



Barcode

GENETIC TEST REQUISITION FORM

Test Code: **SLS162800** Test Name: **Germine Homologous Recombination Repair Test (gHRR)** TR ID:
 Test Code: Test Name: TR ID:
 Test Code: Test Name: TR ID:
 Test Code: Test Name: TR ID:

PATIENT DETAILS

Patient MRN/UHID #:
 Patient Name: **Mr.D.M.A.D.B Dissanayake** DOB: Age: **54** Sex: **M**
 Marital Status: Nationality: Contact No.:
 Address: **Kandy Sri Lanka**
 Email ID: Pincode:
 ID Proof: Driving License Aadhaar Card Voter ID card Ration card Others:

ADDITIONAL DETAILS IF REQUIRED

Transfusion Date (if available): Blood Group:

CLINICIAN INFORMATION

Referring Clinician: **Dr. Senaka Kandegedara** Clinician Contact:
 Referring Hospital: **General Hospital Kandy Sri Lanka**
 Email Id:
 Address:

ADDITIONAL FAMILY MEMBERS' DETAILS

Disease Status - Affected: Age at Diagnosis:
 Affected Sibling/Family members: Yes No (if yes, provide the details in the table below)

Name	Relation with Patient	Type of cancer	Age at Diagnosis	Sex

SAMPLE COLLECTION INFORMATION

Date & Time of Collection: Sample collected by:
 Clinical History/Pathology Report Attached: **Y** H and E Slides sent:

FOR OFFICIAL USE ONLY

Region: Bill type (for internal use only):
 Sales person: GC done by: GC date:

20th May 2024

Mr. D.M.A.D.B Dissanayake,

Colombo

Dear Sir,

QUOTATION FOR CONDUCTING GENETIC TESTING

As requested by Dr. Senaka Kandegedara Clinical Oncologist, we are pleased to offer you the following services for Diagnostic purpose through Strand Life Sciences, India.

Financial Offer

Test Cord	Test Name	Cost for the Toal Package (LKR)
SLS162800	Somatic Homologous Recombination Repair Test (sHRR)	LKR. 175,990/-

- **Terms & Conditions.**
- Diagnose results, reports will be given after 28 working days.
- Full payments should be transferred before the execution of the process.

Account Details.

Aegle Omics (Private) Limited
Bank - Commercial Bank
Branch- Narahenpita
Acc No- 1000756928
Swift Cord- CCEYLKLX

Thanking you!



.....
Amila Herath
Manager Operations

011-4496286



Comprehensive Cancer Care Centre

20 MAY 2024

Dr. Don Adams Doyle
M.D.

~~Dr. Don Adams Doyle~~

Dr. Adams - Received

Medicine Dept -

KSL-162800

RV

Ref.
Date

Dr. S. L. Rajaguru
CLINICAL NOTES

Dr. Mahendra Perera
Consultant Oncologist

Dear Sir

This is the pt
that you discuss in your
club is being seen
in our clinic

All the reports are with the
pt.

Yours



Dr. S.L. RAJAGURU
MBBS, MR. Clinical Oncology
Consultant Clinical Oncology
National Hospital
Kandy

14/12/20



Barcode

GENETIC TEST REQUISITION FORM

Test Code: <input type="text" value="SLS162800"/>	Test Name: Somatic Homologous Recombination Repair Test (sHRR)	TR ID: <input type="text"/>
Test Code: <input type="text"/>	Test Name: <input type="text"/>	TR ID: <input type="text"/>
Test Code: <input type="text"/>	Test Name: <input type="text"/>	TR ID: <input type="text"/>
Test Code: <input type="text"/>	Test Name: <input type="text"/>	TR ID: <input type="text"/>

PATIENT DETAILS

Patient MRN/UHID #:

Patient Name: DOB: Age: Sex:

Marital Status: Nationality: Contact No.:

Address:

Email ID: Pincode:

ID Proof: Driving License Aadhaar Card Voter ID card Ration card Others:

