Hypertension Suite of Templates Tutorial

The **Hypertension Suite of Templates** have been designed to make it possible to efficiently and excellently document the proper care of hypertension and to simultaneously provide the opportunity for healthcare providers to improve their understanding of the disease, while evaluating the quality of care they personally give to patients with hypertension. At present, over 50% of the patients who are treated by SETMA have a diagnosis of hypertension making its treatment a key element in our delivery of quality healthcare.

There is no condition which is not worsened by the concomitant presence of hypertension and there are few chronic conditions which are as devastating to the health and well being of a patient as that of elevated blood pressure. Finally, the complexity of the treatment of hypertension is compounded by the fact that so many special cases exist each of which requires different considerations for the proper treatment of high blood pressure.

On the other hand, few chronic conditions offer the promise as does the effective treatment of hypertension. The consequences of untreated hypertension are grave, but the results of effective control of hypertension are gratifying. However, the reality is that the overwhelming majority of patients with hypertension are inadequately treated.

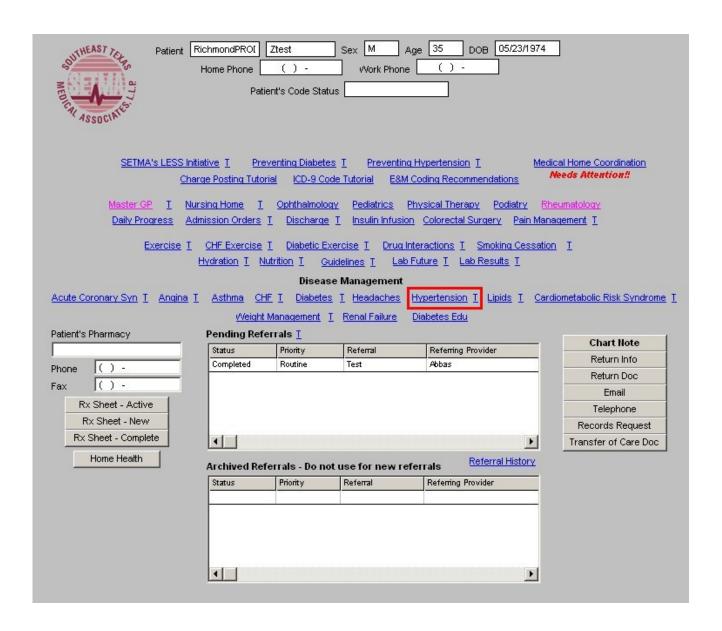
It is for all of these reasons that SETMA's **Hypertension Suite of Templates** not only offers a tool for the documentation of the patient's hypertension treatment but also provides information, insight and algorithms for the analysis of a patient's blood pressure treatment and control in:

- The evaluation of secondary hypertension
- The presence of Dippers and Non-Dippers
- White Coat Syndrome and its cardiovascular risk
- Diabetes
- Depression
- The elderly
- Insulin Resistance
- Isolated Systolic Blood Pressure
- Kidney Disease
- Medication selection in dozens of comorbid conditions

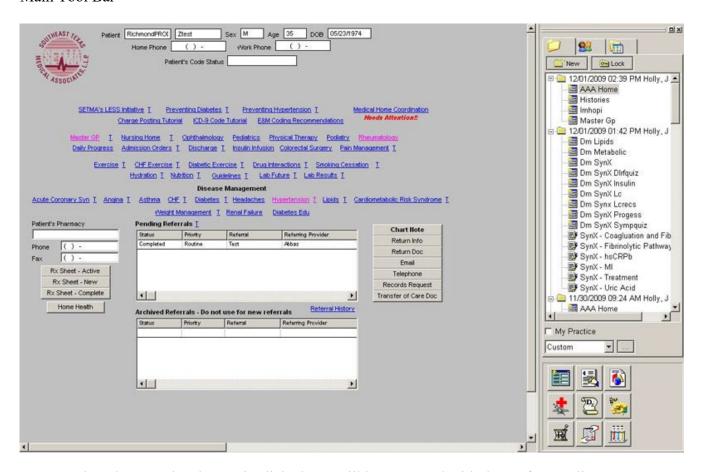
The following is not only an explanation of how to use **SETMA's Hypertension Suite of Templates**, but it is also a structure for logically approaching the treatment of hypertension along with a wealth of information and insight about that treatment.

The Hypertension Suite of Templates can be accessed from:

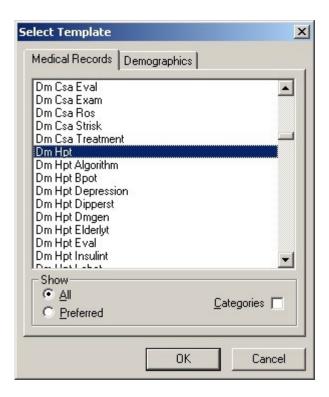
AAA Home



Main Tool Bar



- When the Template button is clicked you will be presented with the preference list.
- If the Hypertension Master Template is listed as one of your preferences, select it.
- If it is not one of your preferences, select the All radio button and then scroll down until you find it in the list.



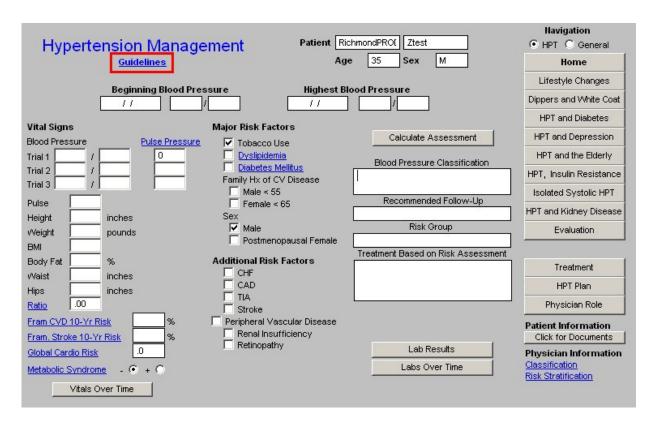
NOTE: For more on how to set up your preferences, <u>Click Here</u>

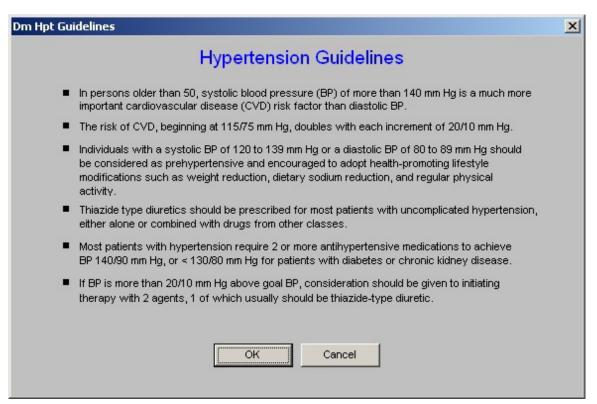
Other Disease Management Tools Hyperlinks

The **Master Hypertension Template** provides the entry point into the management of hypertension

Information at the top of the Master Hypertension Template

- Title
- **Guidelines** this is a hyperlink which launches a list of General Hypertension Guidelines about blood pressure management. These are listed below.





- The **General Hypertension Guidelines** provide the imperative for effective blood pressure control. The **Guidelines** state:
 - o In persons older than 50, systolic blood pressure (BP) of more than 140 mm Hg is a much more important cardiovascular disease (CVD) risk factor than diastolic BP.
 - The risk of CVD, beginning at 115/75 mm Hg, doubles with each increment of 20/10 mm Hg.
 - Individuals with a systolic BP of 120 to 139 mm Hg or a diastolic BP of 80 to 89 mm Hg should be considered as pre-hypertensive and encouraged to adopt health-promoting lifestyle modifications such as weight reduction, dietary sodium reduction, and regular physical activity.
 - o Thiazide type diuretics should be prescribed for most patients with uncomplicated hypertension, either alone or combined with drugs from other classes.
 - Most patients with hypertension require 2 or more antihypertensive medications to achieve BP 140/90 mm Hg, or < 130/80 mm Hg for patients with diabetes or chronic kidney disease.
 - o If BP is more than 20/10 mm Hg above goal BP, consideration should be given to initiating therapy with 2 agents, 1 of which usually should be thiazide-type diuretic.
- Patient Name, Age, Sex

Hypertensio	n Management	Patient RichmondPROI Ztest	Navigation ⊙ HPT ○ General
<u>Gui</u>	<u>delines</u>	Age 35 Sex M	Home
Dit	i Planed Paraness	Winh and Discoul Description	Lifestyle Changes
Begii	nning Blood Pressure	Highest Blood Pressure	Dippers and White Coat
Vital Circus	Maior Dials Forts		HPT and Diabetes
Vital Signs Blood Pressure	Major Risk Facto Pulse Pressure Tobacco Us	Calculate Assessmen	HPT and Depression
Trial 1 /	Pulse Pressure ✓ Tobacco Usi O Dyslipidemia		HPT and the Elderly
Trial 2 /	Diabetes Me	illitus I	HPT, Insulin Resistance
Trial 3 /	☐ Male < 55	2000	Isolated Systolic HPT
Pulse	Female < 6	5 Recommended Follow-I	HPT and Kidney Disease
Weight pounds	Sex ▼ Male Fostmenop	Risk Group	Evaluation
Body Fat %	Additional Risk F	actors Treatment Based on Risk Ass	essment Treatment
Waist inches	☐ CHF ☐ CAD		HPT Plan
Ratio .00			Physician Role
Fram CVD 10-Yr Risk	% ☐ Peripheral Vaso		Patient Information
Fram. Stroke 10-Yr Risk	———— /″	Lab Results	Click for Documents
Global Cardio Risk Metabolic Syndrome - G	.0 Teamopouny	Labs Over Time	Physician Information Classification
Vitals Over Time			Risk Stratification

• **Beginning Blood Pressure** - this is the blood pressure from the first visit when the Master Hypertension Template was accessed, not the blood pressure from the first time the patient was seen by a SETMA Provider.

Hypertension Mar	nagement Patien	t RichmondPRO[Ztest	Havigation ● HPT ○ General
Guidelines		Age 35 Sex M	Home
Danimina Dia	d December 18 also	A Planet Processor	Lifestyle Changes
Beginning Bloc	od Pressure Highe	est Blood Pressure	Dippers and White Coat
Vital Signs	Major Risk Factors		HPT and Diabetes
Blood Pressure Pulse Pre		Calculate Assessment	HPT and Depression
Trial 1 / O	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2	Diabetes Mellitus Family Hx of CV Disease		HPT, Insulin Resistance
Trial 3 /	☐ Male < 55		Isolated Systolic HPT
Pulse	☐ Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Height inches Weight pounds	✓ Male	Risk Group	Evaluation
BMI Podrido	Postmenopausal Fema		
Body Fat %	Additional Risk Factors	Treatment Based on Risk Assessment	Treatment
Waist inches	☐ CHF ☐ CAD		HPT Plan
Hips inches	TIA		Physician Role
Ratio J.UU Fram CVD 10-Yr Risk %	Stroke Peripheral Vascular Disea:	0.0	
Fram. Stroke 10-Yr Risk %	Renal Insufficiency		Patient Information Click for Documents
Global Cardio Risk .0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome - © + C		Labs Over Time	Classification Risk Stratification
Vitals Over Time			ition on annoanon

• **Highest Blood Pressure** - this is the highest blood pressure measured at a time when the Master Hypertension Template was accessed. If a higher blood pressure was recorded on a patient when the Master Hypertension Template was not accessed, it will not be recorded here.

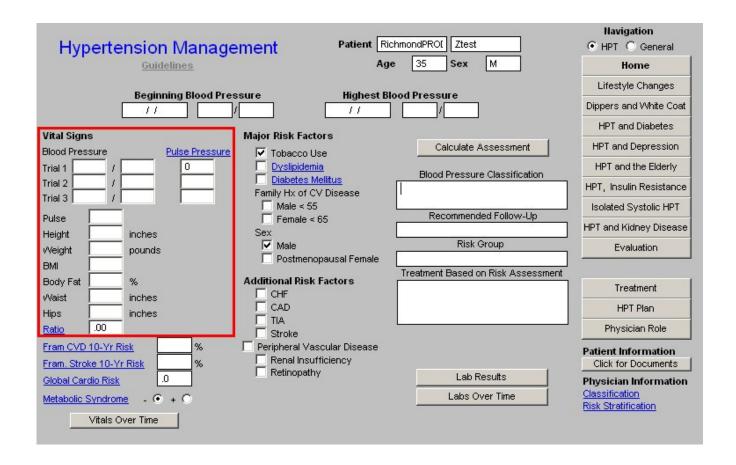
Hypertension Manage	ement Patient Ri	ichmondPROI Ztest	Havigation ⊙ HPT ○ General
Guidelines	Ag	je 35 Sex M	Home
Beginning Blood Pre	naura Highaat P	llood Pressure	Lifestyle Changes
/ /	/ nignest b		Dippers and White Coat
Vital Signs	Major Risk Factors		HPT and Diabetes
Blood Pressure Pulse Pressure	▼ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / 0	□ Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /	Diabetes Mellitus Family Hx of CV Disease	Blood Pressure Classification	HPT, Insulin Resistance
Trial 3 /	Male < 55		Isolated Systolic HPT
Pulse	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Height inches Weight pounds	Sex ✓ Male	Risk Group	Evaluation
BMI Dearlies	Postmenopausal Female		
Body Fat %	Additional Risk Factors	Treatment Based on Risk Assessment	Treatment
Waist inches	☐ CHF ☐ CAD		HPT Plan
Hips inches	TIA		Physician Role
Ratio J.00 Fram CVD 10-Yr Risk %	Stroke Peripheral Vascular Disease		
Fram. Stroke 10-Yr Risk %	Renal Insufficiency		Patient Information Click for Documents
Global Cardio Risk .0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome - ① + ①		Labs Over Time	Classification
Vitals Over Time			Risk Stratification

Beneath the above information, the remainder of the **Hypertension Master Template** is organized into four columns:

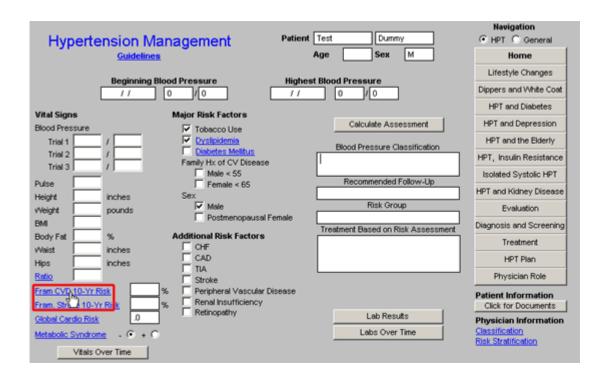
Column 1 -

Vital signs

- Blood Pressure
 - 1. Trial 1
 - 2. Trial 2
 - 3. Trial 3
- Pulse
- Height
- Weight
- BMI
- Body Fat
- Waist
- Hips
- Ratio

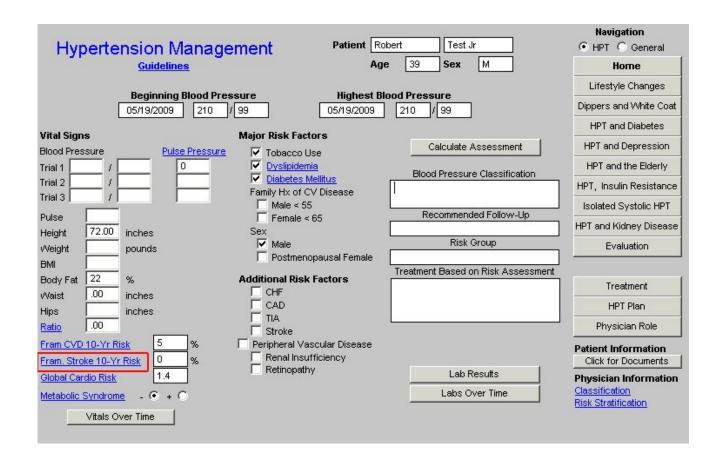


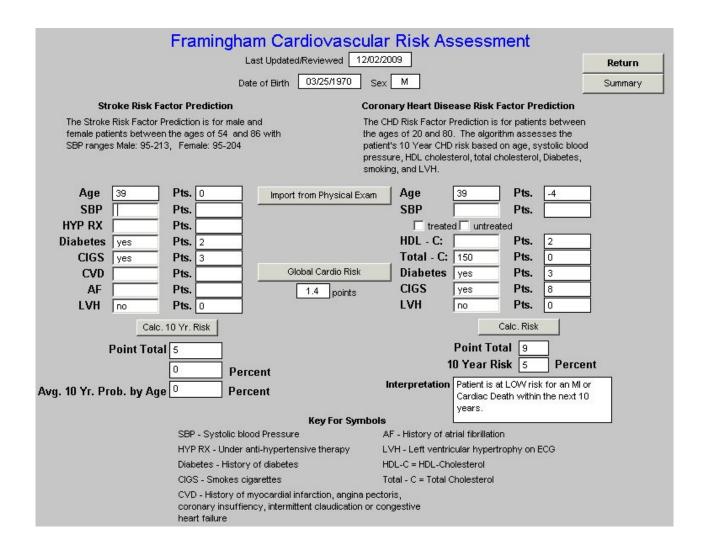
Fram CVD 10-Yr Risk - this is a hyperlink which takes you to the Framingham Risk Template where the Cardiovascular and Stroke Risk are calculated from the Framingham Data and where the Global Cardio Risk is also calculated. The results of all three are displayed here. This enables a provider to address the cardiovascular and cerebrovasculer risk burden carried by the patient and thus to design a treatment program based on that risk.



	Framingham Cardiovaso	cular Risk Assessment
	Last Updated/Reviewed	12/01/2009 Return
	Date of Birth 05/23/1974	Sex M Summary
Stroke Risk Fac	tor Prediction	Coronary Heart Disease Risk Factor Prediction
The Stroke Risk Factor Pro female patients between t SBP ranges Male: 95-213	the ages of 54 and 86 with	The CHD Risk Factor Prediction is for patients between the ages of 20 and 80. The algorithm assesses the patient's 10 Year CHD risk based on age, systolic blood pressure, HDL cholesterol, total cholesterol, Diabetes, smoking, and LVH.
Age 35	Pts. 0 Import from Physical	Exam Age 35 Pts4
SBP	Pts.	SBP Pts.
HYP RX	Pts.	☐ treated ☐ untreated
Diabetes	Pts.	HDL - C: Pts.
CIGS	Pts.	Total - C: Pts.
CVD	Pts. Global Cardio Ris	Didiveres 1 to.
AF	Pts	
LVH	Pts.	LVH Pts.
Calc. 1	0 Yr. Risk	Calc. Risk
		Point Total
		10 Year Risk Percent
	VF	Interpretation
	Key For Sy SBP - Systolic blood Pressure	AF - History of atrial fibrillation
	HYP RX - Under anti-hypertensive therapy	LVH - Left ventricular hypertrophy on ECG
	Diabetes - History of diabetes	HDL-C = HDL-Cholesterol
	CIGS - Smokes cigarettes	Total - C = Total Cholesterol
	CVD - History of myocardial infarction, angir coronary insuffiency, intermittent claudication heart failure	

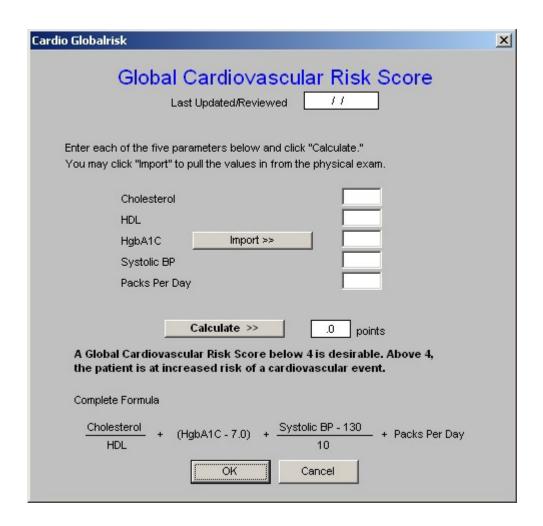
Fram. Stroke 10-Yr Risk - same as above.





Global Cardio Risk - this is a new risk calculation based on the Framingham data but without the non-modifiable risk factors - age, sex, etc - included in the equation. A score of 4 or more represents an increased cardiovascular risk.

Hyperte	nsion Manage	ment Patient Ri	chmondPROI Ztest	Navigation • HPT • General
	Guidelines	Ag	e 35 Sex M	Home
	Danissis - Diagd Danis	anna Minka at Di	lood Pressure	Lifestyle Changes
1	Beginning Blood Pres	sure nignest bi		Dippers and White Coat
Vital Signs		Maior Diels France		HPT and Diabetes
Blood Pressure	Pulse Pressure	Major Risk Factors Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslipidemia Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /		☐ <u>Diabetes Mellitus</u> Family Hx of CV Disease	Diodu Pressure Classification	HPT, Insulin Resistance
Trial 3 /		☐ Male < 55	Bassing ded Faller Library	Isolated Systolic HPT
Pulse	inches	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Weight	pounds	✓ Male	Risk Group	Evaluation
вмі 🗀		Postmenopausal Female	Treatment Based on Risk Assessment	
Body Fat	%	Additional Risk Factors CHF	Trouble Based of Hisk Assessment	Treatment
Waist Hips	inches inches	□ CAD		HPT Plan
Ratio .00	lilicites	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Ri	<u>sk</u> %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr		Renal Insufficiency Retinopathy	1	Click for Documents
<u>Global Cardio Risk</u>	.0	i Reunopatry	Lab Results	Physician Information
Metabolic Syndrom	<u>e</u> - ⊙ + ⊙		Labs Over Time	Classification Risk Stratification
Vitals Ov	ver Time			



Metabolic Syndrome

Hypertension Manage	ment Patient Ric	chmondPROI Ztest	Navigation • HPT • General
Guidelines	Age	e 35 Sex M	Home
Parissian Bland Barr	Windowsk Di	and December	Lifestyle Changes
Beginning Blood Pres	isure nighest bi	ood Pressure	Dippers and White Coat
Vital Signs	Major Risk Factors		HPT and Diabetes
Blood Pressure Pulse Pressure	✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / O	Dyslipidemia Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 / /	Diabetes Mellitus Family Hx of CV Disease	Diodu Fressure Classification	HPT, Insulin Resistance
Trial 3	Male < 55		Isolated Systolic HPT
Pulse	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Height inches Weight pounds	✓ Male	Risk Group	Evaluation
BMI podrids	Postmenopausal Female		
Body Fat %	Additional Risk Factors	Treatment Based on Risk Assessment	Treatment
vVaist inches	☐ CHF ☐ CAD		HPT Plan
Hips inches	□ TIA		
Total Control of the	Stroke		Physician Role
Fram CVD 10-Yr Risk	Peripheral Vascular Disease Renal Insufficiency		Patient Information Click for Documents
	Retinopathy	Lab Results	
		Labs Over Time	Physician Information Classification
Metabolic Syndrome - ⊙ + ○		Labo Over Time	Risk Stratification
Vitals Over Time			

Last Updated/Reviewed 12/02/2009	WHO Diagnostic Criteria	ATP III Diagnostic Criteria ← + ← -	Return
Triglycerides mg	v/dL >= 150 mg/dL	>= 150 mg/dL	
Central Obesity			
vValst 12341 inc	hes Ratio	vVaist	
Hip 1234J inc	hes Men > 0.90	Men > 40 inches	Minor Features
Ratio 1.00	vNomen > 0.85	Women > 35 inches	Pro-inflammatory State: Increased CRP Polycystic Ovarian Syndrome
BMI mg	u/m^2 BMI > 30		Coronary Heart Disease
Blood Pressure			Small, dense LDL particle size
	nHa > 140/90 mmHa	> 130/85 mmHg	Increased Apolipoprotein B Hypercoagulability/Prothrombotic State: increased fibrinogen and PAI-1.
		- 15000 mm g	Vascular Endothelial Dysfunction: The inside of the artery doesn't work correctly
Glucose Abormalities	BOD VAN ES		Microalbuminuria: The kidneys fall and allow protein to go into urine inappropriatel
(0.000)	i/dL Fasting > 110 mg/dL 2 Hr GTT > 140 mg/dL	Fasting > 110 mg/dL	
The state of the s	yol. 2 Hr G11 > 140 mg/di. Diabetes		
Diabetes C + C - Insulin Resistance C + C -	Insulin Resistance		
HDL mg	v/dL Men < 35 mg/dL	Men < 40 mg/dL	
Microalbuminuria	· V/omen < 39 mg/dL	/Vomen < 50 mg/dL	
Alb/Creat mg	ylg > 30 mg/g		
Spot A/C m	ydL > 2.9 mg/dL		

Vitals over Time - this function displays the vital signs over time. This is a very useful tool to

look at patterns and possible seasonal changes in blood pressure, etc.

Hyperter	nsion Manage	ment Patient F	RichmondPROI Ztest	Havigation • HPT • General
	Guidelines	A	ge 35 Sex M	Home
	Barrianian Bland Barri	anna Hintarat I	Diagram December	Lifestyle Changes
Г	Beginning Blood Pres	sure Highest I	Blood Pressure	Dippers and White Coat
W 10:				HPT and Diabetes
Vital Signs Blood Pressure	Pulse Pressure	Major Risk Factors ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /		Diabetes Mellitus Family Hx of CV Disease	Biood Pressure Classification	HPT, Insulin Resistance
Trial 3 /		Male < 55	December Sellen, the	Isolated Systolic HPT
Pulse Height	inches	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Weight	pounds	✓ Male	Risk Group	Evaluation
ВМІ		Postmenopausal Female	Treatment Based on Risk Assessment	
Body Fat	%	Additional Risk Factors	Treatment based of risk Assessment	Treatment
Waist Hips	inches inches	☐ CAD		HPT Plan
Ratio .00	lilories	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Ris	<u>sk</u> %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr	Risk %	Renal Insufficiency Retinopathy		Click for Documents
Global Cardio Risk	.0	The inopatity	Lab Results	Physician Information
Metabolic Syndrome	<u> - </u>		Labs Over Time	Classification Risk Stratification
Vitals Ov	er Time			

			Trial 1		Trial 2		Trial 3				
Visit Date	Avgerage Systolic BP	Avgerage Diastolic BP	Systolic	Diastolic	Systolic	Diastolic	Systolic	Diastolic	Weight	Body Fat %	BMI
12/01/2009 01:42 PM			1234	152345				. 180			

Column 2 --

Major Risk Factors (Cardiovascular Disease Risk)

Hypertension Manage	ement Patient Ri	chmondPROI Ztest	Navigation • HPT • General
Guidelines	Ag	e 35 Sex M	Home
Beginning Blood Pre	oouro Highoot Di	lood Pressure	Lifestyle Changes
/ /	/ nightest bi		Dippers and White Coat
Vital Signs	Major Risk Factors		HPT and Diabetes
Blood Pressure Pulse Pressure	Wajor Risk Factors ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /	☐ <u>Diabetes Mellitus</u> Family Hx of CV Disease	Diodu Pressure Classification	HPT, Insulin Resistance
Trial 3 /	Male < 55	December and Sellen Us	Isolated Systolic HPT
Pulse inches	☐ Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Weight pounds	✓ Male	Risk Group	Evaluation
вмі	Postmenopausal Female	Treatment Based on Risk Assessment	
Body Fat %	Additional Risk Factors	Treatment based off thisk Assessment	Treatment
Waist inches	□ CAD		HPT Plan
Ratio .00	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Risk %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk %	Renal Insufficiency Retinopathy		Click for Documents
Global Cardio Risk .0	I Neuropatry	Lab Results	Physician Information
Metabolic Syndrome - ● + ●		Labs Over Time	Classification Risk Stratification
Vitals Over Time			

- Tobacco UseDyslipidemia

Hypertension Mana	igement Patient Ric	chmondPRO[Ztest	Navigation • HPT • General
<u>Guidelines</u>	Ag	e 35 Sex M	Home
Beginning Blood	Dragoura Highoot D	lood Pressure	Lifestyle Changes
/ /	/ nighest bi		Dippers and White Coat
Wital Ciana	Maior Biola Fortuna		HPT and Diabetes
Vital Signs Blood Pressure Pulse Press	Major Risk Factors	Calculate Assessment	HPT and Depression
Trial 1 / O	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 / /	□ <u>Diapetes Meilitus</u> Family Hx of CV Disease	Diodu Pressure Classification	HPT, Insulin Resistance
Trial 3 /	☐ Male < 55	December of Fellows I be	Isolated Systolic HPT
Pulse Inches	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Weight pounds	✓ Male	Risk Group	Evaluation
вмі	Postmenopausal Female	Treatment Based on Risk Assessment	·
Body Fat %	Additional Risk Factors	Treatment based of risk Assessment	Treatment
Waist inches	☐ CAD		HPT Plan
Hips inches	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Risk %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk %	Renal Insufficiency Retinopathy		Click for Documents
Global Cardio Risk .0	i Reunopatriy	Lab Results	Physician Information
Metabolic Syndrome - ● + ●		Labs Over Time	Classification Risk Stratification
Vitals Over Time			100000000000000000000000000000000000000

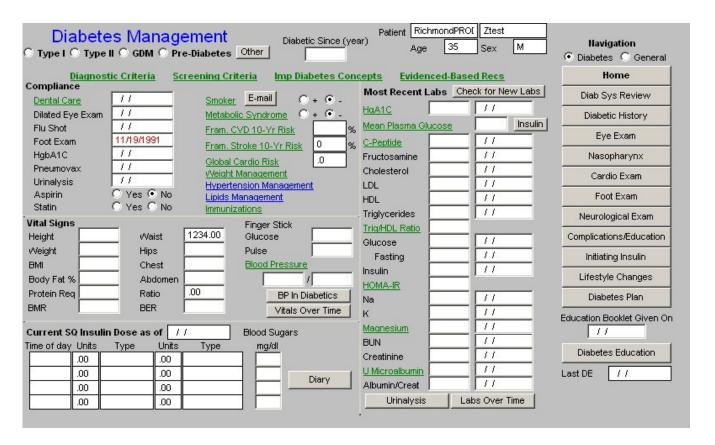
Depressing the Dyslipidemia button launches the Lipid Management Suite of Templates beginning with the Master Lipid Management Template. For more information on the Master Lipid Template, <u>Click Here.</u>

Lipids Managemer	nt Patient RichmondPRO[Z	test	Havigation
SETMA's Lipid Philosophy	Age 35 Sex M	1	Return
Compliance	Most Recent Labs Goals	Risk Factors Coronary Heart Disease	Lipids System Review
Last CRP //	Check for New Labs		Extremity Exam
Last CRP //	Cholesterol //	_ Angina	
Height inches	HDL //	CABG Non-Coronary Atherosclerosis	Eye Exam
Weight pounds	HDL 2	Peripheral Artery Disease	Cardio Exam
BMI	HDL 3 Cholesterol/HDL	Cerebrovascular Disease	Lifestyle Changes
Body Fat %	Triglycerides //	Aortic Aneurysm	Lipids Plan
BMR cal/day	Trig/HDL	Fram. CVD 10-Year Risk %	Lipius Fiam
Protein Req grams/day		Fram. Stroke 10-Year Risk	
vVaist 1234.00 inches	CPK / / /	Global Cardio Risk .0	Lipoprotein Metabolism
Blood Pressure	Lp(a)	— ✓ Male Age > 45	Summary of Lipid Studies
/ mmHg	LDL //	Female Age > 55	Lipoproteins
/ mmHg	IDL IDL	Hypertension > 140/90	Significance
/ mmHg	<u>VLDL</u>	Blood Pressure Medications	Composition
Diabetes Mellitus + C - ©	LDL-Remnant	✓ Smoking HDL	Classification
Metaboilc Syndrome + C - ©	Pattern A Pattern B Info	Male < 40	Hyperlipoproteinemias
Fredrickson Classification	Pattern A/B	Female < 50	Hypolipoproteinemias
Assess from Labs	Homocystiene //	FHx Premature HD	VLDLs
O I O IIa O IIb	hsCRP //	Male First Degree < 55	IDLs
CIII CIV CV	Apo A1	Female First Degree < 65	LDLs
Help Info	Apo B	Assessment Update	HDLs
	Apo E2	Measures should be taken to lower LDL to below 70.	LDL Receptors
Last Updated/Reviewed	Apo E4	Last Updated/Reviewed	Chylomicrons
12/01/2009	Labs Over Time	[]]	Chylomicrons and Triglycerides
		Sec	ondary Causes of Abnormal Lipids
			Hypercholesterolemia
			Hypocholesterolemia
			Low HDL
			Hypertriglyceridemia
			

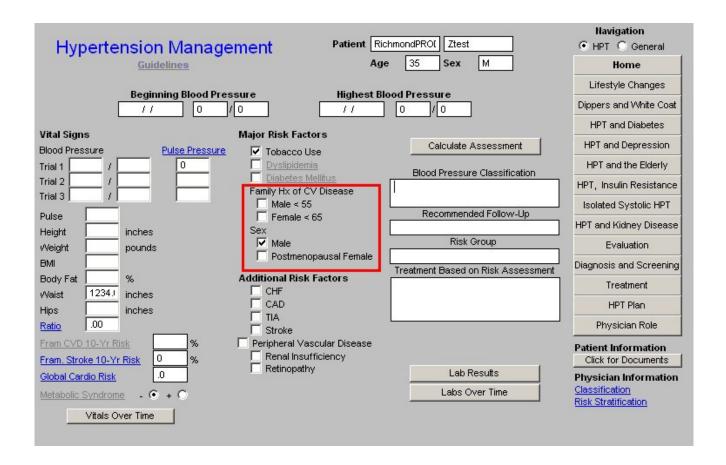
• Diabetes Mellitus

Hyperte	nsion Manage	ment Patient Ric	chmondPROI Ztest	Havigation
Пурсто	<u>Guidelines</u>	Age	9 35 Sex M	Return
	Danissis a Plant Danis	Hinto et Di	and Barrers	Lifestyle Changes
Г	Beginning Blood Pres	O //	ood Pressure	Dippers and White Coat
				HPT and Diabetes
Vital Signs Blood Pressure	Pulse Pressure	Major Risk Factors ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslinidemia	Di il	HPT and the Elderly
Trial 2 /		☐ <u>Diabetes Mellitus</u>	Blood Pressure Classification	HPT, Insulin Resistance
Trial 3 /		Family HX of CV Disease Male < 55	<u>'</u>	Isolated Systolic HPT
Pulse		Female < 65	Recommended Follow-Up	HPT and Kidney Disease
Height	inches	Sex ✓ Male	Risk Group	Evaluation
Weight	pounds	Postmenopausal Female		
BMI	%	Additional Risk Factors	Treatment Based on Risk Assessment	Diagnosis and Screening
Body Fat 1234.		CHF		Treatment
Waist 1234.	inches inches	CAD		HPT Plan
Ratio .00	littles	□ TIA		Physician Role
Fram CVD 10-Yr Ris	sk 7%	Stroke Peripheral Vascular Disease		
Fram. Stroke 10-Yr		Renal Insufficiency		Patient Information Click for Documents
Global Cardio Risk	.0	Retinopathy	Lab Results	
			Labs Over Time	Physician Information Classification
Metabolic Syndrome	≞ - ⊙ + ○		Labs Over Time	Risk Stratification
Vitals Ov	ver Time			
Vitals Ov	ver Time			

Depressing the Diabetes Mellitus Button launches the Diabetes Suite of Templates beginning with the Master Diabetes Management Template. For more information on the Master Diabetes Management Template, Click Here.

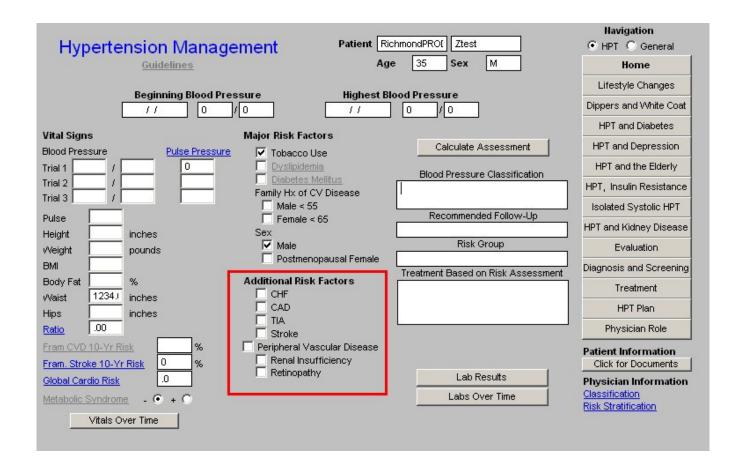


- Family History of CV Disease
 - 1. Male < 55
 - 2. Female < 65
- Sex
 - 1. Male
 - 2. Postmenopausal Female



Additional Risk Factors -- The presence of any of these risk factors increases the cardiovascular risk burden carried by a patient and therefore increases the imperative for lowering the blood pressure.

- CHF
- CAD
- TIA
- Stroke
- Peripheral Vascular Disease
- Renal Insufficiency
- Retinopathy



Column 3 -

Calculate Assessment - when this button is depressed the following calculations are automatically made and displayed in the boxes named hypertension classification, recommended follow-up, risk group and treatment recommendation based on this risk assessment.

- **Blood Pressure Classification** This indicate the risk associated with certain levels of blood pressure.
- **Recommended Follow-up** This indicates how soon the patient should be seen for this blood pressure classification.
- **Risk Group** This classification provides a risk stratification for determining the treatment of the patient. The Risk Group is determined by the factors listed in the document entitled "Risk Stratification" which is launched from the button of that name at the bottom right of this template.
- **Treatment Based on Risk Assessment** Based on the Risk Group identified above, a treatment plan will be displayed. This will range from lifestyle modifications to drug therapy.

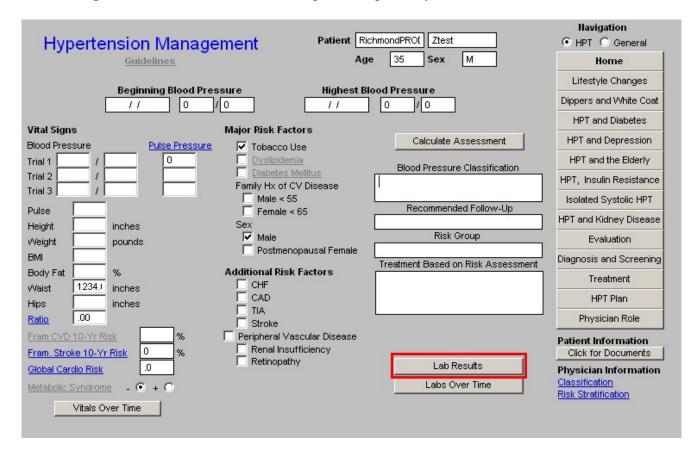
Note: For details see the **Risk Stratification** section below.

Hypertension N	Management Patient Ric	chmondPROI Ztest	Havigation ⊙ HPT ○ General
Guidelin		e 35 Sex M	Home
Diii	Pland Parane	and Programs	Lifestyle Changes
Beginning	Blood Pressure Highest Bl	ood Pressure	Dippers and White Coat
Vital Ciana	Maior Dials France		HPT and Diabetes
Vital Signs Blood Pressure Puls	Major Risk Factors se Pressure ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	O Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /	Diabetes Mellitus Family Hx of CV Disease	Blood Pressure Classification	HPT, Insulin Resistance
Trial 3	☐ Male < 55		Isolated Systolic HPT
Pulse Inches	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Weight pounds	✓ Male	Risk Group	Evaluation
ВМІ	Postmenopausal Female	Treatment Based on Risk Assessment	Diagnosis and Screening
Body Fat %	Additional Risk Factors	Treatment based on Nisk Assessment	Treatment
Waist 1234. inches	☐ CAD		HPT Plan
Ratio .00	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Risk	% Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk 0	% Renal Insufficiency		Click for Documents
Global Cardio Risk .0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome - © +	•	Labs Over Time	Classification Risk Stratification
Vitals Over Time			**************************************

Lab Results - when this button is depressed, a pop-up is launched which has the following laboratory values displayed.

- Na
- K
- BUN
- Creatinine
- Ca
- Mg
- Ca/Mg this ratio reflects the possible presence of a prothrombotic condition in a patient. Calcium promotes blood clotting, while Magnesium inhibits platelet aggregation. If the Ca/Mg ratio is above 5, the patient is likely to have a prothrombotic condition. This ratio is most often regulated by increasing the Magnesium level rather than decreasing the Calcium levels in the blood.
- Glucose
- Fasting Glucose
- Insulin
- **HOMA-IR** Homeostasis Model Assessment of Insulin Resistance. This is a calculated value based on the fasting plasma glucose and the fasting insulin. The calculation is done automatically when these two values are present in NextGen's laboratory module. If this value is greater than 2, the patient is insulin resistant.
- MS Strip

- **Microalbumin** as with diabetic nephropathy, so with hypertensive renal disease, urinary protein is the earliest indicator of renal damange.
- Uric Acid increased uric acid is associated with a prothrombic state as well.
- Cholesterol
- HDL
- LDL
- Triglycerides
- Trig/HDL if this ratio is over 2, the patient is probably insulin resistant.



Labs Over Time - when this button is depressed, the "Hypertension Labs Over Time" template is launched.

Hypertension Manage	ement Patient R	ichmondPROI Ztest	Havigation • HPT • General
<u>Guidelines</u>	Ag	ge 35 Sex M	Home
Beginning Blood Pres	nours Highoot F	Blood Pressure	Lifestyle Changes
	0 //	0 /0	Dippers and White Coat
Vital Signs	Major Risk Factors		HPT and Diabetes
Blood Pressure Pulse Pressure	✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / 0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 / / /	Diabetes Mellitus Family Hx of CV Disease	Bioda i ressai e ciassinoation	HPT, Insulin Resistance
Trial 3 /	Male < 55		Isolated Systolic HPT
Pulse	Female < 65	Recommended Follow-Up	HPT and Kidney Disease
Height inches	Sex ✓ Male	Risk Group	Evaluation
Weight pounds	Postmenopausal Female		
Body Fat %	Additional Risk Factors	Treatment Based on Risk Assessment	Diagnosis and Screening
	CHF		Treatment
Trainet Inches	CAD		HPT Plan
	☐ TIA		Dhusisian Bala
TORIO 1.55	_ Stroke		Physician Role
Fram CVD 10-Yr Risk %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk 0 %	Renal Insufficiency Retinopathy		Click for Documents
Global Cardio Risk .0	Reunopatry	Lab Results	Physician Information
Metabolic Syndrome - C + C		Labs Over Time	Classification
Varia Ocean Time			Risk Stratification
Vitals Over Time			

ncounter	Na	lĸ	BUN	Creatinine	Ca	Mg	Ca/Mg	Time Glucose	Fasting Glucose	Insulin	HOMA IR	MSS
ate:Time	140	IN.	DOIN	Creatifilite	Ca	IMIG	Caring	Giucose	r asting Glucose	IIISUIIII	HOWA IN	IVIS S
			13		g		10					9 - 1
												Þ

Column 4 -

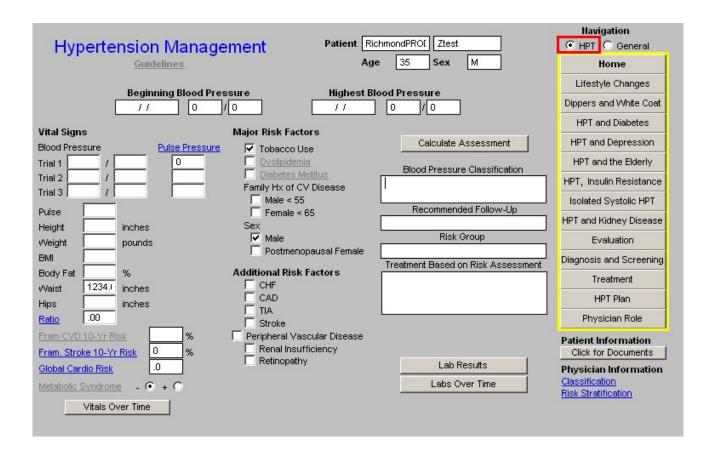
Navigation Buttons - there are two lists of navigation buttons which are displayed depending upon which of two boxes are checked:

- HPT
- General

Below are listed the templates which are displayed when each is checked in turn:

HPT - if this box is checked the templates which are specific to the disease management of the patient with hypertension are displayed:

- Home
- Lifestyle Changes
- Dippers and White Coat
- HPT and Diabetes
- HPT and Depression
- HPT and the Elderly
- HPT, Insulin Resistance
- Isolated Systolic HPT
- HPT and Kidney Disease
- Evaluation
- Diagnosis and Screening
- Treatment
- HPT Plan
- Physician Role

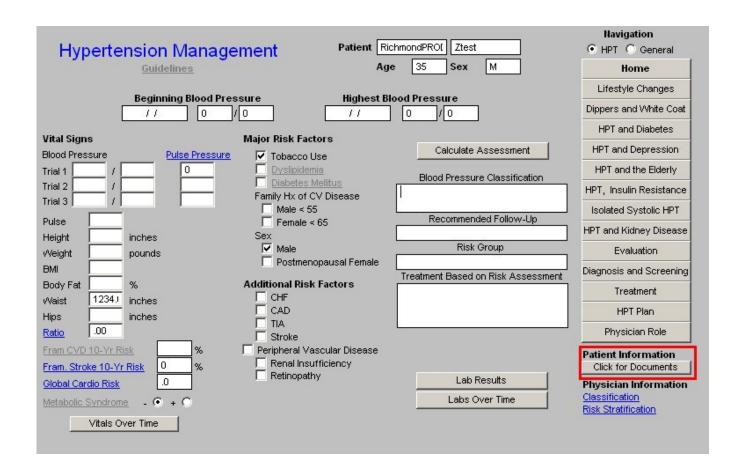


General - if this box is checked the templates which are drawn from the Master GP Suite of templates are displayed. Their use with the Hypertension Suite allows a patient's entire encounter to be documented from this suite, if no other issues are being addressed in this encounter.

- Return
- Chief/Chronic
- HPI
- Histories
- System Review
- Physical Exam

Hypertension Mana	agement Patient Ric	chmondPROI Ztest	Navigation C HPT General
Guidelines	Ag	e 35 Sex M	Return
Davissis - Diagram	December 115 of District Distr		Chief/Chronic
Beginning Blood	/ O / /	ood Pressure	HPI
			Histories
Vital Signs	Major Risk Factors	Calculate Assessment	System Review
Blood Pressure Pulse Press	1 10000000000	Calculate F 100000 Horiz	
Trial 1 0	☐ <u>Dyslipidemia</u> ☐ Diabetes Mellitus	Blood Pressure Classification	Physical Exam
Trial 2	Family Hx of CV Disease		
Trial 3 /	☐ Male < 55		
Pulse	☐ Female < 65	Recommended Follow-Up	1
Height inches	Sex	Risk Group	
Veight pounds	✓ Male ☐ Postmenopausal Female	Risk Group	1
BMI		Treatment Based on Risk Assessment	
Body Fat %	Additional Risk Factors	Trouble Business Common Troubles]
vVaist 1234.1 inches	☐ CHF ☐ CAD		
Hips inches	T TIA		
Ratio .00	Stroke		4
Fram CVD 10-Yr Risk %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk 0 %	Renal Insufficiency		Click for Documents
Global Cardio Risk .0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome - 6 + C		Labs Over Time	Classification
			Risk Stratification
Vitals Over Time			

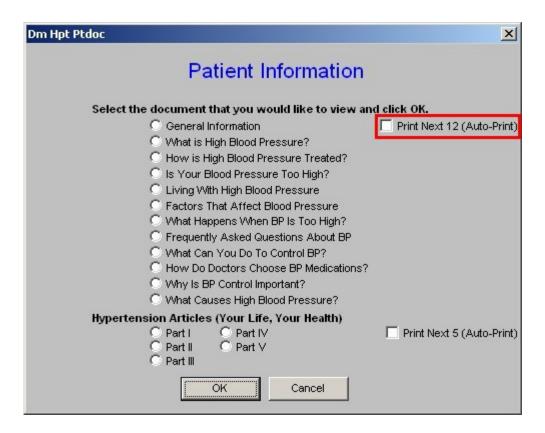
Beneath the navigation buttons are several patient information documents. When the button entitled "click for documents" is depressed the following list of documents appears.



The **first twelve** can be printed as a group and the **last five** can be printed as a group or they can be printed one at a time by following the instructions given.

- 1. General Information
- 2. What is High Blood Pressure?
- 3. How is High Blood Pressure Treated?
- 4. Is Your Blood Pressure too High?
- 5. Living with High Blood Pressure
- 6. Factors That Affect Blood Pressure
- 7. What Happens When BP is Too High?
- 8. Frequently Asked Questions About BP
- 9. What Can You Do to Control BP?
- 10. How Do Doctors Choose BP Medications?
- 11. Why is BP Control Important?
- 12. What Causes High Blood Pressure

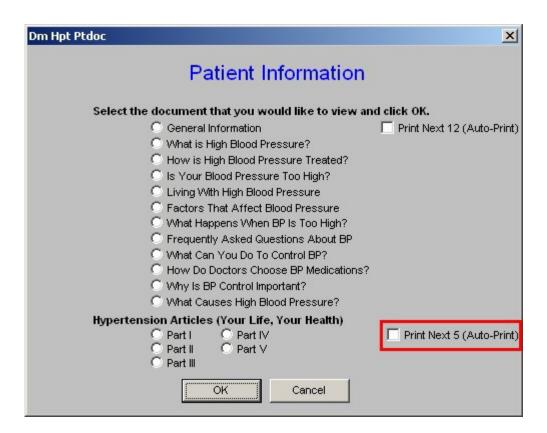
The function entitled "Print Next 12 (auto-print)" prints the above 12 documents



Hypertension Articles (Your Life, Your Health)

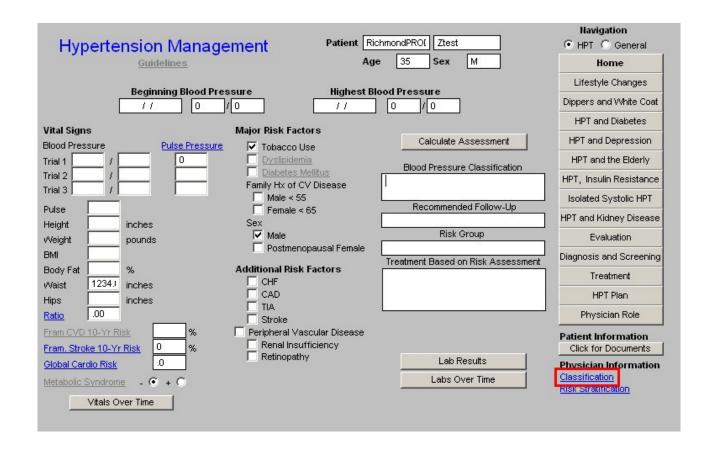
- 1. Part 1
- 2. Part II
- 3. Part III
- 4. Part IV
- 5. Part V

The function entitled "Print Next 5 (auto print)" prints these five articles.



Physician Information - This is beneath the Patient Information function described above.

• **Classification** - this displays a table with the blood pressure classifications which are automatically calculated as described above.





Classification of Blood Pressure Adults > 18 Years of Age

<u>Category</u>	Systolic (mmHg)		Diastolic (mmHg)
Optimal	<120	and	<80
Pre-Hypertension	121 - 130	or	81 - 85
High	131 - 139	or	86 - 89
Hypertension			
Stage 1	140 - 159	or	90 - 99
Stage 2	160 - 179	or	100 - 109
Stage 3	>179	or	>109

• **Risk Stratification** - this launches a document with a table which shows the calculation of the Risk Categories. In addition the following information is given:

Hypertens	ion Manage	ment Patient R	ichmondPROI Ztest	Havigation ⊙ HPT ○ General
	Guidelines	Ag	ge 35 Sex M	Home
_				Lifestyle Changes
Be	eginning Blood Pres	Sure Highest E	Blood Pressure	Dippers and White Coat
				HPT and Diabetes
Vital Signs Blood Pressure	Pulse Pressure	Major Risk Factors ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslipidemia	DI ID OLIGINA	HPT and the Elderly
Trial 2		Diabetes Mellitus Family Hx of CV Disease	Blood Pressure Classification	HPT, Insulin Resistance
Trial 3 /		Male < 55		Isolated Systolic HPT
Pulse		Female < 65	Recommended Follow-Up	HPT and Kidney Disease
	hes unds	Sex ✓ Male	Risk Group	Evaluation
BMI D	an 100	Postmenopausal Female	Transfer of Board or Birth Assessment	Diagnosis and Screening
Body Fat %		Additional Risk Factors	Treatment Based on Risk Assessment	Treatment
	hes	☐ CHF ☐ CAD		HPT Plan
Hips incl	hes	□ TIA		Physician Role
Fram CVD 10-Yr Risk	~	Stroke Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk	<u>k</u> 0 %	Renal Insufficiency		Click for Documents
Global Cardio Risk	.0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome	- 💿 + 🔘		Labs Over Time	Classification Risk Stratification
Vitals Over T	Time			



our Blood Pressure	Your Risk Group (Se	ee Below)	
	Risk Group A	Risk Group B	Risk Group C
130-139/85-89		Lifestyle Modification	Drug therapy
high-normal			
140-159/90-99		Lifestyle Modification	Drug therapy
(called stage 1 hypertension)		(up to 12 months)	(up to 6 months)
> or = 160/> or = 100	Drug Therapy	Drug Therapy	Drug therapy
(Called stages 2 & 3 hypertension)			

1. Risk Group A

- a. No Risk Factors
- b. No Target Organ Damage
- c. No clinical cardiovascular disease

2. Risk Group B

- a. At least 1 risk factor, not including Diabetes
- b. No target organ damage
- c. No clinical cardiovascular disease

3. Risk Group C

- a. Target organ damage/clinical cardiovascular disease and/or Diabetes
- b. Other Risk Factors

4. Risk Factors

- a. Tobacco use
- b. Dyslipidemia,
- c. Diabetes Mellitus,
- d. Sex (male and postmenopausal women),
- e. Family history of cardiovascular disease

5. Target Organ Damage -

- a. LVH,
- b. CHF,
- c. CAD,
- d. Stroke,
- e. TIA,
- f. Peripheral Vascular Disease,
- g. Renal Insufficiency,
- h. Retinopathy

Lifestyle Changes Template

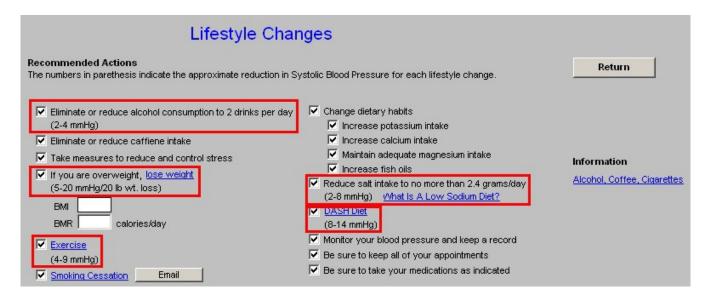
Hyperten	sion Manage	illetit –	RichmondPROI Ztest ge 35 Sex M	Havigation HPT General Home
	Beginning Blood Pres	sure Highest E	Blood Pressure	Lifestyle Changes
	// 0 /	0 //	0 /0	Dippers and White Coat
Vital Signs		Major Risk Factors		HPT and Diabetes
Blood Pressure	Pulse Pressure	▼ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /		Diabetes Mellitus Family Hx of CV Disease	Diodu Pressure Classification	HPT, Insulin Resistance
Trial 3		☐ Male < 55		Isolated Systolic HPT
Pulse ii	nches	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
	nones oounds	Male	Risk Group	Evaluation
BMI D		Postmenopausal Female	Total Book Biological Control	Diagnosis and Screening
Body Fat 9	%	Additional Risk Factors	Treatment Based on Risk Assessment	Treatment
Waist 1234.	nches	☐ CHF ☐ CAD		
	nches	□ CAD □ TIA		HPT Plan
Ratio .00		Stroke		Physician Role
Fram CVD 10-Yr Risk	<u> </u>	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr R	tisk 0 %	Renal Insufficiency		Click for Documents
Global Cardio Risk	.0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome	- O + O		Labs Over Time	Classification
Vitals Ove	er Time			Risk Stratification

Lifestyle changes are the foundation of the treatment of virtually any chronic disease. On this template there are **12 Recommended Actions** for the control and improvement of hypertension through lifestyle changes. The following describes how this template works.

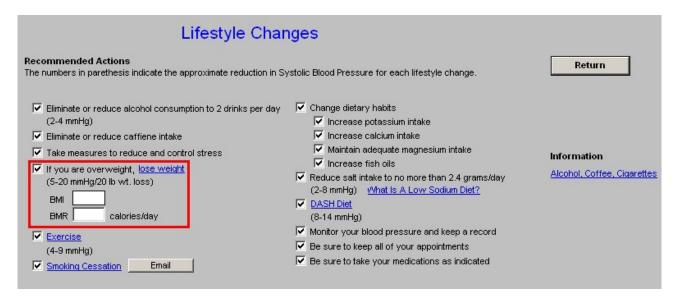
- The 12 Recommended Actions are auto checked as they apply to anyone with hypertension or pre-hypertension.
- When this template is accessed, these recommended actions will appear on the
 Hypertension Follow-up Note which should be given to the patient each time they are
 treated or assessed for hypertension.
- The material described below on a **Low Sodium Diet** and the **DASH Diet** also automatically print with the **Hypertension Follow-up Note**.
- Those actions which have a proven effect on hypertension have a notation concerning the degree of improvement in systolic blood pressure which can be reasonably expected from that lifestyle action.
- The Blood Pressure reductions which can reasonably be expected from the 12 Recommended actions are:
 - 1. A 2-4 mmHg lowering from decreasing or eliminating alcohol use.
 - 2. A 5-20 mmHg lowering for every 20 pounds of weight loss.
 - 3. A 4-9 mmHg lowering from exercise

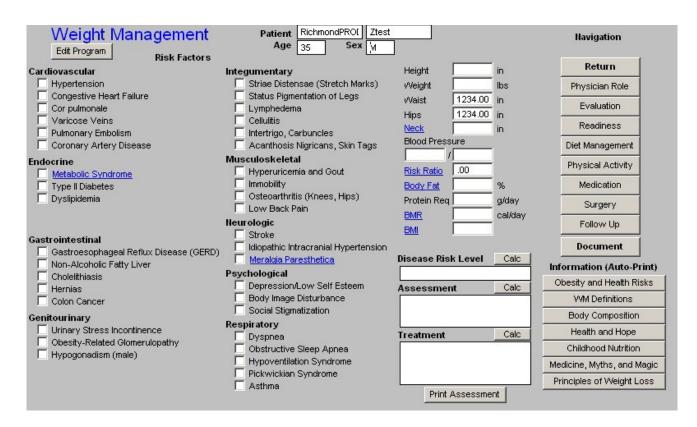
- 4. A 2-8 mmHg lowering from a reduced salt intake
- 5. A 8-14 mmHg lowering from following the DASH diet.

Contrast this with the fact that a single medication will generally provide a maximum of a 10 mmHg lowering of the systolic pressure.

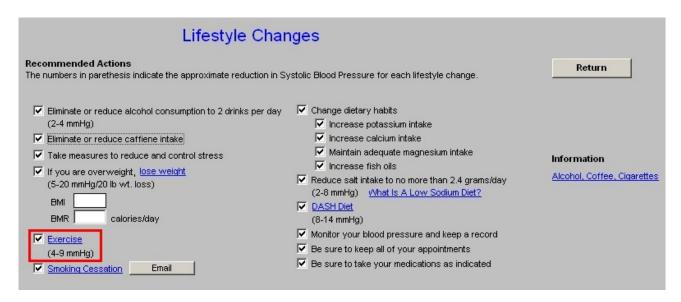


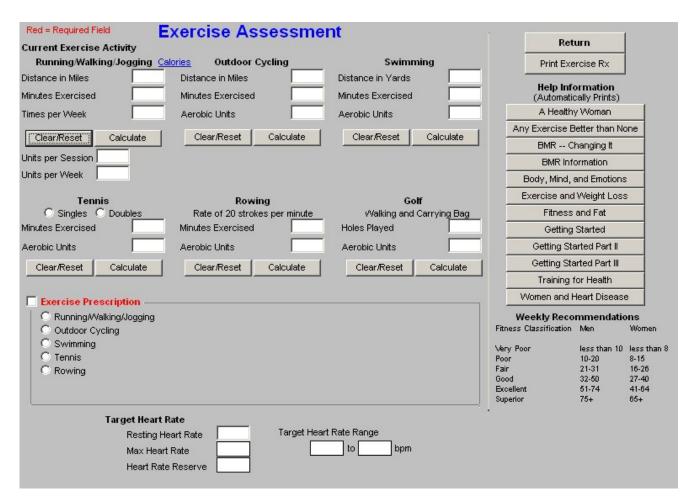
- The 12 Recommended Actions are:
 - 1. Eliminate or reduce alcohol consumption to 1 drink a day (2 4 mmHg)
 - 2. Eliminate or reduce caffeine intake.
 - 3. Take measures to reduce and control stress.
 - 4. **If you are overweight, lose weight** (5-20 mmHg/20 lb. wt. loss) -- there is also a hyperlink to SETMA's Weight Management Suite of Templates. And, there are boxes where the patient's BMI and BMR are automatically displayed.





5. **Exercise** (4-9 mmHg) - there is also a hyperlink to SETMA's Exercise template. For more information on the Exercise Prescription template, Click Here





6. **Smoking Cessation** - there is also a hyperlink to SETMA's Smoking Cessation Template and a button which launches an email for the creation of an electronic tickler file.

Lifestyle Chan	iges	
Recommended Actions The numbers in parethesis indicate the approximate reduction in Sy	stolic Blood Pressure for each lifestyle change.	Return
▼ Eliminate or reduce alcohol consumption to 2 drinks per day (2-4 mmHg) ▼ Eliminate or reduce caffiene intake ▼ Take measures to reduce and control stress ▼ If you are overweight, lose weight (5-20 mmHg/20 lb wt. loss) ■ ■ ■ ■ ■ ■ ■	✓ Change dietary habits ✓ Increase potassium intake ✓ Increase calcium intake ✓ Maintain adequate magnesium intake ✓ Increase fish oils ✓ Reduce salt intake to no more than 2.4 grams/day (2-8 mmHg)	Information Alcohol, Coffee, Cigarettes

Red = Required Field	Smoking Cessation	Last Chest X-Ray	
Patient currently Check here if pai Pipe? Smokeless Toba Date stopped sm Packs per day?	ent has quit! Smoke at home O Yes O No Has the patient co? Yes No	t committed to quit? C Yes C No	Return Pharmacotherapy Document
Ask	At every visit, ask all patients about tobacco use Patients who have never used tobacco or who seed repeated assessments.		Information General Information Process of Guitting Smoking
Advise	☐ Let patients know, in a clear, strong, and person to quit. Men who smoke cut their lives she Women smokers lose 14.5 years o	ort by 13.2 years	Second Hand Smoke
Assess	Find out whether patients are willing to quit now	or at least within the next 30 days.	
Assist	reviewing past attempts to quit nicotin	pating challenges such as ne withdrawal symptoms g total abstinence tion.	
Arrange Follow-Up	Smokers trying to quit are at high risk of relapse, weeks after the quit date. Follow up in person of Email Tickler File Scheduled Date?		

7. Change Dietary habits

- a. Increase potassium intake
- b. Increase calcium intake
- c. Maintain adequate magnesium intake
- d. Increase fish oils
- 8. **Reduce Salt intake to no more than 2.4 grams/day** (2-8 mmHg) there is also a printable document entitled What is a low Sodium Diet?
- 9. **DASH Diet** (8-14 mmHg) there is also a hyperlink which prints an introduction to the DASH Diet **Dietary Approach to Stop Hypertension**.
- 10. Monitor your blood pressure and keep a record
- 11. Be sure to keep all appointments
- 12. Be sure to take your medications as indicated.

Information

A patient information document is launched and printed from this template entitled, "Alcohol, Coffee, Cigarettes."

Dippers and White Coat Syndrome Template

The concepts of "dipping," "non-dipping," and "white coat syndrome" are very important to the proper evaluation and treatment of hypertension. The frequent reviewing of the information on this template will help keep these principles in mind as patients are evaluated for decreasing their stroke risk.

Hypertension Manage	ement Patient Ric	chmondPROI Ztest	Navigation • HPT • General
Guidelines	Age	e 35 Sex M	Home
Beginning Blood Pre	oouro Highoot Pl	ood Pressure	Lifestyle Changes
	/O //	0 /0	Dippers and White Coat
Vital Signs	Major Risk Factors		HPT and Diabetes
Blood Pressure Pulse Pressure	▼ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / 0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2	Diabetes Mellitus Family Hx of CV Disease		HPT, Insulin Resistance
Trial 3 /	Male < 55	Recommended Follow-Up	Isolated Systolic HPT
Pulse Inches	Female < 65 Sex	Recommended Follow-op	HPT and Kidney Disease
Weight pounds	✓ Male	Risk Group	Evaluation
BMI	Postmenopausal Female	Treatment Based on Risk Assessment	Diagnosis and Screening
Body Fat %	Additional Risk Factors	Treatment based of Misk Assessment	Treatment
Waist 1234. inches	☐ CHF ☐ CAD		HPT Plan
Hips inches	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Risk %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk 0 %	Renal Insufficiency Retinopathy		Click for Documents
Global Cardio Risk .0	Retiliopatriy	Lab Results	Physician Information
Metabolic Syndrome - © + C		Labs Over Time	Classification Risk Stratification
Vitals Over Time			The state of the s

There are four pop-ups on this template.

- **Dippers and Non-Dippers** this addresses the very important physiological decreasing of blood pressure in the early morning hours. There three educational documents associated with this phenomenon: **Circadian Rhythms**, **General Information**, **Dippers and Sodium Shifts**. The pop-up displays the following information:
 - 1. **Circadian Rhythms** -- Blood pressure and heart rate tend to be lowest between 2-4 AM. Levels begin to increase after that and before awakening.
 - 2. **Dipping** -- The characteristic decrease in blood pressure at night.
 - 3. **Non-Dippers** -- Some patients do not have this characteristic reduction in blood pressure at night. This has been described consistently in African-Americans with hypertension and in diabetics.
 - 4. Risk
 - a. Non-dipping appears to increase cardiovascular risk.
 - b. CV risk is most closely correlated with total 24-hour pressure load.
 - c. When the pressure "dips" at night, the overall blood pressure load is reduced.
 - d. For non-dippers, the overall blood pressure load is not decreased, thus increasing cardiovascular risk.
 - 5. Other Rhythms with CV Risk

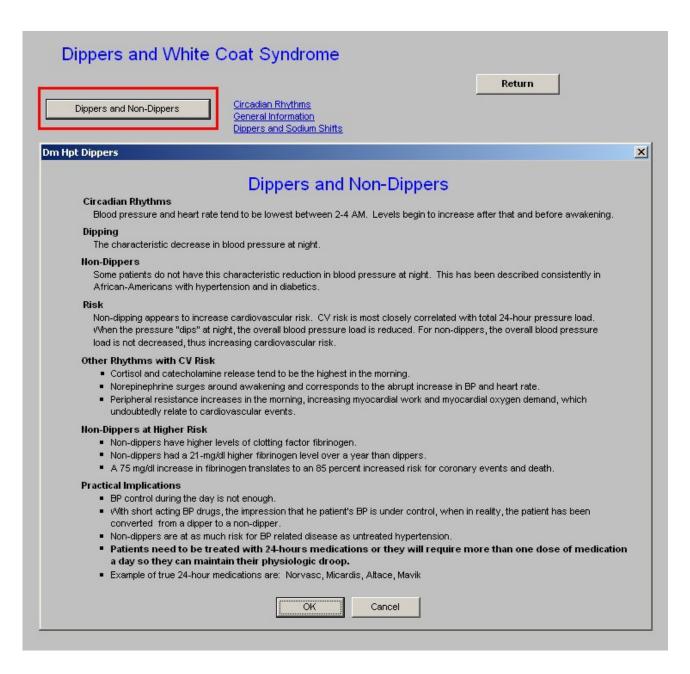
- a. Cortisol and catecholamine release tend to be the highest in the morning.
- b. Norepinephrine surges around awakening and corresponds to the abrupt increase in BP and heart rate.
- c. Peripheral resistance increases in the morning, increasing myocardial work and myocardial oxygen demand, which undoubtedly relate to cardiovascular events.

6. Non-Dippers at Higher Risk

- a. Non-dippers have higher levels of clotting factor fibringen.
- b. Non-dippers had a 21-mg/dl higher fibrinogen level over a year than dippers.
- c. A 75 mg/dl increase in fibrinogen translates to an 85 percent increased risk for coronary events and death.

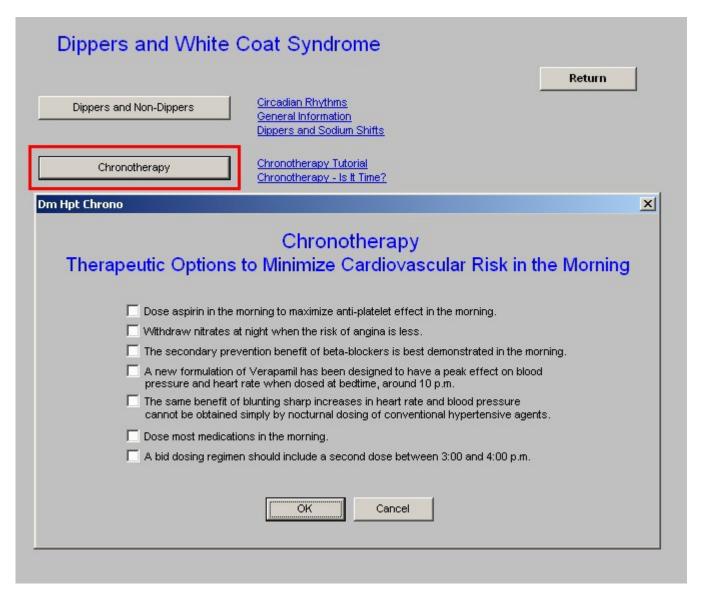
7. Practical Implications

- a. BP control during the day is not enough.
- b. With short acting BP drugs, the impression that he patient's BP is under control, when in reality, the patient has been converted from a dipper to a non-dipper.
- c. Non-dippers are at as much risk for BP related disease as untreated hypertension.
- d. Patients need to be treated with 24-hours medications or they will require more than one dose of medication a day so they can maintain their physiologic droop.
- e. Example of true 24-hour medications are: Norvasc, Micardis, Altace, Mavik



- Chronotherapy the subtitle of this pop-up is "Therapeutic Options to Minimize Cardiovascular Risk." The seven facts about the "timing" of medication administration to improve its beneficial effects is an introduction to the large field of chronotherapy. This pop-up has two documents associated with it: Chronotherapy: A Tutorial and Chronotherapy: Is it Time?
 - 1. Chronotherapy
 - 2. Therapeutic Options to Minimize Cardiovascular Risk in the Morning
 - a. Dose aspirin in the morning to maximize anti-platelet effect in the morning.
 - b. Withdraw nitrates at night when the risk of angina is less.
 - c. The secondary prevention benefit of beta-blockers is best demonstrated in the morning.

- d. A new formulation of Verapamil has been designed to have a peak effect on blood pressure and heart rate when dosed at bedtime, around 10 p.m.
- e. The same benefit of blunting sharp increases in heart rate and blood pressure cannot be obtained simply by nocturnal dosing of conventional hypertensive agents.
- f. Dose most medications in the morning.
- g. A bid dosing regimen should include a second dose between 3:00 and 4:00 p.m.

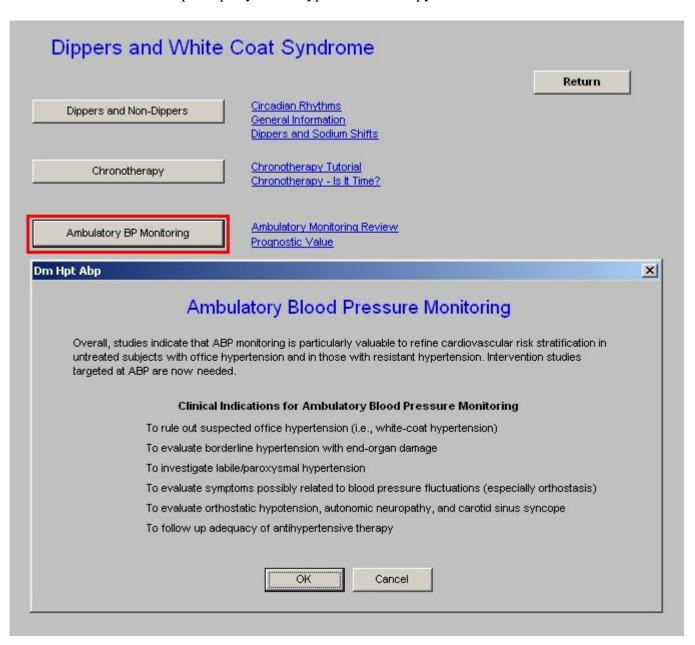


• Ambulatory BP Monitoring - the following information appears on this pop-up. Two documents are associated with this pop-up: Ambulatory Monitoring Review and Prognostic Value.

Overall, studies indicate that ABP monitoring is particularly valuable to refine

cardiovascular risk stratification in untreated subjects with office hypertension and in those with resistant hypertension. Intervention studies targeted at ABP are now needed.

- 1. Clinical Indications for Ambulatory Blood Pressure Monitoring
- 2. To rule out suspected office hypertension (i.e., white-coat hypertension)
- 3. To evaluate borderline hypertension with end-organ damage
- 4. To investigate labile/paroxysmal hypertension
- 5. To evaluate symptoms possibly related to blood pressure fluctuations (especially orthostasis)
- 6. To evaluate orthostatic hypotension, autonomic neuropathy, and carotid sinus syncope
- 7. To follow up adequacy of antihypertensive therapy



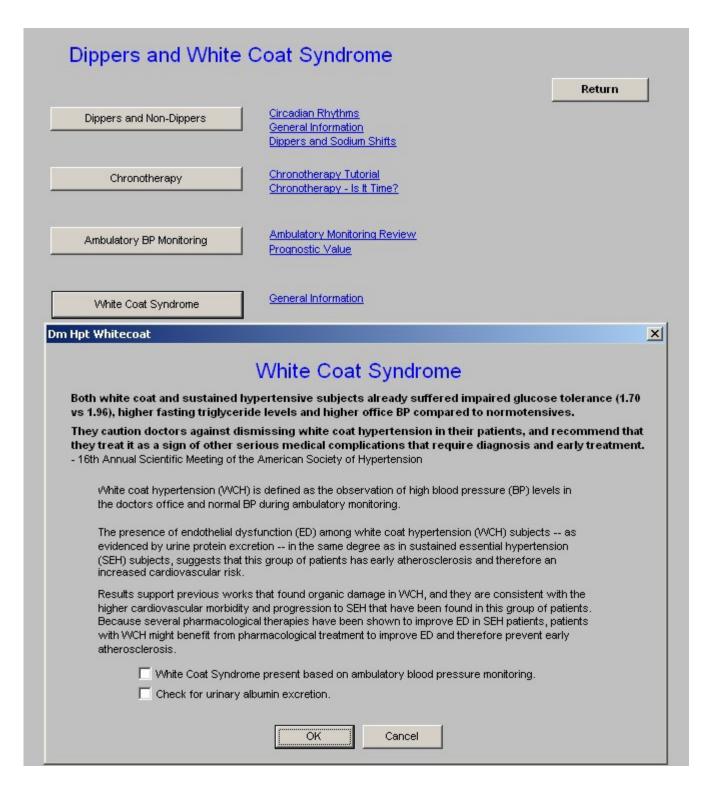
- White Coat Syndrome the following information appears on this pop-up which is associated with a document entitled, General Information. Healthcare providers are cautioned about dismissing patients with "white coat syndrome" as it increases cardiovascular risk.
 - 1. Both white coat and sustained hypertensive subjects already suffered impaired glucose tolerance (1.70 vs 1.96), higher fasting triglyceride levels and higher office BP compared to normotensives.
 - 2. They caution doctors against dismissing white coat hypertension in their patients, and recommend that they treat it as a sign of other serious medical complications that require diagnosis and early treatment.

- 16th Annual Scientific Meeting of the American Society of Hypertension

- 3. White coat hypertension (WCH) is defined as the observation of high blood pressure (BP) levels in the doctors office and normal BP during ambulatory monitoring.
- 4. The presence of endothelial dysfunction (ED) among white coat hypertension (WCH) subjects -- as evidenced by urine protein excretion -- in the same degree as in sustained essential hypertension (SEH) subjects, suggests that this group of patients has early atherosclerosis and therefore increased cardiovascular risk.
- 5. Results support previous works that found organic damage in WCH, and they are consistent with the higher cardiovascular morbidity and progression to SEH that have been found in this group of patients. Because several pharmacological therapies have been shown to improve ED in SEH patients, patients with WCH might benefit from pharmacological treatment to improve ED and therefore prevent early atherosclerosis.

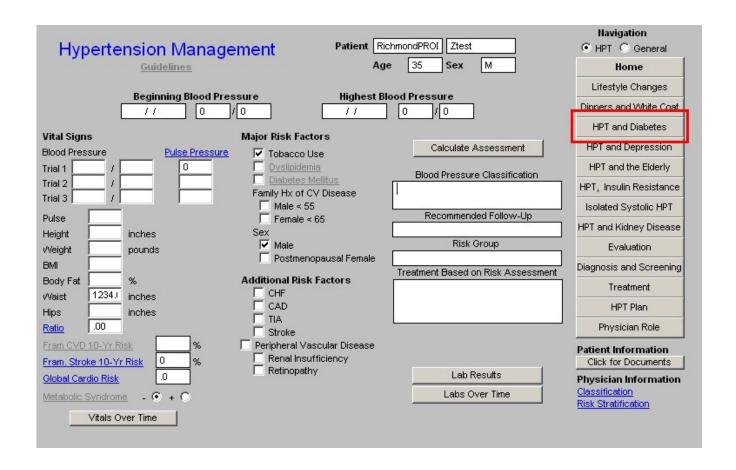
The following two are check boxes for further evaluation of the patient with suspicion of WCS:

- White Coat Syndrome present based on ambulatory blood pressure monitoring.
- Check for urinary albumin excretion.

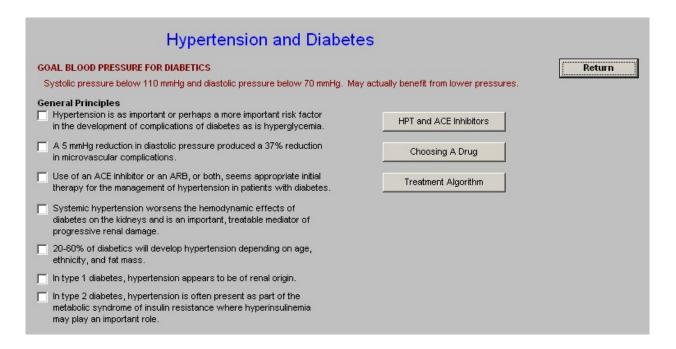


Hypertension and Diabetes Template

This template addresses the goals and principles of the treatment of hypertension in the patient with diabetes. Its review and the knowledge and use of the algorithms contained in it are key to the successful treatment of hypertension in the patient with diabetes.



When the Navigation button entitled HPT and Diabetes is depressed the following template is launched.



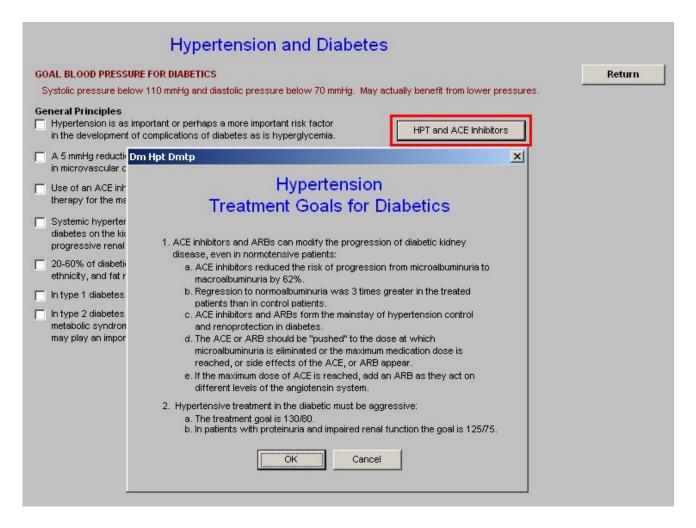
- Goal 110/70 mm Hg
- General Principles
 - 1. Hypertension is as important or perhaps a more important risk factor in the development of complications of diabetes as is hyperglycemia.
 - 2. A 5 mmHg reduction in diastolic pressure produced a 37% reduction in microvascular complications.
 - 3. Use of an ACE inhibitor or ARB, or both, seems appropriate initial therapy for the management of hypertension in patients with diabetes.
 - 4. Systemic hypertension worsens the hemodynamic effects of diabetes on the kidneys and is an important, treatable mediator of progressive renal damage.
 - 5. 20-60% of diabetics will develop hypertension depending on age, ethnicity, and fat mass.
 - 6. In type 1 diabetes, hypertension appears to be of renal origin.
 - 7. In type 2 diabetes, hypertension is often present as part of the metabolic syndrome of insulin resistance where hyperinsulinemia may play an important role.

Three pop-ups appear on this template; they are:

• HPT and Ace Inhibitors - this pop-up displays the following information

Treatment Goals for Diabetics

- 1. ACE inhibitors and ARBs can modify the progression of diabetic kidney disease, even in normotensive patients:
 - a. ACE inhibitors reduced the risk of progression from microalbuminuria to macroalbuminuria by 62%.
 - b. Regression to normoalbuminuria was 3 times greater in the treated patients than in control patients.
 - c. ACE inhibitors and ARBs form the mainstay of hypertension control and renoprotection in diabetes.
 - d. The ACE or ARB should be "pushed" to the dose at which microalbuminuria is eliminated or the maximum medication dose is reached, or side effects of the ACE, or ARB appear.
 - e. If the maximum dose of ACE is reached, add an ARB as they act on different levels of the angiotensin system.
- 2. Hypertensive treatment in the diabetic must be aggressive:
 - a. The treatment goal is 130/80.
 - b. In patients with proteinuria and impaired renal function the goal is 125/75.



• Choosing a Drug - this pop-up displays the following information on five classes of antihypertensive medications:

1. ACE Inhibitors

- a. Renal protective. Diminish proteinuria.
- b. Increase responsiveness to insulin.
- c. No adverse effect on lipids.
- d. May cause increase in potassium.

2. Calcium Channel Blockers

- a. Diltiazem & verapamil decrease proteinuria.
- b. Long term effect on renal function not known.
- c. No adverse effect on lipid or carbohydrate metabolism.

3. Low Dose thiazide/hydrochlorothiazide (12.5 to 25 mg max)

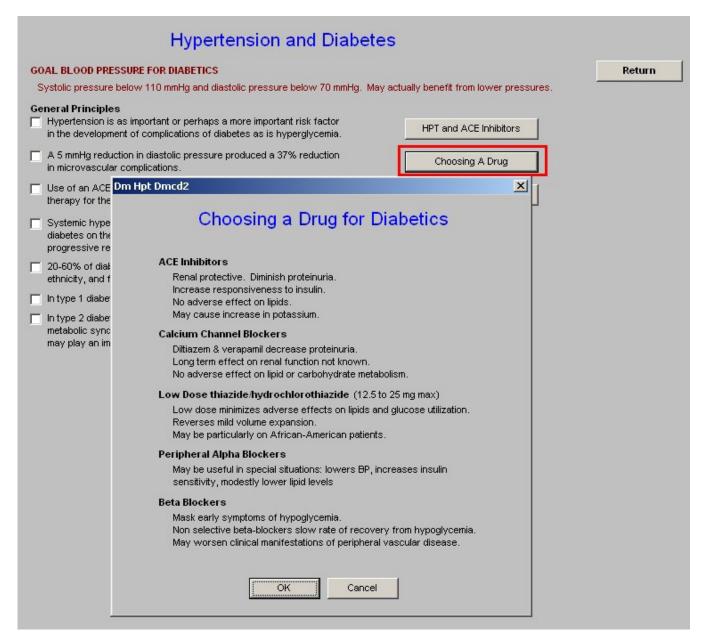
- a. Low dose minimizes adverse effects on lipids and glucose utilization.
- b. Reverses mild volume expansion.
- c. May be particularly on African-American patients.

4. Peripheral Alpha Blockers

May be useful in special situations: lowers BP, increases insulin sensitivity, modestly lower lipid levels

5. Beta Blockers

- a. Mask early symptoms of hypoglycemia.
- b. Non selective beta-blockers slow rate of recovery from Hypoglycemia.
- c. May worsen clinical manifestations of peripheral vascular disease.



• Treatment Algorithm - this pop-up displays information, "Hypertension: Choosing A Drug for Diabetic Patients."

- o First Choice is an ACE-Inhibitor there is a check box for selecting that option.
- o After that

Thiazide Diuretics - there is a check box Beta Blockers - there is a check box ARBs - there is a check box

NOTE: Blood Pressure goal is 110/70 mmHg.

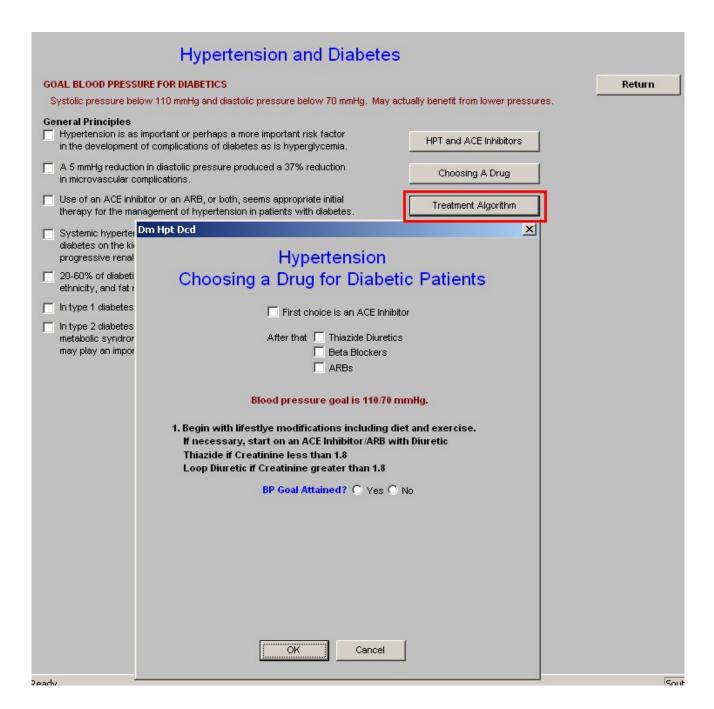
- Steps of treatment of hypertension in patients with diabetes
 - a. Begin with lifestyle modifications including diet and exercise
 - If necessary, start on an ACE Inhibitor/ARB with Diuretic

Thiazide if Creatinine less than 1.8 Loop Diuretic if Creatinine greater than 1.8

- BP Goal Attained? Yes? Stop; No? Proceed
- **b.** Add a long-acting nondihydrophyridine Calcium Channel Blocker such as Amlodipine or Norvasc
 - BP Goal Attained? Yes? Stop; No? Proceed
- c. What is the patient's resting pulse? If above 84 -
 - Add a Beta Blocker; if BP goal still not attained, refer to a hypertension specialist.

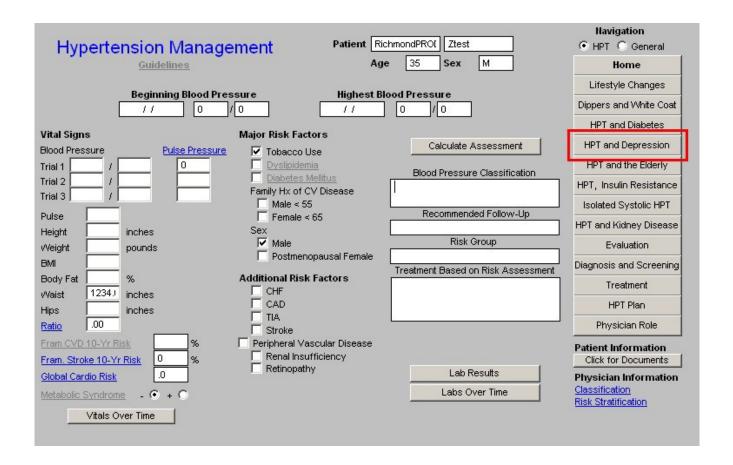
What is the patient's resting pulse? If below 84 -

• Add another subgroup of Calcium Channel Blockers. If BP still not attained, refer to a hypertension specialist.



Hypertension and Depression Template

This template addresses the relationship between both conditions and between the impact which the treatment of one can have on the other. The treatment of hypertension in the elderly is often complicated by the presence of depression particularly since the medications for both conditions often interact with one another.



There are three sections to the information on this template:

- Depression As A Risk Factor for Hypertension
 - 1. Dual Risk of Depression and HPT
 - 2. Mechanisms of Depression Causing HPT

High depression scores are an independent predictor of hypertension

- 3. Vigilance about the coexistence of depression and hypertension is particularly important in:
 - a. Elderly patients
 - b. Women
 - c. Separate or divorced persons
 - d. Those with a family history of depression

epression As A Risk Factor For Hypertension	Return
Dual Risk of Depression and HPT	
Mechanisms of Depression Causing HPT	
High depression scores are an independent predictor of hypertension:	
* In white adults aged 45 to 64, with a relative risk of hypertension of 1.80.	
* In African-American adults aged 25 to 64, with a relative risk of hypertension of 2.99.	
Vigilance about the coexistence of depression and hypertension is particularly important in groups at high risk of depression:	
* Elderly persons	
* Women	
Separated or divorced persons Those with a family history of depression.	
ntihypertensive Drugs Can Worsen Depression	
A link between antihypertensive medications and depression has been suspected for more than four decades.	
Antihypertensive Medications Causing Depression	
Interactions Between Antihypertensives and Antidepressants	
festyle aspects which may worsen depression and hypertension are	
☐ Smoking	
☐ Alcohol	

• Antihypertensive Drugs Can Worsen Depression

- 1. A link between antihypertensive medications and depression has been suspected for more than four decades.
- 2. Antihypertensive Medications Causing Depression
- 3. Interactions Between Antihypertensives and Antidepressants

epression As A Risk Factor For Hypertension	Return
Dual Risk of Depression and HPT Mechanisms of Depression Causing HPT	
High depression scores are an independent predictor of hypertension:	
 In white adults aged 45 to 64, with a relative risk of hypertension of 1.80. In African-American adults aged 25 to 64, with a relative risk of hypertension of 2.99. 	
Vigilance about the coexistence of depression and hypertension is particularly important in groups at high risk of depressi * Elderly persons * Women * Separated or divorced persons * Those with a family history of depression.	ion:
Antihypertensive Drugs Can Worsen Depression	
A link between antihypertensive medications and depression has been suspected for more than four decades.	
Antihypertensive Medications Causing Depression	
Interactions Between Antihypertensives and Antidepressants	
ifestyle aspects which may worsen depression and hypertension are	
□ Smoking	
☐ Alcohol	
☐ Inactivity	
Poor Diet	

- Lifestyle Aspects Which May Worsen Depression and Hypertension
 - Smoking
 Alcohol

 - 3. Inactivity
 - 4. Poor Diet

Hypertension and Depression Depression As A Risk Factor For Hypertension Return Dual Risk of Depression and HPT Mechanisms of Depression Causing HPT High depression scores are an independent predictor of hypertension: * In white adults aged 45 to 64, with a relative risk of hypertension of 1.80. * In African-American adults aged 25 to 64, with a relative risk of hypertension of 2.99. Vigilance about the coexistence of depression and hypertension is particularly important in groups at high risk of depression: * Elderly persons * Women * Separated or divorced persons * Those with a family history of depression. Antihypertensive Drugs Can Worsen Depression A link between antihypertensive medications and depression has been suspected for more than four decades. Antihypertensive Medications Causing Depression Interactions Between Antihypertensives and Antidepressants Lifestyle aspects which may worsen depression and hypertension are... □ Smoking ✓ Alcohol ☐ Inactivity Poor Diet

Hypertension and the Elderly Template

Hyperten	nsion Manage	ment Patient Ri	chmondPROI Ztest	Havigation HPT General
	Guidelines	Ay	e 55 Sex [W]	nome
	Beginning Blood Pres	eure Highest R	lood Pressure	Lifestyle Changes
		0 //	0 /0	Dippers and White Coat
Vital Signs		Major Risk Factors		HPT and Diabetes
Blood Pressure	Pulse Pressure	✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /		Diabetes Mellitus Family Hx of CV Disease		HPT, Insulin Resistance
Trial 3 / /		Male < 55		Isolated Systolic HPT
Pulse		Female < 65	Recommended Follow-Up	HPT and Kidney Disease
	inches	Sex ✓ Male	Risk Group	Evaluation
	pounds	Postmenopausal Female		Evaluation
BMI			Treatment Based on Risk Assessment	Diagnosis and Screening
	%	Additional Risk Factors		Treatment
Waist 1234.I	inches	☐ CHF ☐ CAD		HPT Plan
	inches	E TIA		net Plati
Ratio .00		Stroke		Physician Role
Fram CVD 10-Yr Risk	<u>k</u> %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr R	Risk 0 %	Renal Insufficiency		Click for Documents
Global Cardio Risk	.0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome	- 0 + 0		Labs Over Time	Classification
1/2-1-0	Time 1			Risk Stratification
Vitals Ove	er ime			

Blood pressure is a significant problem in the elderly, particularly when you realized that if you are 55 years of age and do not have hypertension, your lifetime risk of developing hypertension is 90%.. Controlling blood pressure, and particularly systolic blood pressure in the elderly, even those beyond 80 will decrease morbidity and mortality for cerebrovascular accidents and myocardial infarctions.

Hypertension in the Elderly

Treatment Algorithm JHC7 Treatment Recommendations Therapy Comparisons

Return

Information

Morning Blood Pressure

General Principles

- Antihypertensive therapy has been shown to reduce morbidity and mortality in older patients with elevated systolic or diastolic blood pressures.
- This benefit appears to persist in patients older than 80 years, but less than one third of older patients have adequate blood pressure control.
- Systolic blood pressure is the most important predictor of cardiovascular disease.
- Blood pressure measurement in older persons should include an evaluation for orthostatic hypotension.
- · Low-dose thiazide diuretics remain first-line therapy for older patients.
- Beta blockers, angiotensin-converting enzyme inhibitors, angiotensin-receptor blockers, and calcium channel blockers are second-line medications that should be selected based on comorbidities and risk factors.

Treatment Goals

- . The goal blood pressure recommended by JNC7 is
 - * less than 140/90 mmHq
 - * less than 120/70 mmHg in patients with diabetes or chronic kidney disease
- JNC7 recommends treating older patients with stage 1 isolated systolic hypertension (SBP between 140 and 159 mmHg) equally agressively.
- · Active treatment has been shown to reduce
 - * total mortality
 - * cardiovascular mortality
 - fatal or nonfatal cardiovascular events
 - * fatal or nonfatal stroke

The Elderly & Thiazide Diuretics

The Elderly & Beta Blockers

The Elderly & Calcium Channel Blockers

The Elderly & ACE Inhibitors and ARBs

There are three pop-ups at the top of this template; they are:

• Treatment Algorithm - this pop-up displays the following information

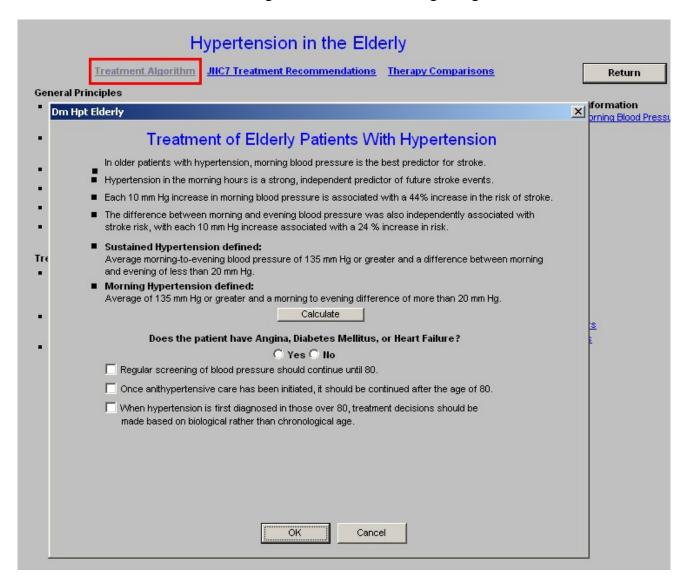
Treatment of Elderly Patients with Hypertension

- 1. In older patients with hypertension, morning blood pressure is the best predictor for stroke.
- 2. Hypertension in the morning hours is a strong, independent predictor of future stroke events.
- 3. Each 10 mm Hg increase in morning blood pressure is associated with a 44% increase in th risk of stroke.
- 4. The difference between morning and evening blood pressure was also independently associated with stroke risk, with each 10 mm Hg increase associated with a 24\$ increase in risk.
- 5. **Sustained Hypertension defined:** -- Average morning-to-evening blood pressure of 135 mm Hg or greater and a difference between morning and evening of less than 20 mm Hg
- 6. **Morning Hypertension defined:** Average of 135 mm Hg or greater and a morning to evening difference of more than 20 mm Hg. At this point on the Treatment of Elderly Patients with Hypertension algorithm there appears a button entitled Calculate.
- 7. When depressed the Calculate button launches a pop-up with the following title Sustained vs. Morning Hypertension

- a. Enter the patient's average morning and evening blood pressure and lcikc calculate.
- b. In the box present the conclusion of morning or sustained blood pressure will appear.

A question is then asked: "Does the patient have angina, diabetes mellitus, or heart failure?" Yes/No

- 1. Check boxes are then provided to guide the following:
 - a. Regular screening of blood pressure should continue until 80.
 - b. Once antihypertensive care has been initiated, it should be continued after the age of 80.
 - c. When hypertension is first diagnoses in those over 80, treatment decisions should be made based on biological rather than chronological age.



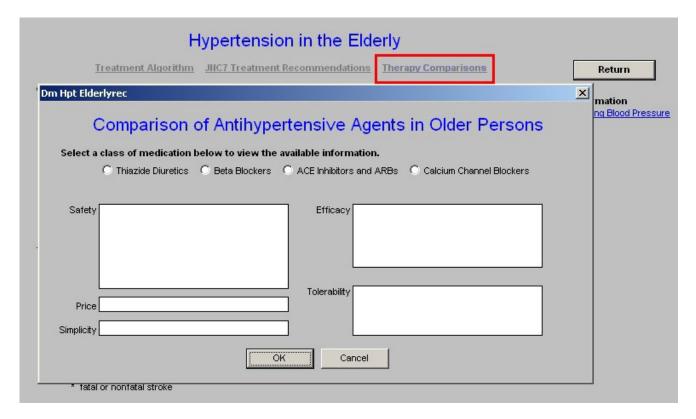
• **JNC7 Treatment Recommendations** - this pop-up displays the following principles of the treatment of hypertension in the elderly:

Note: JNC 7 recommendations for treating hypertension are similar in the general population and older persons. The key points include:

- 1. Treat isolated systolic hypertension
- 2. Thiazide diuretics should be first-line treatment.
- 3. Second-line treatment should be based on comorbidites and risk factors.
- 4. Patients with systolic blood pressure higher than 160 mm Hg or diastolic blood pressure higher than 100 mm Hg usually will require two or more agents to reach goal.
- 5. Treatment should be initiated with a low dose of the hosen antihypertensive agent, and titrated slowly to minimize side effects such as orthostatic hypertension
- 6. Weight loss and sodium reduction have been shown to be feasible and effective interventions in older patients with hypertension. JNC 7 recommends adoption fo the Dietary Approaches of Stop Hypertension (DASH) diet, which has been show to produce blood pressure reductions similar to single-drug therapy.
- 7. To improve adherence with antihypertensive regimens, involve patients in goal setting, and ensure that the patients' cultural beliefs and previous experiences are incorporated in a treatment plan. Simply the medication regimen, keeping in mind how much it costs.



- **Therapy Comparisons** this pop-up compares the safety, price and simplicity of use of four classes of medications in the elderly:
 - 1. Thiazide Diuretics,
 - 2. Beta Blockers,
 - 3. ACE Inhibitors and
 - 4. Calcium Channel Blockers.



The remainder of the template details **General Principles and Goals of treatment** in the elderly:

• General Principles:

- 1. Antihypertensive therapy has shown to reduce morbidity and mortality older patients with elevated systolic and diastolic blood pressures
- 2. This benefit appears to persist in patients older than 80 years, but less than one third of older patients have adequate pressure control.
- 3. Systolic Blood pressure is the most important predictor of cardiovascular disease.
- 4. Blood pressure measurement in older persons should include an evaluation for orthostatic hypotension.
- 5. Low-dose thiazide diuretics remain first-line therapy for older patients
- 6. Beta blockers, angiotensin-converting enzyme inhibitors, angiotensin-receptor blockers and calcium channel blockers are second-line medications that should be selected based on comorbidities and risk factors.

• Treatment Goals

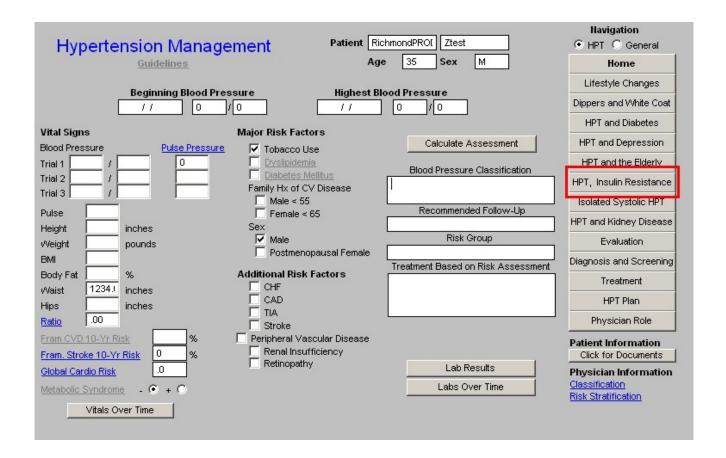
- 1. The goal blood pressure recommended by JNC7 is:
 - a. Less than 140/90 mmHg
 - b. Less than 110/70 mmHg in patients with diabetes or chronic kidney disease.
- 2. JNC 7 recommends treating older patents with stage 1 systolic hypertension (SBP between 140 and 150 mmHg) equally aggressively
- 3. Active treatment has been shown to reduce
 - a. Total mortality
 - b. Cardiovascular mortality
 - c. Fatal or nonfatal cardiovascular events
 - d. Fatal or nonfatal stroke.

There is a series of four documents linked to this template; they are:

- The Elderly and Thiazide Diuretics
- The Elderly and Beta Blockers
- The Elderly and Calcium Channel Blockers
- The Elderly and Ace Inhibitors and ARBs

One last information document is launched from a link entitled, "Morning Blood Pressure." This document addresses the following concept "Early to Rise - Morning Blood Pressure Predicts Stroke."

HPT, Insulin Resistance Template



In SETMA's **Metabolic Syndrome Suite of Templates**, on the **Insulin Resistance Template**, the following statements are made:

- Insulin resistance contributes to decreased renal function reserve and to the development of hypertension
- Insulin resistance is one of the causes of treatment-resistant essential hypertension

The medical literature is not unanimous in these conclusions at present but the association between insulin resistance and the other markers of the metabolic syndrome - obesity, dyslipidemia, hyperglycemia, renal damage as evidenced by microalbuminuria - all of which are both causes of and results of endothelium dysfunction, is strong. So much so, that in the face of treatment resistant hypertension, it is worthwhile to address the issue of insulin resistance and endothelium dysfunction.

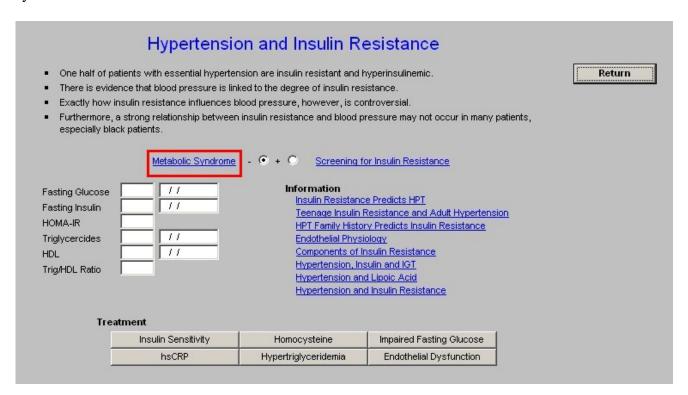
 One half of pati 	ents with essential hyperte	ension are insulin resistant and l	hyperinsulinemic	Return
		nked to the degree of insulin res		
		blood pressure, however, is co		
 Furthermore, a especially black 		n insulin resistance and blood p	oressure may not occur in many patient	s,
	Metabolic Syndrome	e - 🗨 + 🔘 Screening f	for Insulin Resistance	
Fasting Glucose	11	Information	Possibility	
Fasting Insulin	11	Insulin Resistano Teenage Insulin	Resistance and Adult Hypertension	
HOMA-IR			ory Predicts Insulin Resistance	
Triglycercides	1.1	Endothelial Phys	iology	
HDL	11	The same and the s	Insulin Resistance	
Trig/HDL Ratio		Hypertension, In	The state of the s	
		Hypertension an	id Lipoic Acid id Insulin Resistance	
		Trypertensional	id in Suin 1 (CSIStanCC	
Treat	ment			
11040				
	Insulin Sensitivity	Homocysteine	Impaired Fasting Glucose	

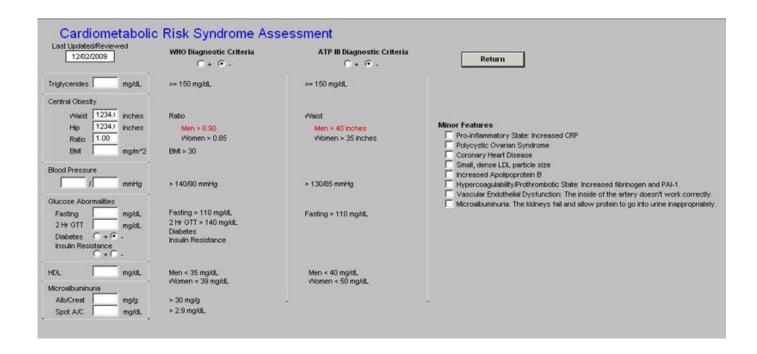
At the top of HPT, Insulin Resistance template are the following General Concepts:

- One half of patents with essential hypertension are insulin resistant and hyperinsulinemic.
- There is evidence that blood pressure is linked to the degree of insulin resistance.
- Exactly how insulin resistance influences blood pressure, however, is controversial.
- Furthermore, a strong relationship between insulin resistance and blood pressure may not occur in many patients, especially black patients.

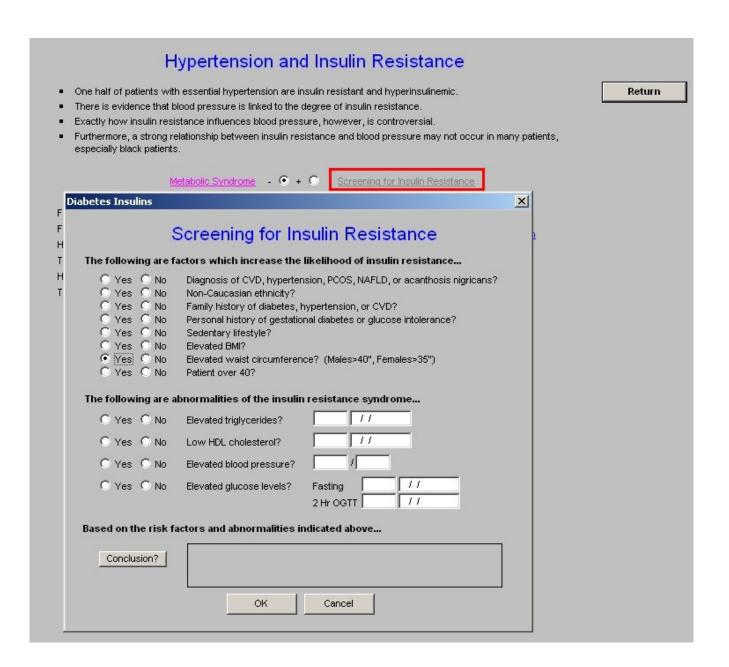
There is evider Exactly how in	ce that blood pressure is I sulin resistance influences strong relationship betwee		sistance.	Return tients,
Fasting Glucose Fasting Insulin HOMA-IR	11	Information Insulin Resistan Teenage Insulin		1
Triglycercides HDL Trig/HDL Ratio	11	Hypertension, In Hypertension ar	nsulin Resistance sulin and IGT	
Triglycercides HDL	II	Components of Hypertension, In Hypertension ar	nsulin Resistance sulin and IGT id Lipoic Acid	

Beneath these principles is a link to the **Metabolic Syndrome Assessment Template** which results will populate the check boxes which display whether or not the patient has the metabolic syndrome.





Next is a link to the **Screening for Insulin Resistance** template which is part of the **Less Initiative**.

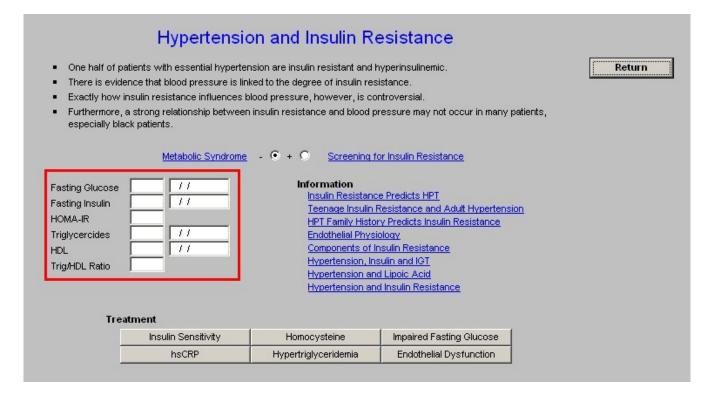


The elements of this template are automatically populated from the EMR data base. When the button entitled Conclusion? is depressed a summary of whether the patient is at risk for insulin resistance, and consequently the potential blood pressure problems associated with it, will appear.

The following laboratory values are then displayed:

- Fasting Glucose
- Fasting Insulin
- **HOMA-IR** Homeostasis Model Assessment of Insulin Resistance. A value above 2 indicates insulin resistance.
- Triglycerides

- HDL
- Trig/HDL Ratio a value above 2 for this ratio indicates Insulin Resistance.



Beside the laboratory values are buttons which launch the following documents:

- Insulin Resistance Predicts HPT
- Teenage Insulin Resistance and Adult Hypertension
- HPT Family Hisory Predicts Insulin Resistance
- Endotehlial Physiology
- Components of Insulin Resistance
- Hypertension, Insulin and IGT
- Hypertension and Lipoic Acid
- Hypertension and Insulin Resistance

 One half of pati 	ents with essential hyperte	ension are insulin resistant and l	nyperinsulinemic.	Return
 There is eviden 	ce that blood pressure is lin	nked to the degree of insulin res	sistance.	<u> </u>
 Exactly how ins 	ulin resistance influences	blood pressure, however, is co	ntroversial.	
 Furthermore, a especially black 			ressure may not occur in many patie or Insulin Resistance	nts,
Fasting Glucose Fasting Insulin HOMA-IR Friglycercides		HPT Family Histo Endothelial Physi Components of I	Resistance and Adult Hypertension ry Predicts Insulin Resistance	
HDL Frig/HDL Ratio		Hypertension, In: Hypertension an Hypertension an		
IDL [Hypertension an	d Lipoic Acid	
HDL ∫ rig/HDL Ratio ∫		Hypertension an	d Lipoic Acid	

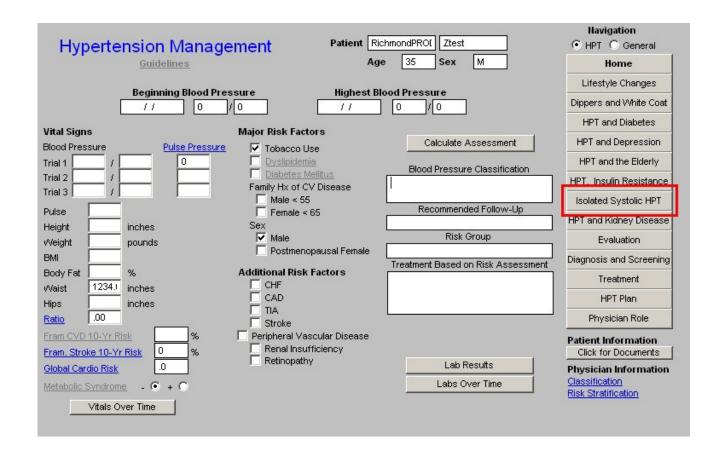
At the bottom of this template are six buttons which launch **Treatment Recommendations** for the following, (for details on these functions, which are self explanatory. see the tutorial for the **Metabolic Syndrome Treatment Recommendations**.):

- Insulin Sensitivity
- hsCRP
- Homocysteine
- Hypertriglyceridemia
- Impaired Fasting Glucose
- Endothelial Dysfunction

	Hypertension	on and Insulin Re	esistance	
There is evident Exactly how in:	ice that blood pressure is lin sulin resistance influences b strong relationship betweer		istance.	patients,
Fasting Glucose Fasting Insulin HOMA-IR Triglycercides HDL Trig/HDL Ratio		Information Insulin Resistanc Teenage Insulin F HPT Family Histor Endothelial Physic Components of In Hypertension, Ins	e Predicts HPT Resistance and Adult Hypertens ry Predicts Insulin Resistance ology nsulin Resistance sulin and IGT	<u>sion</u>
Treat	ment	,	,	7
	Insulin Sensitivity hsCRP	Homocysteine Hypertriglyceridemia	Impaired Fasting Glucose Endothelial Dysfunction	

Isolated Systolic Hypertension Template

Perhaps the most under treated and damaging form of hypertension is Isolated Systolic Hypertension (ISH), particularly in the elderly. The key to ISH treatment is its recognition and diagnosis and the understanding of the increased cardiovascular and cerebrovascular risk associated with it.



<u>x</u>
lated Systolic Hypertension
pressure (> 140 mm Hg) with normal diastolic blood pressure (< 90 mm Hg). of those over 60 years of age regardless of race. The majority of these have ISH. /e societies whose members maintain lean body mass and are physically active.
Evaluation
Exclude diabetes, hypokalemia, and renal disease with blood chemistry Urinalysis to exclude proteinuria EKG to evaluate for MI, arrhythmias, or LVH Echocardiogram for LVH, particularly if signs or symptoms of CHF Treatment Lifestyle Weight Reduction Exercise Decreased Salt Intake Decreased Alcohol Intake Drugs Low-dose, long-acting diuretic, such as 12.5 mg daily of hydrochlorothiazide Low-dose, long-acting, dihydropyridine calcium channel blocker If patient also has angina, a beta-blocker is appropriate If patient also has benign prostatic hypertrophy, an alpha blocker is a reasonable choice

This template contains the following material:

- **Definition** Elevated systolic blood pressure (> 140 mm Hg) with normal diastolic blood pressure (< 90 mm Hg).
- **Incidence** Hypertension occurs in 50% of those over 60 years of age regardless of race. The majority of these have ISH.
- **Prevention** No hypertension in primitive societies whose members maintain lean body mass and are physically active.

• Contributing Causes

- 1. Increased body fat
- 2. Sedentary lifestyle
- 3. Increased sodium intake
- 4. Loss of elasticity of aorta and peripheral vessels compensatory for maintaining oxygen delivery

• Associated Features

- 1. Low plasma volume consistent with general body water loss
- 2. Renal excretion of salt decreased making elder more sensitive to salt

Risk

- 1. Three-fold increase of stroke
- 2. Increased risk of overall mortality

- 3. Increased risk of cardiovascular mortality
- 4. Increased risk of congestive heart failure

Evaluation

- 1. Exclude diabetes, hypokalemia, and renal disease with blood chemistry
- 2. Urinalysis to exclude proteinuria
- 3. EKG to evaluate for MI, arrhythmias, or LVH
- 4. Echocardiogram for LVH, particularly if signs or symptoms of CHF

• Treatment

1. Lifestyle

- 1. Weight Reduction
- 2. Exercise
- 3. Decreased Salt Intake
- 4. Decreased Alcohol Intake

2. Drugs

- 1. Low-dose, long-acting diuretic, such as 12.5 mg daily of hydrochlorothiazide
- 2. Low-dose, long-acting, dihydropyridine calcium channel blocker
- 3. If patient also has angina, a beta-blocker is appropriate
- 4. If patient also has **renal insufficiency** or cardiomyopathy, an ACE is a good choice
- 5. If patient also has **benign prostatic hypertrophy**, an alpha blocker is a reasonable choice

Hypertension and Kidney Disease Template

Hyperte	nsion Manage	ement Patient R	ichmondPROI Ztest	Navigation • HPT • General
	Guidelines	Ag	je 35 Sex M	Home
	Danimaina Dia ad Dana	Uinkand D	n	Lifestyle Changes
Γ	Beginning Blood Pres	o III	Nood Pressure	Dippers and White Coat
Vital Cinna		Maior Diale Forton		HPT and Diabetes
Vital Signs Blood Pressure	Pulse Pressure	Major Risk Factors ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 /	0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /		Diabetes Mellitus Family Hx of CV Disease	Biood Pressure Classification	HPT, Insulin Resistance
Trial 3 /		Male < 55	L	Isolated Systolic HPT
Pulse		☐ Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Height Veight	inches pounds	✓ Male	Risk Group	Evaluation
ВМІ		Postmenopausal Female	Treatment Based on Risk Assessment	Diagnosis and Screening
Body Fat	%	Additional Risk Factors	Treatment based on Risk Assessment	Treatment
Waist 1234.	inches	☐ CHF ☐ CAD		HPT Plan
Hips .00	inches	□ TIA		Physician Role
Fram CVD 10-Yr Ri	sk 5%	Stroke Peripheral Vascular Disease		
Fram. Stroke 10-Yr		Renal Insufficiency		Patient Information Click for Documents
Global Cardio Risk	.0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome	<u> </u>		Labs Over Time	Classification Risk Stratification
Vitals Ov	ver Time			

The association between hypertension in both primary and secondary conditions is well defined. The unique aspect of Kidney disease and hypertension is that hypertension can both be a cause of and a result of kidney disease.

This is a duplicate of the **Hypertension template** in the **Renal Disease Suite of Templates**. The template begins with the following concepts:

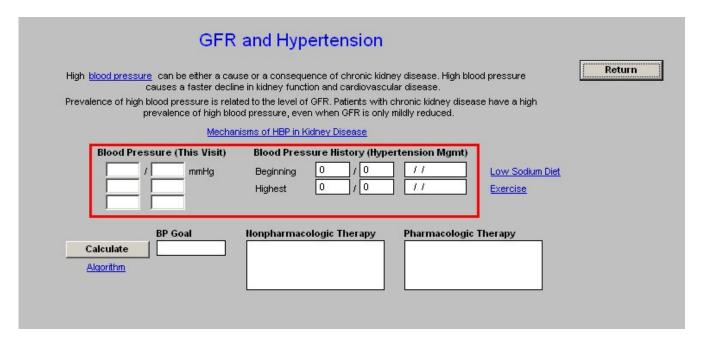
- High blood pressure can be either a cause or a consequence of chronic kidney disease.
- High blood pressure causes a faster decline in kidney disease and cardiovascular disease.
- Prevalence of high blood pressure is related to the level of GFR.
- Patients with chronic kidney disease have a high prevalence of high blood pressure even when GFR is only mildly reduced.

Mechanism of HBP in Kidney Disease - this launches a document which addresses this issue.

Beneath this link, there are boxes which document:

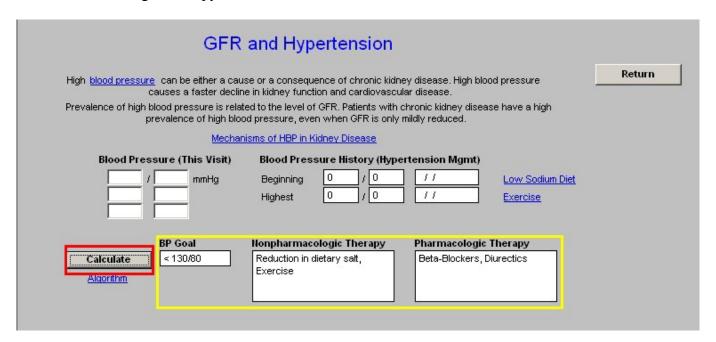
- Blood Pressure (this visit)
- Blood Pressure History (Hypertension Mgmt)

- 1. Beginning Blood Pressure
- 2. Highest Blood Pressure

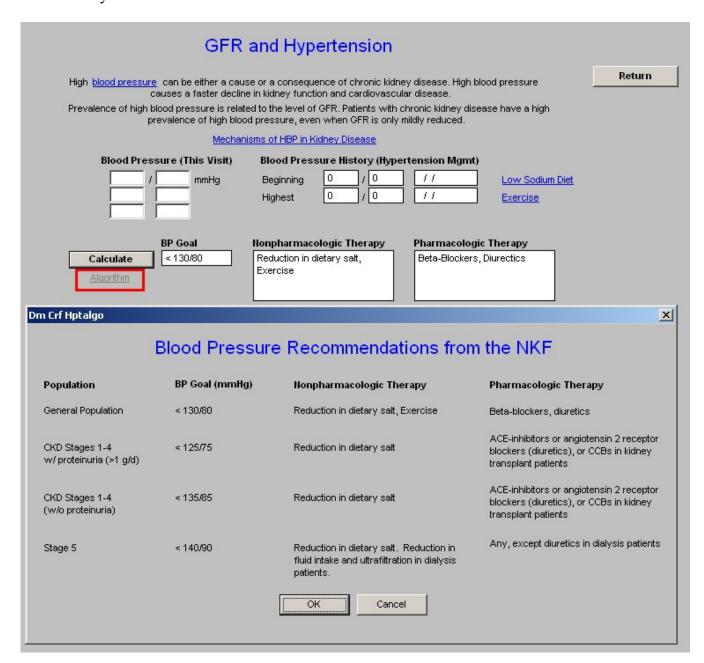


There is then a button entitled "Calculate" when depressed it fills in the following boxes:

- BP Goal
- Nonpharmacologic Therapy
- Pharmacologic Therapy



Algorithm - this button launches a pop-up which lists the **National Kidney Foundation Blood Pressure Recommendations** on the basis of which the above boxes are completed automatically.



There are two links related to lifestyle changes which improve hypertension particularly in relationship to kidney disease; they are:

- Low Sodium Diet
- Exercise

Evaluation Template

Hypertension Ma	nagement Patient	RichmondPROI Ztest	Navigation • HPT • General
Guidelines	A	ge 35 Sex M	Home
Danimaia a Di	Dishest	Blood Pressure	Lifestyle Changes
/ /	ood Pressure Highest	0 //0	Dippers and White Coat
Vital Ciana	Maior Dials Fordana	-	HPT and Diabetes
Vital Signs Blood Pressure Pulse P	Major Risk Factors ressure ▼ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / 0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 /	Diabetes Mellitus Family Hx of CV Disease	Diood Pressure Classification	HPT, Insulin Resistance
Trial 3 /	☐ Male < 55		Isolated Systolic HPT
Pulse Inches	☐ Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Height inches Weight pounds	✓ Male	Risk Group	Evaluation
ВМІ	Postmenopausal Female	Treatment Based on Risk Assessment	Diagnosis and Screening
Body Fat %	Additional Risk Factors	Treatment based on Nisk Assessment	Treatment
Waist 1234.1 inches	☐ CHF ☐ CAD		HPT Plan
Hips inches	□ TIA		Physician Role
	Stroke Stroke Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk 0	% Renal Insufficiency	,	Click for Documents
Global Cardio Risk .0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome - € + €		Labs Over Time	Classification Risk Stratification
Vitals Over Time			

The Evaluation Template has the following content:

Displays the two major types of hypertension:

- Primary Hypertension has no clearly defined etiology
- Secondary Hypertension often has no symptoms

Just below this information is a button entitled "Primary Pulmonary Hypertension".

Primary Pulmonary Hypertension is not a part of systemic hypertension but is presented here because of the close association of the name.

Primary Hypertension			Return
	ry hypertension has no clearly defined etiolog	gy.	
Secondary Hypertension			Information
Most co	ommonly hypertension has no symptor	ms!!!	Arterial Hypertension Renovascular Hypertension
	Primary Pulmonary Hypertension		Total account 11 portorior
Signs & Symptoms	Causes		
☐ Headache	The state of the s	Drugs & Toxins	
Dizziness	Primary Aldosteronism Cushing's Syndrome	☐ Alcohol ☐ Cocaine	
☐ Fatique	Pheochromocytoma	Cyclosporin	
Palpatations	Congenital Adrenal Hyperplasia	☐ Erythropoietin	
	Endocrine	Adrenergic Medications	
Malaise		Renal	
Nosebleeds	☐ Hypercalcemia	Renal Parenchymal Disease	
☐ Weight Loss	☐ Hyperparathyroidism	Polycystic Kidney Disease	
Swelling in the Legs	Hyperthyroidism	Urinary Tract Obstruction	
☐ Weakness	Hypothyroidism	Renin-producing Tumor	
	Oral Contraceptives	Liddle Syndrome	
Frequent Urination	Pregnancy-induced Hypertensio	The second secon	
☐ Increased Thirst		Vascular	
Enlarged, Round, or "Moon" Face	Brain Tumor	Coarctation of Aorta	
Excess Body and Facial Hair	Bulbar Poliomyelitis Intracranial Hypertension	☐ Vasculitis ☐ Collagen Vascular Disease	

- **Definition** -- Pulmonary hypertension is when the blood pressure in the arteries of the lungs is abnormally high. "Primary" means that there are no other diseases of the heart or lungs causing the high blood pressure.
- Alternative Names -- Pulmonary arterial hypertension; Sporadic primary pulmonary hypertension; Familial primary pulmonary hypertension
- Causes And Risk -- The cause of primary pulmonary hypertension is unknown. Some cases are caused by a genetic defect.
 - 1. The small arteries of the lung narrow throughout the lungs. Pulmonary hypertension is the result of greater resistance to blood flow. As a result of the increased workload caused by this resistance, the right side of the heart becomes enlarged.
 - 2. This disease is rare. It affects more women than men.
 - 3. Pulmonary hypertension is the narrowing of the pulmonary arterioles within the lung. The narrowing of the arteries creates resistance and an increased work load for the heart. The heart becomes enlarged from pumping blood against the resistance.
 - 4. The goal of treatment is control of the symptoms, although the disease usually develops into congestive heart failure.

Symptoms

- 1. Shortness of breath during activity
- 2. Chest pain

- 3. Weakness
- 4. Fatigue
- 5. Fainting
- 6. Lightheadedness during exercise
- 7. Dizziness
- 8. Leg Swelling

Signs and Tests

- 1. A physical examination shows enlargement of the veins in the neck, normal lungs, a heart murmur, enlargement of the liver, and swelling due to fluid retention in the tissues.
- 2. Tests
 - a. ECG
 - b. Chest X-Ray
 - c. Echocardiogram
 - d. Pulmonary Function Tests
 - e. Nuclear Lung Scan
 - f. CT Scan of the Chest
 - g. Cardiac Catheterization
 - h. Pulmonary Arteriogram

• Call Your Doctor If...

- 1. Chest pain develops
- 2. You develop shortness of breath when active
- 3. Shortness of breath worsens
- 4. You experience other symptoms

Treatment

- 1. There is no known cure. The goal of treatment is to control symptoms. Some people respond to treatment with oral medications, such as calcium channel blockers and diuretics. Others may need drugs like intravenous epoprostenol.
- 2. Recently, a new oral drug that blocks abnormal constriction of blood vessels has become available and shows promise in treating pulmonary hypertension.
- 3. As the disease progresses, oxygen may be necessary. If all else fails, suitable candidates may be helped by lung or heart-lung transplantation.
- 4. Most patients with primary pulmonary hypertension are treated at centers which specialize in the care of these patients.

n Hpt Pph		x
Definition Pulmonary hypertension is when the blood of the heart or lungs causing the high blood	· · · · · · · · · · · · · · · · · · ·	ertension ally high. "Primary" means that there are no other diseases
Causes And Risk The cause of primary pulmonary hyperter The small arteries of the lung narrow thro the increased workload caused by this re This disease is rare, it affects more wome Pulmonary hypertension is the narrowing	sistance, the right side of the heart becomes en en than men. of the pulmonary arterioles within the lung. The	genetic defect. result of greater resistance to blood flow. As a result of larged. Eventually, progressive heart failure may develop. narrowing of the arteries creates resistance and an
The goal of treatment is control of the syn Symptoms Shortness of breath during activity Chest pain Weakness Fatigue Fainting Lightheadedness during exercise	art becomes enlarged from pumping blood again proms, although the disease usually develops in Signs and Tests A physical examination shows enlargement of enlargement of the liver, and swelling due to ECG Chest X-Ray Echocardiogram Pulmonary Function Tests	of the veins in the neck, normal lungs, a heart murmur,
channel blockers and diuretics. Others may Recently, a new oral drug that blocks abnor As the disease progresses, oxygen may be	need drugs like intravenous epoprostenol. mal constriction of blood vessels has become a	You develop shortness of breath when active Shortness of breath worsens d to treatment with oral medications, such as calcium vailable and shows promise in treating pulmonary hypertension. may be helped by lung or heart-lung transplantation. in the care of these patients.

There are two series of checkboxes in the last section on the "Hypertension Evaluation" screen entitled "Signs & Symptoms" and "Causes". While most patients with hypertension have no signs or symptoms, these are sometimes associated with secondary hypertension:

Signs & Symptoms

- Headache
- Dizziness
- Fatigue
- Palpitations
- Malaise
- Nosebleeds
- Weight Loss
- Swelling in the legs
- Weakness
- Frequent Urination
- Increased Thirst
- Enlarged, Round, or "Moon" Face
- Excess Body and Facial Hair

H	Hypertension E∨aluation	ı	
Primary Hypertension			Return
Prima	ry hypertension has no clearly defined etiolog	gy.	Kettarii
Secondary Hypertension			Information
	ommonly hypertension has no symptor	ns!!!	Arterial Hypertension Renovascular Hypertensio
Signe & Cumptome	Primary Pulmonary Hypertension		
Signs & Symptoms Headache Dizziness Fatigue Palpatations Malaise Nosebleeds Weight Loss Swelling in the Legs Frequent Urination	Primary Aldosteronism Cushing's Syndrome Pheochromocytoma Congenital Adrenal Hyperplasia Endocrine Acromegaly Hypercalcemia Hyperparathyroidism Hyperthyroidism Hypothyroidism Oral Contraceptives Pregnancy-induced Hypertension		
☐ Increased Thirst ☐ Enlarged, Round, or "Moon" Face ☐ Excess Body and Facial Hair	Neurologic Brain Tumor Bulbar Poliomyelitis Intracranial Hypertension	Vascular ☐ Coarctation of Aorta ☐ Vasculitis ☐ Collagen Vascular Disease	

Causes

1. Adrenal

- a. Primary Aldosteronism
- b. Cushing's Syndrome
- c. Pheochromocytoma
- d. Congenital Adrenal Hyperplasia

2. Endocrine

- a. Acromegaly
- b. Hypercalcemia
- c. Hyperparathyroidism
- d. Hypothyroidism
- e. Oral Contraceptives
- f. Pregnancy-induced hypertension

3. Neurological

- a. Brain Tumor
- b. Bulbar Poliomyelitis
- c. Intracranial Hypertension

4. Drugs and Toxins

- a. Alcohol
- b. Cocaine

- c. Cyclosporin
- d. Erythropoietin
- e. Adrenergic Medications

5. Renal

- a. Renal Parenchymal Disease
- b. Polycystic Kidney Disease
- c. Urinary Tract Obstruction
- d. Renin-producing Tumor
- e. Liddle Syndrome
- f. Renovascular Hypertension

6. Vascular

- a. Coarctation of Aorta
- b. Vasculitis
- c. Collagen Vascular Disease

Primary Hypertension Primary hypertension has no clearly defined etiology. Secondary Hypertension Most commonly hypertension has no symptoms::: Primary Pulmonary Hypertension Signs & Symptoms Adrenal Primary Aldosteronism Primary Aldosteronism Primary Aldosteronism Primary Aldosteronism Primary Aldosteronism Pheochromocytoma Cocaine Pheochromocytoma Phypertension Malaise Nosebleeds Nosebleeds Nosebleeds Hypertension Phypertension Phypertension Phypertension Phypertension Phypertension Phypertension Phypertension Phypertension Pregnancy-induced Hypertension Pregnancy-induced Pregn	Н	ypertension E∨aluation	
Primary hypertension has no clearly defined etiology. Secondary Hypertension Most commonly hypertension has no symptoms!! Primary Pulmonary Hypertension Primary Aldosteronism	Primary Hypertension		Paturn
Most commonly hypertension has no symptoms:!! Primary Pulmonary Hypertension Primary Pulmonary Hypertension	Primary	hypertension has no clearly defined etiology.	Ketuiii
Signs & Symptoms Headache	Secondary Hypertension		Information
Headache	Most co		
Primary Aldosteronism	Signs & Symptoms		
Dizziness Cushing's Syndrome Cyclosporin Palpatations Malaise Nosebleeds Weight Loss Weight Loss Hyperparathyroidism Hyperthyroidism Hyperthyroidism Hyperthyroidism Hyperthyroidism Hyperthyroidism Prequent Urination Frequent Urination Increased Thirst Enlarged, Round, or "Moon" Face Congenital Adrenal Hyperplasia Erythropoietin Adrenergic Medications Hypercalcemia Renal Parenchymal Disease Polycystic Kidney Disease Urinary Tract Obstruction Renin-producing Tumor Liddle Syndrome Renovascular Hypertension Neurologic Vascular Brain Tumor Coarctation of Aorta Bulbar Poliomyelitis Vasculitis	☐ Headache		
Fatigue Palpatations Congenital Adrenal Hyperplasia Endocrine Adrenergic Medications Adrenergic Medications Malaise Nosebleeds Hypercalcemia Renal Parenchymal Disease Weight Loss Hyperparathyroidism Polycystic Kidney Disease Hyperthyroidism Urinary Tract Obstruction Hyperthyroidism Renin-producing Tumor Weakness Prequent Urination Pregnancy-induced Hypertension Renovascular Hypertension Neurologic Vascular Brain Tumor Coarctation of Aorta Bulbar Poliomyelitis Vasculitis	☐ Dizziness		
Malaise	☐ Fatigue		
Malaise Nosebleeds Hypercalcenia Renal Parenchymal Disease Hyperparathyroidism Polycystic Kidney Disease Hyperthyroidism Urinary Tract Obstruction Hypothyroidism Renin-producing Turnor Weakness Oral Contraceptives Frequent Urination Pregnancy-induced Hypertension Increased Thirst Neurologic Vascular Enlarged, Round, or "Moon" Face Bulbar Poliomyelitis Vasculitis	☐ Palpatations		
Nosebleeds	☐ Melaise	Endocrine Adrenergic Medications	
Weight Loss		The state of the s	
Swelling in the Legs			
Swelling in the Legs	Weight Loss	_ ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	
Weakness Oral Contraceptives Liddle Syndrome Frequent Urination Pregnancy-induced Hypertension Increased Thirst Neurologic Vascular Enlarged, Round, or "Moon" Face Bulbar Poliomyelitis Vasculitis	Swelling in the Legs		
☐ Increased Thirst Neurologic Vascular ☐ Enlarged, Round, or "Moon" Face ☐ Brain Tumor ☐ Coarctation of Aorta ☐ Bulbar Poliomyelitis ☐ Vasculitis	☐ Weakness		
☐ Enlarged, Round, or "Moon" Face ☐ Brain Tumor ☐ Coarctation of Aorta ☐ Bulbar Poliomyelitis ☐ Vasculitis	Frequent Urination	☐ Pregnancy-induced Hypertension ☐ Renovascular Hypertension	
Enlarged, Round, or "Moon" Face Bulbar Poliomyelitis Vasculitis	☐ Increased Thirst	Neurologic Vascular	
Bulbar Poliomyelitis J Vasculitis		☐ Brain Tumor ☐ Coarctation of Aorta	
Intracranial Hypertension Collagen Vascular Disease			
	Excess Body and Facial Hair	J_ Intracranial Hypertension Collagen Vascular Disease	

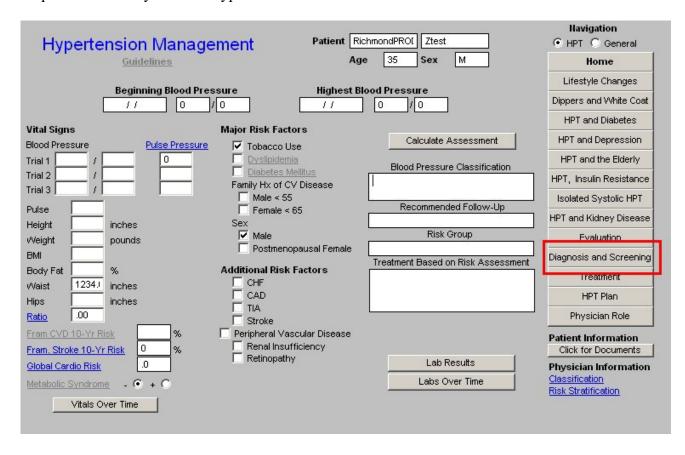
NOTE: All Causes listed in blue are linked to a detailed document describing that cause At the right hand side of the **Evaluation** template is a **Return** button that takes you back to the Master Hypertension template and

Beneath the **Return** button are two Provider education documents

- Arterial Hypertension this button launches a document entitled "Arterial Hypertension: Etiology and Pathogenesis."
- **Renovascular Hypertension** this button launches a document entitled the same as the button.

Diagnosis and Screening Template

This template provides the ability for the provider to establish a plan for evaluating secondary forms of hypertension. In addition to identifying the most common causes of secondary hypertension, this template provides a tool to suggest the proper evaluation for patients with a suspected secondary cause of hypertension.



When the "Diagnosis and Screening" button is clicked, it will display an interactive function entitled "Screening and Diagnostic Studies for Secondary Hypertension." There are two sections on this part of the template, the first is "Suspicions" and the second is "Actions."

Suspicions

- Coaractation of the Aorta
- Cusingh's Syndrome
- Pheochromocytoma
- Renovascular Dsiease

- Thyroid Disease
- Hyperparathyroidism
- Renal Parenchymal Disease

There are check boxes in front of each of these causes of secondary hypertension. If you suspicion one of these conditions to be present, check the box in front of the condition and a list of actions recommended for the evaluation of that condition will be automatically selected.

Suspicions	Actions	Return
Coarctation of Aorta Cushing's Syndrome Primary Aldosteronism Pheochromocytoma Renovascular Disease Thyroid Disease Hyperparathyroidism Renal Parenchymal Disease	24-hour Unrinary Free Cortisol 24-hour Urine Protein and Creatinine Levels 2D Echo, Aorta Abdominal CT Aldosterone Excretion Rate During Salt Loading Adrenal CT Angiography Aortogram Calcium and Phosphorus Levels Captopril Renography Chest Film Clonidine Suppression Test Computed Tomography Dexamethasone Supression Test Iodine 131 Metaiodobenzylguanidine Scan	I lothalamate Glomerular Filtration Rate MRI, Aorta MRI, Renal Artery Plasma Aldesterone:Renin Ratio Plasma Catecholamines Renal Biopsy Renal Duplex Sonography Renal Ultrasound Renal Vein Renin Ratio Serum Calcitonin Level Serum Parathyroid Hormone Level Thyrotropin Urine Catecholamines
Primary Pulmonary Hypertension	Cardiac Catheterization Chest CT ECG Echocardiogram	☐ Nuclear Lung Scan ☐ Pulmonary Arteriogram ☐ Pulmonary Function Testing

• **Primary Pulmonary Hypertension** - this is not pathophysiologically a part of hypertension but it is included here because it does result in hypertension in the pulmonary vasculature and it is an increasingly diagnosed condition.

Suspicions	Actions	Retur
Coarctation of Aorta Cushing's Syndrome Primary Aldosteronism Pheochromocytoma Renovascular Disease Thyroid Disease Hyperparathyroidism Renal Parenchymal Disease		☐ Iothalamate Glomerular Filtration Rate ☐ MRI, Aorta ☐ MRI, Renal Artery ☐ Plasma Aldesterone:Renin Ratio ☐ Plasma Catecholamines ☐ Renal Biopsy ☐ Renal Duplex Sonography ☐ Renal Uthrasound ☐ Renal Vein Renin Ratio ☐ Serum Calcitonin Level ☐ Serum Parathyroid Hormone Level ☐ Serum Thyroid Hormone Level ☐ Thyrotropin ☐ Urine Catecholamines
✓ Primary Pulmonary Hypertens	☐ ☐ Cardiac Catheterization ☐ Chest CT ☐ ECG ☐ Echocardiogram	 ✓ Nuclear Lung Scan ✓ Pulmonary Arteriogram ✓ Pulmonary Function Testing

Actions

- 24-hour urinary Free Cortisol
- 24-hour Urine Protein and Creatinine Levels
- 2D Echo, Aorta
- Abdominal CT
- Aldosterone Excretion Rate during Salt loading
- Adrenal CT
- Angiography
- Aortogram
- Calcium and Phosphorous Levels
- Captopril Renography
- Chest Film
- Clonidine Suppression Test
- Computed Tomography
- Dexamethasone Suppression Test
- Iodine 131 Metaiodobenzylguandine Scan
- Iothalamate Glomerular Filtration Rate
- MRI, Aorta
- MRI, Renal Artery
- Plasma Catecholamines
- Renal Biopsy
- Renal Duplex Sonography
- Renal Ultrasound
- Renal Vein Renin Ratio

- Serum Calcitonin Level
- Serum Parathyroid Hormone Level
- Thyrodropin
- Urine Catecholamines
- Cardiac Cauterization
- Chest CT
- ECG
- Echocardiogram
- Nuclear Lung Scan
- Pulmonary Arteriogram
- Pulmonary Function Testing

uspicions	Actions	Kei
Coarctation of Aorta Cushing's Syndrome Primary Aldosteronism Pheochromocytoma Renovascular Disease Thyroid Disease Hyperparathyroidism Renal Parenchymal Disease	24-hour Unrinary Free Cortisol 24-hour Urine Protein and Creatinine Levels 2D Echo, Aorta Abdominal CT Aldosterone Excretion Rate During Salt Loading Adrenal CT Angiography Aortogram Calcium and Phosphorus Levels Captopril Renography Chest Film Clonidine Suppression Test Computed Tomography Dexamethasone Supression Test	□ lothalamate Glomerular Filtration Rate □ MRI, Aorta □ MRI, Renal Artery □ Plasma Aldesterone:Renin Ratio □ Plasma Catecholamines □ Renal Biopsy □ Renal Duplex Sonography □ Renal Ultrasound □ Renal Vein Renin Ratio □ Serum Calcitonin Level □ Serum Parathyroid Hormone Level □ Serum Thyroid Hormone Level □ Thyrotropin □ Urine Catecholamines
Primary Pulmonary Hypertension	n Cardiac Catheterization Chest CT ECG Echocardiogram	☐ Nuclear Lung Scan ☐ Pulmonary Arteriogram ☐ Pulmonary Function Testing

For instance, if you suspect a **Pheochromocytoma**, check the box beside that name and the following will automatically have the boxes beside them checked

- Abdominal CT
- Clonindine Suppression Test
- Iodine 131 Metaiodobenzyguanidine Scan
- Plasma Catecholamines
- Urine Catecholamines

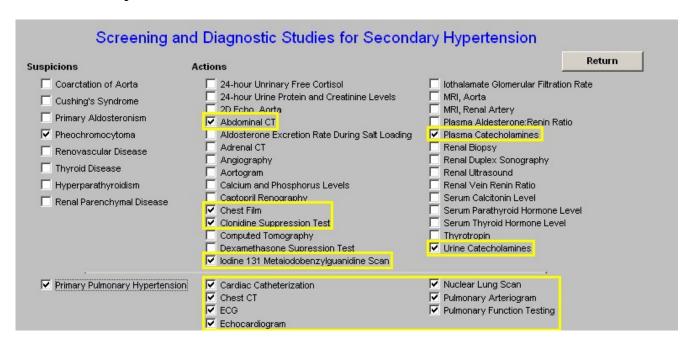
Suspicions	Actions	_	Return
Coarctation of Aorta Cushing's Syndrome Primary Aldosteronism Pheochromocytoma Renovascular Disease Thyroid Disease Hyperparathyroidism Renal Parenchymal Disease	□ 24-hour Unrinary Free Cortisol □ 24-hour Urine Protein and Creatinine Levels □ 2D Echo, Aorta □ Abdominal CT □ Aldosterone Excretion Rate During Salt Loading □ Adrenal CT □ Angiography □ Aortogram □ Calcium and Phosphorus Levels □ Captopril Renography ☑ Chest Film □ Clonidine Suppression Test □ Computed Tomography □ Dexamethasone Supression Test □ Iodine 131 Metaiodobenzylguanidine Scan	lothalamate Glomerular Filtration MRI, Aorta MRI, Renal Artery Plasma Aldesterone:Renin Ratio Plasma Catecholamines Renal Biopsy Renal Duplex Sonography Renal Utrasound Renal Vein Renin Ratio Serum Calcitonin Level Serum Parathyroid Hormone Level Serum Thyroid Hormone Level Thyrotropin Urine Catecholamines	
Primary Pulmonary Hypertension	 ✓ Cardiac Catheterization ✓ Chest CT ✓ ECG ✓ Echocardiogram 	✓ Nuclear Lung Scan ✓ Pulmonary Arteriogram ✓ Pulmonary Function Testing	

If you think the patient has **Primary Pulmonary Hypertension**, check the box beside that name and the following will automatically have the boxes beside them checked:

- Chest Film
- Chest CT
- ECG
- Echocardiogram
- Nuclear Lung Scan
- Pulmonary Arteriogram
- Pulmonary Function Testing

Screening ar	d Diagnostic Studies for Second Actions	ary Hypertension Return
Coarctation of Aorta Cushing's Syndrome Primary Aldosteronism Pheochromocytoma Renovascular Disease Thyroid Disease Hyperparathyroidism Renal Parenchymal Disease	24-hour Unrinary Free Cortisol 24-hour Urine Protein and Creatinine Levels 2D Echo. Aorta ✓ Abdominal CT Aldosterone Excretion Rate During Salt Loading Adrenal CT Angiography Aortogram Calcium and Phosphorus Levels Captopril Renography Chest Film ✓ Clonidine Suppression Test Computed Tomography Dexamethasone Suppression Test I odine 131 Metaiodobenzylguanidine Scan	Iothalamate Glomerular Filtration Rate
Primary Pulmonary Hypertension	Cardiac Catheterization Chest CT ECG Echocardiogram	Nuclear Lung Scan Pulmonary Arteriogram Pulmonary Function Testing

Treatment Template



This template pulls together many of the previously discussed principles of hypertension management.

The **Hypertension Treatment Template** addresses the following issues:

Hypertension Mana	igement Patient Ric	chmondPROI Ztest	Navigation ⊙ HPT ○ General
<u>Guidelines</u>	Ag	e 35 Sex M	Home
Beginning Blood	Dragoura Highoot Pl	lood Pressure	Lifestyle Changes
// 0	/ 0 / /	0 /0	Dippers and White Coat
Vital Signa	Major Diek Fastere		HPT and Diabetes
Vital Signs Blood Pressure Pulse Press	Major Risk Factors ure V Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / O	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2 / /	Diabetes Mellitus Family Hx of CV Disease	Diodu Pressure Classification	HPT, Insulin Resistance
Trial 3 /	☐ Male < 55	Barran and a Sallan Un	Isolated Systolic HPT
Pulse Inches	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
vVeight pounds	✓ Male	Risk Group	Evaluation
вмі	Postmenopausal Female	Treatment Based on Risk Assessment	Diagnosis and Screening
Body Fat %	Additional Risk Factors	Treatment based of Misk Assessment	Treatment
Waist 1234.1 inches	☐ CAD		HPT Plan
Ratio .00	☐ TIA ☐ Stroke		Physician Role
Fram CVD 10-Yr Risk %	Peripheral Vascular Disease		Patient Information
Fram. Stroke 10-Yr Risk 0 %	Renal Insufficiency Retinopathy		Click for Documents
Global Cardio Risk .0	reunopatry	Lab Results	Physician Information
Metabolic Syndrome - © + C		Labs Over Time	Classification Risk Stratification
Vitals Over Time			

- You need one drug for every 10 mm Hg of systolic blood pressure reduction
- The vast majority of patients will need two, three or possibly four medications.
- Drugs that confer an additional cardiovascular benefit beyond direct blood pressure effects are particularly attractive options.
- Data from seven landmark trials found that between 2.5 and 4.0 antihypertensive agents were required to bring blood pressure to target.

Five steps to Hypertensive control

- 1. Begin and/or continue lifestyle modifications
- 2. Goal Blood pressure

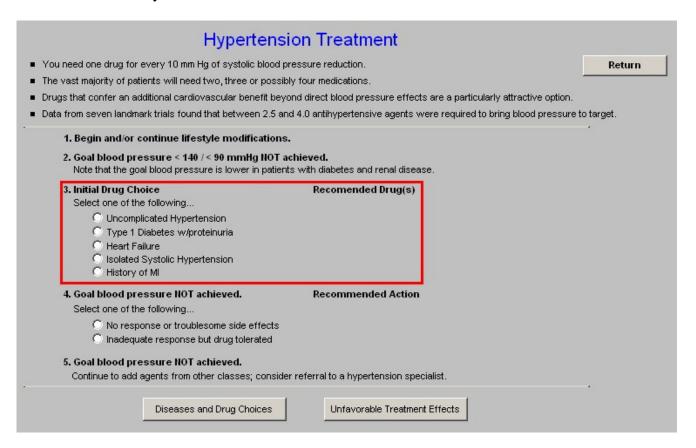
Note that the goal blood pressure is lower in a patient with diabetes and renal disease: 110/70 or lower.

3. Initial Drug Choice

Select one of the following (when one of the below is selected by checking in the box next to it), Recommended Drug(s) will appear on the template.

- a. Uncomplicated Hypertension
- b. Type 1 Diabetes w/proteinuria
- c. Heart Failure
- d. Isolated Systolic hypertension

e. History of MI



4. Goal Blood pressure NOT achieved

Select one of the following

No response or troublesome side effects Inadequate response but drug tolerated

e vast majority of patients will need two, three or	plood pressure reduction. Return possibly four medications.
	fit beyond direct blood pressure effects are a particularly attractive option.
ta from seven landmark trials found that between	2.5 and 4.0 antihypertensive agents were required to bring blood pressure to target.
1. Begin and/or continue lifestyle modific	ations.
2. Goal blood pressure < 140 / < 90 mmHg Note that the goal blood pressure is lower in	
3. Initial Drug Choice Select one of the following C Uncomplicated Hypertension Type 1 Diabetes w/proteinuria Heart Failure Isolated Systolic Hypertension History of MI	Recomended Drug(s)
4. Goal blood pressure HOT achieved. Select one of the following No response or troublesome side ef Inadequate response but drug tolers	NO. COLO
5. Goal blood pressure NOT achieved.	consider referral to a hypertension specialist.

1. Goal Blood Pressure NOT achieved

Continue to add agents from other classes, consider referral to a hypertension specialist.

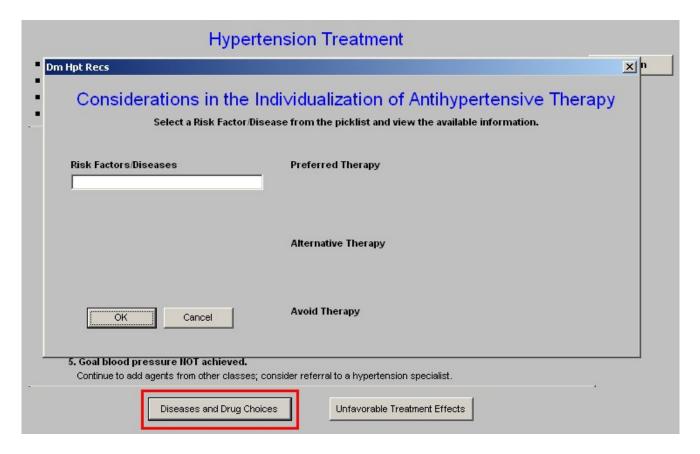
There are two pop-ups on this template:

• **Diseases and Drug choices** - there is a box which when accessed displays the following pick list. When one of the disease states is selected, the second box displays the optimal medicine for treating hypertension in this circumstance.

Risk/Factors Diseases

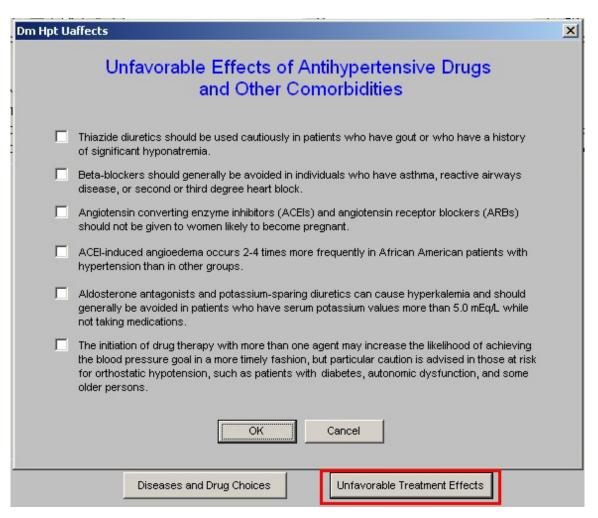
- 1. African-American
- 2. Angina
- 3. Atrial Fibrilation
- 4. Caucasian
- 5. CHF
- 6. Chronic Renal Failure
- 7. Cough Caused by ACE Inhibitors
- 8. Diabetes w/Nephropathy
- 9. Diabetes w/o Nephropathy
- 10. Diabetes w/o Nephropathy w/Systolic HPT
- 11. Dyslipidemia
- 12. History of MI

- 13. Hyperkinetic Circulation
- 14. Hypertension, Uncomplicated
- 15. Impotence From Other Drugs
- 16. Isolated Systolic Hypertension
- 17. Left Ventricular Failure Due To Systolic Dysfunction
- 18. Left Ventricular Hypertrophy
- 19. Migraine Headaches
- 20. Obesity
- 21. Old Age
- 22. Paroxysmal Supraventricular Tachycardia
- 23. Peripheral Artery Disease
- 24. Pregnancy
- 25. Prostatism
- 26. Proteinuria in Chronic Renal Disease
- 27. Renal Disease
- 28. Senile Tremor
- 29. Systolic Dysfunction
- 30. Youth



- Unfavorable Treatment Effects this addresses side effects to hypertensive medications which are not desirable.
 - 1. Thiazide diuretics should be used cautiously in patients who have gout or who have a history of significant hyponatremia.

- 2. Beta-blockers should generally be avoided in individuals who have asthma, reactive airways disease, or second or third degree heart block.
- 3. Angiotensin converting enzyme inhibitors (ACEIs) and angiotensin receptor blockers (ARBs) should not be given to women likely to become pregnant.
- 4. ACEI-induced angioedema occurs 2-4 times more frequently in African American patients with hypertension than in other groups.
- 5. Aldosterone antagonists and potassium-sparing diuretics can cause hyperkalemia and should generally be avoided in patients who have serum potassium values more than 5.0 mEq/L while not taking medications.
- 6. The initiation of drug therapy with more than one agent may increase the likelihood of achieving the blood pressure goal in a more timely fashion, but particular caution is advised in those at risk for orthostatic hypotension, such as patients with older persons.



Hypertension Plan Template

Hypertension Manage	ement Patient Ric	chmondPROI Ztest	Navigation • HPT • General
<u>Guidelines</u>	Ag	e 35 Sex M	Home
Beginning Blood Pres	Uinhaat D	lood Pressure	Lifestyle Changes
	o III	0 /0	Dippers and White Coat
Val Cinn	Maior Diele Fontone		HPT and Diabetes
Vital Signs Blood Pressure Pulse Pressure	Major Risk Factors ✓ Tobacco Use	Calculate Assessment	HPT and Depression
Trial 1 / 0	Dyslipidemia	Blood Pressure Classification	HPT and the Elderly
Trial 2	Diabetes Mellitus Family Hx of CV Disease	Blood Pressure Classification	HPT, Insulin Resistance
Trial 3 /	Male < 55	<u> </u>	Isolated Systolic HPT
Pulse	Female < 65 Sex	Recommended Follow-Up	HPT and Kidney Disease
Height inches Weight pounds	✓ Male	Risk Group	Evaluation
вмі 🗔	Postmenopausal Female	Tarakarah Barah a Biah darah	Diagnosis and Screening
Body Fat %	Additional Risk Factors	Treatment Based on Risk Assessment	Treatment
Waist 1234J inches	☐ CHF ☐ CAD		HPT Plan
Hips inches	□ TIA		Physician Role
Fram CVD 10-Yr Risk %	Stroke Peripheral Vascular Disease		
Fram. Stroke 10-Yr Risk 0 %	Renal Insufficiency		Patient Information Click for Documents
Global Cardio Risk .0	Retinopathy	Lab Results	Physician Information
Metabolic Syndrome - C + C		Labs Over Time	Classification Risk Stratification
Vitals Over Time			The strain out of the strain o

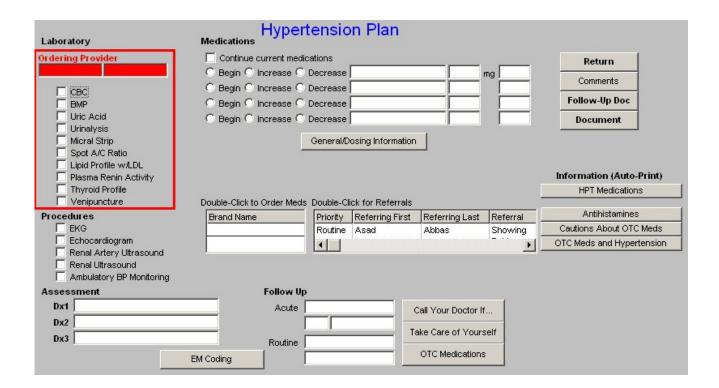
This template facilitates the documentation of the plan which has been developed by the working through of the Hypertension Suite of Templates.

The **Hypertension Plan Template** is divided into three columns:

Column 1 -

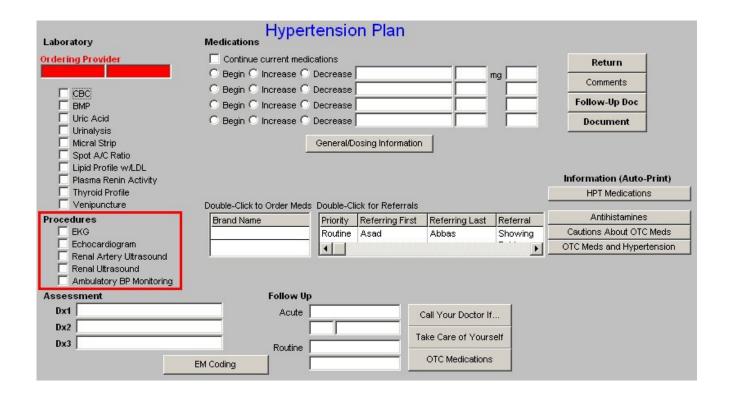
Laboratory - these labs can be ordered from the Hypertension Plan Template and charge posted as well

- CBC
- BMP
- Uric Acid
- Urinalysis
- Micral Strip
- Spot A/C Ratio
- Lipid Profile
- Plasma Renin Activity
- Thyroid Profile



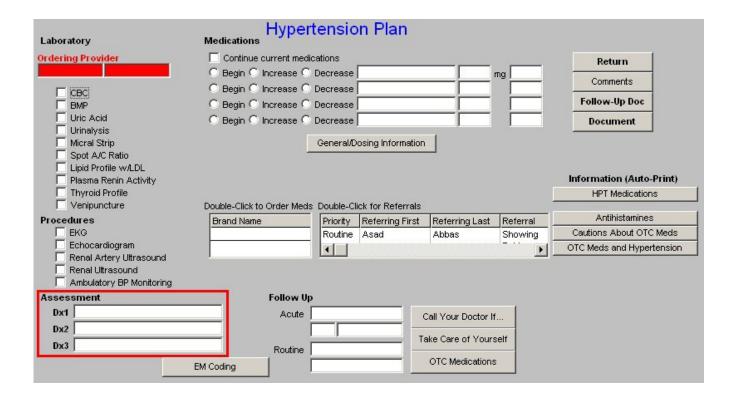
Procedures - these procedures and tests can be ordered and charge posted from the Hypertension Plan Template.

- EKG
- Echocardiogram
- Renal Artery Ultrasound
- Renal Ultrasound
- Ambulatory BP Monitoring



Assessment - this makes it possible to select the appropriate ICD-9 codes from the Hypertension Plan Template.

Dx1 - this first diagnosis box has a pick list made up only of hypertension ICD-9 Codes
DX2 - these next two have pick list made up of all of SETMA's ICD-9 codes
Dx3

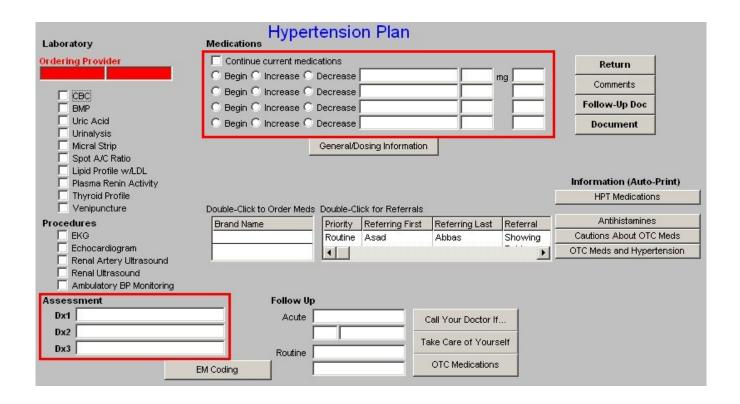


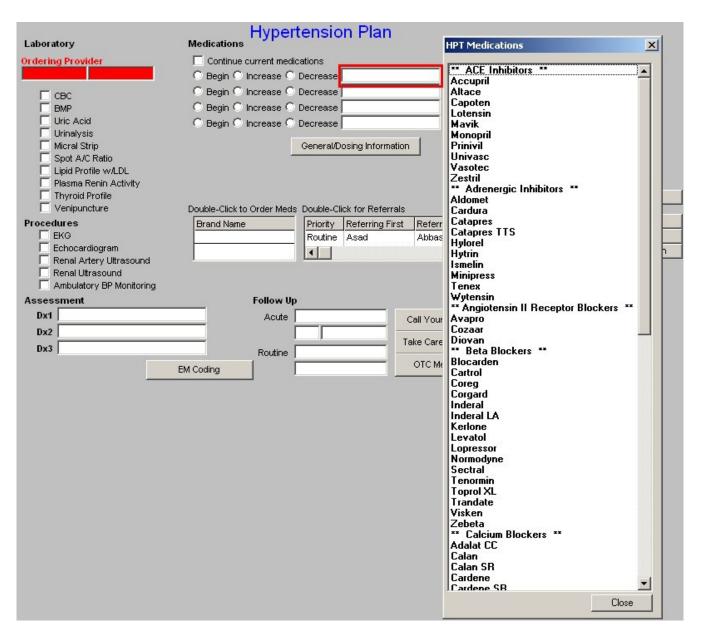
Submit Charge Posting - Once the Assessment is completed and the laboratory and procedures have been selected, depressing this buttion completes charge posting, sending the orders to the lab, x-ray, etc., and putting this data on the patient's chart note.

E & M Coding - This links the provider to the E&M Charge posting template. For more information on Charge posting go to Charge Posting Tutorial or E & M Coding go to E & M E & M Coding.

Column 2 -

Medications -- There are four sets of boxes where medications can be added, increased, decreased. The pick list which is attached to these boxes is:





ACE Inhibitors

- a. Accupril
- b. Altace
- c. Capoten
- d. Lotensin
- e. Mavik
- f. Monopril
- g. Prinivil
- h. Univasc
- i. Vasotec
- i. Zestril

• Adrenergic Inhibitors

- a. Aldomet
- b. Cardura
- c. Catapres
- d. Catapres TTS
- e. Hylorel
- f. Hytrin
- g. Ismelin
- h. Minipress
- i. Tenex
- j. Wytensin

• Angiotensin II Receptor Blockers

- a. Avapro
- b. Cozaar
- c. Diovan

• Beta Blockers

- a. Blocarden
- b. Cartrol
- c. Coreg
- d. Corgard
- e. Inderal
- f. Inderal LA
- g. Kerlone
- h. Levatol
- i. Lopressor
- j. Normodyne
- k. Sectral
- 1. Tenormin
- m. Toprol XL
- n. Trandate
- o. Visken
- p. Zebeta

• Calcium Blockers

- a. Adalat CC
- b. Calan
- c. Calan SR
- d. Cardene
- e. Cardene SR
- f. Cardizem CD
- g. Cardizem SR
- h. Covera-HS
- i. Dilacor XR
- j. DynaCirc

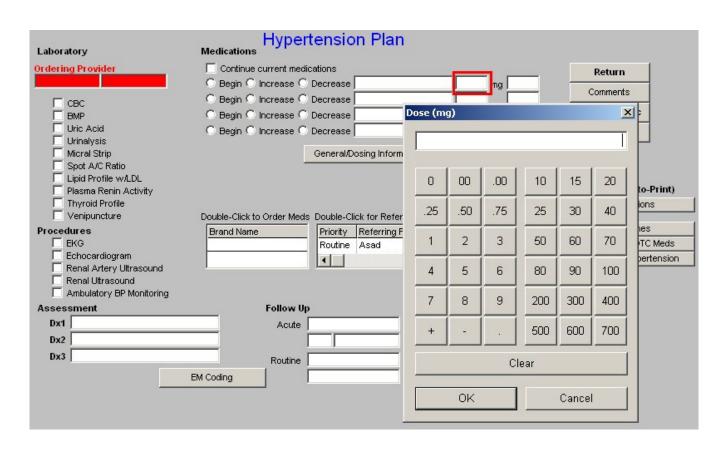
- k. DynaCirc CR
- 1. Isoptin
- m. Isoptin SR
- n. Norvasc
- o. Plendil
- p. Procardia XL
- q. Sular
- r. Tiazac
- s. Verelan

Oral Diuretics

- a. Aldactazide-25
- b. Aldactazide-50
- c. Aldactone
- d. Bumex
- e. Damadex
- f. Diucardin
- g. Diuril
- h. Dyazide
- i. Dyrenium
- j. Edecrin
- k. Enduron
- 1. Esidrix
- m. HydroDiuril
- n. Hygroton
- o. Lasix
- p. Lozol
- q. Maxzide
- r. Maxzide-25
- s. Microzide
- t. Midamor
- u. Moduretic
- v. Mykrox
- w. Naturetin
- x. Oretic
- y. Thalitone
- z. Zaroxolyn

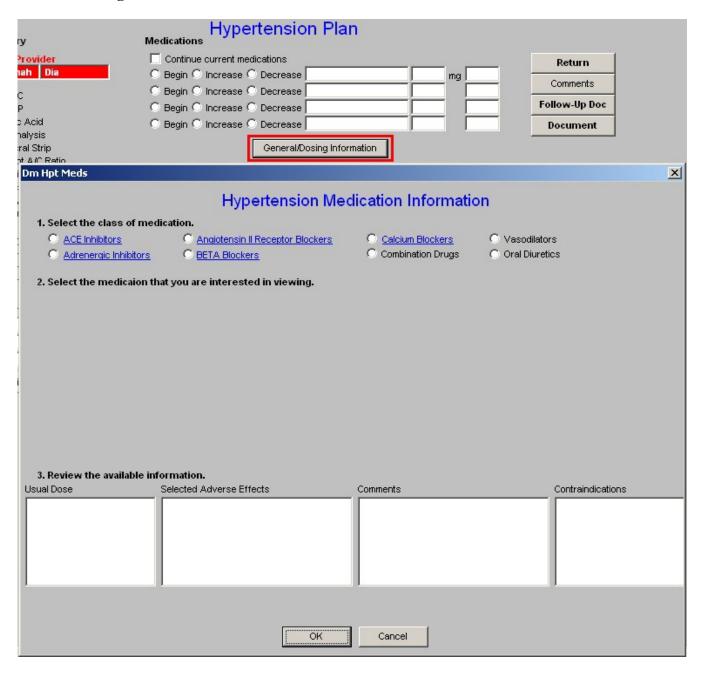
Vasodiltors

- a. Apresoline
- b. Loniten



Laboratory	Hypert Medications	ension Plan			
Ordering Provider	Continue current medic	Decrease	mg	Retur	
CBC BMP Uric Acid Urinalysis Micral Strip Spot A/C Ratio Lipid Profile w/LDL Plasma Renin Activity Thyroid Profile Venipuncture	_	Decrease Decrease General/Dosing Informatio	n l	SMBG BID QD QOD TID	Print)
Procedures EKG Echocardiogram Renal Artery Ultrasound Renal Ultrasound Ambulatory BP Monitoring	Double-Click to Order Meds Brand Name	Priority Referring First Routine Asad		Clo	Meds ension
Assessment	Follow Up				
Dx1 Dx2	Acute		Call Your Doctor If		
Dx3	Routine		Take Care of Yourself		
	EM Coding		OTC Medications		

General/Dosing Information



This is an electronic PDR where a class of drugs can be chosen from:

- Ace Inhibitors
- Adrenergic Inhibitors
- Angiotensin II Receptor Blockers
- Beta Blockers

- Calcium Blockers
- Combination Drugs
- Vasodilators
- Oral Diuretics

Once the class of drugs is chosen, the available drugs from that class are displayed. When you select one of those drugs by check the box beside it, the following information is displayed:

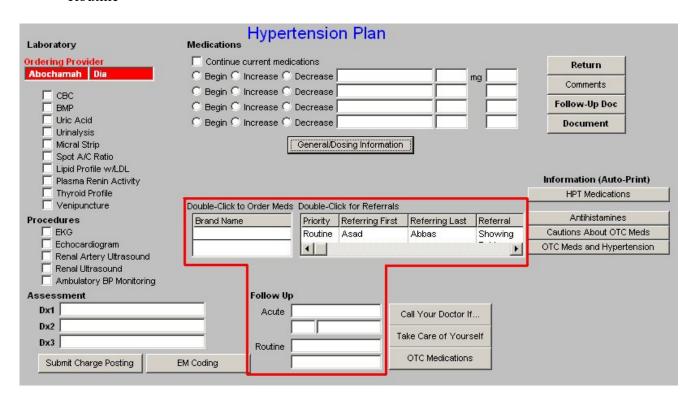
- Usual Dosage
- Selected Adverse Effects
- Comments
- Contraindications

Beneath this function are links to the:

- Medication Module
- Referrals Template

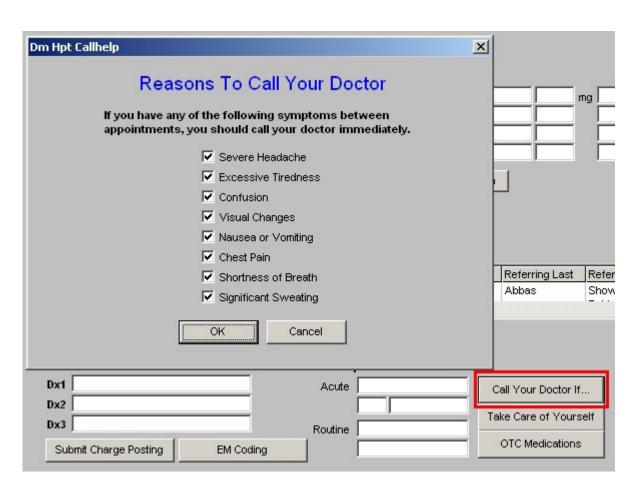
Follow-up

- Acute
- Routine



There are then three Pop-ups with the following content. Each of these prints on the **Hypertension Follow-up Note** automatically.

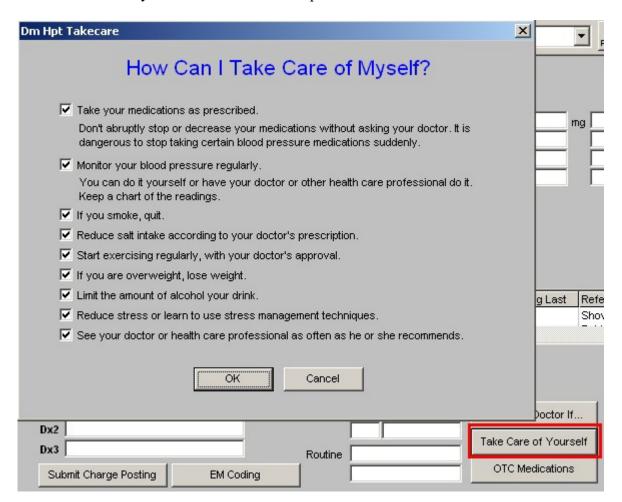
- Call Your Doctor if. -- If you have any of the following symptoms between appointments, you should call your doctor immediately.
 - a. Severe Headache
 - b. Excessive Tiredness
 - c. Confusion
 - d. Visual Changes
 - e. Nausea or Vomiting
 - f. Chest Pain
 - g. Shortness of Breath
 - h. Significant Sweating



• How Can I Take Care of Myself?

- 1. Take your medications as prescribed. -- Don't abruptly stop or decrease your medications without asking your doctor. It is dangerous to stop taking certain blood pressure medications suddenly.
- 2. Monitor your blood pressure regularly. -- You can do it yourself or have your doctor or other health care professional do it. Keep a chart of the readings.
- 3. If you smoke, quit.
- 4. Reduce salt intake according to your doctor's prescription.

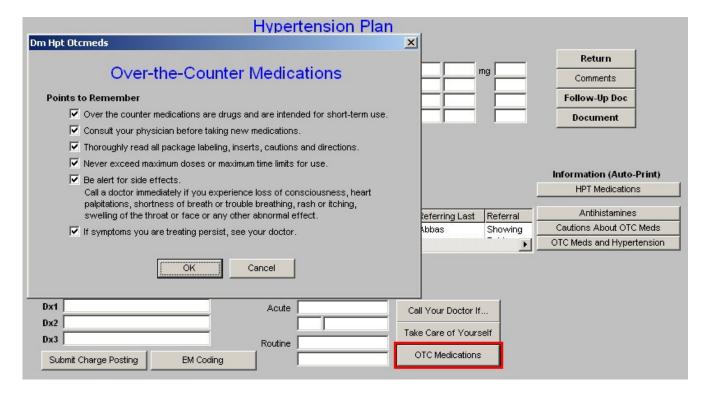
- 5. Start exercising regularly, with your doctor's approval.
- 6. If you are overweight, lose weight.
- 7. Limit the amount of alcohol your drink.
- 8. Reduce stress or learn to use stress management techniques.
- 9. See your doctor or health care professional as often as he or she recommends.



• Over-the-Counter Medications

Points to Remember

- 1. Over the counter medications are drugs and are intended for short-term use.
- 2. Consult your physician before taking new medications.
- 3. Thoroughly read all package labeling, inserts, cautions and directions.
- 4. Never exceed maximum doses or maximum time limits for use.
- 5. Be alert for side effects.
- 6. Call a doctor immediately if you experience loss of consciousness, heart palpitations, shortness of breath or trouble breathing, rash or itching, swelling of the throat or face or any other abnormal effect.
- 7. If symptoms you are treating persist, see your doctor.



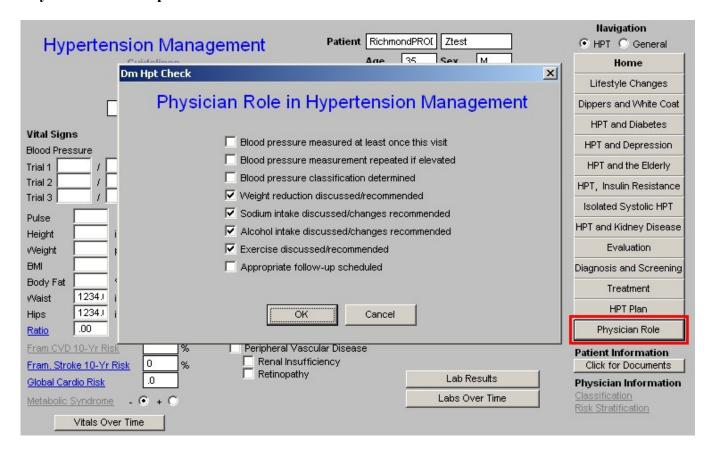
Column 3 -

- **Return** this takes you back to the Hypertension Master Template.
- Comments this allows for free text comments which do not logically fit anywhere else.
- **Follow-up Doc** this creates the Hypertension Follow-up Document which should then be given to the patient.
- **Document** this creates the Hypertension Chart Note

Information

- **HPT Medications** when this button is depressed it launches a list of documents on the following medications, each of which can be printed.
 - 1. ACE Inhibitors and Angiotensin II Blockers
 - 2. Adrenergic Inhibitors
 - 3. Beta Blockers
 - 4. Calcium Channel Blockers
 - 5. Combination Drugs
 - 6. Oral Diuretics
 - 7. Vasodilators
- Antihistamines
- Cautions About OTC Meds
- OTC Meds and Hypertensions

Physician Role Template



As the provider progresses through SETMA's Hypertension Suite of Templates, this template automatically collects the data points for the **Physician Consortium for Performance Improvement Data** for quality of care in hypertension management. A review of this template will allow a provider to see "how he/she is doing," as measured against a national standard of care in hypertensive management. The standard of excellence in the management of hypertension is measured on the following 8 data points.

Physician Role in Hypertension Management

- Blood pressure measured at least once this visit
- Blood pressure measurement repeated if elevated
- Blood pressure classification determined
- Weight reduction discussed/recommended
- Sodium intake discussed/changes recommended
- Alcohol intake discussed/changes recommended
- Exercise discussed/recommended
- Appropriate follow-up scheduled

The Physician Consortium for Performance Improvement

The Physician Consortium for Performance Improvement (Consortium) is a group of clinical and methodological experts convened by the AMA. The Consortium includes representatives from more than 60 national medical specialty and state medical societies, the Agency for Healthcare Research and Quality, and the Centers for Medicare and Medicaid Services.

The Consortium's vision is to fulfill the responsibility of physicians to patient care, public health, and safety by:

- becoming the leading source organization for evidence-based clinical performance measures and outcomes reporting tools for physicians; and
- ensuring that all components of the medical profession have a leadership role in all national forums seeking to evaluate the quality of patient care.

The Consortium's mission is to improve patient health and safety by:

- identifying and developing evidence-based clinical performance measures that enhance quality of patient care and that foster accountability;
- promoting the implementation of effective and relevant clinical performance improvement activities; and
- advancing the science of clinical performance measurement and improvement.

The Consortium works to develop evidence-based clinical performance measures and clinical outcomes reporting tools to support physicians in quality improvement efforts.

The Consortium has published a number of disease management data sets which established quality of care measures with which physicians and other healthcare providers can measure their own performance.