

— TRISTAR TECHNOLOGY GROUP

Quality biospecimens, imaging & data for precision medicine

- ◆ FFPE surgical samples — Primary tumors, Mets, Normal (non-malignant) tissues
- ◆ Clinical follow-up data including treatment, response, OS & PFS
- ◆ Mutations, fusions and other diagnostic markers
- ◆ Donors from UK, Italy, France with informed consent & ethics approval
- ◆ Responders & non-responders to SOC treatment including CKI/IO
- ◆ Expert IHC capabilities — over 100 optimized assays

Imaging + Clinical Data.

Whole-slide imaging tied back to the matched surgical section and the clinical timeline behind every donor. Radiology coming soon.

RADIOLOGY

COMING SOON

Radiological images — capability in development.

PATHOLOGY

Whole-slide imaging (WSI)

CURRENT H&E REPOSITORY

30,000

donors

>100,000

H&E images

- **Aperio GT450 & AT2 scanners** — same platforms used in clinical practice
- **40x standard** scan resolution across the repository
- **.svs files** compatible with all major WSI viewers and AI pipelines
- **Scanned in-house** — no vendor coordination, no provenance gaps
- **Pathologist-reviewed** — every H&E and corresponding image checked by an experienced medical pathologist to confirm image quality is suitable for AI

LINKAGE

Every imaging asset is linked to

IMAGING

Whole-slide imaging

SURGICAL SECTION

Matched FFPE block & slides

Cohorts available

CLINICAL DATA

Treatment, response, OS/PFS, molecular

Donor Profiles

Breadth of donor profiles.

Specimen, treatment, outcomes, and molecular data — structured at the patient level.

— PATIENT TRAJECTORY

Primary & matched distant mets

Matched normal tissue

Neoadjuvant > adjuvant > SOC targeted & CKI/IO

— OUTCOMES, STRUCTURED

Overall Survival (months)

Progression-Free Survival (months)

Time to events (TTE months)

Last follow-up

Event flags (recurrence / death / composite)

— MOLECULAR CHARACTERIZATION

IHC

FISH

RNA-ISH

NGS: SNV, CNV, Fusions

Records are fully anonymized. Not every patient has every field populated, and not every project needs every field. Cohorts are pulled to fit the project; populated fields reflect real-world clinical practice at the source center.

[See an example record →](#)

Cohorts available.

Tumor types by cohort designs we maintain.

[JUMP TO COHORT DETAIL](#)

[Advanced \(III/IV\) →](#)

[Longitudinal →](#)

[Primary + matched mets →](#)

[Pre/Post SOC →](#)

[Pre/Post IO →](#)

TUMOR TYPE	ADVANCED (III/IV)	LONGITUDINAL	PRIMARY + MATCHED METS	PRE / POST SOC	PRE / POST IO
NSCLC	✓	✓	✓	✓	✓
CRC	✓	✓	✓	✓	—
Ovarian	✓	✓	✓	✓	—
Breast	✓	✓	✓	✓	—
Gastric	✓	✓	✓	✓	—
Head & Neck SCC	✓	✓	✓	✓	✓
Prostate (mCRPC)	✓	✓	—	✓	—
Endometrial	✓	✓	✓	✓	—
Pancreatic ADC	✓	—	—	✓	—
Bladder (UC)	✓	✓	✓	✓	—
Melanoma	✓	✓	✓	✓	✓
RCC	✓	✓	✓	✓	✓
HCC	✓	✓	✓	✓	—
Esophageal	✓	✓	✓	✓	—

All cohorts are available as FFPE, TMA, or slides. Treatment, response, OS/PFS data and indication-specific molecular characterization (IHC and mutation panels) are linked at the patient level.

A core partner for translational programs.

Why top pharma and biotechs working on oncology therapeutics work with us.

