

Supplemental Table for:
Attrition of Patients on a Precision Oncology Trial: Analysis of the I-PREDICT Experience
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Supplemental Table 1. Tumor Types of Consented Patients in the I-PREDICT Trial (University of California San Diego site)

Parameter	Evaluable	Inevaluable	Awaiting Treatment
Consented, <i>N</i> = 190	99 (52%)	83 (44%)	8 (4%)
Tumor type			
Blood, <i>n</i> = 3 (1%)	1 (33%)	2 (67%)	0
Breast, <i>n</i> = 5 (3%)	4 (80%)	1 (20%)	0
Central Nervous System, <i>n</i> = 7 (4%)	5 (71%)	2 (29%)	0
Endocrine, Neuroendocrine, <i>n</i> = 6 (3%)	3 (50%)	2 (33%)	1 (17%)
Gastrointestinal ^a , <i>n</i> = 103 (54%)	50 (49%)	49 (47%)	4 (4%)
Genitourinary, <i>n</i> = 5 (3%)	3 (60%)	2 (40%)	0
Gynecological, <i>n</i> = 27 (14%)	13 (48%)	12 (45%)	2 (7%)
Lung, <i>n</i> = 3 (1%)	2 (67%)	1 (33%)	0
Mesothelioma, <i>n</i> = 2 (1%)	0	2 (100%)	0
Oral, Head, and Neck; <i>n</i> = 10 (5%)	6 (60%)	3 (30%)	1 (10%)
Soft Tissue Sarcoma, <i>n</i> = 18 (10%)	11 (61%)	7 (39%)	0
Unspecified site, <i>n</i> = 1 (1%)	1 (100%)	0	0

Note: Data for evaluable, inevaluable, and awaiting treatment are presented as *n* (%) of the tumor type.

^aGastrointestinal tumor type includes hepatobiliary (16%, *n* = 16 of 103 patients) and pancreatic cancers (12%, *n* = 12 of 103 patients).