# Morbidity and Mortality in People with Serious Mental Illness

National Association of State Mental Health Program
Directors
Medical Directors Council
July 2006

#### Overview- THE PROBLEM

- Increased Morbidity and Mortality Associated with Serious Mental Illness (SMI)
- Increased Morbidity and Mortality Largely Due to Preventable Medical Conditions
  - Metabolic Disorders, Cardiovascular Disease, Diabetes Mellitus
  - High Prevalence of Modifiable Risk Factors (Obesity, Smoking)
  - Epidemics within Epidemics (e.g., Diabetes, Obesity)
- Some Psychiatric Medications Contribute to Risk
- Established Monitoring and Treatment Guidelines to Lower Risk Are Underutilized in SMI Populations

#### Overview - PROPOSED SOLUTIONS

- Prioritize the Public Health Problem
  - Target Providers, Families and Clients
  - Focus on Prevention and Wellness
- Track Morbidity and Mortality in Public Mental Health Populations
- Implement Established Standards of Care
  - Prevention, Screening and Treatment
- Improve Access to and Integration of Physical Health and Mental Health Care

### Why Should we be Concerned About Morbidity and Mortality?

■ Recent data from several states have found that people with serious mental illness served by our public mental health systems die, on average, at least 25 years earlier that the general population.

### Recent Multi-State Study Mortality Data: Years of Potential Life Lost

Year	AZ	MO	OK	RI	TX	UT	VA (IP only)
1997		26.3	25.1		28.5		
1998		27.3	25.1		28.8	29.3	15.5
1999	32.2	26.8	26.3		29.3	26.9	14.0
2000	31.8	27.9		24.9			13.5

Compared to the general population, persons with major mental illness typically lose more than 25 years of normal life span

Colton CW, Manderscheid RW. Prev Chronic Dis [serial online] 2006 Apr [date cited]. Available from: URL:http://www.cdc.gov/pcd/issues/2006/apr/05\_0180.htm

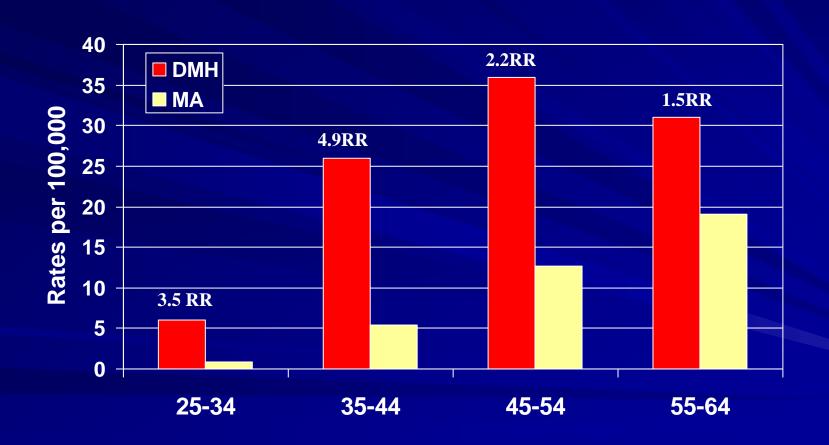
#### Ohio Study-1998-2002

#### **Mean Years of Potential Life lost**

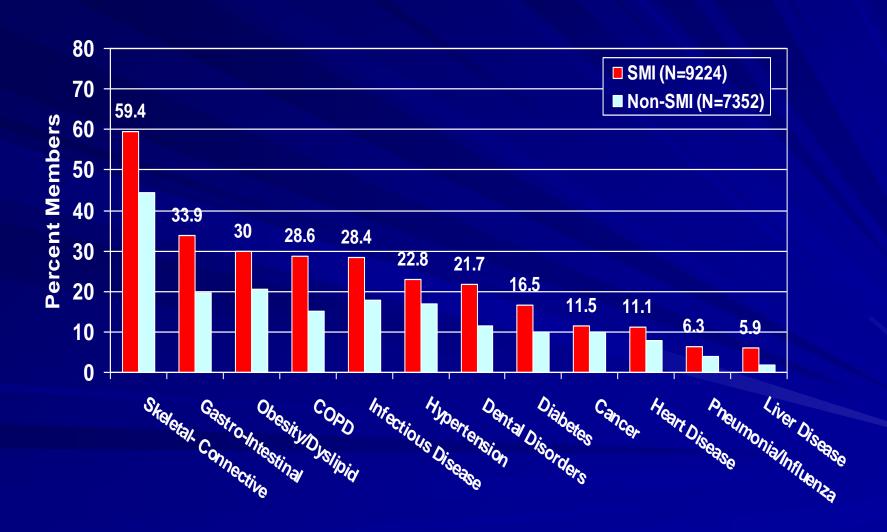
20,018 persons discharged, 608 deaths

Cause	<u>M</u>	<u>F</u>	<u>N</u>
All	31.8	32.5	32.0
Intentional self-harm (suicide)	41.4	42.7	41.7
Assault (homicide)	42.3	35.8	41.6
Accidents (unintentional injuries)	39.5	43.1	40.4
Symptoms, signs, & abnormal	32.8	35.0	33.4
clinical & laboratory findings, NEC			
Diabetes mellitus	25.8	37.2	30.2
Pneumonia & Influenza	29.4	25.0	28.3
Diseases of heart	27.7	26.6	27.3
Cerebrovascular diseases	20.7	32.8	25.5
Malignant neoplasms (cancers)	24.3	26.9	25.3
Chronic lower respiratory diseases	18.6	24.1	21.1

## Massachusetts Study: Deaths from Heart Disease by Age Group/DMH Enrollees with SMI Compared to Massachusetts 1998-2000



