

Genetic Analysis of Homeostasis Iron Regulator (HFE) Gene and Protein in Homo Sapiens and its Future Aspect in Treatment of Classic (Type-I) Hereditary Hemochromatosis

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Abstract:- The HFE gene is present in the genome of various organisms, including Homo sapiens and functions to regulate the amount of iron in the body. With speciation, HFE gene has evolved in the various organisms but organisms having the common ancestry still have the HFE protein structure similar to the ancestral homologues. This has been proved and the applications to this have been extrapolated in this paper. This paper has therefore hypothesized a likely source of the transgenic HFE gene for gene therapy and other therapies that are under development. Besides the hypothesis of transgenic HFE source, the paper also shows the relation of Homo sapiens with its Primate ancestors such as Gorilla gorilla and Pan troglodytes.

The approach to the extrapolation of results was using the Basic Local Alignment Sequence Tool (BLAST) of the Homo sapiens HFE gene (NM_000410.4 Homo sapiens homeostatic iron regulator (HFE), transcript variant 1, mRNA) and then Pairwise alignment of the Closest genes to the Homo sapiens HFE gene. To further strengthen the claim, the Homo sapiens HFE protein was aligned with that of the closest genes that were found using BLAST. The results found that the Pan troglodytes HFE gene was close to 99.35% in identity to the Homo sapiens HFE, whereas Pan troglodytes protein had a 100% match to that of Homo sapiens HFE protein.

Thus the chimpanzee HFE gene can act as a source for the transgenic genes for curing the Classic, Type-I hemochromatosis, wherein the HFE protein synthesized by the body is dysfunctional to regulate the Iron in the body as Iron sensing complex is rendered dysfunctional by the malformed HFE protein which leads to the high amount of iron in the body which causes various problems such as diabetes, liver cirrhosis and etc in the long run.

Keywords:- Hemochromatosis, HFE Gene, HFE Protein, HFE Trans-Genesis.

I. INTRODUCTION

Iron is one of the most essential elements in our body. It plays an important role in the production of hemoglobin and facilitation of oxygen transport. Another role of iron is the enhancement of the immune system, by supporting the functions of all cells including those of the immune system. Further, iron content in our body also affects our cognitive function as well as physical features such as hair, skin and nails ^[1]. Thus, it is crucial for one to properly manage the iron level of one's body.

The HFE gene, or the hemochromatosis gene, is responsible for the production of the Homeostatic Iron Regulator (HFE) protein ^[2].

➤ Features of HFE gene: []

This gene is placed in the short arm of the 6th chromosome. The cytogenetic location of HFE is 6p22.2 (genomic coordinates (GRCh37): (6:26,087,421–26,096,437). The HFE gene contains 7 exons spanning 12 kb. The full-length transcript represents 6 exons ^[2].

➤ The function of HFE protein:

The HFE protein is located on the surface (cell membrane) of intestinal and liver cells. It interacts with the other protein in surrounding cells and monitors the amount of iron in the body along with the regulation and production of another protein called hepcidin ^[2].

For monitoring the iron level of a body, the HFE protein utilizes glycoproteins called transferrin (Tf) and transferrin receptor (TfR). The transferrin transports the iron in the bloodstreams to various tissues. The transferrin receptor, which is a membrane-bound protein, which binds the Tf to itself to receive the iron. When the HFE protein is bound to the TfR protein of a cell, the TfR protein cannot bind the Tf protein, which prevents the iron from entering the cells. If the HFE protein is not bound to HFE, then Tf can attach itself to the TfR protein and thus in this way, HFE protein monitors the iron level of a body ^[2].

Another function of HFE protein is the regulation of the protein called hepcidin, which is produced in the liver. Hepcidin controls how much iron is absorbed into the body and is released from the body (iron metabolism). Deficiency in hepcidin causes excess iron and overproduction of it leads to decreased iron content. When the HFE protein is attached to the TfR, proteins including hepcidin aren't able to enter the cell, which stops the production of hepcidin. When the HFE isn't bound, the production of hepcidin resumes^[2].

➤ *Hereditary Hemochromatosis*

Hereditary Hemochromatosis (HHC) is an autosomal recessive genetic disorder that is caused due to excess accumulation of iron in tissues which leads to organ damage, especially in the liver. It is also referred to as iron overload disorder.

