

(a)

| ID          | Age | Sex | TNM Stage | Treatment prior to PDX  | Patient Tumour Source* | Germline Mutation            |
|-------------|-----|-----|-----------|-------------------------|------------------------|------------------------------|
| <b>Q70P</b> | 47  | M   | IV        | Gemcitabine, FOLFIRINOX | Primary                | BRCA2 c.3167_3171delAAAA     |
| <b>O28</b>  | 53  | F   | IIA       | Gemcitabine - Cisplatin | Primary †              | BRCA2 c.5946delT             |
| <b>O217</b> | 57  | F   | IIA       | Gemcitabine - Cisplatin | Primary †              | BRCA1 c.4327C>T              |
| <b>O232</b> | 50  | M   | IIB       | Naïve                   | Primary                | BRCA2 c.3166delC             |
| <b>Q392</b> | 61  | F   | IIB       | Naïve                   | Primary                | BRCA2 c.8677C>T              |
| <b>S145</b> | 37  | M   | IV        | Naïve                   | Liver Metastasis †     | BRCA1 c.68_69delAG           |
| <b>Q437</b> | 76  | M   | IIA       | Naïve                   | Primary                | BRCA2 c.5065_5066delGCinsAAA |

\* Whole genome sequenced and used to generate PDX for the preclinical trial

† PDX used for WGS when patient sample insufficient

(b)

| ID   | Age | Sex | TNM Stage | Treatment prior to PDX | Patient Tumour Source* | Germline Mutation |
|------|-----|-----|-----------|------------------------|------------------------|-------------------|
| Q66  | 66  | M   | IIB       | FOLFIRINOX             | Primary                | None              |
| Q133 | 60  | M   | IIB       | Naïve                  | Primary                | None              |
| Q155 | 70  | M   | IIB       | Naïve                  | Primary                | None              |

\* Whole genome sequenced and used to generate PDX for the preclinical trial

**Supplementary Table 1. Clinical characteristics of patients from whom patient-derived xenografts were generated and evaluated in the preclinical trial. (a)**

Clinical characteristics of *gBRCA*-mutated PDAC cases. **(b)** Clinical characteristics of HR-proficient PDAC cases.