- ◆ **The translational bar has moved.** Sponsors developing oncology therapeutics need high-quality samples paired with in-depth, standardized data across EHR and orthogonal sources.
- ◆ **Ethically sourced, quality samples and data.** Quality samples and data improve the reliability of results.
- ◆ **Validation for regulators.** Data derived from TriStar samples is frequently included in IND submissions — tissue quality and annotation rigor shape what is defensible.
- ◆ **Validation for the program.** Translational evidence is increasingly what investment committees and partnership negotiations anchor on.
- ◆ **Translational evidence is increasingly the rate-limiting step.** Tissue quality, annotation depth, and provenance decide whether downstream analysis withstands scientific and regulatory scrutiny.

Lab capabilities — validation, extension & characterization.

Run in-house on the same donors and samples behind every cohort — no vendor coordination, no provenance gaps.

100+

OPTIMIZED IHC ASSAYS

DEVELOPED FOR

- Ventana Benchmark
- Dako autostainers

3

MEDICAL PATHOLOGISTS

15+ combined years in pharma

SERVICES

- IHC
- NGS
- Digital Spatial Profiling
- RNAScope
- RNA-Seq
- Digital image analysis

Advanced disease (Stage III/IV) surgical samples

— APPROXIMATE NUMBER OF DONORS **500–900 per tumor type**

— TUMOR TYPES

NSCLC

CRC

Ovarian

Prostate (mCRPC)

Breast

Gastric

Endometrial

Pancreatic ADC

Bladder (UC)

Melanoma

RCC

HCC

Esophageal

Head & Neck SCC

— FOLLOW-UP DATA

Treatment

Response

OS/PFS

— MUTATIONS*

KRAS

NRAS

EGFR

ALK fusion

BRAF

BRCA

MET alteration

HRD status

MSI status

** as available from donor records*

— IHC MARKERS

ER/PR

HER2

MMR IHC

PD-L1

TROP2

Nectin-4

B7-H3

B7-H4

DLL3

p16

p53

— HBS FORMATS

FFPE

TMA

Slides

[Example HBS Cohorts & Data Available](#) [↗](#)

Sequentially collected (longitudinal) surgical samples from the same donor

— APPROXIMATE NUMBER OF DONORS **20–70 sets per indication**

— TUMOR TYPES

NSCLC CRC Ovarian Head & Neck SCC Breast Gastric Endometrial RCC Bladder (UC) Melanoma

— FOLLOW-UP DATA

Treatment Response OS/PFS

— MUTATIONS*

KRAS NRAS EGFR ALK fusion BRAF BRCA MET alteration HRD status MSI status

** as available from donor records*

— IHC MARKERS

ER/PR HER2 MMR IHC PD-L1 TROP2 Nectin-4 B7-H3 B7-H4 DLL3 p16 p53

— HBS FORMATS

FFPE TMA Slides

[Example HBS Cohorts & Data Available](#) [↗](#)

Longitudinal Example: NSCLC.

Per-patient. Per matched metastasis. Treatment-line resolved. Examples below are anonymized records from the repository.

Patient #001

Immunotherapy: **No**

Treatment status: **Treatment naïve**

PRIMARY SPECIMEN

Surgery: Left lung upper lobectomy and lymph node sampling
Site: Lung, left upper lobe

MATCHED MET 1

Site: Superficial cerebrum and dura
Treatment: Immunotherapy naïve

Patient #002

Immunotherapy: **No**

Treatment status: **Treatment naïve**

PRIMARY SPECIMEN

Surgery: Left Thoracotomy and upper lobectomy with wedge of the lower lobe taken en bloc. Sleeve resection of PA
Site: Left lung, upper lobe

MATCHED MET 1

Site: Lymph node
Treatment: Treatment naïve

MATCHED MET 2

Site: Lymph node
Surgery date: Q1-2018
Treatment: Treatment naïve

Patient #007

Immunotherapy: **No**

MATCHED MET 1

Site: Lymph node
Treatment: Treatment naïve

MATCHED MET 2

Site: Lymph node
Treatment: Treatment naïve

MATCHED MET 3

Site: Hilar Lymph node
Surgery date: Q1-2022
Treatment: Treatment naïve
Block #: C7

[Example HBS Cohorts & Data Available](#)

