

### Treatment of Tobacco and Cocaine Use Disorders

Kristina Schnitzer, MD Psychiatrist - Schizophrenia Clinical and Research Program, MGH K12 Scholar - Center for Addiction Medicine, MGH Instructor - Harvard Medical School

### Disclosures

None

### Public Health Burden of Tobacco Dependence

- 34.1 million smokers in the US
- 8 million tobacco-related deaths annually worldwide-- 480,000 in the US
- 14% of Americans currently smoke
- 23% of Americans are former smokers
- 54% of those with SMI smoke
- Numbers of smokers are INCREASING
- 100 million people died in the last century from smoking related causes
- WHO anticipates 1 billion smokers worldwide will die from smoking related causes this century



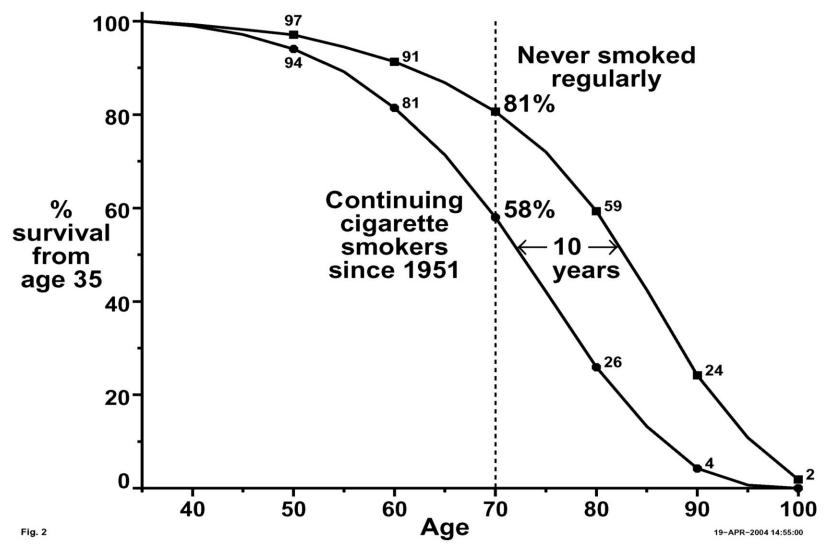
### U.S. Drug Related Deaths

- **Opioid overdoses** killed more than 92,000 people in 2020 and increasing.
- Over 95,000 **alcohol** related deaths per year and increasing.
- Over 480,000 **tobacco** related deaths per year and increasing.



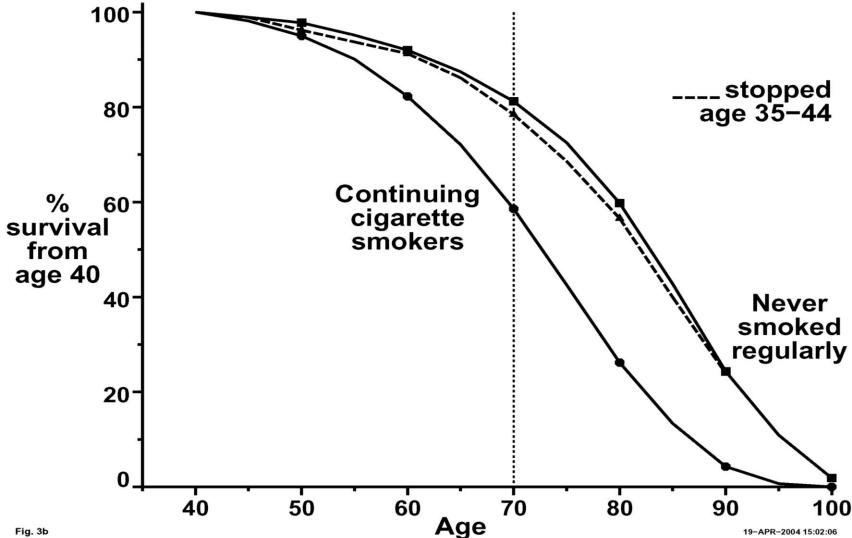
### **Smoking Kills**

UK male doctors born 1900–1930: continuing cigarette vs never smokers. 50-year follow-up of mortality, 1951–2001

