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(71) Applicant: THE JOHNS HOPKINS UNIVERSITY

[US/US]; 3400 North Charles Street, Baltimore, Maryland 21218 (US).

(72) Inventors: VOGELSTEIN, Bert; 3700 Breton Way, Bal-

timore, Maryland 21208 (US). KINZLER, Kenneth W.; 616 Ponte Villas North, Baltimore, Maryland 21230 (US).

DOUVILLE, Christopher; 1650 Orleans Street, Baltimore, Maryland 21231 (US). PAPADOPOULOS, Nickolas; 606 Horncrest, Towson, Maryland 21204 (US).

(74) Agent: LUITJENS, Cameron et al.; Fish & Richardson

P.C., P.O. Box 1022, Minneapolis, Minnesota 55440-1022 (US).

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(54) Title: RAPID ANEUPLOIDY DETECTION

(57) Abstract: This document provides methods and materials for identifying chromosomal anomalies that can be used to identify a mammal as having a disease (e.g., cancer or congenital abnormality). For example, this document provides methods and materials for evaluating sequencing data to identify a mammal as having a disease associated with one or more chromosomal anomalies (e.g., cancer or congenital abnormalities). For example, this document provides methods and materials for evaluating sequencing data that can be used in cancer diagnostics, non-invasive prenatal testing (NIPT), preimplantation genetic diagnosis and evaluation of congenital abnormalities.



From the INTERNATIONAL BUREAU

PCTNOTIFICATION OF THE RECORDING
OF A CHANGE(PCT Rule 92bis.1 and
Administrative Instructions, Section 422)

To:

LUITJENS, Cameron
Fish & Richardson P.C.
P.O. Box 1022
Minneapolis, Minnesota 55440-1022
États-Unis d'Amérique

Date of mailing (day/month/year) 28 April 2021 (28.04.2021)	IMPORTANT NOTIFICATION
Applicant's or agent's file reference 448070322WO1	
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1. The following indications appeared on record concerning:		
☐ the applicant	☒ the inventor	☐ the agent
☐ the common representative		
Name and Address	State of Nationality	State of Residence
	Telephone No.	
	Facsimile No.	
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2. The International Bureau hereby notifies the applicant that the following change has been recorded concerning:		
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3. Further observations, if necessary: The person identified in Box 2 has been added as inventor.		
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