Anti-Obesity Medications: Inside and Outside the Label

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Disclosures

Company	Disease State/Topic	Role	
Novo Nordisk	Obesity	Promotional speaker	
		Advisory board	
Acella	Thyroid	Advisory board	
		Promotional speaker	
Currax	Obesity	Advisory board	
		Promotional speaker	
Lilly	Obesity and Sleep apnea	Advisory Board	
		Promotional speaker	
BI	Obesity	Advisory Board	
WW	Obesity	Advisory board	

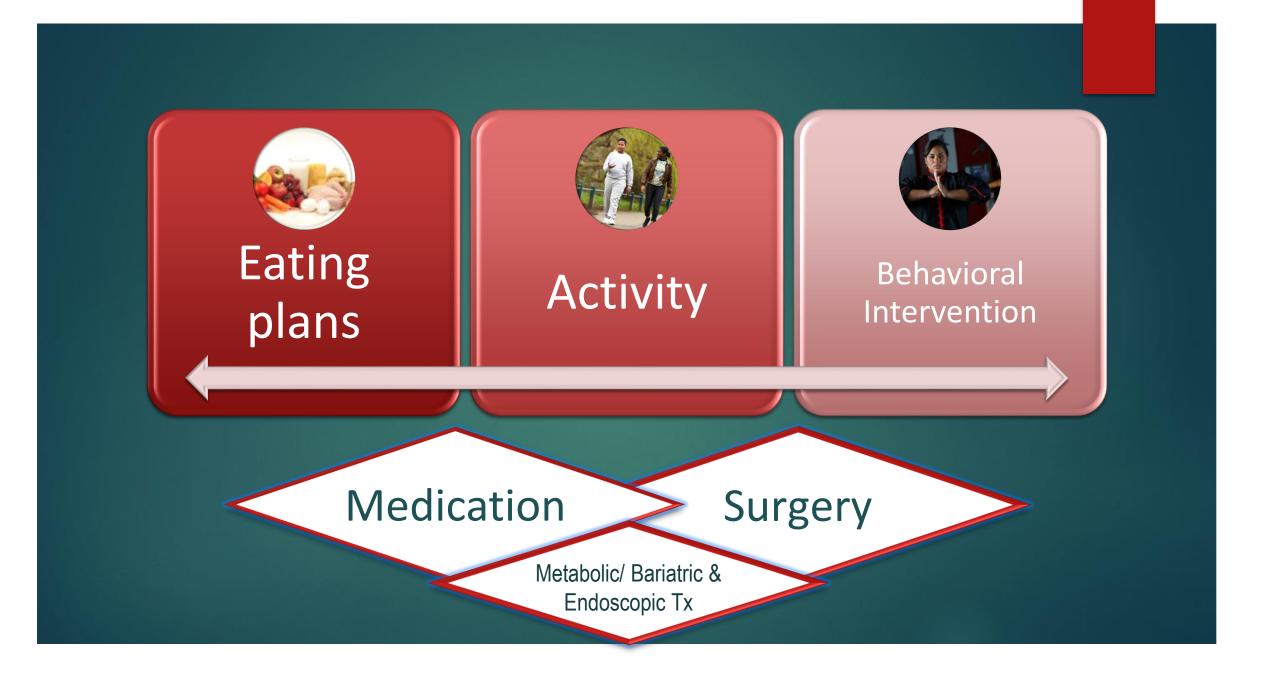
► All relevant financial relationships have been mitigated.

Objectives

Recognize	Recognize the role of pharmacotherapy in obesity treatment.
Evaluate	Evaluate FDA approved medications for the pharmacologic management of chronic disease of obesity.
Identify	Identify available pharmacotherapeutics that can be utilized to support treatment of chronic disease of obesity that are not labeled for obesity by the FDA 20 minutes

Recognize

Recognize the Role of Pharmacotherapy in Obesity Treatment



Case Study



moge: © Obesity Action Coquition

Meet Mariah

48-year-old woman presents today to discuss possible assistance with her weight.

Mariah's Medical History

PMH: Obstructive sleep apnea (OSA) – on BiPAP **Prediabetes** PCOS Seasonal allergies Stage 2 obesity Seizure disorder in teens post MVA, no seizures in 30 years

Medications: Metformin 1000mg XR Certrizine 10mg daily during allergy season Diphenhydramine 25mg nightly during allergy season Multivitamin once daily Pregnancy prevention: IUD NKDA

Mariah's Medical History

Preventative screening:

 Mammogram – 9 months ago

Surgical History: None

Family History:

 DM (mother and older sister) and all are "heavy"; Father CVD with history of MI, no history of cancer

Social History:

- Divorced, one daughter in college
- Works outside the house as an accountant
- ETOH 1x/week (6-ounce glass when out with friends)
- No history of tobacco use or drug abuse



Mariah's First Visit for Obesity

- Vital Signs:
 - 5'4"; 212 Lbs; BP 118/62 mmHg; HR 82 Bpm; RR 16; pOx 98%
- BMI: 36.30 Kg/M²
- Waist Circumference: 45"
- Screening Tools: PHQ-9 (0), BED7 (Neg), PAR-7