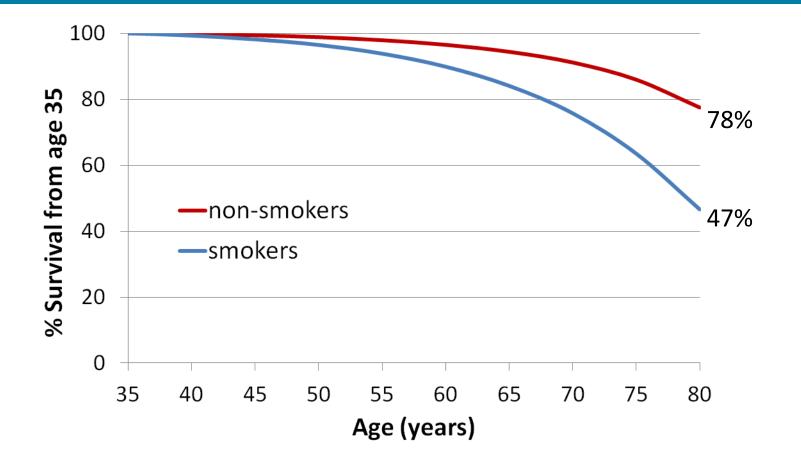


#### **Quitting Helps**

Effect of stopping smoking at age ~40 on survival from age 40



#### One Million Women Study: Effect of 3-fold difference in annual death rates on survival at ages 35-79

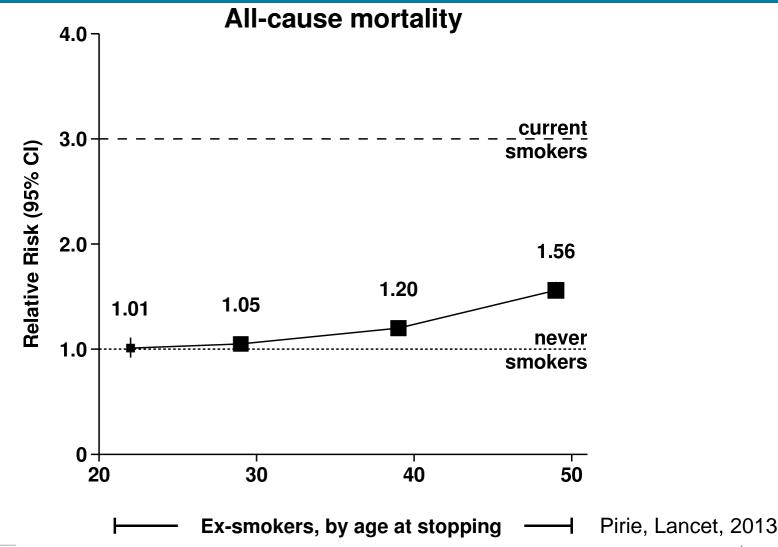


Adapted from the One Million Women Study Pirie, Peto, et al., Lancet 2013



### THE MILLION WOMEN STUDY

#### **Quitting by age 50 halves mortality**





MASSACHUSETTS



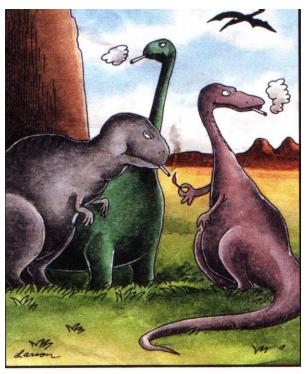
50 Years after the first Surgeon General's report of an association between smoking and cancer, adult smoking has declined 55% in the general US population.

Smoking prevalence among adults with SMI in the US today is 53%.

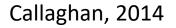
This is higher than in the US general population in 1964.

### Smoking-Related Mortality in Those with Psychiatric Disorders

- In those with one or more lifetime hospitalizations for schizophrenia, bipolar disorder, or MDD:
- HALF died from 1 of 19 diseases identified by CDC as causally linked to tobacco use



The real reason dinosaurs became extinct





## **Quitting Reduces**

- Death
- MI
- Stroke
- Progression of atherosclerosis
- Bronchitis
- Morbidity from Diabetes
- Cancer Risk
- Progression of COPD

#### META-ANALYSIS CONFIRMS: SMOKING CESSATION IMPROVES PSYCHIATRIC SYMPTOMS, QUALITY OF LIFE

• 26 studies



- Change in psychiatric symptoms was compared between continuing smokers and successful quitters
- **Depression, anxiety, stress and quality of life** improved among those who quit smoking significantly compared to those who continued smoking.
- It did not matter whether one had a pre-existing psychiatric diagnosis or not!!!
- Effect sizes comparable to those observed for antidepressant medications!!!

Smoking Cessation Is Associated with Improved Psychiatric Symptoms

Study	SE	Standard mean difference (95% Cl)	Weigh (%)	t Standard mean difference (95% CI)
Anxiety		(95% CI)		
Solomon 2006	0.19	: -	25	-0.06 (-0.42 to 0.30)
Dawkins 2009	0.25		20	-0.19 (-0.68 to 0.30)
McDermott 2013	0.13		29	-0.74 (-1.00 to -0.48)
Becona 2002	0.18		26	-0.37 (-0.72 to -0.02)
Total			100	-0.37 (-0.70 to -0.03)
Test for heterogeneity: $\tau^2=0$				
χ <sup>2</sup> -10.43, df-3, P-0.02, l <sup>2</sup>				
Test for overall effect: z=2.16	, P=0.03			
Depression				
Solomon 2006	0.19		9	0.01 (-0.35 to 0.37)
Berlin 2010	0.22		7	-0.30 (-0.72 to 0.12)
Blalock 2008	0.22		7	-0.58 (-1.00 to -0.16)
Dawkins 2009	0.25		5	-0.39 (-0.88 to 0.10)
Kahler 2011	0.21		7	-0.28 (-0.69 to 0.13)
Vazquez 1999	0.17		11	-0.12 (-0.44 to 0.20)
Busch 2011	0.19		9	-0.30 (-0.67 to 0.07)
Kahler 2002	0.20		8	-0.69 (-1.09 to-0.29)
Munafo 2008	0.09		21	-0.09 (-0.27 to 0.09)
Kinnunen 2006	0.11		17	-0.21 (-0.42 to 0.00)
Total		-	100	-0.25 (-0.37 to -0.12)
Test for heterogeneity: $\tau^2=0$				
χ <sup>2</sup> =12.83, df=9, P=0.17, l <sup>2</sup>				
Test for overall effect: z=3.89				
Mixed anxiety and depressi	on			
Blalock 2008	0.44		4	-0.21 (-1.07 to 0.65)
Kahler 2009	0.29		8	-0.64 (-1.22 to -0.06)
Steinberg 2011	0.14		35	-0.29 (-0.57 to -0.01)
Mino 2000	0.26			-0.44 (-0.95 to 0.07)
Chassin 2002	0.13			-0.23 (-0.48 to 0.02)
Total		-	100	-0.31 (-0.47 to -0.14)
Test for heterogeneity: $\tau^2=0$				
χ <sup>2</sup> =1.94, df=4, P=0.75, I <sup>2</sup> =				
Test for overall effect: z=3.61	, P(0.001			
Stress				
Manning 2005	0.17			-0.25 (-0.58 to 0.08)
Hajek 2010	0.09			-0.22 (-0.40 to -0.04)
Chassin 2002	0.13			-0.36 (-0.61 to -0.11)
Total		+	100	-0.27 (-0.40 to -0.13)
Test for heterogeneity: $\tau^2=0$				
χ <sup>2</sup> =0.77, df=2, P=0.68, I <sup>2</sup> =				
Test for overall effect: z=3.80	, P(0.001			
	-		1	
		avours Favour juitters smoke		