Primary & matched distant metastases (surgical samples)

— APPROXIMATE NUMBER OF DONORS **20–50 sets per indication**

— TUMOR TYPES

NSCLC

CRC

Ovarian

Head & Neck SCC

Breast

Gastric

Endometrial

Bladder (UC)

Melanoma

RCC

— EXAMPLES OF METASTATIC SITES

Lymph nodes

Bone

Liver

Lung

Brain

Peritoneal

**Mets only (without matched primary) samples available*

— FOLLOW-UP DATA

Treatment

Response

OS/PFS

— MUTATIONS*

KRAS

NRAS

EGFR

ALK fusion

BRAF

BRCA

MET alteration

HRD status

MSI status

**As available from donor records.*

— IHC MARKERS

ER/PR

HER2

MMR IHC

PD-L1

TROP2

Nectin-4

B7-H3

B7-H4

DLL3

p16

p53

— HBS FORMATS

FFPE Slides

TMA Block or slides

[Example HBS Cohorts & Data Available](#) [↗](#)

Pre & Post standard of care (SOC) treatment surgical samples

Includes a sub-set of donor matched sets.

— APPROXIMATE NUMBER OF DONORS **25–150 per indication**

— TUMOR TYPES

NSCLC

CRC

Ovarian

Head & Neck SCC

Breast

Gastric

Endometrial

Pancreatic ADC

Bladder (UC)

Melanoma

RCC

— FOLLOW-UP DATA

Treatment

Response

OS/PFS

— MUTATIONS*

KRAS

NRAS

EGFR

ALK fusion

BRAF

BRCA

MET alteration

HRD status

MSI status

** as available from donor records*

— IHC MARKERS

ER/PR

HER2

MMR IHC

PD-L1

TROP2

Nectin-4

B7-H3

B7-H4

DLL3

p16

p53

— HBS FORMATS

FFPE

TMA

Slides

[Example HBS Cohorts & Data Available](#) [↗](#)

Pre & Post IO (Pembro/Nivo) treatment surgical samples

1st line SOC chemo, 2nd line IO. Includes a sub-set of donor matched sets.

— APPROXIMATE NUMBER OF DONORS **10–25 per indication**

— TUMOR TYPES

NSCLC

RCC

Head & Neck SCC

Melanoma

— FOLLOW-UP DATA

Treatment

Response

OS/PFS

— MUTATIONS*

KRAS

NRAS

EGFR

ALK fusion

BRAF

BRCA

MET alteration

HRD status

MSI status

** as available from donor records*

— IHC MARKERS

ER/PR

HER2

MMR IHC

PD-L1

TROP2

Nectin-4

B7-H3

B7-H4

DLL3

p16

p53

— HBS FORMATS

FFPE Slides

TMA Block or slides

[Example HBS Cohorts & Data Available](#) [↗](#)

FFPE blocks & TMAs with matched double-spun plasma.

FFPE block & 4ml matched double-spun plasma or whole blood.

~330

TOTAL SAMPLE SETS

8

INDICATIONS

40 / 60

STAGE I-II / III-IV (%)

5-20_{min}

ISCHEMIC TIME

INDICATION	FFPE BLOCKS	PLASMA / BLOOD	NGS-ELIGIBLE [†]
NSCLC	50	●	●
CRC	100	●	●
Gastric ADC	30	●	●
RCC	30	●	●
Esophageal cancer	40	●	●
Endometrial cancer	50	●	●
Ovarian cancer	20	●	●
Melanoma	10	●	●

COMPATIBLE ASSAYS

IHC

RNA-ISH

NGS (Oncomine)

[†] NGS (Oncomine) panel can be ordered separately on any indication.
Note — these sets do not have outcome data and were collected from Ukraine.
Quantities are approximate and subject to change.

Example HBS Cohorts & Data Available [↗](#)

GET IN TOUCH

Let's talk about your program.

Quality biospecimens, imaging & data for precision medicine — sourced, characterized, and delivered in-house.



HEAD OF PROJECTS & IO APPLICATIONS

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