

RANDOX

EVIDENCE MULTISTAT

Clinical Diagnostics



EVIDENCE
MULTISTAT

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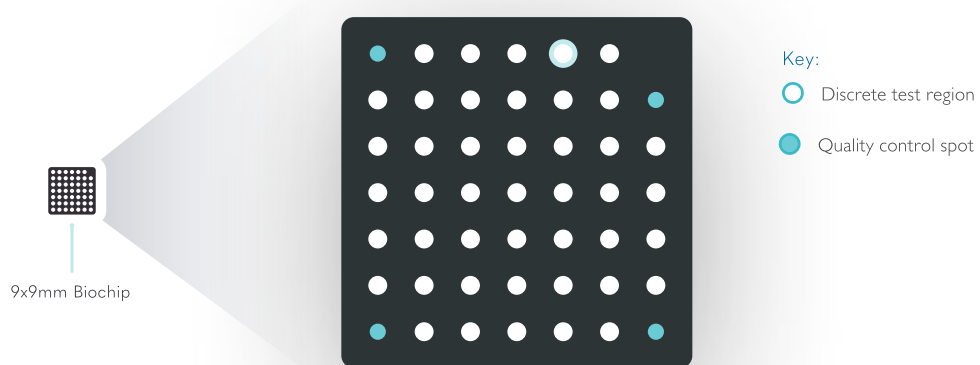
Technical Snapshot

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BIOCHIP ARRAY TECHNOLOGY

Biochip Array Technology enables rapid and precise detection of multiple analytes from a single patient sample.

The biochip is a solid-state device with discrete testing regions onto which antibodies specific to different analytes are immobilised and stabilised. Competitive or sandwich chemiluminescent immunoassays are then employed, offering a highly sensitive screen.



Biochip in Numbers



153 MILLION

Tests performed globally using Biochip Array Technology.



£440 MILLION

Invested into Biochip Array Technology.



203

Number of patents across the Biochip product range.



12,195

Tests within the Biochip portfolio.

Algorithm-Based Testing

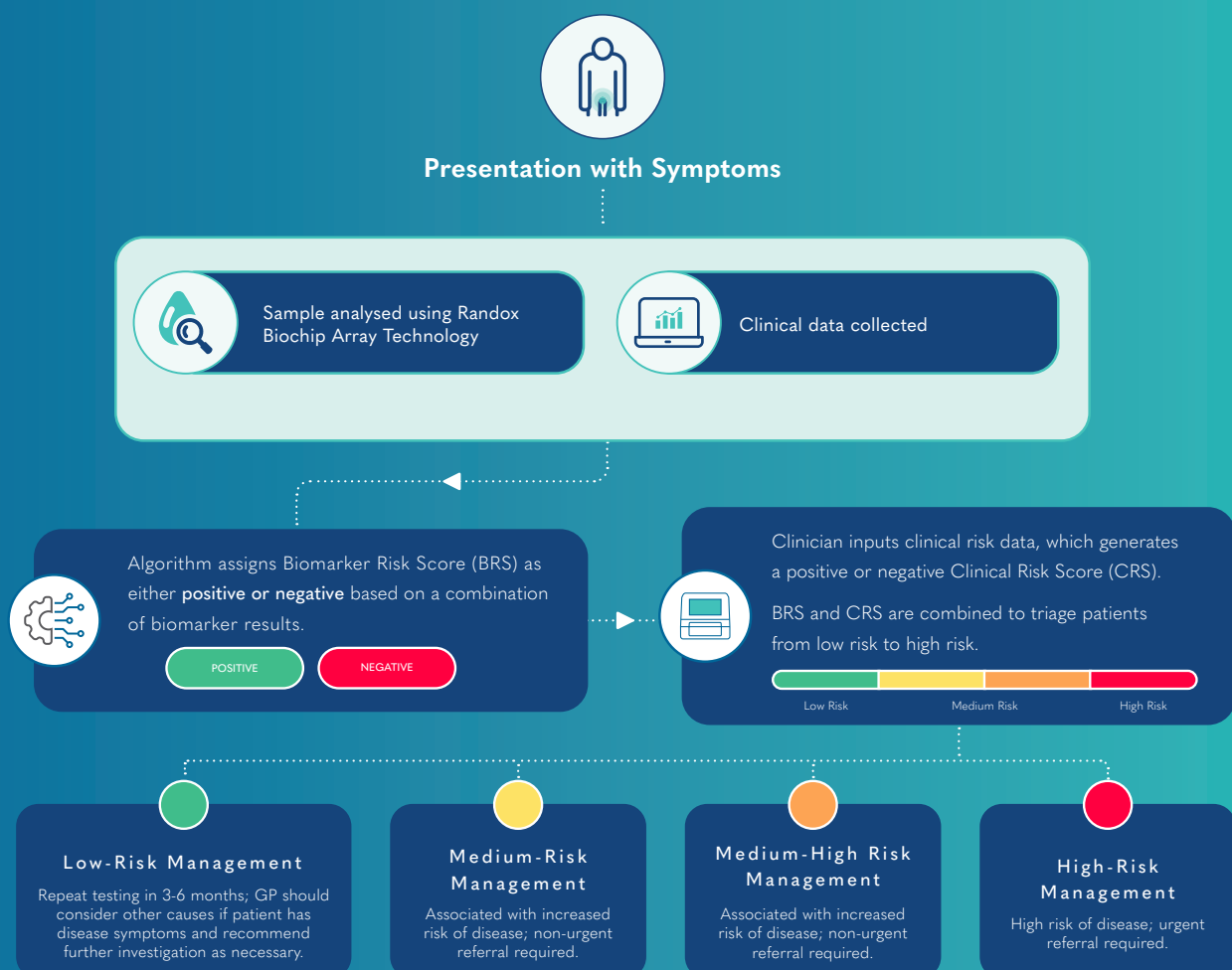
Biomarker algorithms can be applied clinically to some of our Biochip panels to stratify patients from 'low' to 'high' risk of developing certain diseases.

Biomarker Risk Score (BRS)

A BRS is calculated for each patient by applying an algorithm to the quantitative biomarker results. Patients with a score above the cut-off value would be positive, whereas patients below the cut-off would be negative.

Clinical Risk Score (CRS)

A CRS is calculated for each patient; this is a cumulative score utilising clinical and demographic measurements which vary depending on the disease. Patients with a score above the cut-off value would be positive, whereas patients below the cut-off would be negative.



Biochip Benefits



Improve Patient Management

Make better patient management decisions to improve patient outcomes.



Reduce Secondary Care Referrals

Reduce the number of low-risk patients being referred to secondary care, reducing the current healthcare burden.



Monitor Treatment Efficacy and Disease Progression

By measuring biomarkers, Biochip can monitor treatment efficacy and determine if the disease is getting better or worse. This allows for quick adjustments to treatment plans.



Reduce Invasive Procedures

Reduce the need for time consuming scans, as well as risky, invasive and costly biopsies and endoscopies in low-risk patients.



Expedite High-Risk Patients

Expedite those with high-risk scores for urgent investigations, allowing earlier intervention and treatment.



Reduce Costs

Biochip can help utilise healthcare resources more effectively, improve preventative care and reduce associated costs with progressed diseases.

Clinical Diagnostic Biochips

Our risk stratifying biochips use an AI algorithm to triage patients from 'low' to 'high' risk of developing certain diseases, improving patient management at primary care.



Bladder Cancer Risk

Triaging haematuria patients

Risk Stratifying Biochip



Prostate Cancer Risk

Triaging patients with prostate cancer symptoms

Risk Stratifying Biochip



Ovarian Cancer

Quantifying CA 125 to aid detection of epithelial ovarian cancer



Male & Female Hormones

Providing a range of quantitative biochips for fertility, SHBG, Testosterone, AMH, and hCG

BLADDER CANCER RISK

Biochip provides **risk stratification** to triage female patients presenting with haematuria at primary care from low to high risk of bladder cancer. This could allow low-risk patients to be monitored in primary care, while medium and high-risk patients can be referred urgently for secondary care investigations.

Randox investigated 48 urine and 32 serum biomarkers known to be involved in the pathobiology underlying bladder carcinogenesis in a recent study (Duggan et al., 2022). Four markers were demonstrated to be clinically significant in identifying female patients with bladder cancer.

Applications



Primary Care GPs



Urology Departments



Hospital Labs



Oncology Departments

Biochip Markers

Biomarkers	
Interleukin-12p70 (IL-12p70)	Interleukin-13 (IL-13)
Midkine	Clusterin

Product Information



Sample Type
Urine



Time to Result
60 Minutes



Sample Volume
400 µL



Result
Quantitative



Samples per Cartridge
2

49%

Of bladder cancer cases are preventable

\$101k

Could be saved per patient with effective screening and early detection

614,298

New bladder cases globally in 2022

Biochip Benefits

1

Highly Accurate in Identifying Patients at Risk

Our biomarker combination showed a sensitivity of 83.7% and specificity of 79.7% for accurate risk-stratification of female haematuria patients.

2

Reduce Secondary Care Referrals

Reduces the need for costly, risky, and invasive cystoscopy referrals for low-risk patients, decreasing post-procedure complications.

3

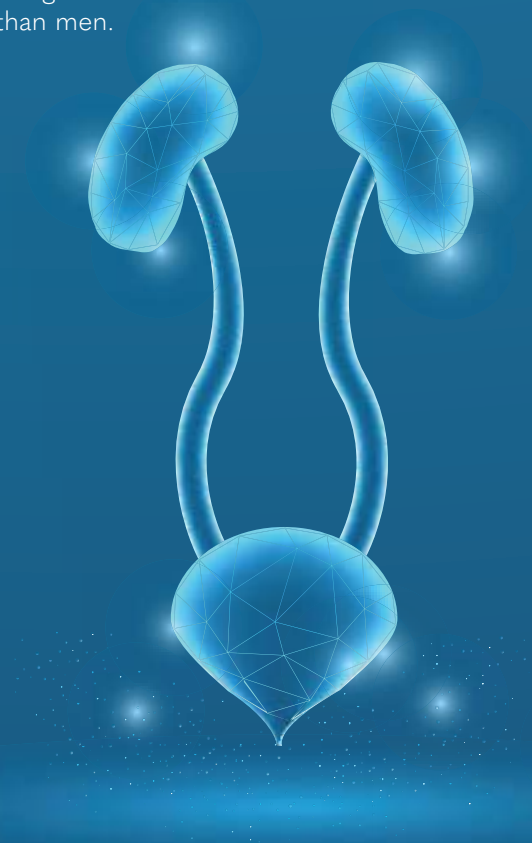
Reduce Underdiagnosis in Women

Biochip's novel combination of biomarkers could help clinicians reduce undiagnosed bladder cancer in women, who are often diagnosed at more advanced stages and have poorer survival rates than men.

4

Reduce Costs

Biochip can help utilise healthcare resources more effectively, improve preventative care and reduce associated costs with progressed diseases.



PROSTATE CANCER RISK

Biochip provides **risk stratification** to triage patients presenting with symptoms at primary care from low to high risk of prostate cancer. This could allow low-risk patients to be monitored in primary care, while medium and high-risk patients can be referred urgently for secondary care investigations.

Randox investigated **19 biomarkers** in a recent study (McNally et al., 2022), identifying **4 novel markers** which were demonstrated to be clinically significant in identifying patients with prostate cancer.

Applications



Primary Care GPs



Urology Departments



Hospital Labs



Oncology Departments

Biochip Markers

Biomarkers	
Epidermal Growth Factor (EGF)	Interleukin-8 (IL-8)
Monocyte Chemoattractant Protein-1 (MCP-1)	Total Prostate-Specific Antigen (tPSA)

Product Information



Sample Type
Serum



Time to Result
60 Minutes



Sample Volume
250 µL



Result
Quantitative



Samples per Cartridge
2

\$28k/year

Is the average cost of early-stage treatment

\$74k/year

Is the average cost to treat advanced prostate cancer

1,414,259

New prostate cancer cases globally in 2022

Biochip Benefits

1

Improved Predictive Potential

Biochip prostate cancer risk score accuracy for detecting prostate cancer is 86% vs 70% for elevated tPSA alone.

2

Independent of Body Mass Index (BMI)

Our prostate cancer risk score is independent of BMI; a high BMI can lower tPSA levels.

3

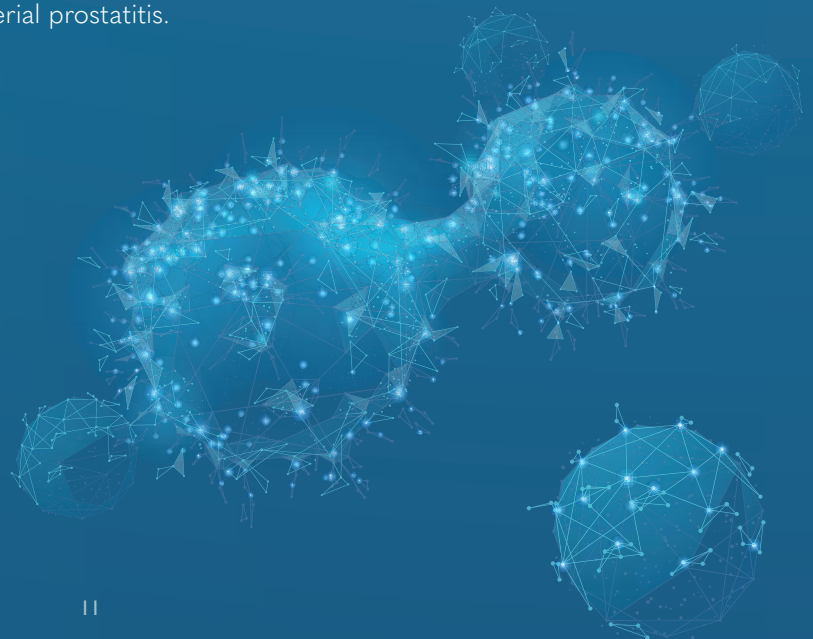
Reduces Unnecessary Biopsies

Reduces unnecessary referrals to secondary care for invasive and costly biopsy investigations - 75% of biopsies from raised tPSA and Digital Rectal Exam referrals are negative.

4

Reduces Hospital Readmissions

Reduces hospital admissions from post-biopsy complications such as infections and bacterial prostatitis.



OVARIAN CANCER

National Institute for Health and Care Excellence (NICE) advises that the best available marker for epithelial ovarian cancer is CA 125 due to the combination of reliability and general availability. Biochip provides the quantitative detection of CA 125 in a serum sample for epithelial ovarian cancer.

Serum CA 125 measurement, an abdominal and pelvic ultrasound, along with the woman's menopausal status, are used to assess whether a referral for histology or cytology is required. CA 125 levels after chemotherapy is one of the strongest available indicators of disease outcome, and the most important application for monitoring patients with epithelial ovarian cancer.

Applications



Primary Care GPs



Gynaecology Departments



Hospital Labs



Oncology Departments

Biochip Markers

Biomarker	
CA 125	

Product Information



Sample Type
Serum



Time to Result
<40 Minutes



Sample Volume
250 µL



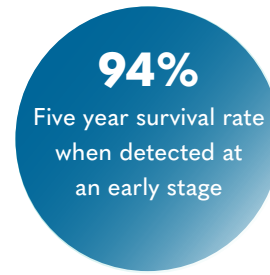
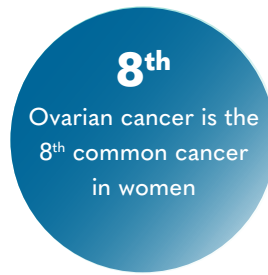
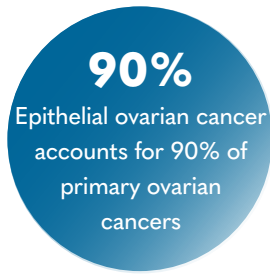
Result
Quantitative



Range
Up to 4,000 U/mL

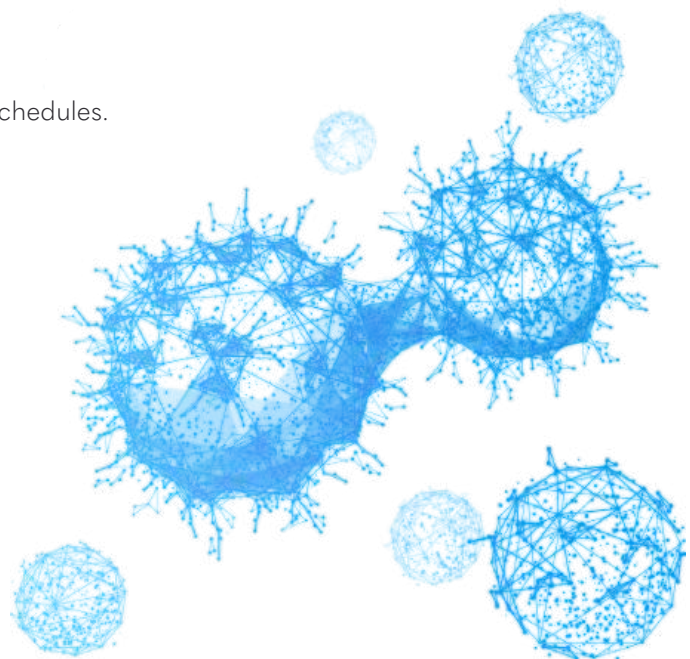


Samples per Cartridge
2



Biochip Benefits

- 1 Wide Measuring Range**
With a measuring range of up to 4,000 U/mL, treatment effectiveness can be monitored. Decreasing biomarker levels indicate efficacy, while stable or increasing levels signal for treatment adjustment.
- 2 Rapid Detection**
Receiving results in <40 minutes aids clinicians in making fast decisions about whether to refer a patient for urgent investigations.
- 3 Ease of Use**
Allowing samples to be processed in primary care, reducing turnaround times, transport issues, and administrative tasks.
- 4 No Sample Batching**
Providing flexibility in testing schedules.



MALE & FEMALE HORMONES

Screening male and female hormones is vital for early detection of health issues. This allows healthcare providers to tailor treatments effectively, improving outcomes for conditions such as infertility, hormonal imbalances and menopause.

Applications



Primary Care GPs



Fertility Health Clinics



Hospital Labs

Fertility Biochip

The Fertility Biochip is an advanced tool that provides a detailed understanding of baseline hormone levels and any imbalances that may be causing adverse effects on a person's overall health or fertility, enabling the tracking of changes over time.

Our Fertility Biochip can provide a full diagnostic hormone profile which can indicate various conditions such as polycystic ovary syndrome (PCOS), menopause, hormone imbalances affecting ovulation and ovulation indication. For patients going through in vitro fertilisation (IVF), the Fertility Biochip can determine the menstrual cycle stage and response to fertility treatment.

Biochip Markers

Biomarkers	
Prolactin	Estradiol
Follicle Stimulating Hormone (FSH)	Progesterone
Luteinizing Hormone (LH)	-

Product Information



Sample Type
Serum



Time to Result
31 Minutes



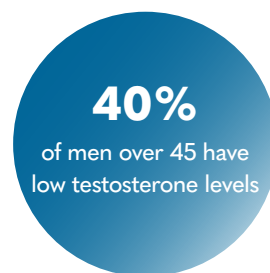
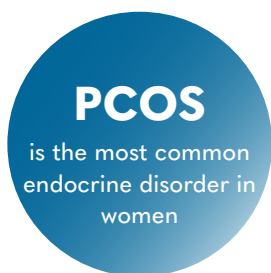
Sample Volume
285 µL



Result
Quantitative



Samples per Cartridge
2



Sex Hormone-Binding Globulin (SHBG) Biochip

The SHBG Biochip is primarily measured alongside a total testosterone test to estimate the amount of free testosterone in the patients' blood. In men it is used to look for testosterone deficiency, whereas in women the primary function for a SHBG test would be to investigate excess production of testosterone.

Biochip Marker

Biomarker
Sex Hormone-Binding-Globulin (SHBG)

Product Information



Sample Type
Serum



Time to Result
34 Minutes



Sample Volume
110 μ L



Result
Quantitative



Samples per Cartridge
2

Testosterone Biochip

The Testosterone Biochip can be used to aid diagnosis and treatment of conditions such as infertility, primary and secondary hypogonadism, testicular failure, androgen resistance and PCOS.

