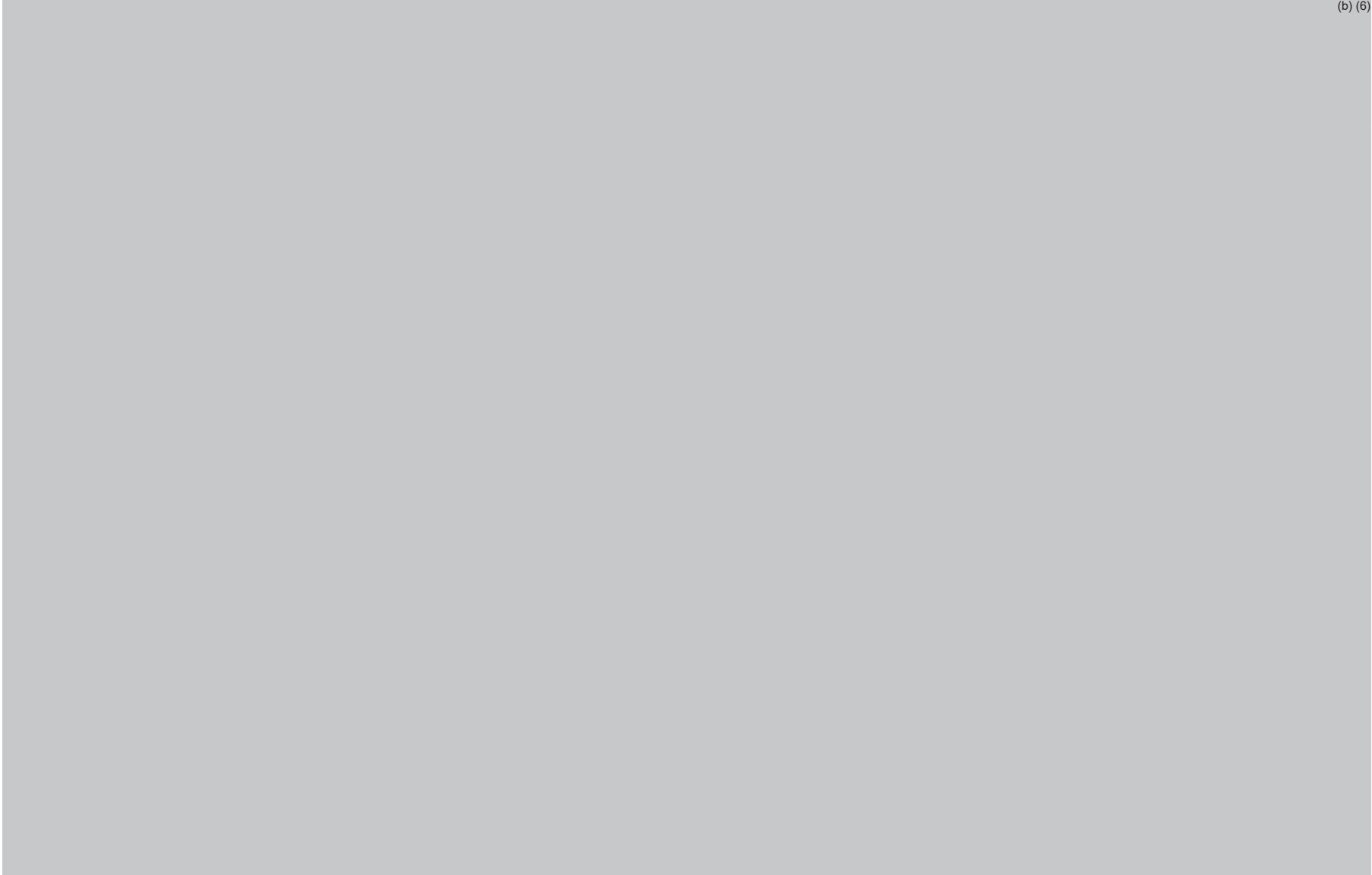




FDA - Adverse Event Reporting System (FAERS)
FOIA Case Report Information

FDACDER3078

Case ID: 10213469



(b) (6)



FDA - Adverse Event Reporting System (FAERS)
FOIA Case Report Information

FDACDER3079

Case ID: 10213469



(b) (6)



FDA - Adverse Event Reporting System (FAERS)
FOIA Case Report Information

FDACDER3080

Case ID: 10213469



(b) (6)