Adverse Events Listing
Example 3: Multi-page with group_by

Subject ID	Study Day	Adverse Event	Severity	Serious
01-701-1363		Headache	MODERATE	N
01-701-1363		Headache	MILD	N
01-701-1363	16.0	Nausea	MILD	N
01-701-1363	48.0	Application Site Pruritus	MILD	N
01-701-1363	137.0	Back Pain	MILD	N
01-701-1363	137.0	Back Pain	MILD	N
01-701-1387	7.0	Diarrhoea	MILD	N
01-701-1387	7.0	Hyperhidrosis	MILD	N
01-701-1392	140.0	Upper Respiratory Tract Infection	MILD	N
01-701-1392	140.0	Upper Respiratory Tract Infection	MILD	N
01-701-1415	29.0	Upper Respiratory Tract Infection	MILD	N
01-701-1415	29.0	Upper Respiratory Tract Infection	MILD	N
01-701-1415	71.0	Micturition Urgency	MILD	N
01-701-1415	121.0	Upper Respiratory Tract Infection	MILD	N
01-701-1415	121.0	Upper Respiratory Tract Infection	MILD	N
01-701-1415	168.0	Diarrhoea	MILD	N
01-701-1360	6.0	Application Site Vesicles	MODERATE	N
01-701-1383	4.0	Application Site Pruritus	MILD	N
01-701-1383	4.0	Application Site Pain	MILD	N
01-701-1383	48.0	Application Site Pruritus	MODERATE	N
01-701-1383	48.0	Application Site Pruritus	MILD	N
01-701-1383	48.0	Application Site Erythema	MILD	N
01-701-1383	68.0	Application Site Irritation	MODERATE	N

Adverse Events Listing
Example 3: Multi-page with group_by

Subject ID	Study Day	Adverse Event	Severity	Serious
01-701-1383	68.0	Application Site Irritation	MILD	N
01-701-1383	68.0	Application Site Erythema	MILD	N
01-701-1383	93.0	Application Site Vesicles	MILD	N
01-701-1383	141.0	Headache	MILD	N
01-701-1383	141.0	Chest Discomfort	MILD	N
01-701-1383	164.0	Cough	MODERATE	N
01-701-1444	15.0	Application Site Pruritus	MILD	N
01-701-1444	15.0	Application Site Erythema	MILD	N
01-701-1444	15.0	Salivary Hypersecretion	MILD	N
01-701-1444	15.0	Application Site Erythema	MODERATE	N
01-701-1444	15.0	Application Site Pruritus	MODERATE	N
01-701-1444	31.0	Application Site Irritation	MODERATE	N
01-701-1444	31.0	Application Site Vesicles	MODERATE	N
01-701-1444	35.0	Paraesthesia	MILD	N
01-701-1442	77.0	Application Site Pruritus	MILD	N
01-702-1082	-19.0	White Blood Cell Count Increased	MILD	N
01-702-1082	-19.0	Neutrophil Count Increased	MILD	N
01-702-1082	-19.0	Urine Analysis Abnormal	MILD	N
01-702-1082	-19.0	Urine Analysis Abnormal	MILD	N
01-702-1082	-19.0	White Blood Cell Count Increased	MILD	N
01-702-1082	-19.0	Neutrophil Count Increased	MILD	N
01-702-1082	39.0	Rectal Haemorrhage	MILD	N
01-702-1082	39.0	Rectal Haemorrhage	MILD	N

Adverse Events Listing
Example 3: Multi-page with group_by

Subject ID	Study Day	Adverse Event	Severity	Serious
01-702-1082	46.0	Application Site Irritation	MILD	N
01-702-1082	79.0	Skin Irritation	MODERATE	N
01-703-1042	3.0	Diarrhoea	MILD	N
01-703-1042	4.0	Insomnia	MILD	N
01-703-1076		Hypercholesterolaemia	MODERATE	N
01-703-1076	23.0	Biopsy Prostate	MODERATE	N
01-703-1076	27.0	Benign Prostatic Hyperplasia	MODERATE	N
01-703-1076	30.0	Application Site Pruritus	MILD	N
01-703-1076	30.0	Application Site Dermatitis	MILD	N
01-703-1076	30.0	Application Site Pruritus	MODERATE	N
01-703-1076	32.0	Hyperhidrosis	MILD	N
01-703-1086	12.0	Application Site Irritation	MILD	N
01-703-1086	12.0	Application Site Irritation	MODERATE	N
01-703-1086	12.0	Application Site Irritation	SEVERE	N

Note: In multi-page listings, group context is automatically restored at the beginning of each new page for better readability.