

Sample Receipt Details:

POD : _____ Temp : _____
 Date & Time : _____ Sample Type : _____
 CS _____ Logistics _____
 Name & Sign: _____ Name & Sign : _____
 Prenatal Sample Yes No Bill type

TEST REQUISITION FORM

Disease Segment* _____
 Each sample must be accompanied by this completed requisition. * Fields are mandatory

Test Details

Tumour HRR (Homologous Recombination Repair) pathway genes analysis by NGS
 Androgen receptor (AR) gene analysis[Expedited TAT]

Test Name:* _____ Test Code:* **MGM1623, MGM599**

Sample type:

<input checked="" type="checkbox"/> Blood (in EDTA tube)	<input type="checkbox"/> Blood (in streck tube)	<input type="checkbox"/> DNA, Specify Source: _____	<input type="checkbox"/> Buccal swab
<input type="checkbox"/> Amniotic Fluid	<input type="checkbox"/> CVS	<input type="checkbox"/> Cultured CV	<input type="checkbox"/> Cultured amniocytes
<input type="checkbox"/> Fetal Blood (PUBS)	<input type="checkbox"/> Maternal blood for MCC (please send for prenatal studies)	<input type="checkbox"/> Products of Conception (POC), specify tissue: _____	<input checked="" type="checkbox"/> FFPE tissue Block (Block no.)
<input type="checkbox"/> Fresh Frozen Tissue	<input type="checkbox"/> Saliva	<input type="checkbox"/> Other sample type (specify site) _____	<input type="checkbox"/> DBS/FTA

PC319B1 , PC319C1
 PC319B2 PC319C2
 PC319B3

Peripheral blood (5 ml in EDTA) 3 Tubs

Patient had a blood transfusion Yes No Date of last transfusion ___/___/___ (minimum 3 days of wait time is required for genetic testing)
 Has he/she undergone allogenic bone marrow transplant: Yes No. **5 Waz Blocks**

Patient Details

Name:* **Mr. Ruwan Lokuge** (In Capital Letters) D.O.B. **DD MM YY** Age:* **67Y/M** Gender:* **M / F**
 Address: _____
 Phone: _____ E-mail I.D: _____

Clinician Details

Clinician's Name:* **Dr. Mahendra Perera** Hospital Affiliation: **Aegle Omics Pvt Ltd**
 Address: _____ Phone : _____
 _____ Email id : _____
 Date of sample collection* **3/3/2025** YY **27/ 1/2025**

I understand that the current analysis is limited to variants which co-relate with disease phenotype/symptoms/terms as mentioned in the clinical details provided by me. Incidental findings which may or may not be actionable are not routinely reported. They can however be provided on request after informed consent from the patient/guardian. As disease phenotype may evolve over time, the appearance of new symptoms/signs may alter test results or their significance: MedGenome laboratories cannot be held responsible for this. A re-analysis or a re-test may be required due to the former; this will be performed (if deemed necessary) at an additional cost. I am authorised to order the above tests as I am the treating physician/consulting physician in this case. I confirm that the patient/guardian (in case of minors) has been provided complete information regarding the test, including its limitations in a language of their understanding.

Dr. MAHENDRA PERERA
 MBBS (Cey), MD (Col), Dip RT
 Consultant in Clinical Oncology
 & Radiotherapy

Medical Professional Signature* _____ Date: _____ Place: _____
 Clinical notes/diagnosis: _____

Disease affection status Parental consanguinity present Age of manifestation: _____
 Affected Siblings Details: _____

GOVERNING LAW, JURISDICTION AND DISPUTE RESOLUTION

These Terms and Conditions and this Test Requisition Form shall be governed by and construed in accordance with Indian law and the courts in Bangalore shall have exclusive injunctive jurisdiction. In the event of any dispute, controversy or claim whatsoever arising from these Terms and Conditions and/or this Test Requisition Form, the parties shall undertake to make every effort to reach an amicable settlement within fifteen (15) days upon reference of the dispute by any party through discussions among the concerned representatives of parties, failing which the dispute, controversy or claim shall be settled by Arbitration by a Sole Arbitrator appointed by the 'President-Arbitration Centre-Karnataka', Bangalore as per Indian Arbitration and Conciliation Act, 1996 as amended from time to time. The venue of arbitration shall be Bangalore and it shall be conducted in English language. The award passed by the Sole Arbitrator shall be final and binding upon the parties.

INDEPENDENT PARTIES

All parties effected hereunder are independent entities and neither of the parties are an agent, employee or joint venture of the other and they shall not represent themselves as such to any third parties.

REFUND

Refund of fees for any reason has to be claimed by the Patient or the guardians of the Patients within 90 days from the date of delivery of report.

