, Level A)
 - Combined estrogen plus progestogen therapy should not be initiated to prevent CVD in postmenopausal women
 - Combined E+P should not be continued to prevent CVD in postmenopausal women
 - Other forms of menopausal hormone therapy (e.g. unopposed estrogen) should not be initiated or continued to prevent CVD pending results from ongoing trials (Class III Level C)

Clinical Recommendations Class III Interventions

- Antioxidant Supplements (Class III, Level A, G1)
 - Should not be used to prevent CVD pending the results of ongoing trials
- → Aspirin (Class III, Level A, G1)
 - Use of low dose ASA is not recommended pending the results of ongoing trials

Clinical Recommendations Atrial Fibrillation/Stroke Prevention

- → Warfarin Atrial Fibrillation (Class I, Level A, G1)
 - Among women with chronic or paroxysmal atrial fibrillation warfarin should be used to maintain the INR at 2.0 – 3.0 unless they are considered to be at low risk of stroke (<1% a year) or high risk of bleeding
- Aspirin (Class I, Level A, G1)
 - Should be used in women with chronic or paroxysmal atrial fibrillation with a contraindication to warfarin or at low risk for stroke (<1% a year)