

**FDA - Adverse Event Reporting System (AERS)
Freedom Of Information (FOI) Report
Standard Report - All Preferred Terms in Cases**

MedDRA Preferred Term Reaction	Count	% Rpts
ANTI-HBC IGM ANTIBODY POSITIVE	1	0.0%
ANTI-ERYTHROCYTE ANTIBODY	1	0.0%
ANTEROGRADE AMNESIA	1	0.0%
ANTERIOR CHAMBER DISORDER	1	0.0%
ANTEPARTUM HAEMORRHAGE	1	0.0%
ANKLE FRACTURE	1	0.0%
ANISOCYTOSIS	1	0.0%
ANTINUCLEAR ANTIBODY NEGATIVE	1	0.0%
ANHIDROSIS	1	0.0%
ANASTOMOTIC COMPLICATION	1	0.0%
ANAL PRURITUS	1	0.0%
ANAL FISSURE	1	0.0%
ANAESTHETIC COMPLICATION	1	0.0%
ANAESTHESIA	1	0.0%
ANAEMIA MACROCYTIC	1	0.0%
AMYOTROPHIC LATERAL SCLEROSIS	1	0.0%
ADJUSTMENT DISORDER	1	0.0%
ADENOMA BENIGN	1	0.0%
ADENOCARCINOMA	1	0.0%
ACUTE STRESS DISORDER	1	0.0%
ACUTE CORONARY SYNDROME	1	0.0%
ACTIVATED PARTIAL THROMBOPLASTIN TIME ABNORMAL	1	0.0%
ACTIVATED PARTIAL THROMBOPLASTIN TIME	1	0.0%
ACQUIRED EPIDERMOLYSIS BULLOSA	1	0.0%
ALCOHOL PROBLEM	1	0.0%
ALCOHOL POISONING	1	0.0%
ALCOHOL INTOLERANCE	1	0.0%
ALCOHOL ABUSE	1	0.0%
ALBUMINURIA	1	0.0%
AGONAL DEATH STRUGGLE	1	0.0%
ACCELERATED IDIOVENTRICULAR RHYTHM	1	0.0%
ACCIDENT AT WORK	1	0.0%
ACCIDENT	1	0.0%
ABSCESS ORAL	1	0.0%
ABSCESS INTESTINAL	1	0.0%
ABORTION INCOMPLETE	1	0.0%
ABDOMINAL HERNIA	1	0.0%

Total Reactions: 61983

Total Death Outcomes by Case: 1015