ADDITIONAL DETAILS IF REQUIRED

Transfusion Date (if available): Blood Group:

CLINICIAN INFORMATION

Referring Clinician: Clinician Contact:

Referring Hospital:

Email Id:

Address:

ADDITIONAL FAMILY MEMBERS' DETAILS

Disease Status - Affected: Age at Diagnosis:

Affected Sibling/Family members: Yes No (If yes, provide the details in the table below)

Name	Relation with Patient	Type of cancer	Age at Diagnosis	Sex
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>

SAMPLE COLLECTION INFORMATION

Date & Time of Collection: Sample collected by:

Clinical History/Pathology Report Attached: H and E Slides sent:

FOR OFFICIAL USE ONLY

Region: Bill type (for internal use only):

Sales person: GC done by: GC date:

PEDIGREE / FAMILY HISTORY / CLINICAL HISTORY (TO BE FILLED BY CLINICIAN / GENETIC COUNSELOR)

(PLEASE ATTACH SEPARATELY INCASE OF DETAILED PEDIGREE)

DETAILS OF SAMPLE SENT TO THE LAB

SEND SAMPLES TO
(Monday to Saturday only)

Strand Life Sciences Pvt. Ltd.
Ground Floor, UAS Alumni Association Building,
Veterinary College Campus, Hebbal, Bangalore-560024
Phone: +91 99 8044 8044, E-mail: support.strandx@strandls.com

INFORMED CONSENT FOR GENETIC TESTING

- If the proband is over 18 years of age, 'You' in this form refers to the proband
- If the proband is a minor or differently-abled, 'You' in this form refers to the Legal Guardian of the proband
- 'We', 'Our', 'Us' refers to Strand Life Sciences Pvt. Ltd.

You hereby consent to undergo testing offered by us and understand that:

1. Your biological sample(s) will be collected using generally accepted techniques. The sample(s) could be blood, saliva, tumor tissue, or any other biological sample as needed.
2. You understand that the sample(s) may be used to determine if you and/or your family members have variants in your gene(s). Results may indicate affected status, increased risk of being affected in future, inherited risk of disease, somatic mutations in tumor tissue, or other such findings.
3. Genetic tests are relatively new and are being improved and expanded continuously. You are aware of the risks and limitations of genetic testing.
4. You may need to share your relevant health records to correlate the findings from the genetic testing.
5. The genetic test report will be usually released within the Turnaround Time (TAT) specified by us for the test.
6. A positive test result is an indication that the individual being tested has a genomic variant that might have implications for their health or their progeny's health. Consultation may be sought from any physician or genetic counselor of your choice. You may also consider independent testing and consultation in addition.
7. The report will be shared with your authorized physician where applicable, or shared to your registered email ID, or shared on our secure portal, or a hard copy of the report can be shared upon request to the address provided in this form.
8. It is possible that knowledge of genetic information of an individual might be misused if it is in the wrong hands. Hence, we cannot reveal the genetic information even to family members without your explicit written authorization to do so. We will maintain complete confidentiality of the test results, as genetic test results can have social implications.
9. Genetic testing has its limitations. A repeat or alternative tests might be recommended by your treating physician accordingly.
10. Genetic testing might identify secondary findings in genes outside of the original test genes as defined by the American College of Medical Genetics. You can opt out from receiving secondary findings by ticking this box here.
11. All laboratory raw data are confidential and will not be released unless a special written request is made by you or the consulting clinician on record, or a valid court order is received by us. VCF, FASTQ and BAM files can be provided to you

