

PATIENT INFORMATION	SAMPLE INFORMATION
Name: Date of Collection: (DD/MM/YYYY) Time: (HH/MM) Last name First name HKID/ Passport no: Date of Birth: (DD/MM/YYYY) Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female Phone number: Clinical Diagnosis:	Sample Type (Please select one): <input type="checkbox"/> Blood (<input type="checkbox"/> EDTA/ <input type="checkbox"/> Plasma) <input type="checkbox"/> Saliva in collection kit <input type="checkbox"/> Swab (<input type="checkbox"/> Buccal / <input type="checkbox"/> Nasal) <input type="checkbox"/> Urine <input type="checkbox"/> Other, please specify:
REPORTING INFORMATION	
Ordering Physician: Institution: Route of report delivery: <input type="checkbox"/> Email / <input type="checkbox"/> Mail / <input type="checkbox"/> Fax	Email: Phone: Fax:
Please put a "✓" in the box(es) to indicate the test(s) to be performed.	
Neurology	
<u>Alzheimer's Disease</u> <input type="checkbox"/> CoGenesis® AD <input type="checkbox"/> APOE Test <input type="checkbox"/> p-Tau 217 Test <input type="checkbox"/> Amyloid- Beta 42/40 Ratio <input type="checkbox"/> NFL Test <input type="checkbox"/> α-Synuclein Test <input type="checkbox"/> GFAP Test <input type="checkbox"/> Precivity AD2™	<u>Neurofibromatosis</u> <input type="checkbox"/> Neurofibromatosis type 1 & 2 (NF1 & NF2) panel <u>Movement & Neurodegenerative Disorders</u> <input type="checkbox"/> Amyotrophic Lateral Sclerosis (ALS) & Frontotemporal Dementia (FTLD) Panel <input type="checkbox"/> CoGenesis® Neuro <input type="checkbox"/> CoGenesis® Myopathy <input type="checkbox"/> Spinocerebellar ataxia (SCA) Panel <input type="checkbox"/> Retinitis Pigmentosa Panel
WES/WGS	
<input type="checkbox"/> Whole Exome Sequencing (WES)	<input type="checkbox"/> Whole Genome Sequencing (WGS)
Paediatrics	
<u>Neurodevelopmental disorders</u> <input type="checkbox"/> CoGenesis® ASD	<u>Recessive Diseases Carrier Test</u> <input type="checkbox"/> CoGenesis® Family (Partner's HKID no. _____)
Cardiovascular Disease	
<input type="checkbox"/> CoGenesis® CVD	
Pharmacogenomics	
<input type="checkbox"/> CoGenesis® Drug Response	
Cancer	
<u>Hereditary Cancer Screening</u> Specify TAT: <input type="checkbox"/> Standard (4 weeks) <input type="checkbox"/> Paid- Accelerated (14 working days)	
<input type="checkbox"/> CoGenesis® Pan-Cancer (Please also fill in the <i>Health Information Questionnaire</i>) <input type="checkbox"/> CoGenesis® BRCA-Pro (Please also fill in the <i>Health Information Questionnaire</i>)	<input type="checkbox"/> CoGenesis® Colo <input type="checkbox"/> CoGenesis® Gastric <input type="checkbox"/> CoGenesis® Pancreas <input type="checkbox"/> CoGenesis® Prostate
<u>Somatic Cancer Diagnosis</u> <input type="checkbox"/> CONCISE ColoScan (Please also fill in the <i>Tissue Handling Authorization Letter</i>) <input type="checkbox"/> CONCISE LungScan (Please also fill in the <i>Tissue Handling Authorization Letter</i>)	
Infectious Diseases	
<input type="checkbox"/> EBV Real-time PCR (<input type="checkbox"/> Qualitative <input type="checkbox"/> Quantitative) <input type="checkbox"/> HBV Real-time PCR	<input type="checkbox"/> HPV Real-time PCR <input type="checkbox"/> STD Combo (<input type="checkbox"/> for 3 <input type="checkbox"/> for 9 <input type="checkbox"/> for 13)
<u>Designed Panel/Other, Please Specify:</u>	

Please call **3008 2560** or fax this form to **2381 8688** to arrange for sample collection.

Requesting Physician:

Print Name:

Signature:

Date (DD/MM/YYYY):

免責聲明及同意書

本免責聲明及同意書由科德施基因有限公司 (Codex Genetics Limited, 以下簡稱「本公司」) 發出。

請仔細通讀此同意書。如果您還有其他疑問，請向本公司查詢。您有權在任何時候改變主意，包括在簽署本同意書後。您亦有權隨時向科德施基因查閱或更改您在科德施基因登記的個人資料。本人確認本表格所提供的資料準確無誤。

本人同意本同意書所述的測試將由本公司進行。

- 本人已明白我所進行的測試的目的、範圍及限制。
- 本人的測試結果將發送至本人的醫生，本人將在結果準備就緒時從醫生處獲取相關資訊。
- 您的個人資料受《個人資料(私隱)條例》(PDPO) 及 ISO27001 國際標準保障，將僅用於診斷用途；您可選擇授權 / 不授權*我們使用經去識別化/匿名化處理的數據進行研究以優化服務，我們會實施嚴格安保措施防止未經授權的存取、使用或披露。您的個人資料將被保存至少四年，或按法律及認證標準 (包括《個人資料 (私隱) 條例》及 ISO 15189) 之要求保留。您可聯絡我們查閱或要求刪除您的個人資料，惟須符合相關法律規定。
- 本人已閱讀並理解本表格中上述信息，特此同意本公司對上述樣品進行相關化驗項目，並代表本人取回上述樣本和病理報告，也了解樣本可能會在測試過程中消耗。
- 若進行測試人士未滿 18 歲，本人作為其法定監護人，代表測試人士同意其接受該基因檢測，並同意按照上述規定處理測試人士的個人資料。

*請刪去不適用選項

Disclaimer and consent form

This Disclaimer and Consent Form is issued by Codex Genetics Limited (hereinafter referred to as "the Company").