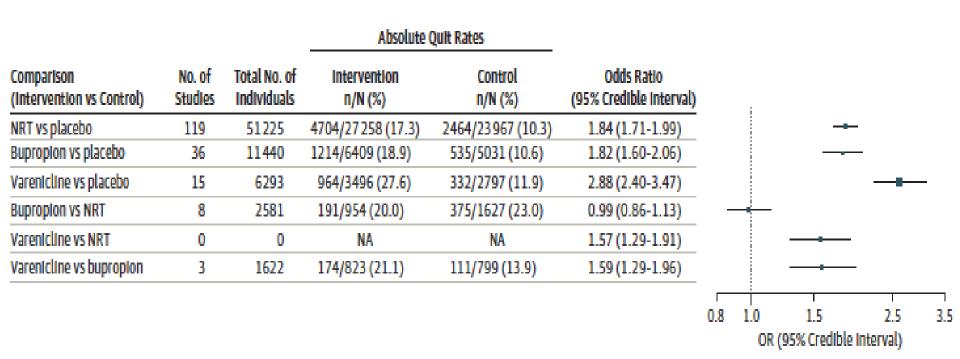
#### Taylor et al., BMJ, 2014

# Addiction to Nicotine: Mechanism and Therapeutic Targets

- •Nicotine binds to nicotinic cholinergic receptors ( $\alpha 4\beta 2$ ,  $\alpha 3\beta 4$ ,  $\alpha 7$ ) > neurotransmitter release
  - –<u>dopamine, glutamate, GABA</u> important in develop of nicotine dependence
  - –<u>Corticotropin-releasing factor</u> contributes to nicotine withdrawal
- •Agonist therapies target nicotinic receptors: NRT, Varenicline
- •Bupropion increases levels of dopamine and norepinephrine, some nicotinic receptor-blocking activity

#### Cessation Works: Pharmacotherapy + Behavioral Therapy Doubles to Triples Abstinence Rates

#### Figure. Odds Ratios for Smoking Abstinence of 6 Months or More



Cahill et al., JAMA 2014



### Addiction Treatment Works: Expect and Treat Relapses

For tobacco dependence: average of 5 attempts at abstinence before long-term abstinence achieved

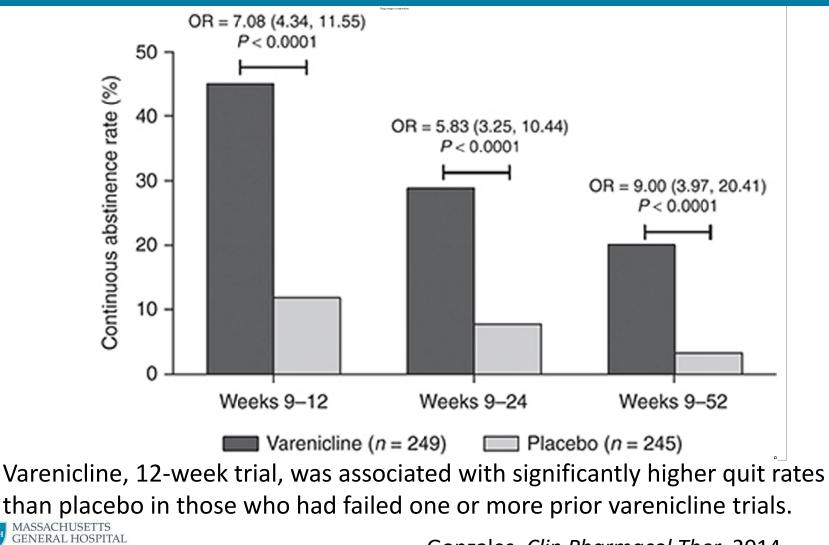
Treatments double to triple abstinence rates and are Underutilized!

#### New clinical practice guidelines: Treat all smokers

Offer pharmacotherapy and behavioral support to all smokers willing to accept such treatment, not just those who report being 'ready to quit'