#### Maine Study Results: Comparison of Health Disorders Between SMI & Non-SMI Groups



## Ohio Study Leading Causes of Death

<u>Cause</u>	<b>ICD-9 Codes</b>	ICD-10 Codes	<u>M</u>	<u>F</u>	<u>N</u>	<u>%</u>
Diseases of heart	390-398, 402,	100-09, 111,	83	43	126	20.7
	404, 410-429	I13, I20-51				
Intentional self-harm (suicide)	E950-959	X60-84,	84	24	108	17.8
		Y87.0				
Accidents (unintentional injuries)	E800-869,	V01-X59,	61	22	83	13.7
	E880-929	Y85-86				
Malignant neoplasms (cancers)	140-208	C00-C97	27	17	44	7.2
Symptoms, signs, & abnormal	780-799	R00-99	23	9	32	5.3
clinical & laboratory findings, NEC						
Chronic lower respiratory diseases	490-494, 496	J40-J47	17	14	31	5.1
Diabetes mellitus	250	E10-14	11	7	18	3.0
Pneumonia & Influenza	480-487	J10-18	12	4	16	2.6
Cerebrovascular diseases	430-434,	160-69	6	4	10	1.6
	436-438					
Assault (homicide)	E960-969	X85-Y09,	9	1	10	1.6
		Y87.1				

## Ohio Study Standardized Mortality Ratios

Cause	<u>Overall</u>	
	<u>N</u>	<u>SMR</u>
All causes of death	608	3.2†
Intentional self-harm (suicide)	108	12.6†
Symptoms, signs, & abnormal	32	9.7†
clinical & laboratory findings, NEC		
Pneumonia & Influenza	16	6.6†
Chronic lower respiratory diseases	31	5.5†
Accidents (unintentional injuries)	83	3.8†
Diseases of heart	126	3.4†
Diabetes mellitus	18	3.4†
Assault (homicide)	10	1.7
Cerebrovascular diseases	10	1.5
Malignant neoplasms (cancers)	44	0.9
† P<0.001		

## What are the Causes of Morbidity and Mortality in People with Serious Mental Illness?

- While suicide and injury account for about 30-40% of excess mortality, about 60% of premature deaths in persons with schizophrenia are due to "natural causes"
  - Cardiovascular disease
  - Diabetes
  - Respiratory diseases
  - Infectious diseases

### Schizophrenia: Natural Causes of Death

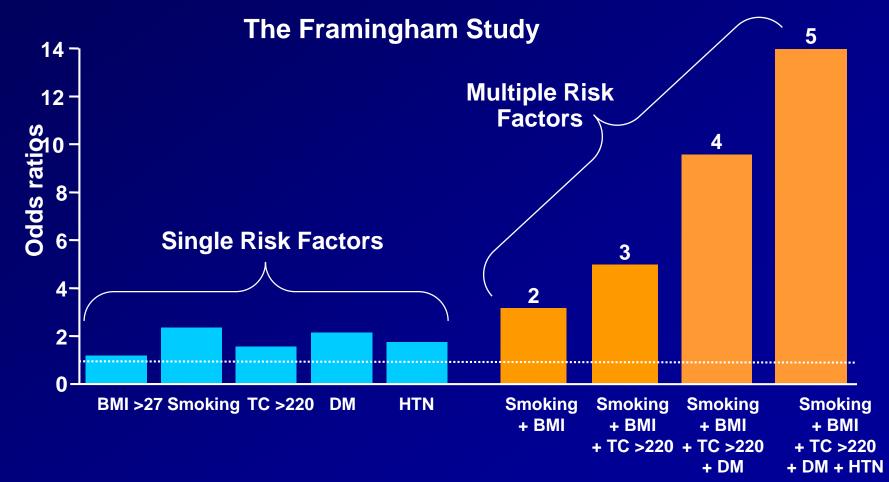
Higher standardized mortality rates than the general population from:

<ul><li>Diabetes</li></ul>	2.7x
<ul> <li>Cardiovascular disease</li> </ul>	2.3x
<ul> <li>Respiratory disease</li> </ul>	3.2x
<ul> <li>Infectious diseases</li> </ul>	3.4x

- Cardiovascular disease associated with the largest number of deaths
  - 2.3 X the largest cause of death in the general population



### Cardiovascular risk factors – overview



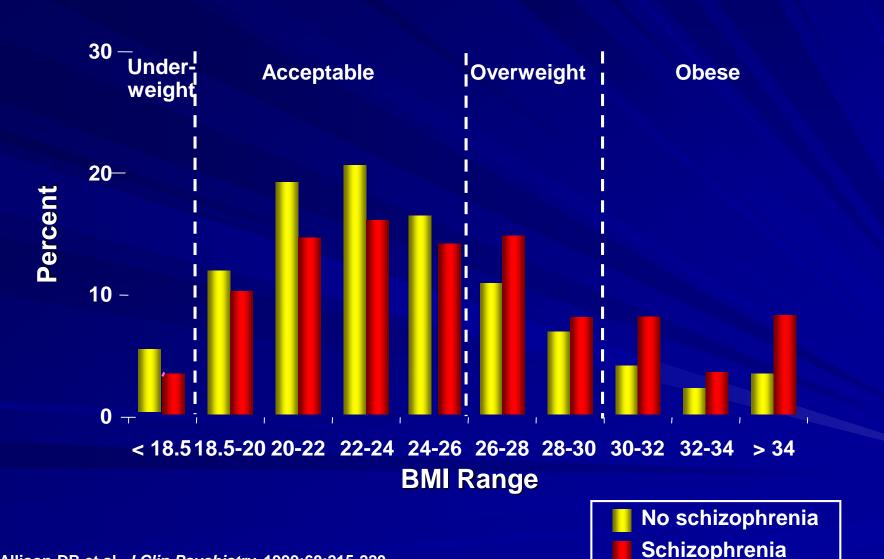
BMI = body mass index; TC = total cholesterol; DM = diabetes mellitus; HTN = hypertension. Wilson PWF et al. Circulation. 1998;97:1837–1847.

