# MarketVue® Pancreatic Ductal Adenocarcinoma (PDAC)

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## MarketVue®: Pancreatic Ductal Adenocarcinoma

UNDERSTAND THE PDAC MARKET

MarketVue market landscape reports combine primary (KOL interviews and survey data) and secondary market research to empower strategic decision-making and provide a complete view of the market.

Every MarketVue includes a disease overview, epidemiology (US and EU5), current treatment, unmet needs, pipeline and access and reimbursement chapter.

Methodology: Research is supported by 7 qualitative interviews with key opinion leaders (4 U.S. oncologists, 1 UK oncologist, 1 French oncologist, and 1 Italian oncologist), a quantitative survey with 23 U.S. physicians and secondary research.

Geographies covered: United States plus epidemiology for EU5 (France, Germany, Italy, Spain, United Kingdom)

EPIDEMIOLOGY: Understand prevalence, diagnosed and drug-treated prevalence of the population and key market segments

CURRENT TREATMENT: Understand the treatment decision tree and strengths and weaknesses of current on-label and off-label treatment

UNMET NEEDS: Identify opportunities to address treatment or disease management gaps

PIPELINE ANALYSIS: Compare current and emerging therapy clinical development strategy; their performance on efficacy, safety, and delivery metrics; and their potential to address unmet needs

VALUE AND ACCESS: Review the evidence needed to assess and communicate value to key stakeholders (e.g., providers, payers, regulators) and learn what competitors have done or are doing

#### Why MarketVue?

- PMR-Driven Insights informed by qualitative interviews and/or quantitative surveys
- Senior Team Experienced team members (10+ years in pharma market research) lead the research
- **Strategic –** Delivered in a concise and strategic report template vetted by pharmaceutical industry professionals
- Fresh New reports or report refreshes delivered in as little as 15 business days





UNDERSTAND THE PDAC MARKET

#### **COMPANIES MENTIONED**

- Novartis
- Pancreatic Cancer Action Network
- Panbela Therapeutics
- BioLineRx
- Astellas Pharma
- TME Pharma
- Merck Sharp & Dohme
- SynerGene Therapeutics

- Ability Pharma
- AstraZeneca
- ERYTECH
- Cornerstone Pharmaceuticals
- Eli Lilly
- Pharmacyclics
- Halozyme Therapeutics
- Sumitomo Pharma

#### DRUGS MENTIONED

- Capecitabine
- FOLFIRINOX
- Gemcitabine
- Cisplatin
- Olaparib (Lynparza)
- Larotrectinib (Vitrakvi)
- Entrectinib (Rozlytrek)
- Pembrolizumah (Keytruda)
- Pamrevlumab
- Canakinumab (Ilaris)
- Spartalizumab
- SM-88
- Ivospemin

- Zolbetuximat
- Olaptesed pegol
- SGT-53
- ABTL0812
- Selumetinib (Koselugo)
- Durvalumab (Imfinzi)
- Paclitaxel (Abraxane)
- Eryaspase
- Devimistat
- Pegilodecakin
- Ibrutinib (Imbruvica)
- PEGPH20
- Napabucasir



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#### Meet the REACH Team







MELISSA CURRAN is the Director of Product Management at REACH. Melissa has over 10 years of life sciences market research and consulting experience spanning from bespoke strategy consulting to syndicated market research product development and management. Prior to joining REACH, she worked at Decision Resources Group (DRG) for 7 years assisting pharmaceutical and biotechnology commercial teams across the product lifecycle to inform strategic decision making. Melissa is particularly passionate about new product planning and portfolio management, especially in the rare disease space where data can be scarce, and decision-making can be challenging. Specific types of strategic assessments Melissa specializes in include market landscape assessments, commercial opportunity assessment, patient journey mapping, product positioning and TPP optimization, portfolio prioritization, and competitive intelligence. She also has extensive experience working across various market research methodologies including qualitative interviews, quantitative surveys, patient chart audits, real world claims and EHR data, conjoint analysis and secondary research. Melissa received her bachelor's degree in Biology and minor in Business from Providence College.



MICHAEL HUGHES, MSc, Ph.D., Dr. Hughes is the Director of Research at REACH. He has worked in academia, regulatory affairs (NICE) and in RWE and epidemiology consultancies, leading the global epidemiology team at Clarivate (previously Decision Resources Group) for many years. Over that period, he has built numerous new approaches to epidemiological forecasting and imputation, which now form industry best-practice. He has built syndicated and custom epidemiological models and forecasts for many blockbuster drugs across many therapeutic areas, often using a hybrid approach sourcing data from multiple types of dataset and primary market research. He has recently worked on projects in prostate cancer, amyloidosis, anaphylaxis and multi-drug resistant UTIs, among others. He has supported the needs of both big pharma, including Novartis, GSK and Johnson and Johnson, as well as smaller companies and biotechs.



#### Meet the REACH Team



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