Barua, et al., Am Col Cardiology 2018

#### Repeat Cessation Attempts Are Effective

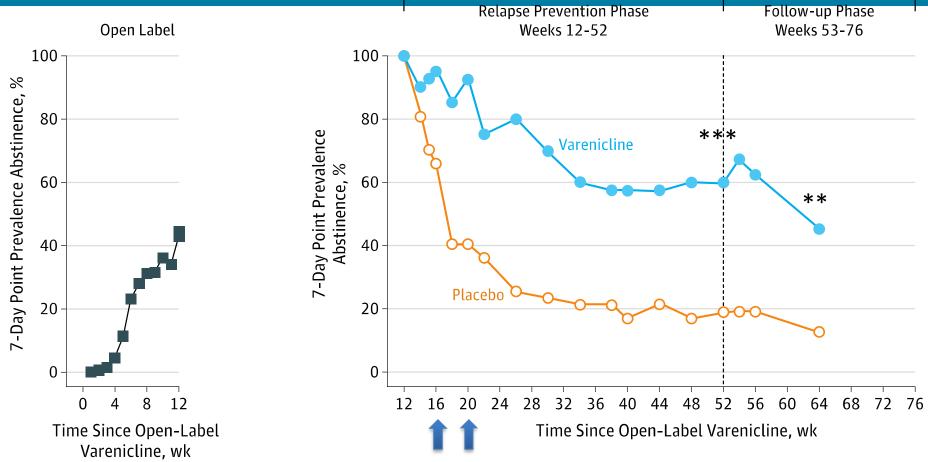


PSYCHIATRY ACADEMY

Gonzales, Clin Pharmacol Ther, 2014

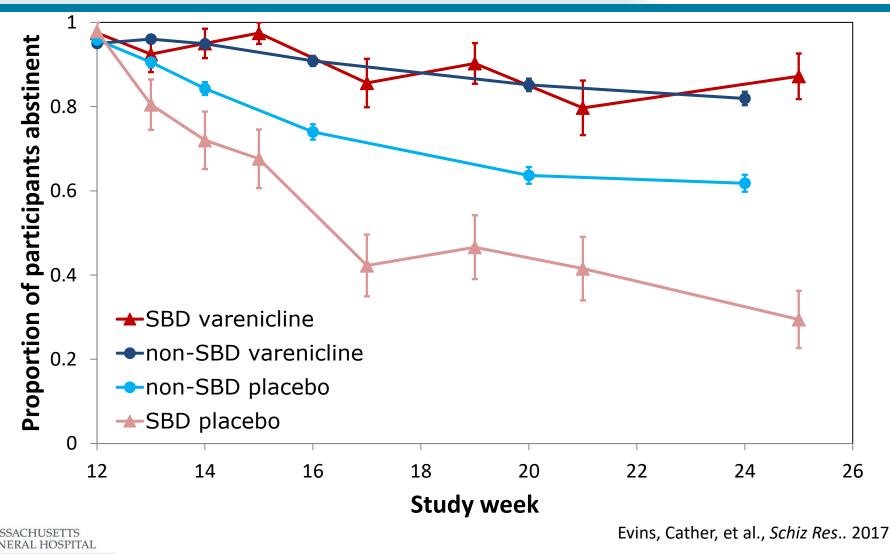
### Varenicline Maintenance Treatment for One Year Triples Abstinence Rates at One Year in

#### **Smokers with Schizophrenia**



43% attained abstinence with 12 weeks open label varenicline MASSACHUSETTS GENERAL HOSPITAL PSYCHIATRY ACADEMY A3% attained abstinence with 12 weeks open label varenicline and were randomized to 40 weeks varenicline or placebo + Group CBT Evins, Cather, et al., JAMA. 2014 WWW.mghcme.org

#### Maintenance Tx Normalizes the Relapse Curve for Smokers with Schizophrenia and Bipolar Disorder



PSYCHIATRY ACADEMY

#### Smoking Cessation Tx Recommended but Underutilized, Particularly in Psychosis

- 2010 PORT guidelines for tx of schizophrenia recommend physician advice to quit and medication with bupropion with or without NRT for all smokers with schizophrenia.
- Smoking rates are not declining in those with psychosis
- Psychiatrists rarely offer counseling to quit smoking. In one study, only 12.4% of smoking patients were advised to quit.
- Smokers with SMI are even less likely to receive a medication to help them to quit.
- Varenicline especially underutilized in smokers with psychosis

#### **Treatment is effective in the long run and is under-prescribed**

Buchanan et al., 2010; Himloch and Daumit, 2003; Thorndike et al., 2001; Huang et al., 2014; Cook 2014 www.mghcme.org

#### **EAGLES** Trial

- Compare risk of clinically significant neuropsychiatric AEs & efficacy of varenicline, bupropion, NRT patch, placebo
- >8000 Smokers aged 18 to 75 years; ≥10 cigs/day
- > 4000 smokers, no lifetime psychiatric diagnosis
- > 4000 smokers, 1+ clinically stable, lifetime diagnoses

Mood Disorders	Major depressive disorder (MDD), bipolar I, bipolar II
Anxiety Disorders	Panic disorder with or without agoraphobia, post- traumatic stress disorder, obsessive-compulsive disorder, social phobia, generalized anxiety disorder
Psychotic Disorders	Schizophrenia, schizoaffective disorder

Anthenelli RM, Benowitz NL, West R, St. Aubin L, McRae T, Lawrence D, Ascher J, Russ C, Krishen A, Evins AE. Effects of varenicline and bupropion in smokers with and without psychiatric disorders. *Lancet*. 2016 Apr 22

### Clinical Characteristics of the EAGLES Psychiatric Cohort

- Included:
  - Stable but symptomatic
  - Half on psychotropic medication at baseline (>95% with psychotic disorder)
  - Half with major depressive disorder had recurrent depression
  - One third had a second psychiatric diagnosis / comorbidity
  - One fourth had a prior substance use disorder
  - One eighth had made a prior suicide attempt
  - Excluded those with active self-injurious behaviors, imminent suicide risk, or active SUD



### **Primary Endpoint**: Composite Neuropsychiatric Adverse Event

**Primary Safety Endpoint:** Percent of subjects reporting worsening or new onset of one or more of the following during treatment and up to 30 days after last dose:

≥1 "severe" AE of:							
Anxiety	Depression	Feeling abnormal	Hostility				
And/or ≥1 "moderate" or "severe" AE of:							
Agitation	Aggression	Delusions	Hallucinations				
Homicidal ideation	Mania	Panic	Paranoia				
Psychosis	Suicidal ideation	Suicidal behavior	Completed suicide				
AE, adverse event; NPS, neu	ropsychiatric						

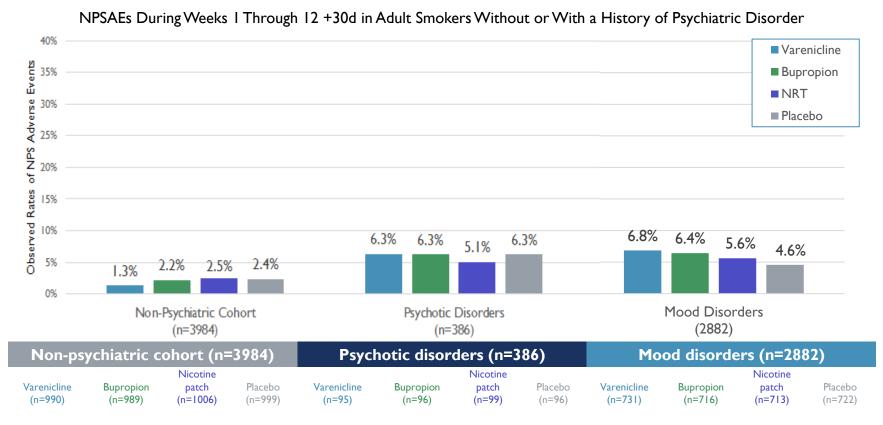
Designed in collaboration with FDA and EMA to be broad and capture an array of events

#### Severity assessment

Moderate = interferes to some extent with subject's usual function Severe = interferes significantly with subject's usual function

