# DNA Profile Report / Allele Sizing <br> T029470 

| Specimen Sample: | Jan Wigestrand |
| :--- | :--- |
| Date of Birth: | $03 / 02 / 1965$ |
| Ethnicity: | Unspecified/Mixed |
| Collection Date: | $11 / 30 / 2004$ |

## Explanation of DNA Profile Analysis:

Several discrete locations on different chromosomes were used to construct the DNA profile for Jan Wigestrand. Table 1 shows Jan Wigestrand to have identification markers 10 and 11 at the CSF1PO STR chromosome location. The parents contributed these identification markers to Jan Wigestrand, one marker from each parent. Results from the other STR chromosome locations are also listed in Table 1. Each marker has an established probability of occurrence in the general population and among different ethnic groups.

## Conclusion:

DNA profiling (also known as DNA fingerprinting) was performed by PCR, and has been completed on a sample in the name of Jan Wigestrand. Based on the observed scientific evidence, it is concluded, for all practical purposes, and in reference to the submitted samples, that the above DNA profile is that of Jan Wigestrand. The chance is 1 in 9,640,000,000,000,000 that some (particular) person other than the individual tested would have a genetic profile identical to the profile provided.

Table 1: The DNA profile established for Jan Wigestrand.

|  |  |
| :--- | :---: |
|  |  |
| Jan Wigestrand <br> CODIS profile <br> Unspecified |  |
| Locus |  |
| AMEL |  |
| CSF1PO |  |
| D13S317 |  |
| D16S539 |  |
| D18S51 |  |
| D19S433 |  |
| D21S11 |  |
| D2S1338 |  |
| D3S1358 |  |
| D5S818 |  |
| D7S820 |  |
| D8S1179 |  |
| FGA |  |
| FGA |  |
| TH01 |  |
| TPOX |  |
| vWA |  |
| Random Match Probability: |  |

This is to certify that

## Jan Wigestrand

has had a Y-chromosome analysis performed by DNA Heritage.

Passed from father to son every generation, the Y-chromosome is a genetic legacy tracing back up the direct paternal line.

At each Y-chromosome marker, there are regions of DNA that are repeated several times. This DNA sequence is copied exactly when it is passed onto future generations - it is this similarity that is used to trace paternal lineages.

Occasionally however, small changes in that marker do occur resulting in different repeat numbers - it is these differences that make your
 haplotype distinctive from other lineages.

| Marker <br> Name | No. of <br> Repeats | Marker <br> Name | No. of <br> Repeats |
| :---: | :---: | :---: | :---: |
| DYS19 | 14 | DYS452 | 11 |
| DYS385a | 11 | DYS454 | 11 |
| DYS385b | 13 | DYS455 | 11 |
| DYS388 | 12 | DYS456 | 16 |
| DYS389i | 13 | DYS458 | 17 |
| DYS389ii | 29 | DYS459a | 9 |
| DYS390 | 24 | DYS459b | 10 |
| DYS391 | 11 | DYS460 | 11 |
| DYS392 | 13 | DYS461 | 12 |
| DYS393 | 14 | DYS462 | 11 |
| DYS426 | 12 | DYS463 | 22 |
| DYS437 | 15 | DYS464a | 15 |
| DYS438 | 12 | DYS464b | 15 |
| DYS439 | 12 | DYS464c | 17 |
| DYS441 | 14 | DYS464d | 17 |
| DYS442 | 12 | GATAA10 | 13 |
| DYS444 | 12 | GATAC4/ <br> DYS635 | 23 |
| DYS445 | 13 | TAGAH4 | 12 |
| DYS446 | 15 | GGAAT1B07 | 10 |
| DYS447 | 25 | YCAIIa | 19 |
| DYS448 | 19 | YCAIIb | 23 |
| DYS449 | 29 |  |  |
|  |  |  |  |

Your 43-marker results show the number of repeats for any given marker.

Your test involved the use of internationally-recognised standards and testing protocols ensuring that you can confidently compare your results with others. Any comparison should only be made with individuals who share your surname, or have a similarly spelled surname.

Sample number: YN1D547
Analysis date: 1st February 2005


Alastair Greenshields
Principal DNA Heritage



## GenQuest DNA Analysis Laboratory

University of Nevada School of Medicine
Department of Microbiology
Mail Stop 320
Reno, Nevada 89557

## Personal Screening Report

## Case Number: \#20058982 (DTC0027616)

| Party | Race | Date Collected |
| :--- | :--- | :--- |
| Jan Wigestrand | Caucasian | $07 / 08 / 2005$ |

## DNA ANALYSIS RESULTS: Allele and number of repeats

| Alleles | Number of Repeats |
| :---: | :---: |
| CSF1P0 | 10,11 |
| D2S1338 | 20,23 |
| D3S1358 | 17,18 |
| D5S818 | 9,11 |
| D7S820 | 8,9 |
| D8S1179 | 12,13 |
| D13S317 | 12,13 |
| D16S539 | 11,12 |
| D18S51 | 16,18 |
| D19S433 | 15 |
| D21S11 | 27,30 |
| FGA | 20,21 |
| THO1 | 7,9 |
| TPOX | 8,9 |
| vWA | 17,18 |
| Amelogenin | XY |

## Conclusions:

The above profile for Jan Wigestrand was obtained from the analysis of the alleles, CSF1PO, D2S1338, D3S1358, D5S818, D7S820, D8S1179, D13S317, D16S539, D18S51, D19S433, D21S11, FGA, THO1, TPOX, VWA and Amelogenin. This profile occurs in the Caucasian population one in every $4.36 \times 10^{22}$ persons.
Note: Personal Identity Screening Test results cannot be used in a court of law. Samples were not collected in accordance with AABB standards.

[^0]Date $\qquad$ 1 1

## JAN WIGESTRAND HAS HAD A Y-CHROMOSOME

 ANALYSISPERFORMED BY DNA HERITAGE.
SAMPLE NUMBER: YN1D547.
ANALYSIS DATE: 1ST FEBRUARY 2005.
JAN WIGESTRAND'S 43-MARKER.
RESULTS:
DYS19: 14 DYS385a: 11 DYS385b: 13 DYS388: 12
DYS389i: 13 DYS389ii: 29 DYS390: 24 DYS391: 11
DYS392: 13 DYS393: 14 DYS426: 12 DYS437: 15
DYS438: 12 DYS439: 12 DYS441: 14 DYS442: 12
DYS444: 12 DYS445: 13 DYS446: 15 DYS447: 25
DYS448: 19 DYS449: 29 DYS452: 11 DYS454: 11
DYS455: 11 DYS456: 16 DYS458: 17 DYS459a: 9
DYS459b: 10 DYS460: 11 DYS461: 12 DYS462: 11
DYS463: 22 DYS464a: 15 DYS464b: 15 DYS464c: 17
DYS464d: 17
GATAA10: 13 GATAC4/DYS635: 23
TAGAH4: 12 GGAAT1B07: 10
YCAlla: 19 YCAllb: 23

DYS452: 30 (+19), DYS 452: 30 (+19),
Y-GATA-A10: 13, Y-GATA-H4: 12, Y-GGAAT-1B07: 10, GAAT1B07: 10, YGATAA10: 13, YGATAH4: 12, YGGAAT1B07: 10, GATAH4: 12, GATA H4: 12, GATAH4.1: 12,
GATAC4: 23, GATA C4: 23, DYS635: 23, DYS 635: 23, Y-GATA-C4: 23, YGATAC4: 23, DYS394: 14, DYS 394: 14,
DYS 389-1: 13, DYS 389-2: 29, GATA H4: 12,
DYS395: 14, DYS 395: 14,
Y-GATA-A4: 12, YGATAA4: 12,
DYS385, DYS 385, DYS389, DYS 389, DYS459, DYS 459, DYS464, DYS 464, YCAll, YCA, lla, llb, i, ii, a, b, c, d, 43marker, Ychromosome, test, Y-DNA.

DYS 19: 14 DYS 385a: 11 DYS 385b: 13 DYS 388: 12
DYS 389i: 13 DYS 389ii: 29 DYS 390: 24 DYS 391: 11
DYS 392: 13 DYS 393: 14 DYS 426: 12 DYS 437: 15
DYS 438: 12 DYS 439: 12 DYS 441: 14 DYS 442: 12
DYS 444: 12 DYS 445: 13 DYS 446: 15 DYS 447: 25
DYS 448: 19 DYS 449: 29 DYS 452: 11 DYS 454: 11
DYS 455: 11 DYS 456: 16 DYS 458: 17 DYS 459a: 9
DYS 459b: 10 DYS 460: 11 DYS 461: 12 DYS 462: 11
DYS 463: 22 DYS 464a: 15 DYS 464b: 15 DYS 464c: 17
DYS 464d: 17
GATA A10: 13 GATA C4 / DYS 635: 23
TAGA H4: 12 GGAAT1B07: 10
YCA Ila: 19 YCA IIb: 23

## FamilyTreeDNA Certificate - mtDNA

Family Tree DNA certifies that a mitochondria DNA sample taken from
Jan Wigestrand
differs from the Cambridge Reference Sequence (CRS)* at the numbered positions indicated, by the presence of the bases designated $\mathrm{A}, \mathrm{C}, \mathrm{G}$ or T :

## Haplogroup T2

Sample \# 368457
The letters designate the base that occurs at each of those positions in place of the entire CRS. These are distinctive of this sample and may be compared to other people to confirm or rule out common descent, providing genetic evidence of genealogical relationships

HVR1: 16126C, 16292T, 16294T, 16296T, 16324C, 16519C
HVR2: 73G, 263G, 315.1C
*The Cambridge Reference Sequence is the accepted mtDNA standard.

\&RQFHWW [\%RLP DQV

| Case | 8478937 |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Patient Name | Jan Wigestrand |  |  |  |  |
| Sample Number | 8478937-50 |  |  |  |  |
| Date Collected |  |  |  |  |  |
| Collected by |  |  |  |  |  |
| Locus | Allele Sizes |  | Locus | Allele Sizes |  |
| D3S1358 | 17 | 18 | FGA | 20 | 21 |
| vWA | 17 | 18 | D22S1045 | 15 |  |
| D16S539 | 11 | 12 | D5S818 | 9 | 11 |
| CSF1PO | 10 | 11 | D13S317 | 12 | 13 |
| TPOX | 8 | 9 | D7S820 | 8 | 9 |
| D8S1179 | 12 | 13 | SE33 | 22.2 | 28.2 |
| D21S11 | 27 | 30 | D10S1248 | 14 |  |
| D18S51 | 16 | 18 | D1S1656 | 11 | 17.3 |
| D2S441 | 10 | 11 | D2S1338 | 20 | 23 |
| D19S433 | 15 |  | Amelogenin | X | Y |
| TH01 | 7 | 9 |  |  |  |

Note: Since the samples were not collected under a strict chain of custody by a third neutral party and the Laboratory cannot verify the origin of the samples, this test result may not be defensible in a court of law for the establishment of paternity and other legally related issues. The tested parties' names that may appear on this report have been provided by the client and cannot be verified. The laboratory assumes no responsibility for incorrect or misspelled patient information.

Based on the samples received from the tested parties whose identities cannot be independently verified, I, the undersigned
Laboratory Director, declare the genetic data is correct as reported on


## Jan Wigestrand Yoursample \# 368457

## FamilyTreeDNA

This Certificate confirms that you have had your DNA analyzed by Family Tree DNA. The outcome from each of the one hundred eleven Loci examined is reported in the table below. If your alleles for the one hundred eleven Loci match another person exactly, then you share the same Haplotype.


Section for Genetic Kinship and Identity

Our ref.: 2017401626
Jan Wigestrand
Postboks 99
4301 Sandnes

Your ref.: 2017401626
Date: 17.08.2017

## Forensic Genetic Expert Report

DNA-profile
Jan Wigestrand
Date of birth: 020365

## DNA-analysis

Investigations are made on different parts of the DNA-molecules (loci), which are hypervariable. For each locus two variants (alleles) appear; one inherited from the mother and one inherited from the father. 23 loci have been investigated of which 22 are inherited independently.

## Results

The attached table shows result for sample labelled Jan Wigestrand 02.03.1965 (see attached sampling documentation).


Marguerethe Stenersen, cand. scient. Berit Myhre Dupuy, Ph.D.
Daniel Kling, Ph.D.
Attachments: Sampling documentation, analysis results