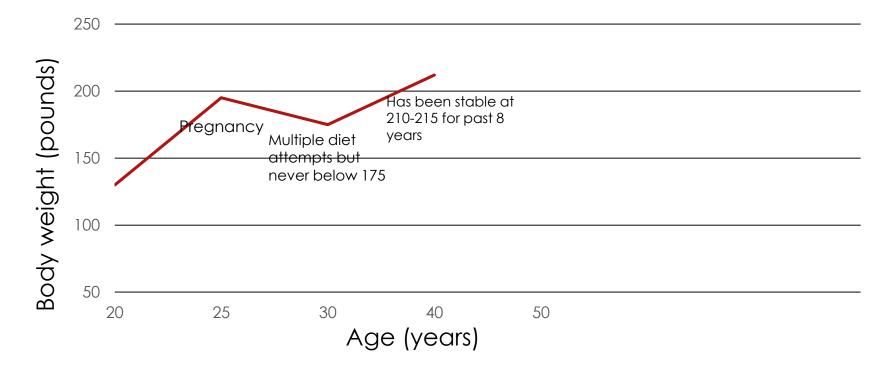
	Yes	No
Has your NP or healthcare provider said you have a heart condition or high blood pressure?		x
Do you feel pain in your chest at rest, during your daily activities of living or when you do physical activity?		x
Do you lose balance because of dizziness, or have you lost consciousness in the last 12 months (answer no if your dizziness was associated with over-breathing, including during vigorous exercise)?		x
Have you ever been diagnosed with another chronic medical condition (other than heart disease or high blood pressure)? Please list conditions here: Depression, Sleep apnea	х	
Are you currently taking prescribed medications for a chronic medical condition? Please list conditions and medications here:		x
Do you currently have (or have had within the past 12 months) a bone, joint, or soft tissue (muscle, ligament, or tendon) problem that could be made worse by becoming more physically active? Please answer no if you had a problem in the past, but it does not limit your current ability to be physically active. Please list conditions here:		x
Has your healthcare provider ever said that you should only do medically supervised physical activity?		x

Form adapted from PAR 7 by Angela Golden for use at NP Obesity Treatment Center.

BED7, Binge Eating Disorder; PAR-7, Physical Activity Readiness. PHQ, Patient Health Questionnaire; STOP-BANG, snoring, tiredness, observed apnea, pressure, BMI, age, neck circumference, and gender.

Body Weight Graph

Use this graph to chart live events, health conditions, times of stress, and other factors that have influenced your weight



Polling Question

Are any of Mariah's medications obesogenic?

- a) Metformin 1000mg XR
- b) Certrizine 10mg daily
- c) Diphenhydramine 25mg
- d) Multivitamin once daily
- e) None of the above

Obesogenic Medications and Alternatives

Common Obesogenic Medications and Alternatives

Medication class	Obesogenic medications	Mechanism of weight gain	Alternatives
Anticonvulsants	Carbamazepine Gabapentin Pregabalin Valproic acid	Hypothalamic mediated increase in appetite and decrease in energy expenditure	Duloxetine, topiramate, zonisamide
Antidepressants/ antianxiety medications	Amitriptyline Fluoxetine Mirtazapine Nortriptyline Paroxetine	Appetite increase stimulated via serotonergic pathways	Sertraline, citalopram, escitalopram, bupropion
Antihistamines	Diphenhydramine Fexofenadine Hydroxyzine Cetirizine	Increase appetite, alter body weight regulation	Loratidine
Antipsychotics	Olanzapine Quetiapine Risperidone Ziprasidone	Increase orexigenic and decrease anorexigenic neuropeptide expression in hypothalamus	Cariprazine, aripiprazole
	Thiazolidinediones	Act as insulin sensitizers, cause water retention	

IUD = intrauterine device; MoA = mechanism of action; SGLT2 = sodium glucose cotransporter 2. Modified from Desalermos A, et al. *Obesity (Silver Spring)*. 2019;27:716-723.

Common Obesogenic Medications and Alternatives

Medication class	Obesogenic medications	Mechanism of weight gain	Alternatives
Beta blockers	Atenolol Metoprolol propranolol	Inhibit sympathetic tone, decrease lipolysis, reduce exercise tolerance, increase fatigue, and reduce resting energy expenditure	Carvedilol; limited given MoA; assess risk vs benefit
Corticosteroids and hormones	Medroxyprogesterone Oral contraceptives Prednisone	Alters energy intake and expenditure of the human body	Progesterone IUD, copper IUD
Diabetes medications	Insulin	Anabolic and adipogenic hormone, decreases daily energy expenditure	GLP-1, SGLT2, metformin
	Sulfonylureas	Increase secretion of insulin and cause water retention	
	Thiazolidinediones	Act as insulin sensitizers, cause water retention	

IUD = intrauterine device; MoA = mechanism of action; SGLT2 = sodium glucose cotransporter 2.

Modified from Desalermos A, et al. *Obesity (Silver Spring)*. 2019;27:716-723.

First Visit for Obesity

- Most recent labs: Triglycerides 263 mg/dl; TC 228 mg/dl; LDL 132 mg/dl; HDL 36 mg/dl; Vitamin D 18 ng/ml
- Fasting insulin 16 mlu/L; glucose 88 mg/dl; HOMA IR 4.5
- Obesity-related complications: prediabetes, PCOS
- Obesity comorbidities: OSA
- Obesogenic medication: certrizine, diphenhydramine
- Staging of obesity
 - ▶ WHO Class 2
 - ► AACE/ACE Stage 2

AACE, American Association of Clinical Endocrinologists; ACE, American College of Endocrinology; HDL, high-density lipoprotein; HOMA-IR, homeostatic model assessment of insulin resistance; LDL, low-density lipoprotein; TC, total cholesterol; WHO, World Health Organization.



Evaluate

Evaluate FDA approved medications for the pharmacologic management of chronic disease of obesity.