Although HHC is already present at birth, the symptoms of this disease are usually experienced later in life, from around 40 years of age^[3].

➤ *Clinical Symptoms and Physical Manifestations in Patients with Hereditary Hemochromatosis^[3]:*

- >Amenorrhea
- >Cardiomyopathies
- >Abdominal Pain
- >Apathy
- >Ascites
- >Cirrhosis
- >Congestive heart failure
- >Diabetes Mellitus
- >Hypogonadism
- >Hepatocellular Carcinoma
- >Loss of Libido
- >Weakness
- >Weight Loss
- >Splenomegaly
- >Lethargy
- >Impotence
- >Osteoporosis
- >Testicular atrophy

II. MUTATIONS

➤ *C282Y*

The C282Y mutation is seen in adults with HHC and is the most common mutation. In C282Y mutation the Nucleotide 845 which usually is Guanine is replaced with Adenine which results in the transcribed protein to have Tyrosine instead of Cysteine at Amino acid number 282^[4].

➤ *H63D*

The H63D mutation has Guanine in the place of Cytosine a Nucleotide number 184 which in turn causes the transcribed protein to contain Aspartic acid in place of Histidine at Amino acid number 63. In this mutation, an increase in the ferritin concentration and hemoglobin level of a body is observed^[4].

III. HFE MECHANISM IN IRON REGULATION

➤ *Role of HFE protein:*

The HFE gene encodes a non-classical MHC (major histocompatibility complex) class I-like protein which can be found on the surfaces of the intestinal cells. By associating with β 2-microglobulin, HFE protein binds with TfR1 (Transferrin Receptor 1) preventing the expression of hepcidin. But, in the case that HFE protein does not bond with TfR1, the receptor gains affinity towards other proteins, primarily, transferrin. This results in hepcidin production, allowing ferrous ions to enter the hepatic cells^[5].

However, mutations can occur in HFE genes. C282Y mutation prevents the HFE proteins from reaching the surface of the hepatic cells. This disrupts the disulfide bridges in the extracellular domains of the protein. Such a mutation prevents the linkage of the HFE protein and the TfR1 leading to an iron overload in many of the vital organs, common symptoms of HH type 1^[5].

➤ *Role of Hepcidin:*

Hepcidin synthesized and secreted by hepatocytes in the human liver matures in hepcidin-25 (due to 25-amino acid peptide). It binds with ferroportin (also referred to as IREG1) where they later endocytosis in lysosomes. It is worthy to note that the Hepcidin-25 expression is indirectly correlated with ferroportin. Hence, in situations where there is a temporary lack of iron in the blood, hepcidin protein is suppressed. Soon when the concentrations stabilize, hepcidin is expressed to prevent further iron export into the blood.^[6]

In normal conditions, ferritin formed and isolated in hepatocytes can sustain a large number of iron ions with the help of the IRP-IRE regulatory system. In HH, however, the iron overload ferritin begins to denature in lysosomes due to excess iron ions, rather than endocytosis. This results in a redox-active NTBI (non-transferrin bound iron).^[6]

This protein releases isolated Fe^{2+} ions which react with hydrogen peroxide. This process is known as hydroxyl radical generation since it produces free radicals of OH. These OH free radicals undergo lipid peroxidation when they bond with lipid peroxides to generate respective radicals.^[7]

These toxic lipid radicals disrupt organelle function by weakening the lysosome membrane which further leads to enzyme leakage and eventually cell death. Lysosome fragility can also induce collagen synthesis and in turn fibrosis. Thus it induces tissue damage in the body.^[8]

IV. TOOLS

➤ *The tools used for the analysis were:*

➤ *BLAST - Basic Local Alignment Sequence tool*^[9]:

BLAST is a bioinformatics tool that is used to find the regions of similarity between biological sequences. The program compares nucleotide or protein sequences to sequence databases and calculates the statistical significance.

➤ *Pairwise Sequence Alignment*^[10]:

A pairwise sequence alignment (MSA) is a sequence alignment of two biological sequences, generally a protein, DNA, or RNA. In many cases, the input set of query sequences are assumed to have an evolutionary relationship by which they share a linkage and are descended from a common ancestor. From the resulting PSA, sequence homology can be inferred and phylogenetic analysis can be conducted to assess the sequences; shared evolutionary origins.

