

**Supplemental Table 1 Treatment-related adverse events in this study and previous trials**

<b>Variables</b>	<b>PRODIGE4 /ACCORD11</b>	<b>FOLFIRINOX% (n = 86)</b>	<b>MPACT</b>	<b>Gem + nabPTX (n = 81)</b>
<i>Hematologic</i>				
Grade ≥ 3 Anemia	7.8% (n = 13)	19.8% (n=17)	13% (n=53)	14.8% (n=12)
Grade ≥ 3 Thrombocytopenia	9.1% (n = 15)	8.1% (n=7)	13% (n=52)	6.2% (n=5)
Grade ≥ 3 Neutropenia	45.7% (n = 75)	74.4% (n=64)	38% (n=153)	46.9% (n=38)
Febrile neutropenia	5.4% (n = 9)	25.6% (n=22)	3% (n=14)	16.0% (n=13)
<i>Nonhematologic</i>				
Grade ≥ 3 Nausea/Vomiting	14.5% (n = 24)	41.9% (n=36)	NA	9.8% (n=8)
Grade ≥ 3 Diarrhea	12.7% (n = 21)	3.5% (n=3)	6% (n=24)	7.4% (n=6)
Grade ≥ 3 Fatigue	23.6% (n = 39)	20.9% (n=18)	17% (n=70)	28.4% (n = 23)
Grade ≥ 3 neuropathy	9.0% (n = 15)	3.5% (n=3)	17% (n=70)	18.5% (n = 15)