## SAFETY

#### Varenicline, Bupropion Do NOT Increase NPSAEs



Evins et al., J Clin Psychopharm 2019; Anthenellii et al., Lancet 2016

Neuropsychiatric (NPS) safety data based on EAGLES (Evaluating Adverse Events in a Global Smoking Cessation Study)<sup>1,2</sup>, an FDA required trial to evaluate NPS safety in over 8000 smokers with and without a psychotic, anxiety or mood disorder<sup>+</sup>

## SAFETY

Neuropsychiatric (NPS) safety data based on EAGLES (Evaluating Adverse Events in a Global Smoking Cessation Study)<sup>1,2</sup>, an FDA required trial to evaluate NPS safety in over 8000 smokers with and without a psychotic, anxiety or mood disorder<sup>+</sup>

EAGLES provides data that can be used to counsel smokers on the likelihood of experiencing a moderate to severe NPS adverse events during a smoking cessation attempt.

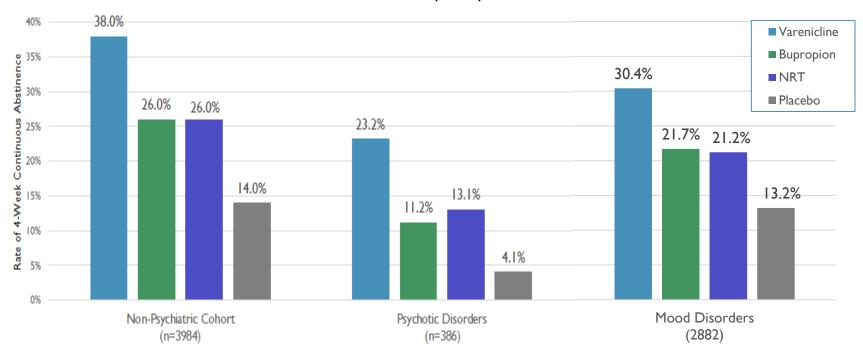
Risk of NPS AEs is independent of treatment

~2% NPS AE rate in smokers without mental illness

- ~5-7% NPS AE rate in smokers with mental illness
- NPS AE rates during a cessation attempt are not different across active treatments or placebo
- No pattern of NPS AEs in the most worrisome NPS AEs
- No psychiatric subgroup appears to be at particularly increased risk

## **EFFICACY** Comparative efficacy data based on EAGLES<sup>2</sup>

#### Varenicline was superior to bupropion, NRT and placebo, while bupropion and NRT were superior to placebo for biochemically-confirmed tobacco abstinence.‡



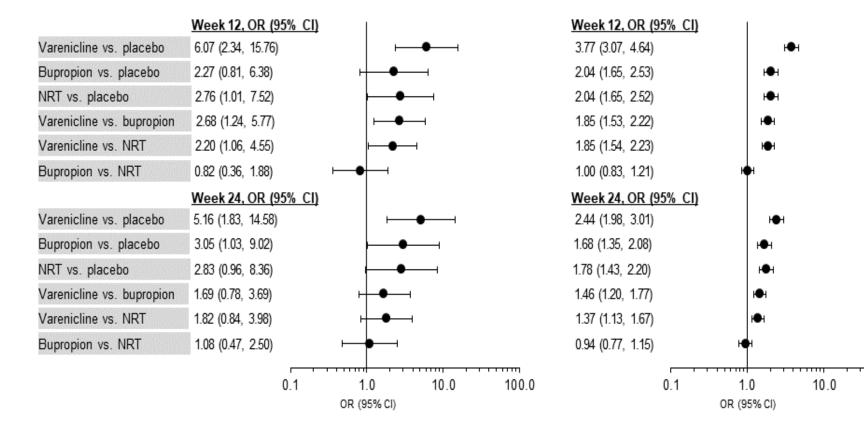
Continuous Abstinence During Weeks 9 Through 12 in Adult Smokers Without or With a History of Psychiatric Disorder

<sup>||</sup>"N" and analyses based on all-randomized populations in the EAGLES trial published in Antheneli et al., *The Lancet* (2016) and Evins et al., J Clin Psychopharm 2019

### **EFFICACY** Comparative efficacy data based on EAGLES<sup>2</sup>

Schizophrenia spectrum disorders subcohort

No psychiatric disorders cohort

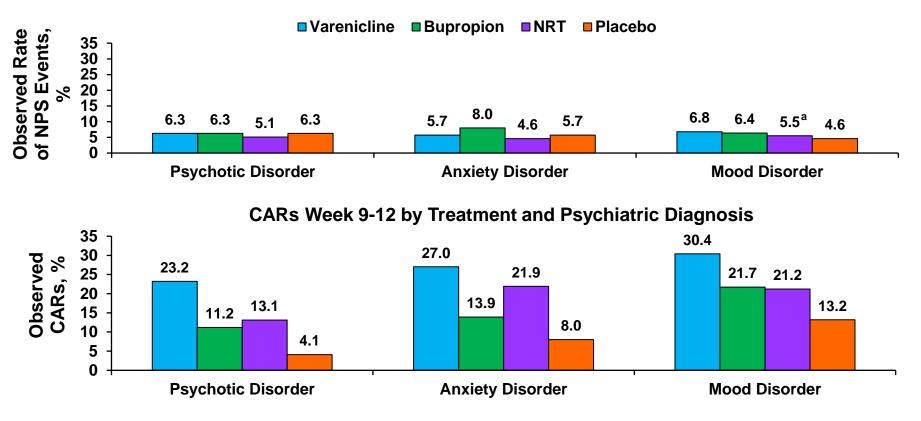


Evins et al., Psych Services 2020

100.0

#### Risk / Benefit: EAGLES Allows Comparison of Neuropsychiatric Safety and Efficacy in Those with Psychiatric Illness

#### Primary NPS Composite Safety Endpoints by Treatment for Those with Primary Psychotic, Anxiety and Mood Disorders



Adapted from Evins, et al., J Clin Psychopharm 2019

a. One additional participant (NRT group/mood subcohort) who reported suicide ideation was identified after clinical database lock and was not included in the analysis www.mghcme.org