➤ *EMBOSS*^[11]:

Dottup is a commonly used tool for drawing dot plots between two sequences. It looks for places where words of a specified length have an exact match in both sequences and draw a diagonal line over the position of these words. Shorter sequences are more sensitive to shorter regions of similarity but also display random points of similarity and run slower as compared to longer sequences which run faster, display minimum random points of similarity but are less sensitive.

V. ANALYSIS

In the analysis, the BLAST was executed on the Homo sapiens DNA code for HFE gene and based on the results of BLAST, the Pairwise alignment was run on sequences closest to the query entered. The Protein transcribed from the closest DNA sequence found using Pairwise alignment was then analyzed against the Human HFE protein and the results were extrapolated. The sequence of the Homo sapiens HFE gene (NM_000410.4) was acquired from NCBI, alongside sequences of HFE genes from several other species.

➤ *The sample used as a query from Homo sapiens:*

Ascension Number: Homo sapiens HFE gene (NM_000410.4)

Number of Amino Acids in transcribed protein: 348

➤ *Database Used for BLAST:*

nr (All non-redundant GenBank CDS translations+PDB+SwissProt+PIR+PRF excluding environmental samples from WGS projects)

➤ *BLAST results:*

BLAST results showed that the genes ascension number XM_016954624.2 of Gorilla gorilla and XM_031011852.1 of Pan troglodytes were the closest variants to the Homo sapiens HFE gene (NM_000410.4).

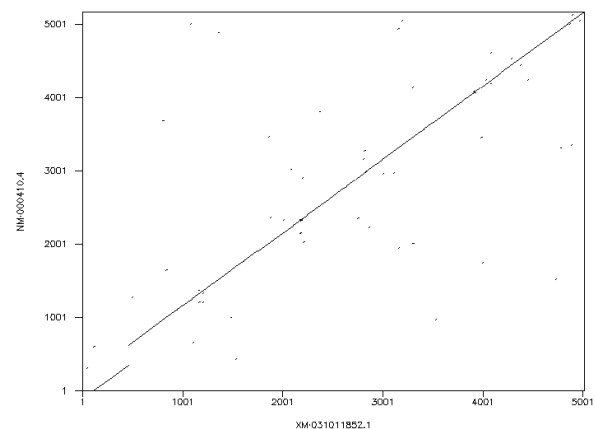
➤ *Pairwise alignment of Genes:*

Pairwise alignment of NCBI Reference Sequence: NC_000006.12 and XM_016954624.2:

The results found out that 3189/3211bp of NC_000006.12 and XM_016954624.2 perfectly aligned meaning the alignment was roughly 99.3 percent and Waterman-Eggert score was 9829.

➤ *EMBOSS:*

Dottup: fasta::emboss:dottup-l20201206-045657-0477-73350...
Sun 6 Dec 2020 04:56:59

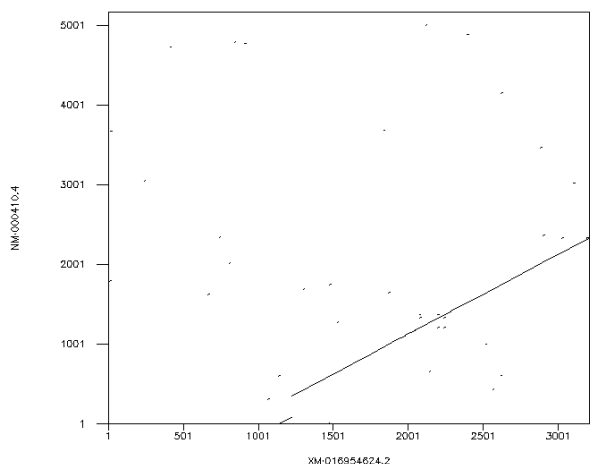


➤ *Pairwise alignment of NCBI Reference Sequence: NC_000006.12 and XM_031011852.1:*

The results found out that 4664/5015 bp of NC_000006.12 and XM_031011852.1 perfectly aligned, meaning that the alignment was roughly 93.0 percent and Waterman-Eggert score was 23206.