### Cardiovascular Disease (CVD) Risk Factors

Modifiable Risk	Estimated Prevalence and Relative Risk (RR)						
Factors	Schizophrenia	Bipolar Disorder					
Obesity	45–55%, 1.5-2X RR <sup>1</sup>	26% <sup>5</sup>					
Smoking	50–80%, 2-3X RR <sup>2</sup>	55% <sup>6</sup>					
Diabetes	10–14%, 2X RR <sup>3</sup>	10% <sup>7</sup>					
Hypertension	≥18%⁴	15% <sup>5</sup>					
Dyslipidemia	Up to 5X RR <sup>8</sup>						

<sup>1.</sup> Davidson S, et al. *Aust N Z J Psychiatry*. 2001;35:196-202. 2. Allison DB, et al. *J Clin Psychiatry*. 1999; 60:215-220. 3. Dixon L, et al. *J Nerv Ment Dis*. 1999;187:496-502. 4. Herran A, et al. *Schizophr Res*. 2000;41:373-381. 5. MeElroy SL, et al. *J Clin Psychiatry*. 2002;63:207-213. 6. Ucok A, et al. Psychiatry Clin Neurosci. 2004;58:434-437. 7. Cassidy F, et al. *Am J Psychiatry*. 1999;156:1417-1420. 8. Allebeck. Schizophr Bull. 1999;15(1)81-89.

#### BMI Distributions for General Population and Those With Schizophrenia (1989)



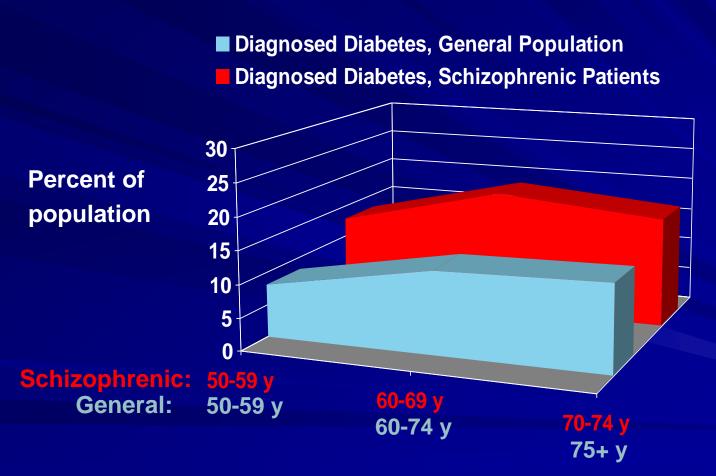
Allison DB et al. J Clin Psychiatry. 1999;60:215-220.

#### **Mental Disorders and Smoking**

- Higher prevalence (56-88% for patients with schizophrenia) of cigarette smoking (overall U.S. prevalence 25%)
- More toxic exposure for patients who smoke (more cigarettes, larger portion consumed)
- Smoking is associated with increased insulin resistance
- Similar prevalence in bipolar disorder

George TP et al. Nicotine and tobacco use in schizophrenia. In: Meyer JM, Nasrallah HA, eds. Medical Illness and Schizophrenia. American Psychiatric Publishing, Inc. 2003; Ziedonis D, Williams JM, Smelson D. Am J Med Sci. 2003(Oct);326(4):223-330

## Prevalence of Diagnosed Diabetes in General Population Versus Schizophrenic Population





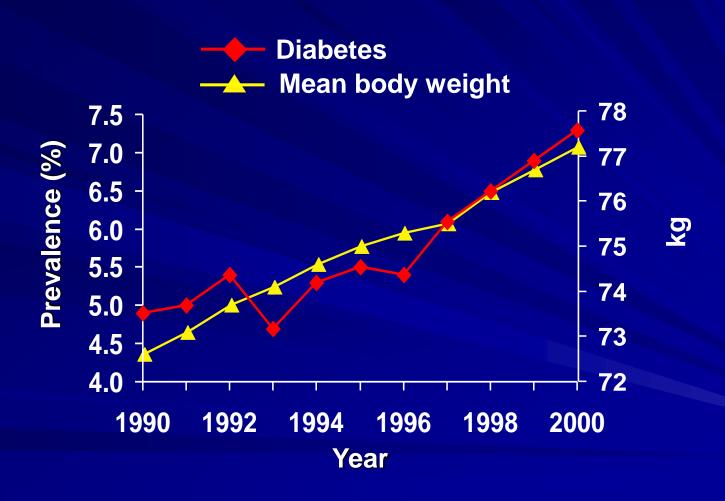
#### Hypothesized Reasons Why There May Be More Type 2 Diabetes in People With Schizophrenia

- Genetic link between schizophrenia and diabetes
- Impact of lifestyle
- Medication effect increasing insulin resistance by impacting insulin receptor or postreceptor function
- Drug effect on caloric intake or expenditure (obesity, activity)

### How Does This Relate to What is Happening in the General Population?

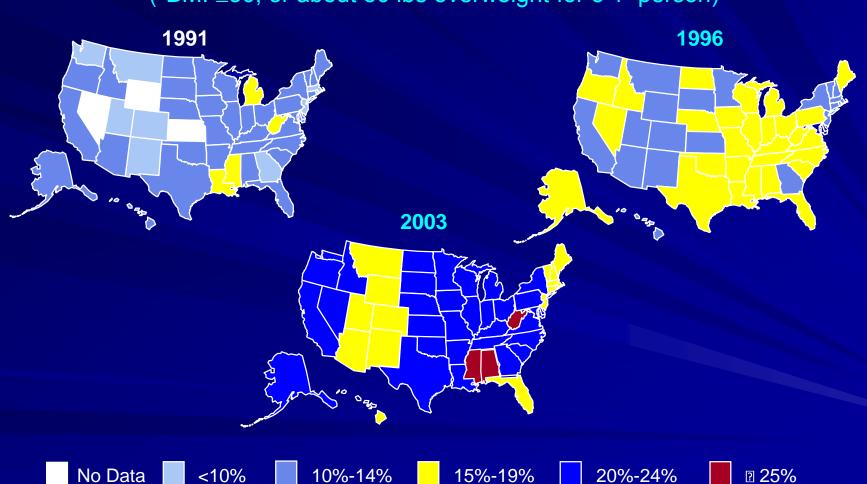
- There is an "epidemic" of obesity and diabetes, increasing risk of multiple medical conditions and cardiovascular disease.
  - Obesity
  - Diabetes
  - Metabolic Syndrome
  - Cardiovascular Disease

### Diabetes and Obesity: The Continuing Epidemic

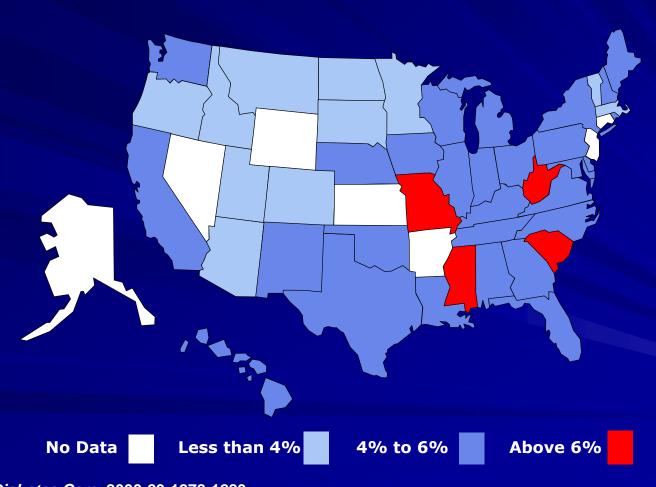


### Obesity Trends\* Among US Adults BRFSS, 1991, 1996, 2003

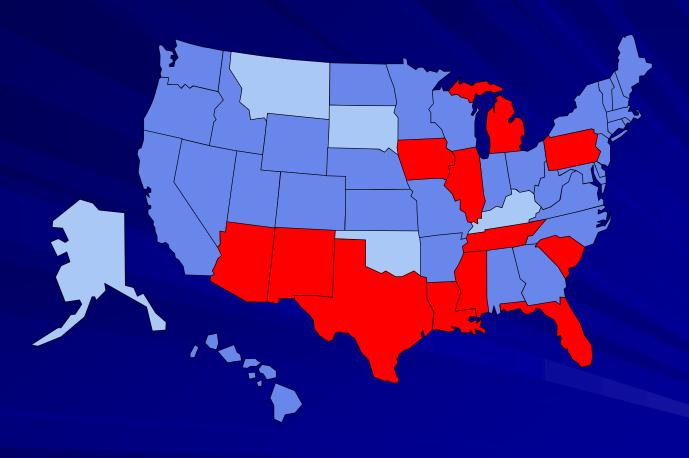
(\*BMI ≥30, or about 30 lbs overweight for 5'4" person)



#### Diabetes and Gestational Diabetes Trends: US Adults, BRFSS 1990

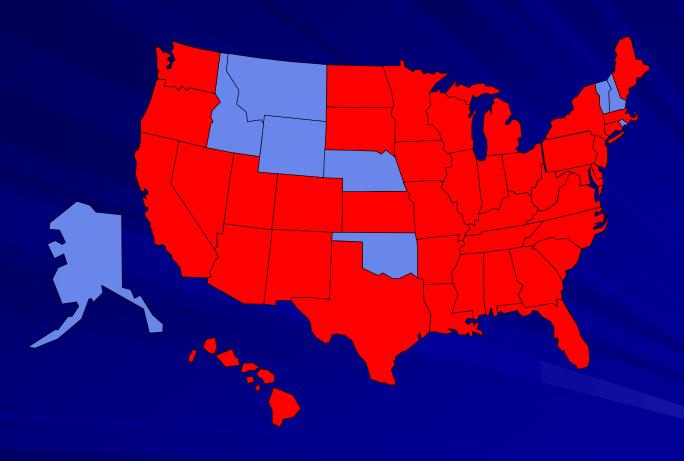


#### Diabetes and Gestational Diabetes Trends: US Adults, BRFSS 1995



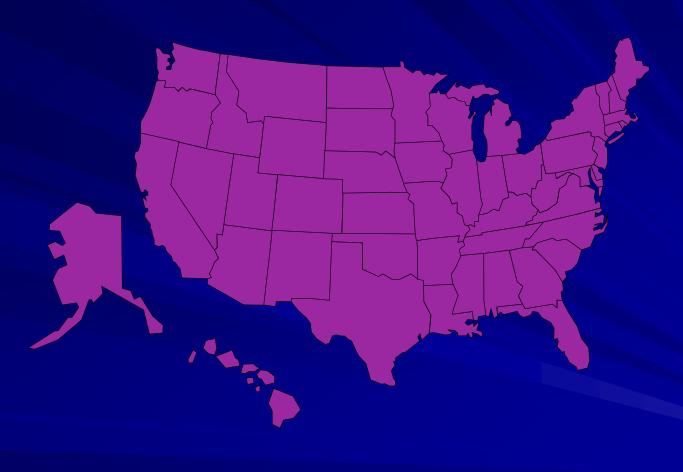
No Data Less than 4% 4% to 6% Above 6%

#### Diabetes and Gestational Diabetes Trends: US Adults, BRFSS 2000



No Data Less than 4% 4% to 6% Above 6%

#### Diabetes and Gestational Diabetes Trends: US Adults, Estimate for 2010



No Data

Less than 4%

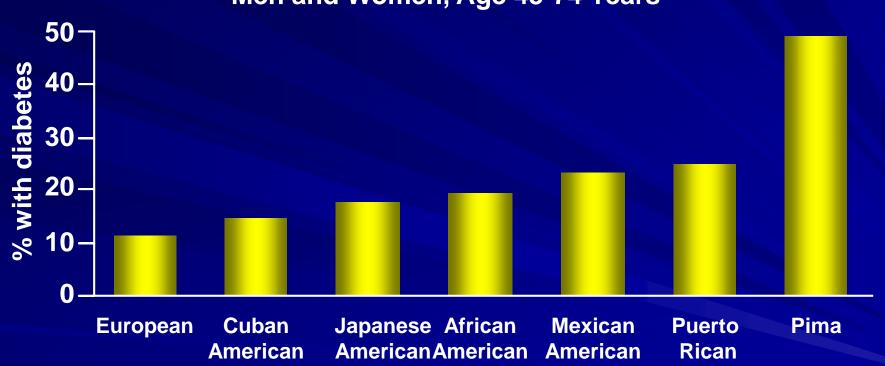
4% to 6%

**Above 6%** 

Above 10%

### US Diabetes Prevalence by Ethnic Group





Harris et al. *Diabetes*. 1987;36:523.

Flegal et al. Diabetes Care. 1991;14(suppl 3):628.

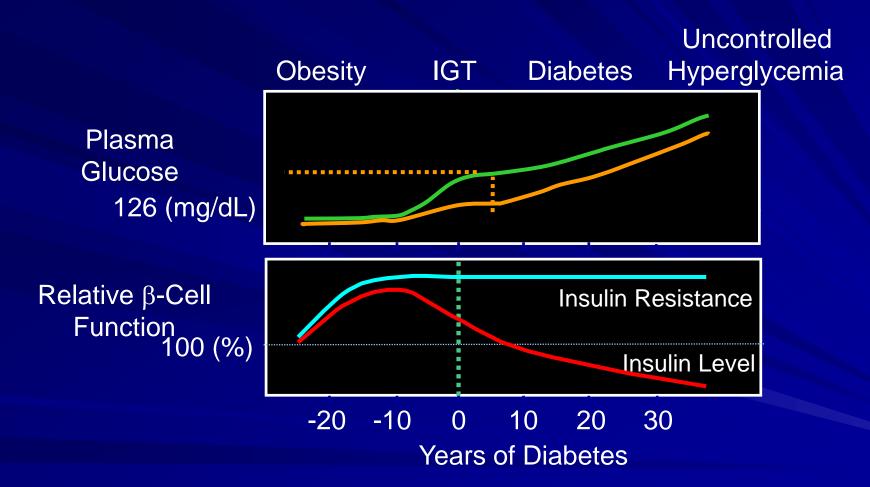
Knowler et al. Diabetes Care. 1993;16(suppl 1):216.

Fujimoto et al. Diabetes Res Clin Pract. 1991;13:119.

Fujimoto et al. Diabetes. 1987;36:721.



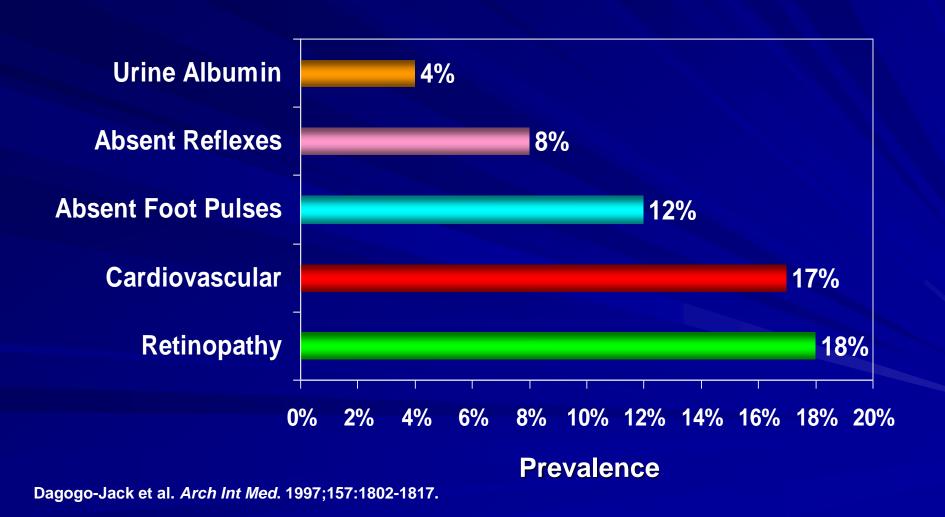
#### Natural History of Type 2 Diabetes



IGT = impaired glucose tolerance.

Adapted from: International Diabetes Center (IDC). Available at: <a href="https://www.parknicollet.com/diabetes/disease/diagnosing.cfm">www.parknicollet.com/diabetes/disease/diagnosing.cfm</a>. Accessed March 26, 2006.

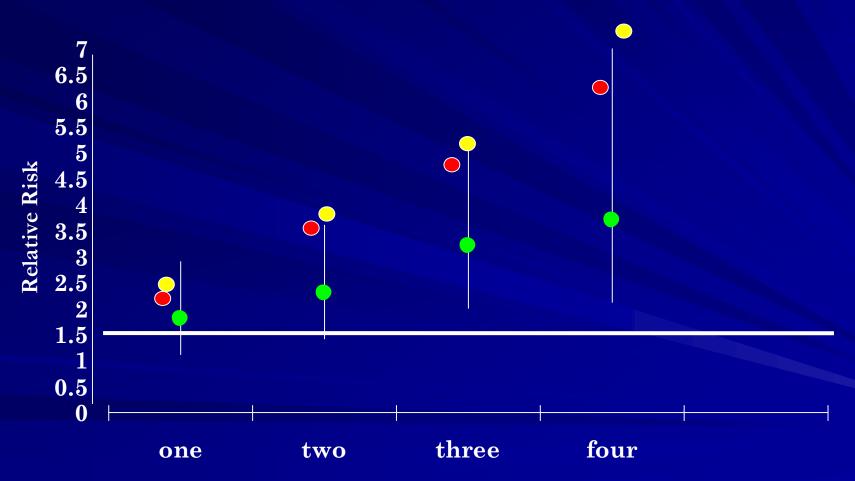
### Prevalence of Diabetic Tissue Damage at Diagnosis of Type 2 Diabetes



#### Identification of the Metabolic Syndrome

≥3 Risk Factors Required for Diagnosis							
Risk Factor	Defining Level						
Abdominal obesity Men Women	Waist circumference >40 in (>102 cm) >35 in (>88 cm)						
Triglycerides	≥150 mg/dL (1.69mmol/L)						
HDL cholesterol Men Women	<40 mg/dL (1.03mmol/L) <50 mg/dL (1.29mmol/L)						
Blood pressure	≥130/85 mm Hg						
Fasting blood glucose	≥110 mg/dL (6.1mmol/L)						

## CHD Risk Increases with Increasing Number of Metabolic Syndrome Risk Factors

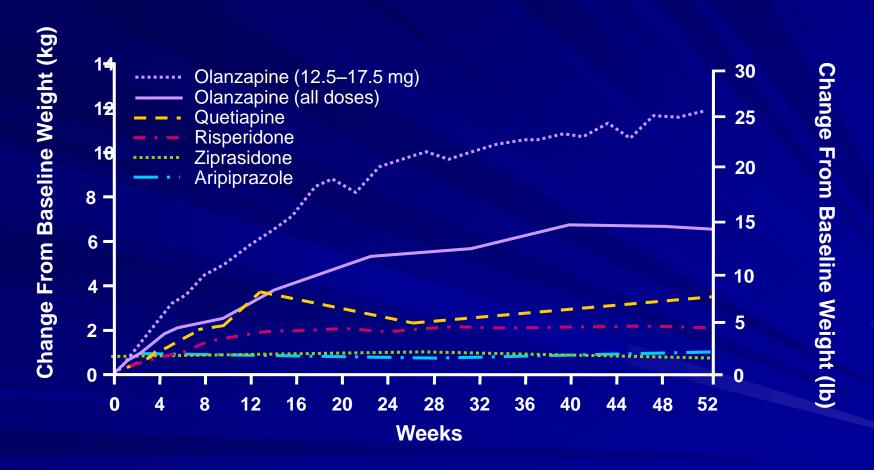


- Sattar et al, Circulation, 2003;108:414-419
  - Whyte et al, American Diabetes Association, 2001 Adapted from Ridker, Circulation 2003;107:393-397

## Modifiable Risk Factors Affected by Psychotropics

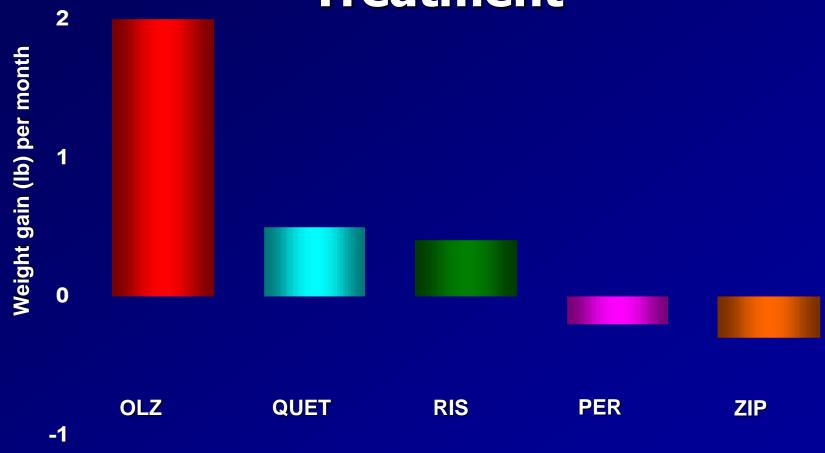
- Overweight / Obesity
- Insulin resistance
- Diabetes/hyperglycaemia
- Dyslipidemia

#### 1-Year Weight Gain: Mean Change From Baseline Weight

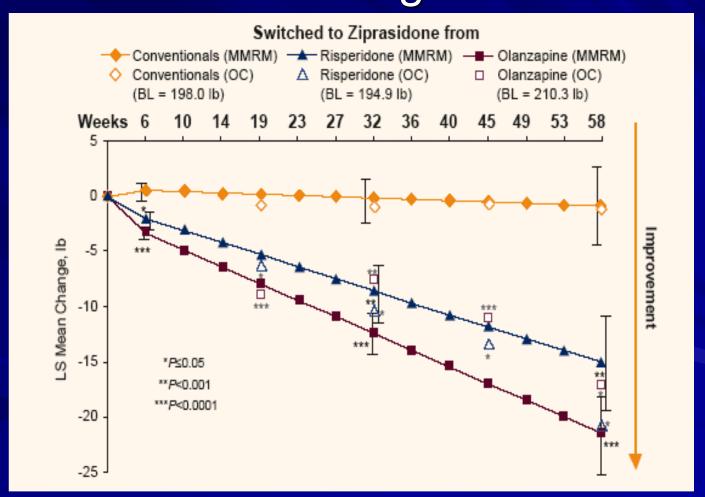


Nemeroff CB. *J Clin Psychiatry*. 1997;58(suppl 10):45-49; Kinon BJ et al. *J Clin Psychiatry*. 2001;62:92-100; Brecher M et al. American College of Neuropsychopharmacology; 2004. Poster 114; Brecher M et al. *Neuropsychopharmacology*. 2004;29(suppl 1):S109; Geodon<sup>®</sup> [package insert]. New York, NY:Pfizer Inc; 2005. Risperdal<sup>®</sup> [package insert]. Titusville, NJ: Janssen Pharmaceutica Products, LP; 2003; Abilify<sup>®</sup> [package insert]. Princeton NJ: Bristol-Myers Squibb Company and Rockville, Md: Otsuka America Pharmaceutical, Inc.; 2005.

## CATIE Trial Results: Weight Gain Per Month Treatment



# Change in Weight From Baseline 58 Weeks After Switch to Low Weight Gain Agent



## ADA/APA/AACE/NAASO Consensus on Antipsychotic Drugs and Obesity and Diabetes: Monitoring Protocol\*

	Start	4 wks	8 wks	12 wk	qtrly	12 mos.	5 yrs.
Personal/family Hx	X					X	
Weight (BMI)	X	X	X	X	X		
Waist circumference	X					X	
Blood pressure	X			X		X	
Fasting glucose	X			X		X	
Fasting lipid profile	Χ			X		Χ←	-X

<sup>\*</sup>More frequent assessments may be warranted based on clinical status

\*Diabetes Care. 27:596-601, 2004\*

#### **METABOLIC SCREENING AND MONITORING FORM**

#### NAME:

There is a growing awareness that some psychiatric illnesses and atypical antipsychotics can increase metabolic risks. Frequency of monitoring for modifiable risk factors depends on level of risk present at baseline screening.

OBESITY SCREENING 1,2	BASELINE Dates/Values From Subsequent Visits						
Consider BMI (weight/height in kg/m²) at each visit.	Height	Date / /	//	//	//	//	//
Normal (18.5-24.9); Overweight (25-29.9); Obese (≥30)	Ü	ВМІ					
		Wt					

LIPID	SCREENIN	G — CHOLES	TEROL, TRIGLYO	ERIDES (TG) <sup>3</sup>		BASELINE		Dates/Value	s From Subse	equent Visits	
	Optimal/ Desirable (mg/dL)	Near/Above Optimal (mg/dL)	Borderline High (mg/dL)	High/ Undesirable (mg/dL)	Very High (mg/dL)	//_	//_	//_	//_	//_	//_
Total	<200		200-239	≥240							
LDL	<100	100–129	130–159	160–189	≥190						
HDL	≥60			<40		Enter values as indicated in the Metabolic Syndrome (MS)					
TG	<150		150–199	200–499	≥500*	Screening section of the form below.					

\* ≥500 for TG requires immediate pharmacotherapeutic intervention without waiting for therapeutic lifestyle changes.

METABOLIC SYNDROME (MS) SCREENING <sup>3</sup>	BASELINE	Dates/Values From Subsequent Visits				
Risk Criteria:	//_	//_	//_	//_	//_	//
Abdominal Obesity measured in waist circumference (men >40 inches, women >35 inches)						
Triglycerides (mg/dL) (≥150; or drug treatment)						
HDL Cholesterol (mg/dL) (men <40, women <50; or drug treatment)						
Blood Pressure (mmHg) (≥130/≥85; or drug treatment)						
Fasting Plasma Glucose (≥100 mg/dL; or drug treatment) <sup>4</sup>						
Total Criteria for each visit (≥3 = MS Diagnosis*)						

\*Risk for cardiovascular disease increases with each criterion present, motivating intervention for any single criterion.<sup>5</sup>

				-				
<b>TYPE 2 DIABETI</b>	ES MELLITUS (T2DM) SCREE	NING'						
Risk Factors:	Age (≥45)     Habitual physical inactivity     Rece/ethnicity*     Polycystic ovary syndrome	☐ History of GD☐ Hypertension	Overweight (BMI ≥25 kg/m²)†					<u>&gt;</u> 250 mg/dL
			BASELINE		Dates/Value	s From Subse	equent Visits	
Diagnostic Criteria for Prediabetes and T2DM‡			//	//	//	//_	//_	//_
Fasting Plasma G Normal: <100 mg/dl	ilucose (FPG)§ L; Prediabetes: 100-125 mg/dL; T2DN	l: <u>&gt;</u> 126 mg/dL						
	ld Glucose (OGTT)§ L; Prediabetes: 140-199 mg/dL; T2DN	l: <u>&gt;</u> 200 mg/dL						
Symptoms of T2	OM [Yes + casual (random) PG ≥200	mg/dL]						
Random Plasma	Glucose	n <sub>6</sub>						

\* Includes African Americans, Hispanic Americans, Native Americans, Asian Americans, Pacific Islanders

† May not be correct for all ethnic groups

‡ Screen at 3-year intervals beginning at age 45, particularly for those with BMI of >25; test at <45 or more frequently when overweight and have 1+ other risk factors.

§ FPG and OGTT are the only measures currently approved by the ADA for diabetes screening/diagnosis; ADA recommends preferential use of FPG due to ease of use/acceptance.

|| Diagnosis must be confirmed on a subsequent day with FPG, 2-h PG, or casual (random) PG if symptoms (e.g., polyuria, polydipsia) are present, unless unequivocal hyperglycemia with acute metabolic decompensation is present.

ATP-III recommends therapeutic lifestyle changes (TLC) for those with prediabetes, hypertension, 0-1 CHD risk factor and LDL ≥160 mg/dL, 2+ CHD risk factors and LDL ≥130, MS, and perhaps subsyndromal MS. Follow-up monitoring of 6- to 12-week intervals to monitor TLC response is recommended and pharmacotherapy intervention if TLC fails after 3 months — unless lipid, blood pressure, or glucose values demand immediate drug treatment.

ADA/APA Consensus Statement recommends considering antipsychotic medication switch for those who gain ≥5% of baseline body weight.<sup>9</sup>

Authored by John W. Newcomer, MD and Dan W. Haupt, MD. Compiled primarily from ADA and ATP-III guidelines.

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### Problem: SMI and Reduced Use of Medical Services

- Fewer routine preventive services (Druss 2002)
- Worse diabetes care (Desai 2002, Frayne 2006)
- Lower rates of cardiovascular procedures (Druss 2000)

### Access and Quality of Care

- SMI may be a health risk factor because of:
  - Patient factors, e.g.: amotivation, fearfulness, homelessness, victimization/trauma, resources, advocacy, unemployment, incarceration, social instability, IV drug use, etc
  - Provider factors: Comfort level and attitude of healthcare providers, coordination between mental health and general health care, stigma,
  - System factors: Funding, fragmentation

### Goals: Lower Risk for CVD

- Blood cholesterol
  - 10%  $\downarrow$  = 30%  $\downarrow$  in CHD (200-180)
- High blood pressure (> 140 SBP or 90 DBP)
  - 4-6 mm Hg  $\downarrow$  = 16%  $\downarrow$  in CHD; 42%  $\downarrow$  in stroke
- Cigarette smoking cessation
  - 50%-70% ↓ in CHD
- Maintenance of ideal body weight (BMI = 25)
  - \_ 35%-55% ↓ in CHD
- Maintenance of active lifestyle (20-min walk daily)
  - 35%-55% ↓ in CHD

# Why Should we be Concerned About Morbidity and Mortality?

■ Recent data from several states have found that people with serious mental illness served by our public mental health systems die, on average, at least 25 years earlier that the general population.

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- Prioritize the Public Health Problem
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### Recommendations NATIONAL LEVEL

- 1. Seek federal designation of people with SMI as a distinct at-risk health disparities population. Establish co-ordinated mental health and general health care as a national healthcare priority.
- 2. Establish a committee at the federal level to recommend changes to national surveillance activities that will incorporate information about health status in the population with SMI.
  - Consider representation from SAMHSA, Medicaid, the Centers for Disease Control and Prevention, state MH authorities / NASMHPD, and experts
  - This may include the IOM project and other national surveys.

### Recommendations

#### **NATIONAL LEVEL**

3. Share information widely about physical health risks in persons with SMI to encourage awareness and advocacy. Educate the health care community. Encourage consumers and family members to advocate for wellness approaches as part of recovery.

### Recommendations STATE LEVEL

- Seek state designation of people with SMI as BOTH an at-risk and a health disparities population.
- Establish co-ordinated mental health and general health care as a state healthcare priority.
- 3. Education and advocacy policy makers funders providers individuals, family, community

### Recommendations

#### STATE LEVEL

4. Require, regulate and lead Behavioral Health provider systems to screen, assess and treat both mental health and general health care issues. Provide for

staffing
time
record keeping
reimbursement
linkage with physical healthcare providers

- 5. Funding
- Promote co-ordinated and integrated mental health and physical health care for persons with SMI.

See 11th NASMHPD Technical Paper: Integrating Mental Health and Primary Care.

# Recommendations LOCAL AGENCY / CLINICIAN

- BH providers shall provide quality medical care and mental health care
  - Screen for general health with priority for high risk conditions
  - Offer prevention and intervention especially for modifiable risk factors (obesity, abnormal glucose and lipid levels, high blood pressure, smoking, alcohol and drug use, etc.)
  - Prescribers will screen, monitor and intervene for medication risk factors related to treatment of SMI (e.g. risk of metabolic syndrome with use of second generation anti-psychotics)
  - Treatment per practice guidelines, e.g heart disease, diabetes, smoking cessation, use of novel anti-psychotics.

# LOCAL AGENCY/CLINICIAN Recommendations

#### 2. Care coordination Models

- Assure that there is a specific practitioner in the MH system who is identified as the responsible party for each person's medical health care needs being addressed and who assures coordination all services.
- Routine sharing of clinical information with other providers (primary and specialty healthcare providers as well as mental health providers
- Care integration where services are co-located

# LOCAL AGENCY/CLINICIAN RECOMMENDATIONS

- 3. Support consumer wellness and empowerment to improve personal mental and physical well-being
  - educate / share information to make healthy choices regarding nutrition, tobacco use, exercise, implications of psychotropic drugs
  - teach /support wellness self-management skills
  - teach /support decision making skills
  - motivational interviewing techniques
  - Implement a physical health Wellness approach that is consistent with Recovery principles, including supports for smoking cessation, good nutrition, physical activity and healthy weight.
  - attend to cultural and language needs

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### Full report available at

http://www.nasmhpd.org/publications.cfm#techpap