NOTICE

All notices, statements or other communication required or permitted to be given or made shall be in writing and in English language. Such notices will deliver by hand or sent by prepaid post with recorded delivery, or facsimile transmission addressed to the intended recipient at the address mentioned in this Test Requisition Form.

Patient/Guardian Authorization

By my signature below I attest to the following:

I have read and I understand the information provided on this form.

Patient Consent (sign here or on the consent document)

I have read the Informed Consent document and I give permission to MedGenome to perform genetic testing as described. I also give permission for my specimen / genetic data to be used in (de-identified) studies at MedGenome to improve genetic testing for other patients.

By agreeing to this informed consent below, I am confirming that I understand the benefits, risks and limitations associated with genetic testing. Furthermore, I am affirming that I recognize the seriousness of conditions for which {I am/my child} being tested, and that disease descriptions, prognoses, and treatment options have been made available to me by {my/my child's} health care provider. Finally, if I have the legal authorization to provide this informed consent on behalf of another person, I am attesting that the sample provided belongs to that person.

Patient/Guardian Name Mr. Ruwan Lokuge
 First Name Middle Name Last Name Date of Birth: mm/dd/yyyy

Patient/Guardian Signature* _____ Date: _____ Place: _____

Father Name _____ Mother Name _____

Signature*  _____ Date and time _____ Signature* _____ Date and time _____

Relationship with the proband _____

Note :

Signature of both parents is requested for prenatal testing.
 For trio testing, each parent should provide separate informed consent for the sequencing of his or her sample.

24 FEB 2025

Co. Sw Lu

Dr. Perera

S. AR
BRACA II,
(HRP Power)

Dr. MAHENDRA PERERA
MBBS (Cey), MD (Coll. Dip RT)
Consultant in Clinical Oncology
& Radiotherapy
Principal Investigator - Clinical Trials

CONFIDENTIAL LABORATORY REPORT

Member of Clinical and Laboratory Standards Institute, U.S.A.

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** OPD/AHH/ALS **

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REFERENCE No. : 01 0652 27/01/25
 SAMPLE DATE & TIME : 27/01/2025 19:37 AGE : 67 Y/M
 REPORT DATE & TIME : 04/02/2025 10:34 AHH2099932 / ahh6070
 PATIENT : MR. R.K. LOKUGE (UHID-120250971) 112C
 REFERRED BY : DR (MRS) LAKMALIE PARANAHEWA

TEST : TRUS BIOPSIES

Clinical features :- Increased PSA (12 ng/ml).

Specimen :- TRUS guided biopsies of the prostate.

Macroscopy :- A. Right lobe anterior : 2 cores of tissue measuring 14 and 12mm in length.

B. Right lobe middle : 3 cores of tissue measuring 19, 16 and 14mm in length.

C. Right lobe posterior : 2 cores of tissue measuring 19 and 18mm in length.

D. Left lobe anterior : 2 cores of tissue measuring 22 and 13mm in length.

E. Left lobe middle : 3 cores of tissue measuring 18, 17 and 17mm in length.

F. Left lobe posterior : 2 cores of tissue measuring 19 and 16mm in length.

Microscopy :- A to F. Sections confirm the presence of an acinar type prostate carcinoma predominantly composed of separate well-formed, small neoplastic glands (Gleason grade 3) This pattern account for about 60% of the tumour. Fused neoplastic glands and ill-formed small neoplastic glands constitute about 40% of the tumour volume (Gleason grade 4). The tumour is seen in 6 out of 7 cores from the right lobe where it occupies 25% to 70% of the surface areas of the cores (average 43.3%). The tumour is not seen in cores from the left lobe. There is no evidence of vascular or perineural invasion.



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Details of tumour grade and percentage involvement

A1 : 3+3 - 25%.
A2 : Not involved.
B1 : 3+4 - 50%
B2 : 3+4 - 70%
B3 : 4+3 - 25%
C1 : 4+3 - 50%
C2 : 3+4 - 40%
D1,D2,E1,E2,E3,F1,F2 : Not involved.

Conclusion :- TRUS guided biopsies of the prostate

1. Tumour type : Acinar type prostatic adenocarcinoma.
2. Gleason sum score : 3 (60%) + 4 (40%) = 7
3. Grade group (ISUP) : Group 2
4. Tumour burden : Percentage of positive biopsy cores = 42.8%
The tumour occupies 25 to 70% of the surface areas of the affected cores (average 43.3%).
Maximum length involved = 11mm
5. Perineural invasion : Absent.
6. Vascular invasion : Absent.

PC - 319 (S.C.T. 27/01/2025 at 9.15 am)

Chandu de Silva

PROF. CHANDU DE SILVA
MBBS, D.Path, MD (Histopathology), FCPATHSL, FCSSL, MIAC
Professor of Pathology