➤ **EMBOSS :**

Dottup: fasta::emboss-dottup-120201206-045806-0102-39074..
Sun 6 Dec 2020 04:58:09



➤ *Pairwise alignment of proteins:*

Pairwise alignment of Homo sapiens HFE protein vs Gorilla gorilla HFE protein transcribed from XM_031011852.1 gene:

As seen from the result mentioned in (fig.1) all the 345/348 amino acids of Gorilla gorilla HFE protein align with the amino acids of Homo sapiens HFE protein. Therefore there is a 99.1% similarity in the two proteins compared.

CLUSTAL O(1.2.4) multiple sequence alignment

```

sp|Q30201|HFE_HUMAN      MGPRARPALLLLMLLQTAVLQGRLLRSHSLHYLFHNGASEQDLGLSLFEALGYVDDQLFVF 60
tr|G3QU39|G3QU39_GORGO  MGPRARPALLLLMLLQTAVLQGRLLRSHSLHYLFHNGASEQDLGLSLFEALGYVDDQLFVF 60
*****

sp|Q30201|HFE_HUMAN      YDHESTRVEPRTPHWSSRISQMLQLSQSLKGDHMFVDFWTIMENHNHKSESHTLQV 120
tr|G3QU39|G3QU39_GORGO  YDHESTRVEPRTPHWSSRISQMLQLSQSLKGDHMFVDFWTIMENHNHKSESHTLQV 120
*****

sp|Q30201|HFE_HUMAN      ILGCEHQEDNSTEGYWKYGDQDHLFCPTLDNRAAEPRAMPKLEWERHKIRARQNR 180
tr|G3QU39|G3QU39_GORGO  ILGCEHQEDNSTEGYWKYGDQDHLFCPTLDNRAAEPRAMPKLEWERHKIRARQNR 180
*****

sp|Q30201|HFE_HUMAN      AYLERDCPAQLQQLLELGRGLDQQVPLVKVTHHTVSSVTLRACRALNYYPQNIHMKWL 240
tr|G3QU39|G3QU39_GORGO  AYLERDCPAQLQQLLELGRGLDQQVPLVKVTHHTVSSVTLRACRALNYYPQNIHMKWL 240
*****

sp|Q30201|HFE_HUMAN      KDKQPHDAKEFEKDVLPNGDGTQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIEWPS 300
tr|G3QU39|G3QU39_GORGO  KDKQPHDAKEFEKDVLPNGDGTQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIEWPS 300
*****

sp|Q30201|HFE_HUMAN      PSGLTVIGVISGIAVVFVILFIFILFIIILRKRQGSRGANGHYVLAERE 348
tr|G3QU39|G3QU39_GORGO  PSGLTVIGVISGIAVVFVILFIFILFIIILRKRQGSRGANGHYVLAERE 348
*****
    
```

Fig : Pairwise alignment result of Homo sapiens HFE protein V/S Gorilla gorilla HFE protein

Pairwise alignment of Homo sapiens HFE protein vs Pan troglodytes HFE protein transcribed from XM_016954624.2 gene:

As seen from the result mentioned in (fig.1) all the 348/348 amino acids of Pan troglodytes HFE protein align with the amino acids of Homo sapiens HFE protein. Therefore there is a 100% similarity in the two proteins compared

CLUSTAL O(1.2.4) multiple sequence alignment

```

sp|Q30201|HFE_HUMAN      MGPRARPALLLLMLLQTAVLQGRLLRSHSLHYLFHNGASEQDLGLSLFEALGYVDDQLFVF 60
sp|P60018|HFE_PANTR      MGPRARPALLLLMLLQTAVLQGRLLRSHSLHYLFHNGASEQDLGLSLFEALGYVDDQLFVF 60
*****

sp|Q30201|HFE_HUMAN      YDHESTRVEPRTPHWSSRISQMLQLSQSLKGDHMFVDFWTIMENHNHKSESHTLQV 120
sp|P60018|HFE_PANTR      YDHESTRVEPRTPHWSSRISQMLQLSQSLKGDHMFVDFWTIMENHNHKSESHTLQV 120
*****

sp|Q30201|HFE_HUMAN      ILGCEHQEDNSTEGYWKYGDQDHLFCPTLDNRAAEPRAMPKLEWERHKIRARQNR 180
sp|P60018|HFE_PANTR      ILGCEHQEDNSTEGYWKYGDQDHLFCPTLDNRAAEPRAMPKLEWERHKIRARQNR 180
*****

sp|Q30201|HFE_HUMAN      AYLERDCPAQLQQLLELGRGLDQQVPLVKVTHHTVSSVTLRACRALNYYPQNIHMKWL 240
sp|P60018|HFE_PANTR      AYLERDCPAQLQQLLELGRGLDQQVPLVKVTHHTVSSVTLRACRALNYYPQNIHMKWL 240
*****

sp|Q30201|HFE_HUMAN      KDKQPHDAKEFEKDVLPNGDGTQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIEWPS 300
sp|P60018|HFE_PANTR      KDKQPHDAKEFEKDVLPNGDGTQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIEWPS 300
*****

sp|Q30201|HFE_HUMAN      PSGLTVIGVISGIAVVFVILFIFILFIIILRKRQGSRGANGHYVLAERE 348
sp|P60018|HFE_PANTR      PSGLTVIGVISGIAVVFVILFIFILFIIILRKRQGSRGANGHYVLAERE 348
*****
    
```

Fig : Pairwise alignment result of Homo sapiens HFE protein V/S Pan troglodytes HFE protein

VI. POSSIBLE METHODS OF CORRECTION FOR CLASSIC TYPE-I HHC

The analysis that was run, found out that there is immense scope for interspecies transgenesis of HFE gene to cure the Type-I hemochromatosis. The existing technologies such as embryonic gene therapy, vector-based infusion of the gene and stem cell therapy for it to be expressed in the Human DNA can be possible cures for the treatment of classic type-I HHC in humans. The recent studies pertaining to the use of adenoviruses, Lentivirus and Epstein Barr Virus as vectors for insertion of foreign therapeutic genes have shown great success in the correction of other genetic anomalies such as Hemophilia which can be cured by the expression of correct genetic sequences.

Moreover, the current studies being conducted on the use of adenoviral vector for the correction of Hemophilia^[12] gives rise to the fundamental argument of it being used for curing classic type-I Hemochromatosis by insertion of therapeutic sequences of HFE gene as analyzed above in the cells of Homo sapiens, thus reversing the classic type-I hemochromatosis.

Why insertion of HFE gene using Lentiviral and Adenoviral vectors is an impractical approach?

The approach to curing HHC Type-I is impractical with the use of conventional Lentiviral and Adenoviral vectors due to their limitation of the length of DNA they can carry. The adenovirus and Lentivirus vectors are limited to less than 12KB whereas the complete CDS length of the HFE gene in the Homo sapiens is found to be more than 10KB, 12KB^[13].

Due to the length of DNA being a constraint we have to use other vectors that are being used for transgenesis of hepatic cells.

➤ *Use of Epstein Barr Virus*

The Epstein Barr Virus holds a large transgenic capacity unlike other virus vectors such as retroviruses and lentiviruses (that are often used as a vector). The EBV virus

can hold up to 120kb as a vector compared to other vectors^[14]. This allows EBV to sufficiently accommodate and deliver genomic transgenes for gene therapy.

This can be possible cure for the treatment of classic type-I HHC in humans. The recent studies pertaining to the use of adenoviruses, Lentivirus and Epstein Barr Virus as vectors for insertion of foreign therapeutic genes have shown great success in the correction of other genetic anomalies such as Hemophilia which can be cured by the expression of correct genetic sequences.

Therefore the similar methodologies using the EBV virus as a vector for transgenic HFE gene can be used for the correction of type-I classic HHC in Homo sapiens.

➤ *Proposal and the case for the use of CRISPR-cas based gene-editing technology and stem cell therapy to reverse C282Y mutation and cure Classic Type-I HHC:*

CRISPR-cas based gene-editing technology has shown immense and tremendous amounts of scope in reversing the single point mutation related genetic diseases such as hemophilia^[15].

Therefore based on this model the in vivo experiment for transgenesis of the hepatic cells and the hematopoietic stem cells for reversing the C282Y mutation should be possible by using the CRISPR-cas technology.

VII. CONCLUSION

In our study, we conclude that Homo sapiens variant of the HFE gene, **NM_000410.4 Homo sapiens homeostatic iron regulator (HFE), transcript variant 1, mRNA** can be substituted with either Gorilla gorilla gene, **XM_031011852.1** or Pan troglodytes gene, **XM_016954624.2**.

The basis of this hypothesis is that the exons of the human gene that was entered as a query against the gene of the Gorilla gorilla and Pan troglodytes had a high degree of similarities as given in the results above. This causes the proteins being transcribed out of the proposed substitute genes to have identical HFE proteins as Homo sapiens (Query protein), 348/348 amino acids similar with Pan troglodytes and 347/348 amino acids similar with Gorilla gorilla.

Thus with the proposed methods of correction combined with the use of transgenic material analyzed using BLAST software we will be able to devise a genetic therapy to cure Classic Type-I HHC.

REFERENCES

- [1]. Abbaspour N, Hurrell R, Kelishadi R. Review on iron and its importance for human health. *J Res Med Sci*. 2014 Feb;19(2):164-74. PMID: 24778671; PMCID: PMC3999603.
- [2]. Barton JC, Edwards CQ, Acton RT. HFE gene: Structure, function, mutations, and associated iron abnormalities. *Gene*. 2015 Dec 15;574(2):179-92. DOI: 10.1016/j.gene.2015.10.009. Epub 2015 Oct 9. PMID: 26456104; PMCID: PMC6660136.
- [3]. Brissot P, Cavey T, Ropert M, Guggenbuhl P, Loréal O. Genetic hemochromatosis: Pathophysiology, diagnostic and therapeutic management. *Presse Med*. 2017 Dec;46(12 Pt 2):e288-e295. doi: 10.1016/j.lpm.2017.05.037. Epub 2017 Nov 20. PMID: 29158016.
- [4]. Kelley M, Joshi N, Xie Y, Borgaonkar M. Iron overload is rare in patients homozygous for the H63D mutation. *Can J Gastroenterol Hepatol*. 2014 Apr;28(4):198-202. doi: 10.1155/2014/468521. PMID: 24729993; PMCID: PMC4071918.
- [5]. Maja Vujić. Molecular Basis of HFE-hemochromatosis. *Front Pharmacol*. 2014; 5: 42. Published online 2014 Mar 11. doi: 10.3389/fphar.2014.00042; PMCID: PMC3949417; PMID: 24653703
- [6]. Taro Takami, Isao Sakadai. Iron regulation by hepatocytes and free radicals. *Journal of Clinical Biochemistry and Nutrition*; 2011 Mar; 48(2): 103–106; Published online 2011 Feb 26. doi: 10.3164/jcbrn.1076; PMCID: PMC3045680; PMID: 21373260
- [7]. Mitchell D Khutson. Non-transferrin-bound iron transports. *Free Radical Biology and Medicine* (IF 6.170) Pub Date : 2018-10-12 , DOI: 10.1016/j.freeradbiomed.2018.10.413
- [8]. R. Fleming, W. Sly. Mechanisms in iron accumulation in hereditary hemochromatosis. Published 2002. DOI :10.1146/ANNUREV.PHYSIOL.64.081501.15588 ; Corpus ID: 517972
- [9]. National Center for Biotechnology Information (NCBI)[Internet]. Bethesda (MD): National Library of Medicine (US), National Center for Biotechnology Information; [1988] – [cited 2020 DEC 31]. Available from:<https://www.ncbi.nlm.nih.gov/>
- [10]. Xiaoqiu Huang, Webb Miller, A time-efficient, linear-space local similarity algorithm, *Advances in Applied Mathematics*, Volume 12, Issue 3, 1991, Pages 337-357, ISSN 0196-8858, [https://doi.org/10.1016/0196-8858\(91\)90017-](https://doi.org/10.1016/0196-8858(91)90017-)

D.(<http://www.sciencedirect.com/science/article/pii/S09688589190017D>)

[11]. Rice P., Longden I. and Bleasby A. EMBOSS: The European Molecular Biology Open Software Suite. Trends in Genetics. 2000 16(6):276-277

[12]. Brunetti-Pierri N, Ng P. Adenoviral Vectors for Hemophilia Gene Therapy. J Genet Syndr Gene Ther. 2013 Apr 30;2(Suppl 1):017. PMID: 24883229; PMCID: PMC4039643