Pharmacologic Therapy

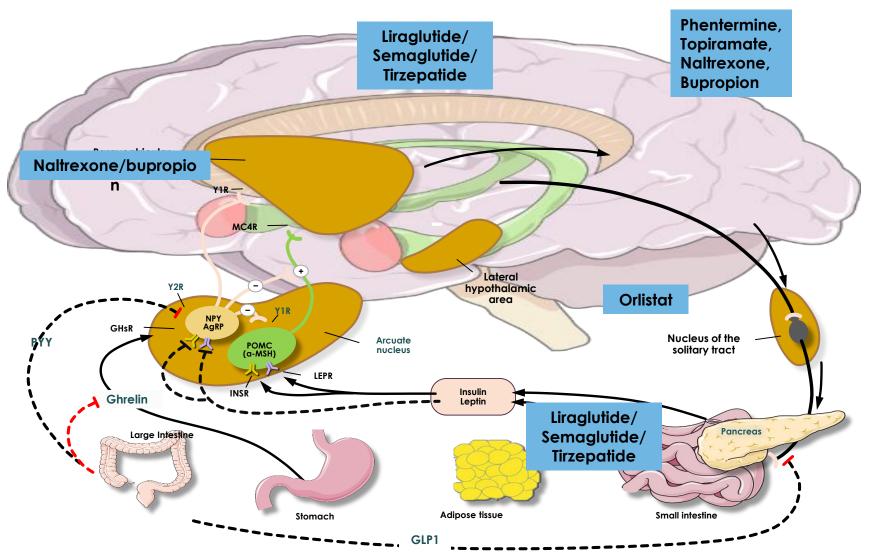
Therapy Options, Factors to Consider When Selecting Therapy, and Efficacy/Safety Evidence

Why Use Medication With Obesity Treatment?

- Therapy aimed at the dysregulated weight regulated biology
- Metabolic adaptation difficult to treat with diet and behavior modification alone
- Anti-obesity medications can assist in managing the > 200 obesity induced complications/comorbidities



Medication Mechanisms



a-MSH, a-melanocyte-stimulating hormone; GHsR, growth hormone secretagogue receptor; INSR, insulin receptor; LEPR, leptin receptor; MC4, melanocortin-4 receptor; POMC, pro-opiomelanocortin; Y1R, NPY Y1 receptor; Y2R, NPY Y2 receptor. Apovian CM, et al. J Clin Endocrinol Metab. 2015;100(2):342-362. Servier Medical Art by Servier is licensed under a <u>Creative Commons Attribution 3.0 Unported License</u>

Polling Question

What percentage of weight reduction should be the first therapeutic goal?

- a. <5%
- b. 5-10%
- c. 11-15%
- d. Depends on the individual's baseline BMI

General Considerations in Pharmacologic Initiation

Pharmacologic interventions may be helpful as adjuvant therapy with lifestyle interventions for patients 18 years and older* with BMI ≥30 kg/m² or ≥27 kg/m² with comorbidities.

- Different patients respond to different medications
 - If one option does not work, consider others
- Discontinue medication in patients who do not respond with weight loss of at least 5% at 12 weeks after maximum dose[†]
- Avoid in pregnancy
 - Pregnancy tests at baseline
 - Consider a disclosure signature

*December 2020: liraglutide label change to include 12–17-year-olds with body weight of >60 kg and initial BMI corresponding to 30 kg/m² or greater for adults. †Liraglutide label suggests only 4% weight loss at 12 weeks after maximum dose. Apovian CM, et al. J Clin Endocrinol Metab 2015;100(2):342-362.

Medications to Treat Obesity

FDA Approved

Orlistat

Phentermine

Phentermine/Topiramate ER #

Bupropion/Naltrexone XL

Liraglutide 3mg #

Semaglutide 2.4mg. #

Tirzepatide 15 mg

Setmelanotide monogenic obesity*

approved for adults and adolescents

Topiramate Zonisamide Bupropion Naltrexone Metformin Dulaglutide RA Exenatide Liraglutide GLP-1 Lixisenatide Semaglutide GIP/GLP-1 RA Tirzepatide Amylin analogue Pramlintide Canagliflozin SGLT2I Dapagliflozin Empagliflozin

Additional Medications that

May Cause Weight Loss and Used Off Label

FDA-Approved Short-Term (Anti) Obesity Medications

Generic Drug*	Dose	Contraindications	Side Effects	
Phentermine	8-37.5 mg	Anxiety disorder, CVD, hypertension, MAO inhibitors, glaucoma, hyperthyroidism, seizures, pregnancy/ breastfeeding, drug abuse history	Insomnia, palpitations,	Insomnia, palpitations,
Diethylpropion	25 mg or 75 mg, SR		tachycardia, dry mouth, taste alterations, dizziness,	
Phendimetrazine	17.5-70 mg or 105 mg <i>,</i> SR		tremors, headache, diarrhea, constipation, vomiting,	
Benzphetamine	25-50 mg		breastfeeding, drug abuse bistory	gastrointestinal distress, anxiety, restlessness, increased blood pressure

*Mechanism of action = sympathomimetic-noradrenergic causing appetite suppression.

MAO, monoamine oxidase; SR, sustained release.

DailyMed. https://dailymed.nlm.nih.gov/dailymed/index.cfm. Accessed February 23, 2021. Bray GA, et al. Circulation. 2012;125(13):1695-1703. Apovian CM, et al. J Clin Endocrinol Metab. 2015;100(2):342-362.

Phentermine

- US Drug Enforcement
 Agency scheduled IV drug
 - Risk for addiction
- Not indicated for longterm use
 - 13 weeks by label

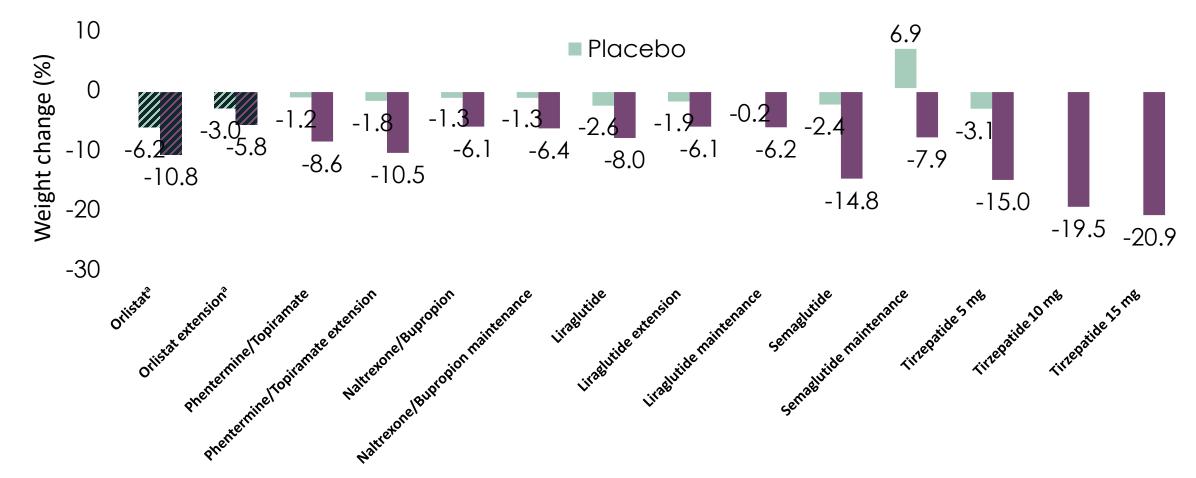
Endocrine Society allows for possible long-term use:

- No CVD
- No psychiatric/substance abuse history
- Has been informed about therapies that are approved for long-term use
- Document off-label use in patient's medical record
- No clinically significant increase in pulse/BP when taking phentermine
- Demonstrates significant weight loss with phentermine
- Start at 7.5 or 15 mg/d—dose escalate if not achieving significant weight loss
- Monitor monthly during dose escalation

FDA-Approved (Anti) Obesity Medication

Generic (listed alphabetically)	Mechanism of Action
Liraglutide (subcutaneous injection)	GLP-1 receptor agonist
Naltrexone/bupropion ER (oral)	Opioid receptor antagonist; dopamine and noradrenaline reuptake inhibitor
Orlistat (oral)	Pancreatic lipase inhibitor—impairs gastrointestinal energy absorption, causing excretion of approximately 30% of ingested triglycerides in stool
Phentermine/topiramate-ER (oral)	Noradrenergic + GABA-receptor activator, kainite/AMPA glutamate receptor inhibitor causing appetite suppression
Semaglutide (subcutaneous injection)	GLP-1 receptor agonist
Tirzepatide (subcutaneous injection)	GLP-1/GIP receptor agonist

Mean Percent (%) Weight Change Reported in the Main Phase 3 and Extension Trials of Antiobesity Medications



Orlistat: XENDOS trial (years 1 and 4). Phentermine/topiramate: CONQUER and SEQUEL trials. Naltrexone/bupropion: COR-I and COR-II trials. Liraglutide: SCALE obesity, SCALE obesity, and prediabetes extension, and SCALE maintenance trials. Semaglutide: STEP 1 and STEP 4 trial. All trials are listed in order as seen in the figure from left to right. The grey color represents placebo arms; the red color represents intervention arms. Athe mean weight change in the orlistat group is in kg not in percent (striped bar charts).

Chakhtoura M, et al. EClinicalMedicine. 2023;58:101882.

Orlistat

Dose Frequency	Efficacy			indications/ tions/Warnings	Side Effects
60 mg OTC 120 mg TID within 1 h of fat-containing meal	at Year 1 i (120 mg T • ↓ BP, TC, fasting glu year • Slows risk	om 3.9-10.2% n 17 RCTs ID) LDL-C, Icose at 1	syndro breastf some n warfari	c malabsorption me, pregnancy, eeding, cholestasis, nedications (eg, n, antiepileptic levothyroxine, oorine)	Oily spotting, cramps, flatus with discharge, fecal urgency, fatty oily stool, increased defecation, fecal incontinence
	Practical Considerat		ions	 Consider fat-soluble Limit fat intake to 30^o Counsel on risk of GI 	% of calories

Bragg R, et al. J Am Assoc Nurse Pract 2016;28(2):107-115. Kahan S. Am J Manag Care. 2016;22(7 Suppl):S186-S196.

Phentermine/Topiramate ER

Dose Frequence	су	Efficacy		Contraindications/ Precautions/Warnings	Side Effects
 Initiate treatment at 3.75 mg/23 mg for 2 weeks Increase to 7.5 mg/ 46 mg Escalate to 11.25 mg/ 69 mg for 2 weeks then to max 15 mg/92 mg Inote a treatment at 10% weight loss with treatment vs 2% with placebo Improved cardiometabolic markers Reduced progression to T2DM 		Pregnancy and breastfeeding, hyperthyroidism, glaucoma, use of MAO inhibitors	Paresthesia, dizziness, taste alterations, insomnia, constipation, dry mouth, elevation in heart rate, memory or cognitive changes		
	Practical Conside	ations • Counsel about ris suicidal thoughts • Taper highest dos		nt Agency Schedule IV • 7/2022 sk for mood disorders, years ol	of childbearing age: pregnancy ion plan and monthly ncy testing approved for adolescents >12 d with BMI of 95 th percentile or

Liraglutide

Dose Frequency	Efficacy		Contraindications/ Precautions/Warnings	Side Effects		
Weekly titration by 0.6 mg over 5 weeks to target dose of 3.0 mg	 Mean weight loss 9% at 1 year Reduced progression to T2DM in patients with prediabetes Reduced risk of weight regain at 1 year 		Medullary thyroid cancer history, multiple endocrine neoplasia type 2 history, history of pancreatitis, pregnancy, breastfeeding	Nausea, vomiting, diarrhea, constipation, hypoglycemia in patients with T2DM, increased lipase, increased heart rate, pancreatitis		
	Practical Considerations	 FDA a with B with a Gener 	able administration pproved for use in adults MI ≥30kg/m ² or ≥27 kg/m ² t least one complication ic summer 2024, price ted at \$700	Approved 12/2020 label cha treatment of obesity in adolescents aged 12 to 17 ye with a body weight of at leas kg and an initial BMI corresponding to 30 kg/m ² of greater for adults	ears st 60	

Naltrexone/Bupropion ER

Dose Frequency	Efficacy	Contraindications/ Precautions/Warnings	Side Effects
 Initiate 8 mg/90 mg x 1 week Weekly escalation to target dose of 32 mg/360 mg (2 tablets BID) 	 Weight loss of 8.2% vs 1.4% (placebo) Improved cardiometabolic parameters Fewer cravings Lowered HbA1c in patients with T2DM 	Uncontrolled hypertension, seizure disorder, anorexia or bulimia, drug or alcohol withdrawal, chronic opioid use, MAO inhibitors, caution with renal/ hepatic impairment	Nausea, constipation, headache, dizziness, vomiting, insomnia, dry mouth Transient increase in BP
	Practical Consideratio	Titrate dose on initiation Monitor BP Monitor closely for depres	ssion

Semaglutide

Dose Frequency	Efficacy	Contraindications/ Precautions/Warnings	Side Effects	
Weekly injections with titration every four weeks, 0.25 mg, 0.5 mg, 1 mg, 1.7 mg or 2.4 mg dose of 2.4 mg	 Mean weight loss 14.9% at 68 weeks 	Medullary thyroid cancer history, multiple endocrine neoplasia type 2 history, suicidal behavior and ideation, pregnancy, breastfeeding, acute gallbladder disease, diabetic retinopathy, acute kidney injury	Nausea, vomiting, diarrhea, constipation, hypoglycemia in patients with T2DM, increased lipase, increased heart rate, pancreatitis	
Practical Consider	$>30 \text{ kg/m}^2 \text{ or } >27 \text{ kg}^2$	use in adults with BMI with or with a	22 approved for 12–17-year-old besity a most common issue – slow on ; and 2.4 mg as maintenance dos	
2024 Label update: reduce the risk of MACE in adults with established CV disease and overweight or obesity				

https://www.novo-pi.com/wegovy.pdf

MACE: Major Adverse Cardiovascular events (cardiovascular death, non-fatal MI, or non-fatal stroke

Tirzepatide

Dose Frequency		Efficacy		Contraindications/ Precautions/Warnings	Side Effects
Weekly injections with titration every four weeks, 2.5mg, 5mg, 7.5mg, 10mg, 12.5mg, 15mg		 Mean weight loss 20.9% at 72 weeks 		Medullary thyroid cancer history, multiple endocrine neoplasia type 2 history, suicidal behavior and ideation, pregnancy, acute gallbladder disease, diabetic retinopathy, acute kidney injury	Nausea, vomiting, diarrhea, constipation, hypoglycemia in patients with T2DM
	Practical Considerations		 Injectable administration FDA approved for use in adults with BMI ≥30kg/m² or ≥27 kg/m² with at least one obesity related comorbid condition Nausea most common issue – slow titration Smg, 10mg, and 15mg as maintenance doses 		

Rare Genetic Cause of Obesity Treatment

Uncommon Obesity



>20 rare genetic disorders

Common symptoms

- Early onset of severe obesity, often less than one year of age
- Insatiable hunger (hyperphagia)

Genetic testing is critical (free)

Patient eligibility criteria:

- ≤18 years of age, BMI ≥97th percentile or
- ≥19 years of age, BMI ≥40 kg/m², and a history of childhood obesity before age 10

Setmelanotide – Imcivree[™]

- Approved in November 2020 for patients with obesity due to POMC, PCSK1, or LEPR deficiency
 - Impaired MC4 receptor pathway
 - Adults and pediatric patients 6 years of age and older with deficiency confirmed by genetic testing
- Action: MC4 receptor agonist
 - Restore impaired MC4 receptor pathway activity arising due to genetic deficits upstream of the MC4 receptor
- Rare pediatric disease priority review voucher, breakthrough therapy designation, orphan drug designation

PCSK1, proprotein convertase subtilisin/kexin type 1.

Uncommon Obesity. https://www.uncommonobesity.com/. Accessed February 23, 2021. Rhythm Pharmaceuticals. https://www.rhythmtx.com/science-overview/. Accessed September 20, 2024.

Identify

Identify Available Pharmacotherapeutics That Can Be Utilized to Support Treatment of Chronic Disease of Obesity That Are Not Labeled for Obesity by the FDA.

	Medication	Information	Weight loss	Cost
	GLP1RA's		Semaglutide (lower doses with Ozempic or Rybelsus) weight reduction from baseline – Semaglutide 0.5mg 2.3 kg, Semaglutide 1.0 mg 6.5 kg Semaglutide 14mg orally 4.4kg Liraglutide (lower doses with Victoza) 1.0mg dose 3.1kg Dulaglutide 1.5mg 3.0kg Liraglutide 1.8mg generic launched June 2024	\$500-1400 30 days for \$334
v	Metformin	Go as high as patient can tolerate – 2500mg max	500 mg 120 tabs	Good RX ~ \$12

32 mg max. It is sco down to 25 mg pret often too much for	ng so well beyond the	Range of weight loss is about 4-7# after 2 months	You can get it compounded to get the 8mg and 16 mg doses (then can increase to 25 mg from the cheaper		
Bupropion XL	ness, fatigue, nausea, g up.		version) ~ \$25/month.		
		can expect about 5 pounds of weight loss with 150mg to 300mg	~\$14		
	**so you could do off label with naltrexone and bupropion together and likely have a lower cost BUT doses would be different so nothing in literature to support amount of				

	Medication	Information	Weight loss	Cost
	Phentermine	I suggest using low dose – 8mg tablets and start with one dose in am and ½ to 1 tablet about 3- 4pm.	Approximately 5% weight loss	90 tablets ~ \$35-50
	Topiramate (not extended release)	Often works nicely for people with evening eating alone.	When used alone approximately 6% of patients at 50mg/day lose weight	25mg (using ½ to 1 tablet per day) ~ \$12. NOTE Topiramate ER is capsule only and close to ~\$75 so no savings.
	so you could do off label with phentermine and topiramate together and likely have a lower cost BUT doses would be different so nothing in literature to support amount of weight loss			

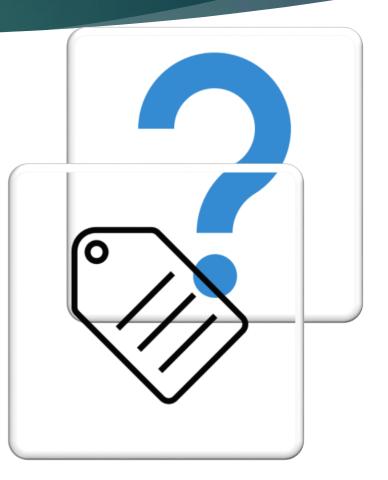


Medication	Information	Weight loss	Cost
SGLT2Is – Canagliflozin (Invokana)	There is some literature to use phentermine with SGLT2I – 7.5% weight	Approximately 1.9% weight loss	~\$550

Off Label Combinations (WHY)

Central mechanisms of actions

- GLP-1RA's and topiramate decrease signaling of AGRP and NPY (these stimulate food intake)
- GLP1-RA's, phentermine, bupropion, and naltrexone increase signaling of POMC and CART (these decrease food intake)



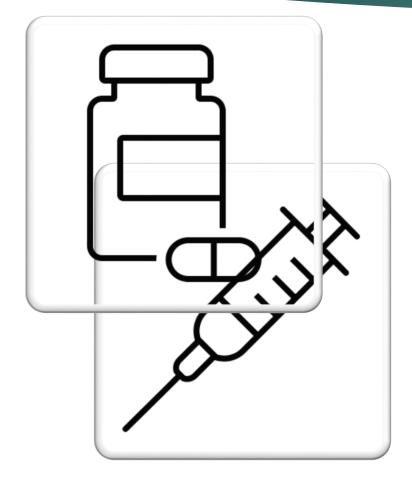
Off Label Combinations (WHAT)



- ▶ GLP-1 plus any medication
 - Watch for nausea
- Orlistat plus any medication
 - Especially if constipation noted as side effect of GLP-1
- Phentermine plus GLP-1 or SGLT-2
- Metformin plus any medication

Shared Decision Making

Medication Selection - Shared



- Complete RXAO
- List medications still appropriate
- Present to the patient
- Make the selection

Polling Question

Which medication would you recommend for Mariah?

- A. Liraglutide
- B. Naltrexone/bupropion ER
- **C**. Orlistat
- D. Phentermine
- E. Phentermine/topiramate ER
- F. Semaglutide
- G. tirzepatide
- H. None at this point

Mariah: Shared Decision-Making

AOM	Additional Benefit	eXcluded
Liraglutide (subcutaneous injection)	Insulin Resistance	None
Naltrexone/bupropion EB (oral)	None	History of seizure post
	None	concussion MVA
Orlistat (oral)	None	None
Phentermine (oral)	Cost	None
Phentermine/topiramate-ER (oral)	Efficacy	None
Semaglutide (subcutaneous injection)	Insulin Resistance	None
Tirzepatide (subcutaneous injection)	Insulin Resistance	None

Mariah: Measuring Efficacy

Begin therapy with semaglutide



Initiate at 0.25mg once a week for 4 weeks, increase to 0.5mg x 4 wks, 1.0 mg x 4 wks, 1.7mg (and 2.4mg if needed)

Effective response to therapy



>5% weight loss from baseline 20 weeks after starting medication

Improvement in CV risk markers



Improvement in insulin resistance

Continue to See the Patient Every 2-4 Weeks

Intensive Lifestyle Intervention

Can be done by other providers as well, e.g., dietitians, physical therapy/exercise physiologist, health coaches

Monitoring: weight, hunger, satiety



Mariah's Visit at 20 Weeks

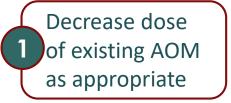


- •BP 112/58 mmHg; HR 76 bpm; RR 16 breaths/min; pOx 97%; weight 199 lbs (6% weight loss)
- •Currently at 1.7mg with hunger controlled and satiety signals well heard, holding at this dose
- Continues to eat reduced carbohydrate
- •Continues activity with swimming three days a week and not lifting at gym twice a week

Ineffective Response to Therapy

- <4-5% weight loss at 12 weeks of maximum dose
 - Medications with escalating doses could be 16 weeks or longer
 - Unable to tolerate maximum doses
- < ? 3% weight loss but with improvement in ORCs



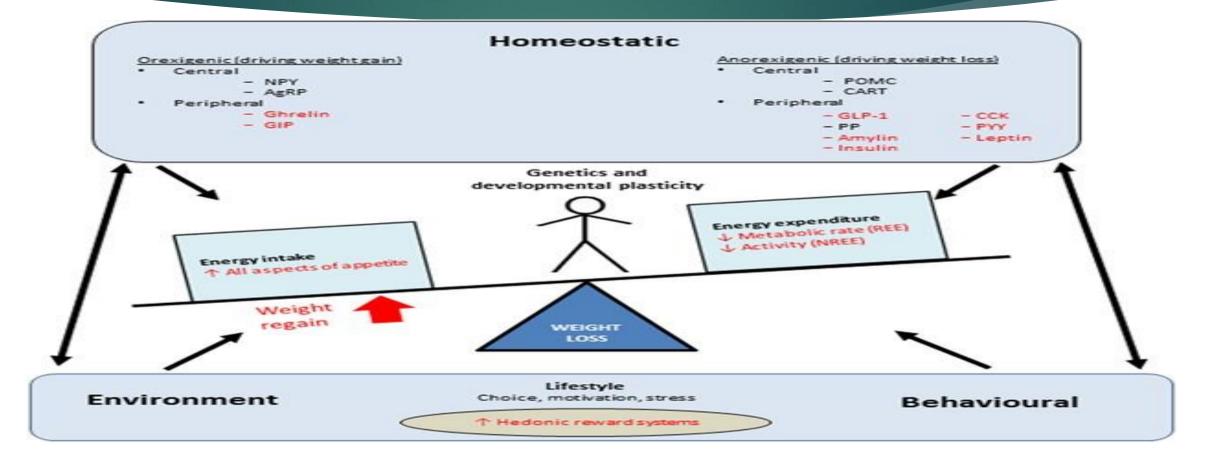




AOM, anti-obesity medication.