Please read this consent form carefully. If you have any questions, please contact the Company. You have the right to change your mind at any time, including after signing this form. You may also access or amend your personal information registered with the Company at any time. I confirm that the information provided in this form is accurate.

I agree that the test described in this consent form will be conducted by the Company.

- I understand the purpose, scope, and limitations of this test.
- My test results will be sent to my doctor, and I will obtain the relevant information from my doctor when the results are ready.
- Your information is protected under the Personal Data (Privacy) Ordinance (PDPO) and ISO27001 standards; it will be used for diagnostic purposes, and you may **authorize/ not authorize*** the use of deidentified/anonymized data for research to improve our services, with strict security measures in place to prevent unauthorized access. Your personal data will be kept for at least four years, or as required by law and accreditation standards (including the Personal Data (Privacy) Ordinance and ISO 15189). You may contact us to request access to or deletion of your data, subject to legal requirements.
- I consent to Codex Genetics Limited conducting the tests indicated above and retrieving the sample and pathology report on my behalf, understanding that the sample may be consumed during testing.
- If the individual is under 18 years of age, I, as the legal guardian, consent on behalf of the individual to undergo this test and agree to the handling of their personal data as outlined above.

*Please delete the option that does not apply.

Patient:

Print Name: _____ Signature: _____ Date (DD/MM/YYYY): _____

Codex Genetics Laboratory use only

Receipt date & time:

Received by:

Data Privacy and Patient Authorization

By submitting this test request and corresponding patient samples, the referring physician confirms that the patient's consent or authorization has been duly obtained for the collection, use, and disclosure of their personal data — including name, date of birth, gender, biological samples, and test results — for the purpose of performing and reporting the requested medical tests. The referring physician further acknowledges that such data will be handled in accordance with applicable data protection and privacy laws and the laboratory's confidentiality and information security policies.

[Fill this form ONLY when tissue specimen retrieval from third-party laboratory is needed]

Date: _____

To: Histopathology Laboratory, _____ (Hospital Name)

RE: Sample Retrieval and Preparation for Cancer Tissue Testing for Patient _____ (Name), HKID: _____

To Whom It May Concern,

Please accept this letter as formal authorization to your laboratory to:

1. **Retrieve** archived tissue specimen(s) of the indicated patient,
2. **Prepare** the specimen(s) based on the specimen requirements of the selected test request, and
3. **Release** the prepared specimen(s) to the **Codex Genetics** in handling all test requisition documents to the third-party service provider and local logistics for sample pick-up and transportation.

<input type="checkbox"/> For Guardant360 TissueNext + PDL1 Dako22C3: 8 FFPE unstained coated slides (*NOTE: Tissue biopsy with tumor content of 5-10%, additional slides up to 10-12 slides is preferred)	<p>Guardant IHC FFPE Slide Requirement:</p> <ul style="list-style-type: none"> 5µm thick each Surface Area <ul style="list-style-type: none"> Minimum: ≥ 2 mm² Optimum: 25mm² Tumor Content <ul style="list-style-type: none"> Minimum: ≥ 5% Optimum: ≥ 20% <p>Pathology report is required.</p>														
<input type="checkbox"/> Additional Guardant IHC Testing: (*Please check the box for each additional test request)															
<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr style="background-color: #cccccc;"> <th style="padding: 5px;">Additional Biomarker(s)</th> <th style="padding: 5px;">No. of additional FFPE slides required</th> </tr> </thead> <tbody> <tr> <td style="padding: 5px;"><input type="checkbox"/> c-MET</td> <td style="text-align: center; padding: 5px;">+1</td> </tr> <tr> <td style="padding: 5px;"><input type="checkbox"/> HER2</td> <td style="text-align: center; padding: 5px;">+1</td> </tr> <tr> <td style="padding: 5px;"><input type="checkbox"/> Ki-67</td> <td style="text-align: center; padding: 5px;">+1</td> </tr> <tr> <td style="padding: 5px;"><input type="checkbox"/> ER/PR</td> <td style="text-align: center; padding: 5px;">+2</td> </tr> <tr> <td style="padding: 5px;"><input type="checkbox"/> FOLR1</td> <td style="text-align: center; padding: 5px;">+2</td> </tr> <tr> <td style="padding: 5px;"><input type="checkbox"/> CLDN18</td> <td style="text-align: center; padding: 5px;">+3</td> </tr> </tbody> </table>	Additional Biomarker(s)	No. of additional FFPE slides required	<input type="checkbox"/> c-MET	+1	<input type="checkbox"/> HER2	+1	<input type="checkbox"/> Ki-67	+1	<input type="checkbox"/> ER/PR	+2	<input type="checkbox"/> FOLR1	+2	<input type="checkbox"/> CLDN18	+3	
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<input type="checkbox"/> For CONCISE LungScan / ColoScan 10 FFPE unstained coated slides															

Bill to: Codex Genetics Limited, Unit 220, 2/F, Hong Kong Science Park 16W, Shatin, N.T.

For any enquiries, please contact Ms. Sherie Lee (Codex Genetics) at 3008 2560 for specimen collection and logistics.

I, Dr. _____ (Dr. Name), hereby to confirm that the indicated patient has understood and accepted the recommendation for the genetic testing service, that this specimen retrieval and preparation is medically indicated. I authorize both your laboratory and Codex Genetics to proceed as outlined.

Yours Sincerely,

Clinician Name:

Patient Name:

Clinician Signature:

Patient Signature: