

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

BEFORE THE PATENT TRIAL AND APPEAL BOARD

GENEDX, INC.

Petitioner

v.

MYRIAD GENETICS, INC.

Patent Owner

U.S. Patent No. 6,051,379 to Lescallett *et al.*

Issue Date: April 18, 2000

Title: Cancer Susceptibility Mutations of BRCA2

Inter Partes Review No: Unassigned

GENEDX, INC.'S EXHIBIT LIST

Mail Stop "PATENT BOARD"

Patent Trial and Appeal Board

U.S. Patent and Trademark Office

P.O. Box 1450

Alexandria, VA 22313-1450

GeneDx Exhibit List
Petition for Inter Partes Review of USPN 6,051,379

<i>GDX Exhibit #</i>	<i>Description</i>
1001	U.S. Patent No. 6,051,379, issued April 18, 2000
1002	Declaration of Dr. Madhuri Hegde, PH.D., FACMG
1003	<i>Curriculum Vitae</i> of Dr. Madhuri Hegde, PH.D., FACMG
1004	File History of U.S. Patent No. 6,051,379
1005	Miki, Y., <i>et al.</i> , "Mutation analysis in the <i>BRCA2</i> gene in primary breast cancers," <i>Nature Genetics</i> 13: 245-247 (June 13, 1996)
1006	Bowcock, A.M., "Molecular cloning <i>BRCA1</i> : a gene for early onset familial breast and ovarian cancer," <i>Breast Cancer Research and Treatment</i> 28: 121-135 (1993)
1007	Hacia, J.G., <i>et al.</i> , "Detection of heterozygous mutations in <i>BRCA1</i> using high density oligonucleotide arrays and two-colour fluorescence analysis," <i>Nature Genetics</i> 14: 441-447 (December 14, 1996)
1008	Grompe, M., "The rapid detection of unknown mutations in nucleic acids," <i>Nature Genetics</i> 5: 111-117 (1993)
1009	Orita, M., <i>et al.</i> , "Detection of polymorphisms of human DNA by gel electrophoresis as single-strand conformation polymorphisms," <i>PNAS USA</i> 86: 2766-2770 (1989)
1010	Scharf, S. J., <i>et al.</i> , "Direct Cloning and Sequence Analysis of Enzymatically Amplified Genomic Sequences," <i>Science</i> 233: 1076-1078 (1986)
1011	Weber, B.H.F., <i>et al.</i> , "A Somatic Truncating Mutation in <i>BRCA2</i> in a Sporadic Breast Tumor," <i>Am. J. Hum. Genet.</i> 59: 962-964 (October 1996)
1012	Lancaster, J.M., <i>et al.</i> , " <i>BRCA2</i> mutations in primary breast and ovarian cancers," <i>Nature Genetics</i> 13: 238-240 (June 13, 1996)
1013	Myriad Genetics 2013 Annual Report <i>downloaded from</i> http://files.shareholder.com/downloads/MYGN/3399011197x0x699241/42B88246-0E90-4F5C-9697-3DF98027C46D/2013_Annual_Report.pdf
1014	Declaration of Dr. Gregory C. Critchfield, filed December 23, 2009, in <i>Assoc. for Molecular Pathology v. U.S. Patent & Trademark Office</i> , No. 09-cv-04515-RWS (S.D.N.Y.)
1015	Declaration of Mark Allan Kay, M.D., Ph.D, filed August 31, 2013, in <i>University of Utah Research Foundation v. Ambry Genetics Corp.</i> , No. 13-cv-00640-RJS (D.Utah)

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1016	Eng, C., <i>et al.</i> , "Interpreting epidemiological research: blinded comparison of methods used to estimate the prevalence of inherited mutations in <i>BRCA1</i> ," <i>J. Med. Genet.</i> 38: 824-833 (2001)
1017	Sambrook, J., <i>et al.</i> , <i>Molecular Cloning, A Laboratory Manual</i> , 2 nd Ed., Cold Spring Harbor Laboratory Press, §§ 5.28-5.32, 6.36-6.48, 13.6-13.77, and 14.5-14.21 (1989)
1018	Conner, B.J., <i>et al.</i> , "Detection of sickle cell β^S -globin allele by hybridization with synthetic oligonucleotides," <i>PNAS USA</i> 80: 278-282 (1983)
1019	Teng, D.H.F., <i>et al.</i> , "Low incidence of <i>BRCA2</i> mutations in breast carcinoma and other cancers," <i>Nature Genetics</i> 13: 241-244 (June 13, 1996)
1020	Cotton, R.G.H., "Current methods of mutation detection," <i>Mutation Research</i> 285: 125-144 (1993)
1021	Handelin, B., <i>et al.</i> , "Simultaneous Detection of Multiple Point Mutations Using Allele-Specific Oligonucleotides," <i>Current Protocols in Human Genetics</i> Suppl. 6: 9.4.1-9.4.8, online publication of May 1, 2001 corresponding to August 1995 print publication
1022	Wooster, R., <i>et al.</i> , "Localization of a Breast Cancer Susceptibility Gene, <i>BRCA2</i> , to Chromosome 13q12-13," <i>Science</i> 265: 2088-2090 (1994)
1023	Wooster, R., <i>et al.</i> , "Identification of the breast cancer susceptibility gene <i>BRCA2</i> ," <i>Nature</i> 378:789-792 (1995)
1024	Schutte, M., <i>et al.</i> , "Identification by representational difference analysis of a homozygous deletion in pancreatic carcinoma that lies within the <i>BRCA2</i> region," <i>Proc. Natl. Acad. Sci. USA</i> 92: 5950-5954 (1995)
1025	Alberts, <i>et al.</i> "Molecular Biology of the Cell," 3 rd Ed., pp. 98-99, 104-106, 242-243, 292-293, 314-317, 337, 339, 1072-1073, G-5-G-6, G-10, G-17 (1994)
1026	Tavtigian, S.V., <i>et al.</i> , "The complete <i>BRCA2</i> gene and mutations in chromosome 13q-linked kindreds," <i>Nature Genetics</i> 12: 333-337 (March 12, 1996)
1027	GenBank Accession No. U43746, "Human breast cancer susceptibility (<i>BRCA2</i>) mRNA, complete cds," modification date of September 3, 1996, available at http://www.ncbi.nlm.nih.gov/nuccore/1161383?sat=13&satkey=6559532 (last accessed August 6, 2014)

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1028	GenBank Accession No. AAB07223, "BRCA2 [Homo sapiens]," modification date of September 3, 1996, available at http://www.ncbi.nlm.nih.gov/protein/1161384?sat=13&satkey=6559532 (last accessed August 6, 2014)
1029	Shattuck-Eidens <i>et al.</i> , "In Vivo Mutations and Polymorphisms in the 17q-linked Breast and Ovarian Cancer Susceptibility Gene," WO 96/05306 (filed August 11, 1995; published on February 22, 1996)
1030	PACER List of Associated Cases for 14-md-02510, available at https://ecf.ugd.uscourts.gov/cgi-bin/qryAscCases.pl?89779 (last accessed on August 15, 2014)

Respectfully submitted,

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CERTIFICATION OF SERVICE (37 C.F.R. §§ 42.6(e), 42.105(a))

The undersigned hereby certifies that the above-captioned "GENEDX, Inc.'s Exhibit List" and Exhibits 1001 – 1030 were served in their entirety on August 18, 2014, upon the following party via FEDEX:

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