## **EFFICACY** Comparative efficacy data based on EAGLES<sup>2</sup>

Varenicline was superior to bupropion, NRT and placebo, while bupropion and NRT were superior to placebo for biochemically-confirmed tobacco abstinence.‡

#### "FDA removes warnings on smoking cessation medication"

Pharmacy Times, December 16, 2016

FDA and EMA removed boxed warnings for varenicline and bupropion in 2016 based on results of EAGLES, a required, randomized, double-blind, triple dummy, active-and placebo-controlled clinical trial conducted by Pfizer in collaboration with GlaxoSmithKline, designed in consultation with the FDA and the European Medicines Agency (EMA). It is the largest smoking cessation clinical trial ever conducted and the largest samples of smokers with psychotic, anxiety, and mood disorders ever conducted.

### EAGLES is a Confirmatory Trial for Efficacy

- Efficacy conclusions replicate and extend findings from smaller trials and meta-analyses in those with and without mental illness
- The efficacy data are clear

Varenicline > bupropion and nicotine patch > placebo

• Agreement with overall, growing body of evidence, raising confidence in the findings



### EAGLES is a Landmark Study of Clinical and Public Health Importance

- The EAGLES trial is the first:
  - To compare safety and efficacy of all 3 FDA approved smoking cessation therapies in large samples of patients with and without a history of psychiatric disorder
  - To allow for comparison of safety and efficacy of smoking cessation aids in smokers with different mental illnesses



### Neuropsychiatric Adverse Events During Smoking Cessation Are Independent of Treatment

- NPS AEs are seen in trials regardless of treatment
- Clinicians who prescribe a treatment and observe a NPS AE likely attribute this AE to the treatment.
- This happened in our large maintenance treatment trial of varenicline, in trials of bupropion, and in clinical practice.



### Why Might There be Significant NPS AEs Among Smokers, Independent of Treatment (and Abstinence)?

- Smoking is an addiction; like all drug addictions, there are:
  - Well documented brain changes
  - Increased neuropsychiatric events, e.g. suicide
  - Suicide risk reduced in smokers who quit
- People with psychiatric illness are more likely to smoke
- Attempts to quit smoking are not risk free, with or without pharmacologic support and independent of abstinence
  - Well replicated in smokers with history of depression

Volkow et al., Am. J. Psych, 1999; Fehr et al., Am J Psych 2008; Li, et al., J Psych Res 2012; Berlin et al., NTR 2011; Brown 1996; Tsoh, et al., Am J Psych 2000; Torres, et al., Psychol Med 2010; Evins, et al., JAMA 2014, Evins et al.. Psychol Med 2017



### Varenicline Safety in 17 RCTs and 5 Large Observational Studies

- Pooled Analysis of ALL Psychiatric Adverse Effects in 17 RCT's of Varenicline Varenicline increased incidence of nausea but not psychiatric adverse events
- Varenicline increased abstinence rates by 124% vs. placebo and 22% vs. bupropion
- Having a psychiatric illness increased the risk for psychiatric adverse events in smokers trying to quit equally in those assigned to varenicline and placebo
- In a large observational study in 35,800 outpatients trying to quit smoking, there were fewer psychiatric adverse events in those prescribed varenicline than those prescribed NRT

Results replicated now in multiple studies in different practice populations: DoD, VA, UK NHS

Gibbons and Mann 2013; Tonstad et al., 2010; Kotz et al., 2015

### Implication of EAGLES: Offer Treatment to ALL Smokers, Especially Those with SMI

- Confirms NPS safety and efficacy of smoking cessation treatments for smokers with mental illness, a group that is:
  - More likely to smoke, to smoke heavily, and be dependent
  - Less likely to quit without a cessation aid
  - More likely to relapse after discontinuation of cessation aids
  - More likely to benefit from maintenance treatment
- Smokers with mental illness are less likely to receive a pharmacotherapeutic cessation aid from a medical provider
  - This contributes to the 25 year mortality gap in those with mental illness from diseases causally related to smoking
  - 28 year mortality gap for those with schizophrenia

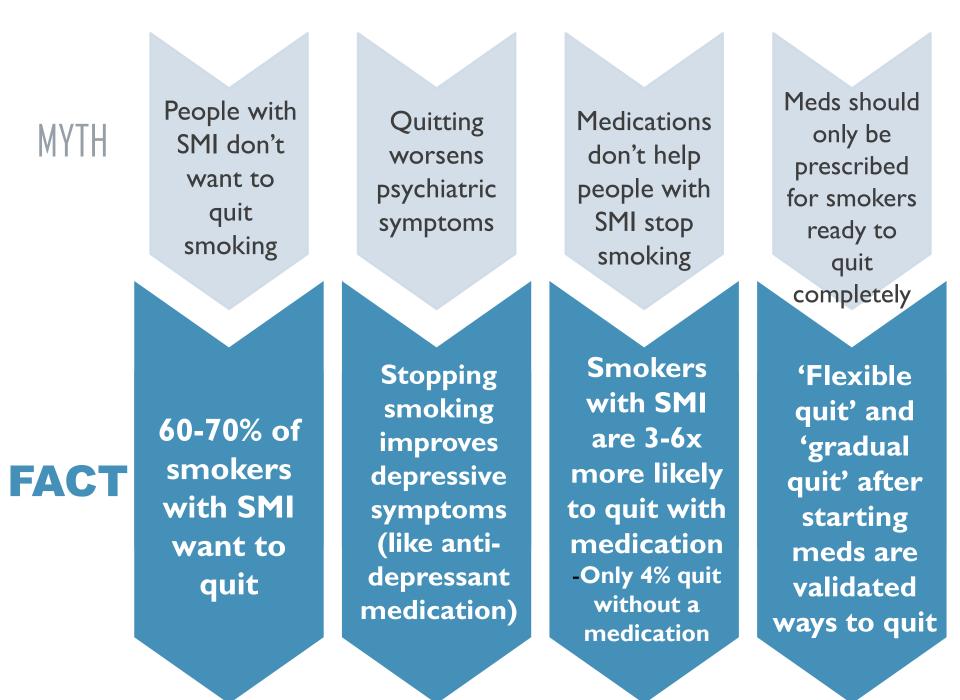


## **Risk/Benefit Considerations**

- Clinicians **overestimate** the risk of NPS AEs with varenicline and bupropion, particularly in those with psychotic illnesses
- And underestimate the benefit of varenicline and bupropion on improving quit rates
- It is imperative we find ways to increase use of the most effective smoking cessation treatment for our patients who try time and again to quit smoking



Huang et al., BMC Public Healthm2014org



2018 American College of Cardiology Expert Consensus Decision Pathway on Tobacco Cessation Treatment

- Recommend/Prescribe Pharmacotherapeutic Cessation Aid to ALL Smokers, not just those who state they are ready to quit.
- Prescribe to all smokers willing to start a pharmacotherapy
- Follow up in 2-4 weeks for tolerability
- For those who decline, continuous engagement to quit at each clinic visit



Barua et al., JACC 2018

#### THREE WAYS TO QUIT SMOKING: All Start with Smoking Cessation Medication



#### FIXED QUIT

For people who want to quit smoking in a week

- Set a target quit date I week after starting smoking cessation med
- Can keep smoking for the first week while they prepare to quit
- Take smoking cessation medication for 12-24 weeks



#### FLEXIBLE QUIT

- \*\*Recommended\*\*
- Start taking smoking cessation medication and pick a quit date 8 to 35 days after starting treatment
- Can keep smoking for up to a month on smoking cessation medication while preparing to quit
- Take smoking cessation meds for 12-24 weeks

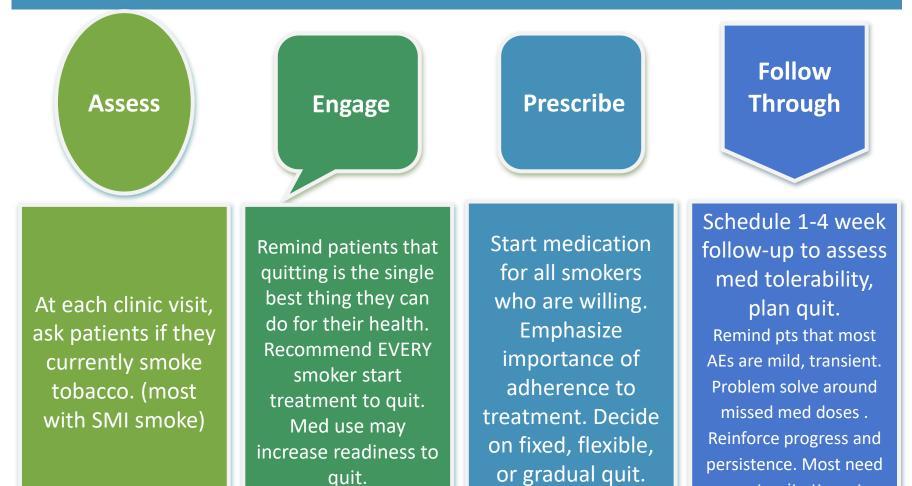


#### GRADUAL QUIT

For people not able/willing to quit abruptly

Start taking smoking
cessation med and reduce
smoking by 50% over 4
wks, by another 50% in the
next 4 wks, etc. Goal of
quitting by 12 weeks.
Continue smoking
cessation med for an
additional 12 weeks, for a
total of 24 weeks

# **STEPS TO ADDRESS SMOKING**



repeat quit attempts.

# How To:

ARENICLINE

Available as 0.5 and 1.0 mg tabs

- 0.5 mg/d at hs x 3 d
- 0.5 mg bid x 4 d
- I.0 mg bid x II weeks
- Additional 3-9 months Tx recommended in those who achieve abstinence
- I2-month safety data published: well tolerated

Renal excretion, used in chronic renal disease with dose reduction

No significant drug-drug interactions or effect on cytochrome enzymes

Nausea, headache, insomnia, and vivid dreams are common

# How To:

Ω **DUAL NICOTINI** П L U. 

	NICOTINE PATCH +	NICOTINE GUM, LOZENGE	
	Dosing: 21 mg/d x 4-6 weeks then - 14 mg/d x 4 weeks then - 7 mg/d x 3-4 weeks	Dosing: up to 20 mg/d x 4-5 weeks then - Up to 14 mg/d x 4 weeks then - Up to 10 mg/d x 3-4 weeks	
	Apply one new patch every 24 hours (preferable am) to dry, clean	Do not chew, break, crush, or swallow whole	
	skin Move site with each new patch to avoid skin irritation	Gum: Chew a few times then 'park it' between cheek and gum	
		Move around mouth until it melts (lozenge) or loses flavor (gum)	
	Remove patch at night if bothered by insomnia or vivid dreams	Do not eat or drink for 15 minutes before or during use	

# How To:

**JPROPION** 

Dosing: 150mg QD x 3 days, then 150mg BD

#### Insomnia common

Meta-analysis of 182 studies with 70,000 smokers:<sup>9</sup> Varenicline triples chances of quitting vs. placebo and increases odds of quitting by 50% over NRT and bupropion

NRT and bupropion nearly **double** odds of quitting vs. placebo (80% increase)

## Varenicline and Bupropion Improved Health Related Quality of Life

- Treatment with Varenicline (n=696) and Bupropion (n=671) Significantly Improved Self Rated Quality of Life Over Placebo (n=685) at 12, 24, and 52 Weeks
- Significant positive association between smoking cessation and self rating of vitality, self-control, anxiety, and overall mental health profile
- Replication of several studies demonstrating reduced self report of anxiety after smoking cessation...

Hays et al., 2010

www.mghcme.org

## **Combination Pharmacotherapy for Nicotine Dependence**

May improve abstinence rates

For smokers who have relapsed after treatment with single agent, consider maintenance treatment or combination treatment:

- NRT: long acting (patch) + short acting (gum, inhaler or nasal spray) + CBT
- Bupropion 150 mg bid + NRT + CBT
- Varenicline + NRT

### **Behavioral Interventions**

- Current guidelines recommend behavioral tx + pharmacotherapy
  - Motivational enhancement
  - Relapse prevention
  - Partner support
- Guidelines are based on several large meta-analyses of controlled trials
- Telephone counseling provides a modest benefit in quit rates vs minimal intervention, patient follow through can be low
  - www.trytostop.org or 1-800-TRY-TO-STOP
- Physical exercise can decrease cravings and attenuate weight gain

# **Nicotine Withdrawal Syndrome**

- Peaks in 4 days
- Lasts for several weeks
- Can be severe, not life threatening
  - Anxiety
  - Awakening during sleep
  - Depression
  - Difficulty concentrating
  - Impatience
  - Irritability/anger
  - Restlessness
  - Decreased heart rate
  - Weight gain



### Tobacco Abstinence: Effects on Metabolism

- Smoking speeds hepatic metabolism of many medications
- Serum concentrations of medications that are stable in smokers may rise following abstinence, allowing lower doses
- CYP 1A1, 1A2, and 2E1
  - Abstinence associated with 30-42% reduction in 1A2 activity over the first 1-3 days of abstinence
  - Therapeutic drug monitoring and 10% dose reduction has been recommended
- Take care when prescribing bupropion to those on clozapine because of additive seizure risk

Seppala NH, et al., 1999; Desai HD, et al., 2001; Faber & Fuhr, 2004.

### Summary – Nicotine Dependence

- Give physician advice to quit smoking to all smokers at every visit
- Prescribe/Offer Pharmacotherapeutic Cessation Aid to All Smokers
  - (as for Other Chronic Illnesses with Behavioral Component eg. Type 2 Diabetes, Hypertension)
- Choose a Quit Plan: Fixed, Flexible, or Gradual.
- Develop a "quit day" plan, refer or review coping skills, build in selfrewards, and provide written cues to reinforce abstinence
- Long-term pharmacotherapy may be warranted, both to sustain abstinence and to improve symptoms

# **Cocaine Use Disorder**

- Major epidemic since 1980
- Availability of cheap, high-potency drug
- Includes freebase/crack
- 30 million in US have used cocaine
- < 20% become regular users
- 17% risk of dependence
- Lacing common
  - Levamisole in up to 80% of samples in some locations
  - 3-13% risk of agranulocytosis with sustained exposure

#### Pharmacology of Cocaine Use Disorder

- Dopamine stimulation of neurons in nucleus accumbens normally limited by dopamine reuptake
- Cocaine blocks dopamine reuptake
- Assoc. with excessive dopamine stimulation in reward system of brain - "HIGH"
- Also assoc. with depletion of dopamine in nerve terminals of dopaminergic neurons "LOW"
- Compensatory down-regulation of post-synaptic dopamine receptors
  - Protracted syndrome of refractoriness to reward

#### **Cocaine Use Patterns**

- Binge symptoms:
  - Intense euphoria
  - Paranoia, anxiety, dysphoria, tremor, hyperactivity
  - Panic attacks, depression, mania
- Withdrawal:
  - Onset: <24 hrs, peak: 2-4 days</p>
  - Duration: 7-10 days
  - Protracted depression: 1-3 months
  - Intense cravings: 1-3 months

## **Treating Acute Cocaine Intoxication**

- Acute cocaine intoxication:
  - Onset: seconds
  - Duration: 30-60 min
  - Dysphoria: within hours
  - Recovery: < 48 hrs</p>
  - OD requires life support, airway
- Cocaine delusional disorder
  - Diazepam for agitation
  - Antipsychotics for delusions
- Hospitalize if suicidal or delusional

## **Treating Cocaine Withdrawal**

- Pharmacotherapy not required in mild withdrawal states
- For severe cocaine withdrawal:
  - <u>Amantadine</u> indirect dopamine agonist, increases dopamine levels
  - <u>Propranolol</u> B-adrenergic blocker reduces anxiety / severe adrenergic symptoms - 1 mg IV q min, up to 8 min
- Seizures: IV diazepam

#### Treating Cocaine Use Disorder: Pharmacotherapy not Standard of Care

#### **Relapse prevention: Pharmacotherapy**

- Disulfiram: effective in 3 trials
  - Inhibits DA-beta hydroxylase
  - Reduced craving & relapse
  - Overall evidence does NOT support efficacy
- <u>Baclofen:</u> GABA-B agonist: 20 mg tid
- <u>Topiramate:</u> increases GABA & inhibits glutamate:
  - 25 mg po qd, slowly increase to 200 mg qd (Kampman, 2004)
  - Overall evidence does NOT support efficacy
- <u>Modafinil:</u> enhances glutamate levels: 200-400 mg po qd

#### **Overall:**

- Disulfiram: evidence not supportive
- Topiramate, other anticonvulsants: evidence not supportive
- Anticonvulsants: evidence not supportive
- Antipsychotics: evidence not supportive

#### **Treating Cocaine Use Disorder**

<u>Relapse prevention: Psychotherapy</u>

- Contingency Management
- Manual-guided CBT
- 12-step facilitation
- Individual plus group therapy
- Behavioral reinforcement:
  - Urine testing with contingencies
  - Restrict access to money & friends
- High-intensity support to disrupt binge cycles

#### **Treating Cocaine Use Disorder**

As with any substance use disorder, treat anxiety and depressive symptoms in those suspected of having an independent mood or anxiety disorder, especially if these symptoms appear to be interfering with attainment of abstinence

#### Co-morbid depression:

- SSRIs effective if depressed
- "May" also reduce cocaine use
- Avoid TCAs, may be associated with cardiac arrhythmia when combined with cocaine

#### Co-morbid bipolar disorder: No adequate med trials

Consider combination therapy if rapid cycling