Biochip Markers

Biomarker
Total Testosterone

Product Information



Sample Type
Serum



Time to Result
31 Minutes



Sample Volume
275 μ L



Result
Quantitative



Samples per Cartridge
2

Anti-Müllerian Hormone (AMH) Biochip

Anti-Müllerian Hormone (AMH) testing offers critical insights into a woman's ovarian reserve and egg count. Biochip quantifies the concentration of AMH, a hormone produced by ovarian follicles. Elevated AMH levels indicate a higher ovarian reserve, whereas diminished levels indicate a reduced level. AMH testing can aid clinicians in assessing the body's responsiveness to ovarian stimulation protocols.

Biochip Marker

Biomarker
Anti-Müllerian Hormone (AMH)

Product Information



Sample Type
Serum



Time to Result
60 Minutes



Sample Volume
275 µL



Result
Quantitative



Samples per Cartridge
2

Human Chorionic Gonadotropin (hCG) Biochip

Human Chorionic Gonadotropin (hCG) testing measures the levels of hCG, a hormone produced by the placenta during pregnancy. This hormone is crucial for maintaining pregnancy as it supports the uterine lining and prevents menstruation. Testing for hCG can be used to confirm pregnancy, estimate age of the foetus, identify abnormal pregnancies, and screen for potential miscarriages. Biochip uses serum samples as blood tests are more precise and can detect hCG levels at an early stage after conception.

Biochip Marker

Biomarker
Human Chorionic Gonadotropin (hCG)

Product Information



Sample Type
Serum



Time to Result
37 Minutes



Sample Volume
275 µL



Result
Quantitative



Samples per Cartridge
2





QC



RANDOX

—evidence—
MULTISTAT

Evidence MultiSTAT

In the Non-Critical Care Setting

Using our revolutionary Biochip Array Technology, the Evidence MultiSTAT is a fully automated analyser that enables the detection of up to 48 targets simultaneously from a single patient's sample.



ANALYSER OVERVIEW

The Cartridge



- 1 Well One**
Cut-off material is added (qualitative kits) or adjuster/QC/sample is added (quantitative kits).
- 2 Well Two**
Adjuster/QC/sample is added.
- 3 Foil Cover & Fluid Reservoirs**
All additional fluids required are stored here.
- 4 Biochip Wells**
Two biochips are located here. Each biochip has up to 48 discrete testing regions.

Three Step Process



Prepare sample &
add to cartridge



Load reagent & tip
cartridge to MultiSTAT



Press Play



Rapid Screening

Minimal sample preparation is required, and results for 2 samples can be provided in under 30 minutes, allowing for quicker clinical decision and timely patient management

The Analyser



- 1 Touch Screen**
A large touchscreen interface allows the user to easily navigate through the analyser and view results.
- 2 Tip Cartridge Drawer**
The user will insert the prefilled tip cartridge here prior to testing.
- 3 Reagent Cartridge Drawer**
The user will insert the reagent cartridge here prior to testing.
- 4 2 x USB Ports**
USB Ports allow the user to add accessories, for example, barcode scanner, printer, or USB to export test results.

Benefits



No-Fuss Procedure

Pre-filled reagent cartridges and a simple interface mean that minimal laboratory training is required. This versatile benchtop analyser can achieve accurate, quantitative results in minutes.



Multi-Panel

The Evidence MultiSTAT can run a variety of panels, and test for multiple markers, facilitating comprehensive near-patient testing

TECHNICAL SNAPSHOT

Dimensions	585 (H) x 535 (D) x 570 (W) mm
Weight	48 kg, 106 lbs
Analyser Description	Fully automated touchscreen biochip array analyser
Biochip Format	Cartridge based system – assay reagents sealed in a pre-filled cartridge
Data Back-up Methods	Data export functionality via USB
Measurement Principal	Competitive and sandwich techniques with chemiluminescent reaction
Accreditation	FDA Approved, CE marked, Health Canada approved, ANVISA, GIVD, TGA and SFDA MDMA authorised
Sample Loading	Single cartridge loading bay

Also Available

Critical Care Biochips



Hyperinflammation



InflamiSTRAT



Neurovascular Dysfunction



Kidney Dysfunction



Clinical Drug Testing

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