Resultatrapport Autosomale markører - vedlegg til 2017401626
Oppdragsgivers saksnummer: 2017401626

| Genetisk <br> markør | JW65 | Bevisvekt <br> (LR) |
| :--- | :---: | :---: |
| AMELOGENIN | X-Y |  |
| D3S1358 | $17-18$ | 17.582 |
| TH01 | $7-9$ | 18.0181 |
| D21S11 | $27-30$ | 50.7761 |
| D18S51 | $16-18$ | 55.8771 |
| D10S1248 | $14-14$ | 10.4314 |
| D1S1656 | $20-23$ | 44.2506 |
| D2S1338 | $11-12$ | 38.28 |
| D16S539 | $15-15$ | 5.71209 |
| D22S1045 | $17-18$ | 9.0052 |
| vWA | $20-13$ | 11.04622 |
| D8S1179 | $10-11$ | 18.6829 |
| FGA | $19-21$ | 6.67399 |
| D2S441 | $15-15$ | 33.1044 |
| D12S391 | $22.2-28.2$ | 213.557 |
| D19S433 | $7-8$ | 228.685 |
| SE33 | $9-11$ | 35.2308 |
| Penta E | $12-13$ | 15.387 |
| D5S818 | $8-9$ | 18.1648 |
| D13S317 | $10-11$ | 6.76838 |
| D7S820 | $9-12$ | 9.90377 |
| CSF1PO | $8-9$ | 8.73147 |
| Penta D | $3.29242 e+\mathbf{0 3 0}$ |  |
| TPOX |  |  |
| Bevisvekt |  |  |
| (total LR) |  |  |
|  |  |  |

Oslo Universitetssykehus HF
Rettsmedisinske fag
Rettsgenetisk slektskap og identitet
Postboks 4950, Nydalen
0424 OSLO
Utskriftsdato: 17.08.2017:
E-post: rettsgenetikk@ous-hf.no

16 August 2017
Jan Wigestrand is a Caucasian male Y-DNA test results Y-111 (111 markers test) (analyzed by Family Tree DNA).

DYS393 14 DYS390 24 DYS19 14 DYS391 11 DYS385 11-13 DYS426 12 DYS388 12 DYS439 12 DYS389-I 13 DYS392 13 DYS389-II 29

DYS458 17 DYS459 9-10 DYS455 11 DYS454 11 DYS447 25 DYS437 15 DYS448 19 DYS449 29 DYS464 15-15-17-17

DYS460 11 GATA-H4 11 YCAII 19-23 DYS456 16 DYS607 15 DYS576 18 DYS570 17 CDY 37-39 DYS442 12 DYS438 12

DYS531 11 DYS578 9 DYF395S1 15-16 DYS590 8 DYS537 10 DYS641 10 DYS472 8 DYF406S1 10 DYS511 10

DYS425 12 DYS413 23-23 DYS557 16 DYS594 10 DYS436 12 DYS490 12 DYS534 18 DYS450 8 DYS444 12 DYS481 22 DYS520 20 DYS446 15

DYS617 12 DYS568 11 DYS487 13 DYS572 11 DYS640 11 DYS492 12 DYS565 12
DYS710 31 DYS485 15 DYS632 9 DYS495 16 DYS540 11 DYS714 27 DYS716 26 DYS717 19
DYS505 11 DYS556 11 DYS549 12 DYS589 12 DYS522 10 DYS494 9 DYS533 12 DYS636 12 DYS575 10 DYS638 11

DYS462 11 DYS452 30 DYS445 13 Y-GATA-A10 13 DYS463 24 DYS441 14 Y-GGAAT-1B07 10 DYS525 10

DYS712 20 DYS593 15 DYS650 18 DYS532 13 DYS715 24 DYS504 17 DYS513 12 DYS561 15 DYS552 24

DYS726 12 DYS635 23 DYS587 18 DYS643 10 DYS497 14 DYS510 17 DYS434 9 DYS461 12 DYS435 11

17 August 2017
Jan Wigestrand is a Caucasian male autosomal DNA test results (23 loci and Amelogenin) (analysed by Department of Genetic Kinship and Identity (Avdeling for rettsgenetisk slektskap og identitet), Oslo University Hospital).

## AMELOGENIN X-Y

CSF1PO 10-11 D3S1358 17-18 D5S818 9-11 D7S820 8-9 D8S1179 12-13 D13S317 12-13
D16S539 11-12 D18S51 16-18 D21S11 27-30 FGA 20-21 TH01 7-9 TPOX 8-9 vWA 17-18
D1S1656 11-17.3 D2S441 10-11 D2S1338 20-23 D10S1248 14-14 D12S391 19-21 D19S433 15-15 D22S1045 15-15

Penta D 9-12 Penta E 7-8 SE33 22.2-28.2


[^0]:    Stephen St.Jeor, Ph.D.
    Elmer Otteson, Ph. D.

