Psychiatry Drug Alerts 2019 Self-Assessment Module 3: Peer Comparison

You recently participated in an ABPN-approved Self-Assessment activity relevant to your specialty and/or subspecialty. This peer comparison report provides you with feedback on your performance, relative to your peers, on the test module. In order to recognize your current knowledge base and to identify specific topics where further study may be needed, please review your answers to the following questions and compare them with those of your peers.

	rexanolone (Zulresso) as the first agent specifically indicated for treatment of
· · · · · · · · · · · · · · · · · · ·	and sudden loss of consciousness, the drug will only be available
	on Strategy (REMS) program with restricted distribution.
Dizziness	0.00 %
Excessive sedation	100.00 %
Injection site reaction	0.00 %
All of the above	0.00 %
2) According to a retrospective chart rev	view, veterans had modestly better PTSD outcomes when treated with:
An SSRI	0.00 %
An opioid	0.00 %
Buprenorphine-naloxone	100.00 %
None of the above	0.00 %
3) In these patients, PTSD symptom scor scores increased slightly with SSRI treat	res decreased by 24% with buprenorphine—naloxone and by 16% with opioids; ment.
True	100.00 %
False	0.00 %
buzzing sound-found accounte prescribing frequency. Fluoxetine	within the skull sometimes accompanied by dissociation, vertigo, and a d for about one-fourth of the occurrences, which is disproportionate to its 0.00 %
Bupropion	0.00 %
Venlafaxine	100.00 %
Duloxetine	0.00 %
antidepressant activity diminishes in the	s unknown, but they appear to be related in part to how rapidly brain after discontinuation. Patients reported using many methods to get relaxation, and various supplements); seemed effective.
Most	
None	100.00 %
methadone and most antiretroviral (AR metabolism. While most combinations of	tant concern in patients with HIV. Clinically significant interactions between T) classes are uncommon, however, individual agents can affect methadone do not require methadone dosage adjustments, clinical guidelines recommend
Ritonavir	vithdrawal symptoms when it is used in combination with: 8.33 %
Abacavir or nelfinavir	0.00 %
Elvitegravir	0.00 %
Efavirenz or nevirapine	91.67 %

7) In patients receiving treatment with buprenorphine for opioid dependence, guidelines recommend against

coadministration of buprenorphine with un regimens that include ritonavir, which can p	roduce a significant in buprenor	
Increase	100.00 %	
Decrease	0.00 %	
8) Of the 4 agents FDA-approved to maintainaltrexone, intramuscular naltrexone), non- generally considered to be safe. However, c	e have significant CYP effects, and coadm	inistration with ART regimens is
disulfiram, and the lopinavir-ritonavir comb could lead to a disulfiram-like reaction.		
Nevirapine	0.00 %	
Atazanavir	100.00 %	
Darunavir	0.00 %	
All of the above	0.00 %	
9) In addition to pharmacokinetic interactio	ns, concurrent use of substance use disor	der medications and ART
regimens can have compounding effects, w	hich can include:	
Liver enzyme elevations	0.00 %	
Hepatotoxicity	0.00 %	
QT prolongation	0.00 %	
All of the above	100.00 %	
10) In a preliminary randomized trial in pati who received the long-acting injectable form	nulation following detoxification	
Remained in treatment longer	41.67 %	
Had fewer opioid-positive screens	0.00 %	
Attended more therapy sessions	0.00 %	
All of the above	58.33 %	
11) A network meta-analysis supports all of disorder except:	the following as first-line pharmacothero	apy for generalized anxiety
Venlafaxine	0.00 %	
Vortioxetine	100.00 %	
Duloxetine	0.00 %	
Escitalopram	0.00 %	
12) The analysis found was most e	-	e for Anxiety (HAM-A) scores but
was associated with high rates of prematur	•	
Vilazodone	0.00 %	
Paroxetine	8.33 %	
Quetiapine	91.67 %	
Pregabalin	0.00 %	

13) According to the results of an observation with schizophrenia for whom monotherapy wi	,, , , , , , , , , , , , , , , , , , ,	 ··		
Benzodiazepine	0.00 %			
Antidepressant	100.00 %			
Mood stabilizer	0.00 %			
Additional antipsychotic	0.00 %			
14) Despite the advantages of long-acting injectable antipsychotics in providing consistent medication exposure, patients may experience breakthrough symptoms. Potential causes for these breakthrough symptoms can include:				
Low plasma drug levels	8.33 %			
Comorbid medical illness	0.00 %			
Improper administration technique	0.00 %			

91.67 %

All of the above