



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

Case ID: 7979016



(b) (6)



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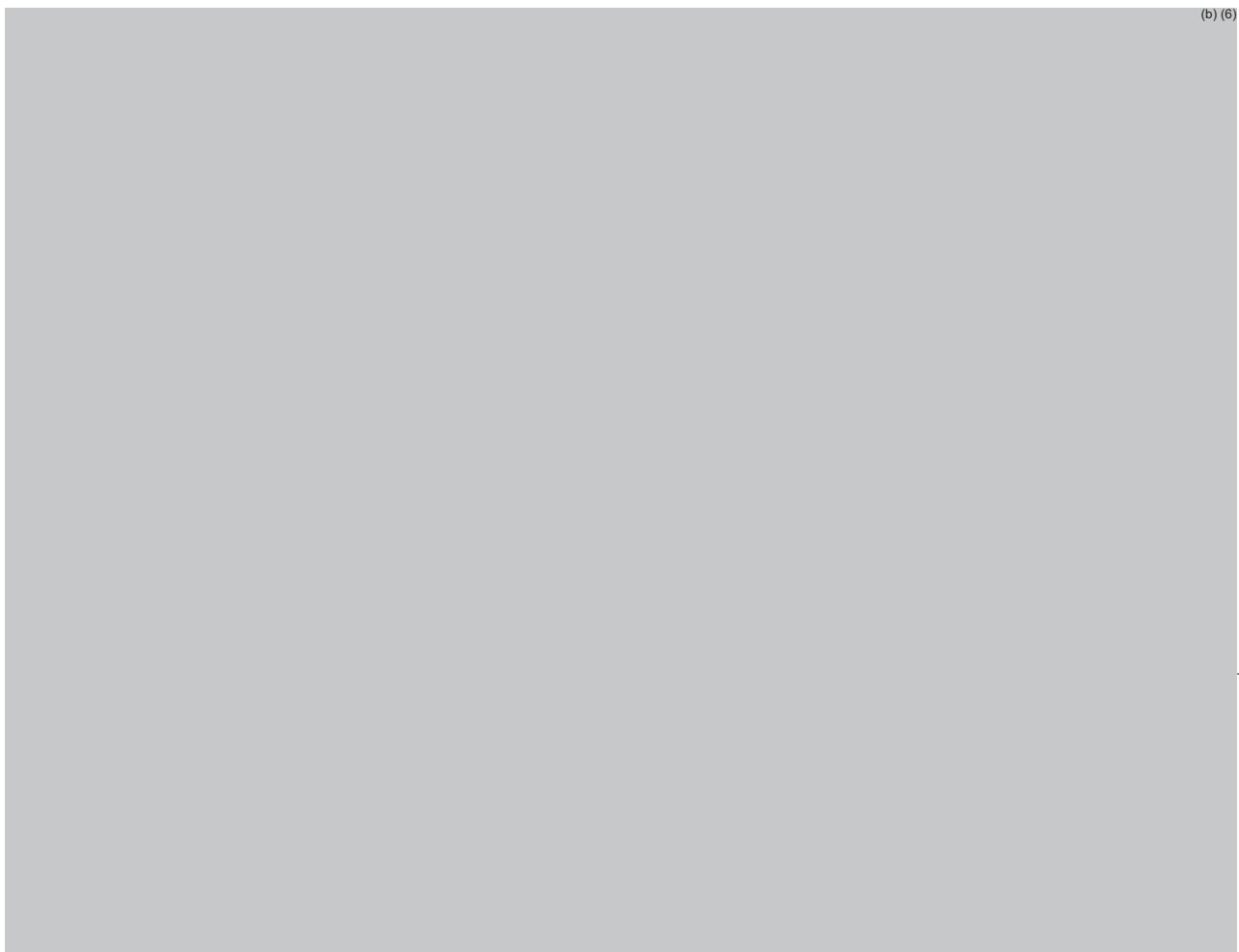
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