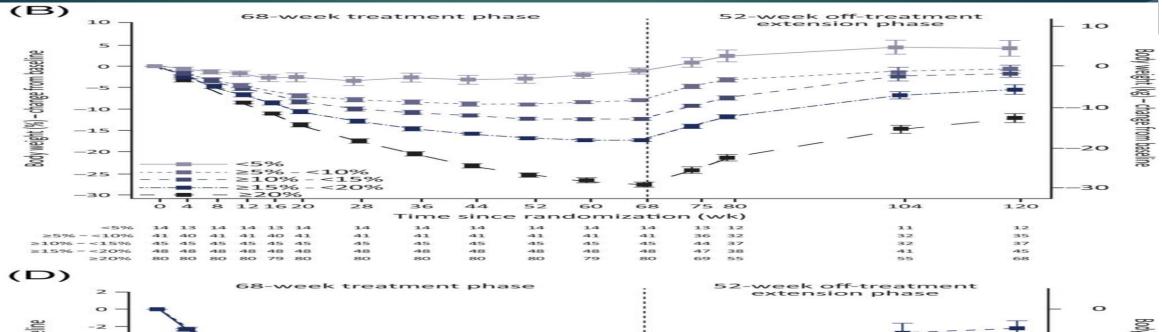
Bray GA, et al. Lancet. 2016;387(10031):1947-1956. Apovian CM, et al. J Clin Endocrinol Metab. 2015;100(2):342-362.

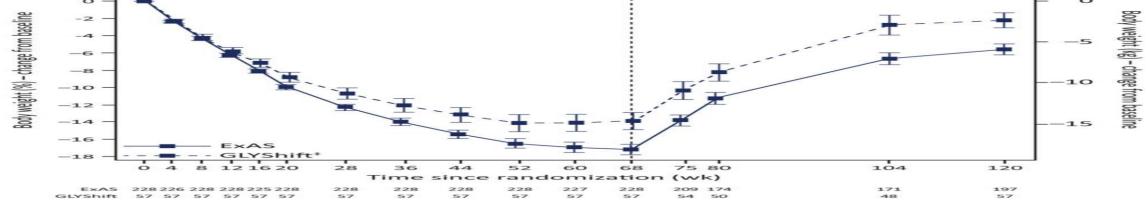
Pathology Of Weight Regain – Metabolic Adaptation



Greenway, F. Physiological adaptations to weight loss and factors favouring weight regain. Int J Obes 39, 1188–1196 (2015). https://doi.org/10.1038/ijo.2015.59 OPEN ACCESS Figure 2

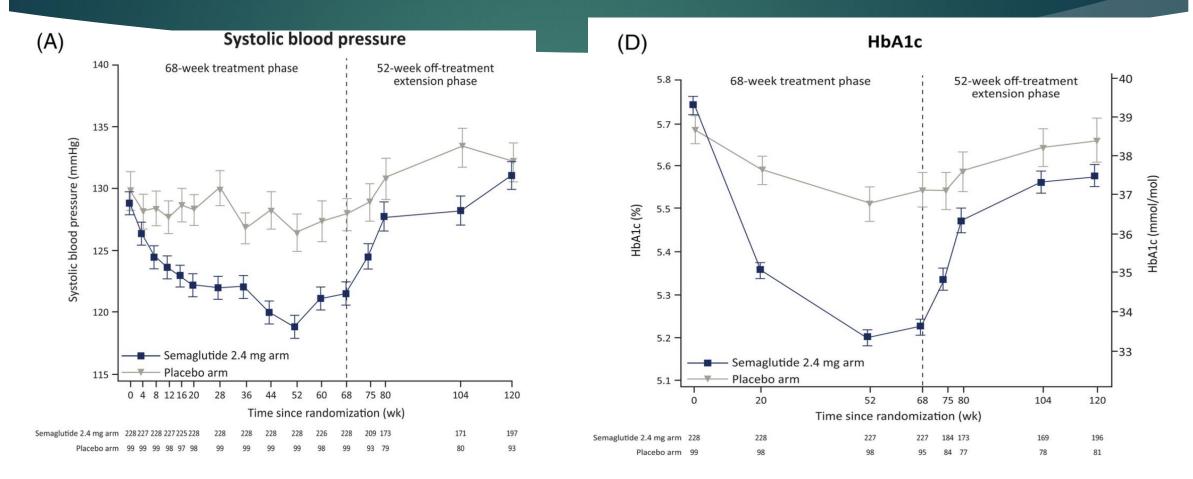
Metabolic Adaptation – STEP 1 Extended





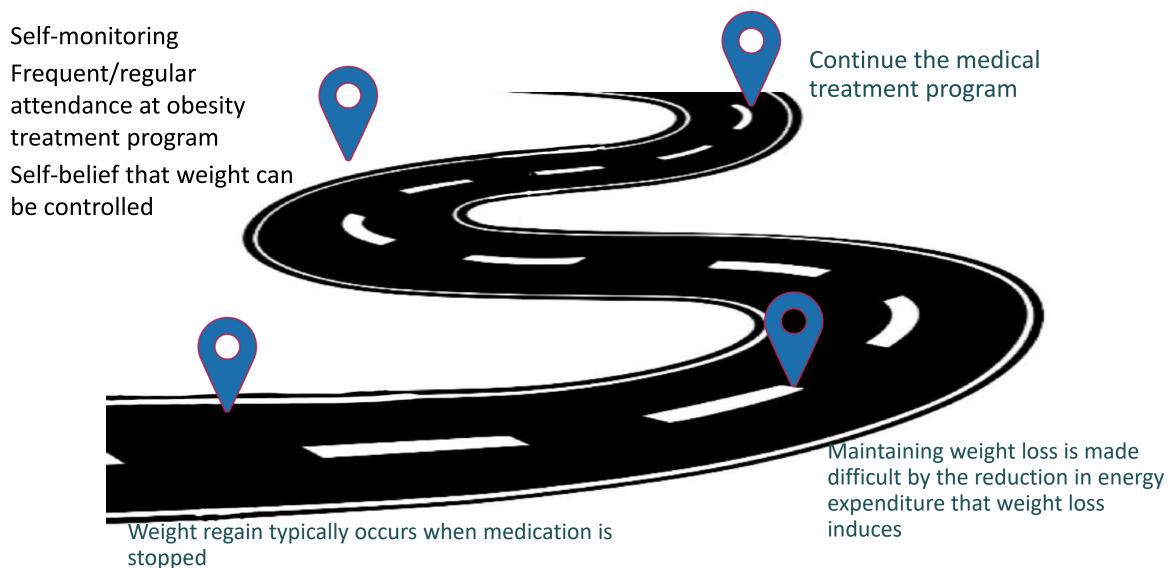
Wilding, J., et al., & STEP 1 Study Group (2022). Weight regain and cardiometabolic effects after withdrawal of semaglutide: The STEP 1 trial extension. *Diabetes, obesity & metabolism*, 10.1111/dom.14725. Advance online publication. <u>https://doi.org/10.1111/dom.14725</u> open access Figure 1

Metabolic Adaptation – Step 1 Extended



Wilding, J., et al, & STEP 1 Study Group (2022). Weight regain and cardiometabolic effects after withdrawal of semaglutide: The STEP 1 trial extension. *Diabetes, obesity & metabolism*, 10.1111/dom.14725. Advance online publication. <u>https://doi.org/10.1111/dom.14725</u> open access Figure 2

Obesity Treatment to Impact ORCs Long Term = Maintaining Weight Loss



Post Bariatric Surgery

- Optimal time for pharmacotherapy
 - At weight plateau
 - Maximize weight loss outcomes
- In the literature
 - Topiramate 7.7% weight loss post surgery (Stanford)
 - Non-GLP1RA's at 9 months 5.6% weight loss, GLP1RA's at 9 months 6.9% (Gazda)

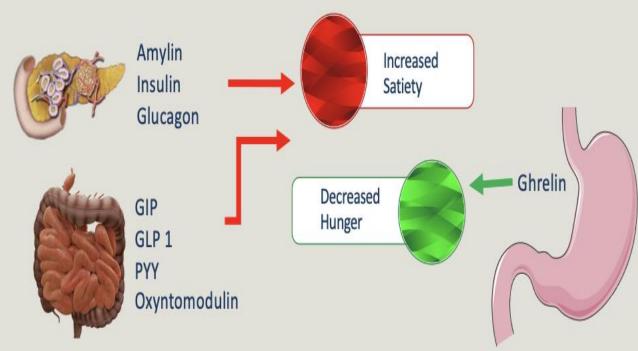
Stanford F. C. (2019). Controversial issues: A practical guide to the use of weight loss medications after bariatric surgery for weight regain or inadequate weight loss. Surgery for obesity and related diseases : official journal of the American Society for Bariatric Surgery, 15(1), 128–132. <u>https://doi.org/10.1016/j.soard.2018.10.020</u> Gazda, C. L., Clark, J. D., Lingvay, I., & Almandoz, J. P. (2021). Pharmacotherapies for Post-Bariatric Weight Regain: Real-World Comparative Outcomes. Obesity (Silver Spring, Md.), 29(5), 829–836. https://doi.org/10.1002/oby.23146

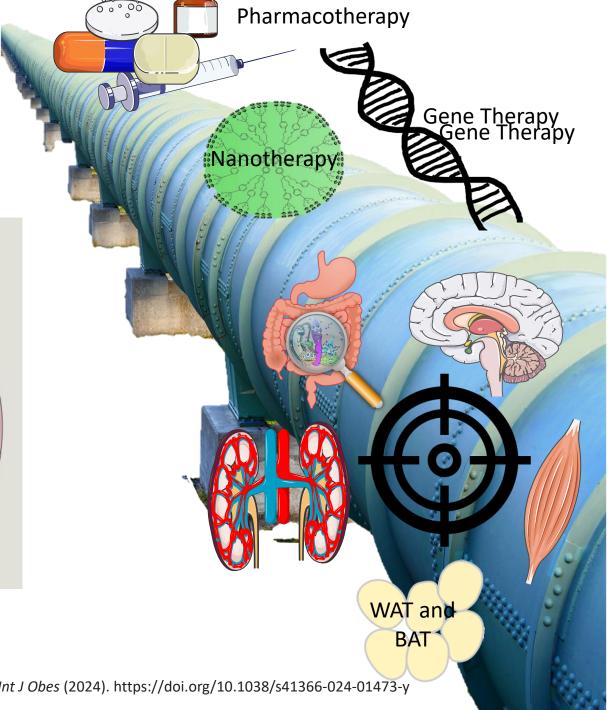
Recognize

Recognize Medications That Are In The Pipeline For Treating Obesity

In The Pipeline

- 50 + companies are developing obesity therapies
- 80 + therapies in trials





Practice Pearls

Obesity is a chronic and often progressive condition

Obesity management is not about simply reducing numbers on the scale

Intensify treatment with pharmacology

Evaluate medication success at "12 weeks"

If one medication doesn't work, try another

With success, continue medical management



Thank you!



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Resources

Image resources

- https://www.worldobesity.org/resources/image-bank
- https://www.obesityaction.org/get-educated/publicresources/oac-image-gallery/
- Canadian Obesity Network Image Bank: <u>https://www.flickr.com/photos/144769815@N06/</u>