- or your authorized personnel for a period of 3 months. For requests beyond that period we may charge an additional cost, as per institutional policy.
12. Our laboratory does not return any leftover sample after completion of testing under any circumstances. The only exception to the above is in the case of FFPE tumor blocks which can be returned upon request by you or your ordering physician within a period of 6 months. Any left over DNA (if available), regardless of the sample type, can be requested by you for up to a period of 3 months, provided you bear the transportation costs.
13. Samples collected as part of routine care with potential for future genetic research will be stored by our laboratory in accordance with ICMR guidelines 2017, clause 10.3.7.
14. Samples can also be shared with collaborators within or outside the country in line with existing relevant guidelines, in accordance with ICMR guidelines 2017, clause 10.3.8.
15. You understand and agree to the use of your data and biological sample for future research by us and our collaborators. We will use your samples and data in anonymised or aggregated form, such that it will be incapable of identifying you. By voluntarily signing this consent form you agree to provide broad or blanket consent for the storage and use of your samples and data as specified by ICMR guidelines 2017. You can opt out of this clause by ticking this box here.
16. Sharing of data with our collaborators will be bound under a data access agreement that will maintain individual confidentiality. Your personal identity will not be revealed in any information shared with third parties or published; your data will be coded accordingly.
17. The future use of your data or sample in research may result in commercial gains. Based on the nature of research outcomes, further investments by us may be needed to commercialize these outcomes. You will not have the right to participate in any direct monetary gains resulting from any future commercial activity.
18. You understand and agree to being re-contacted in the future if there is new information available on your genomic variants or new research envisaged that you could benefit from.
19. You have the full rights to decide whether or not you wish to provide consent, nobody can coerce you into providing consent. You can also choose to withdraw your consent at a later stage if you so wish, you need to notify us regarding the same.
20. In case you have any concerns or perceive any conflict of interest, you may seek clarification on institutional policy from relevant authorities.

AEGLE OMICS (PRIVATE) LIMITED

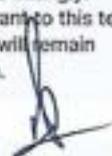
Name of Individual/Legal Guardian: 1211-Level 12, Parkland Building
 No. 33, Park Street,
 Colombo 00200

Date: 26/05/24 Signature: 

STATEMENT OF PHYSICIAN/GENETIC COUNSELOR

- I have explained the genetic testing suitable for this individual and the individual has had the opportunity to ask questions.
- I have addressed the limitations outlined above, and I have answered this person's questions.
- I have obtained consent from the patient or the legal guardian for this testing. I confirm that the individual has given consent willingly.
- I allow Strand Life Sciences to contact the aforementioned patient for clinical history, treatment plan, and other details relevant to this test.
- I undertake to maintain the privacy and confidentiality of the patient's genetic data obtained from Strand Life Sciences and will remain solely responsible for any wrongful acts and/or omissions arising out of, or in relation to my use of the patient's genetic data.

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

Date: 26/05/24 (Physician/Genetic Counselor) Signature: 

www.asirihealth.com

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



ASIRI
LABORATORY
 LIVE MORE

35+
 YEARS OF SERVICE

Asiri Laboratories Kandy
 No. 907, Panabokka Road, Kandy
 T. 081 285 04 04, F. 081 285 04 05

HISTOPATHOLOGY

Block and slides of this
 will be retained ONLY for
 after the date of this report
 will be kept for one week.

** OPD/AKH/AKH **

REFERENCE No. : 05 0246 02/05/23
 SAMPLE DATE & TIME : 02/05/2023 22:39
 REPORT DATE & TIME : 06/05/2023 10:54
 PATIENT : MR. D M A D B DISSANAYAKE (NITF 403945)
 REFERRED BY : DR CHATHURANGA KEPPETIYAGAMA

AGE : 53 Y/M

TEST : **HISTOPATHOLOGY REPORT**

Specimen : Sigmoid colonic biopsy.

Macroscopy : Five fragments of soft tissue - each 2x4x5 mm.

Microscopy : The colonic biopsies show a moderately differentiated
 adeno carcinoma.

No lympho vascular tumour emboli are seen.

Diagnosis : Sigmoid colon (Biopsy): G2 adeno carcinoma.

PRH 3089

Ratnayake

Dr. Palitha Ratnayake
 MBBS, D.path, MD (PATHOLOGY)
 Consultant Pathologist