

FOOD AND DRUG ADMINISTRATION (FDA)
Center for Drug Evaluation and Research (CDER)

***Joint Meeting of the Anesthetic and Analgesic Drug Products Advisory Committee (AADPAC)
and the Drug Safety and Risk Management Advisory Committee (DSaRM)***

FDA White Oak Campus, Building 31 Conference Center, the Great Room (Rm. 1503)
10903 New Hampshire Avenue, Silver Spring, Maryland
June 8, 2016

QUESTIONS

1. **DISCUSSION:** Please discuss whether there are sufficient data to support a finding that Troxyca ER (oxycodone hydrochloride and naltrexone hydrochloride extended-release capsules) has properties that can be expected to deter abuse, commenting on support for abuse-deterrent effects for each of the three possible routes of abuse:
 - a. Oral
 - b. Nasal
 - c. Intravenous

2. **VOTE:** Should Troxyca ER be approved for the proposed indication, management of pain severe enough to require daily, around the clock, long-term opioid treatment and for which alternative treatment options are inadequate?

3. **VOTE:** If approved, should Troxyca ER be labeled as an abuse-deterrent product by the oral route of abuse?

4. **VOTE:** If approved, should Troxyca ER be labeled as an abuse-deterrent product by the nasal route of abuse?

5. **VOTE:** If approved, should Troxyca ER be labeled as an abuse-deterrent product by the intravenous route of abuse?