[13]. Michael White, Roger Whittaker, Carolina Gándara, and Elizabeth A. Stoll. Human Gene Therapy Methods. Aug 2017. 163-176. <http://doi.org/10.1089/hgtb.2017.096>

[14]. Robert E. White, Richard Wade-Martins, Michael R. James, Infectious Delivery of 120-Kilobase Genomic DNA by an Epstein–Barr Virus Amplicon Vector, Molecular Therapy, Volume 5, Issue 4, 2002, Pages 427-435, ISSN 1525-0016, <https://doi.org/10.1006/mthe.2002.0557> (<http://www.sciencedirect.com/science/article/pii/S1525001602905575>)

[15]. Nguyen TH, Anegon I. Successful correction of hemophilia by CRISPR/Cas9 genome editing in vivo: delivery vector and immune responses are the key to success. EMBO Mol Med. 2016;8(5):439-441. Published 2016 May 2. doi:10.15252/emmm.201606325

APPENDIX:

I. The sequence of Homo sapiens HFE gene (NM_000410.4) :

Homo sapiens homeostatic iron regulator (HFE), transcript variant 1, mRNA

AGAGCTGGGAAATGGGCCCGCGAGCCAGGCCGGCGCTTCTCCTCCTGATGCTTTTGCAGACCGCGGTCTGCAGGGG
CGCTTGCTGCGTTCACACTCTCTGACTACCTCTTCATGGGTGCTCAGAGCAGGACCTTGGTCTTTCCCTTGTTTGAAGC
TTTGGGCTACGTGGATGACCAGCTGTTTCGTGTTCTATGATCATGAGAGTCGCCGTGTGGAGCCCCGAACCTCCATGGGT
TCCAGTAGAATTTCAAGCCAGATGTGGCTGCAGCTGAGTCAGAGTCTGAAAGGGTGGGATCACATGTTCACTGTTGACT
TCTGGACTATTATGGAAAATCACAACCACAGCAAGGAGTCCCACACCCTGCAGGTCATCCTGGGCTGTGAAATGCAAG
AAGACAACAGTACCGAGGGCTACTGGAAGTACGGGTATGATGGGCAGGACCACCTTGAATTCTGCCCTGACACACTGG
ATTGGAGAGCAGCAGAACCAGGGCTGGCCACCAAGCTGGAGTGGGAAAGGCACAAGATTCGGGCCAGGCAGAAC
AGGGCCTACCTGGAGAGGGACTGCCCTGCACAGCTGCAGCAGTTGCTGGAGCTGGGGAGAGGTGTTTTGGACCAACAA
GTGCCTCCTTTGGTGAAGGTGACACATCATGTGACCTCTTCAGTGACCACTCTACGGTGTCCGGCCTTGAATACTACCC
CCAGAACATCACCATGAAGTGGTGAAGGATAAGCAGCCAATGGATGCCAAGGAGTTCGAACTAAAGACGTATTGCC
AATGGGGATGGGACCTACCAGGGCTGGATAAACCTTGGCTGTACCCCTGGGGAAGAGCAGAGATATACGTGCCAGGTG
GAGACCCAGGCCTGGATCAGCCCCCTCATTGTGATCTGGGAGCCCTCACCGTCTGGCACCCTAGTCATTGGAGTCATCA
GTGGAATTGCTGTTTTTGTGCTCATCTTGTTCATTGGAATTTTGTTCATAATATTAAGGAAGAGGCAGGGTTCAAGAGGA
GCCATGGGGCACTACGTCTTAGCTGAACGTGAGTGACACGCAGCCTGCAGACTCACTGTGGGAAGGAGACAAAAGTAG
AGACTCAAAGAGGGAGTGCATTTATGAGCTCTTCATGTTTCAGGAGAGAGTTGAACCTAAACATAGAAATTGCCTGAC
GAACTCCTTGATTTTAGCCTTCTCTGTTTCAATTTCCCTCAAAAAGATTTCCCATTTAGGTTTCTGAGTTCCTGCATGCCGGT
GATCCCTAGCTGTGACCTCTCCCCTGGAAGTGTCTCTCATGAACCTCAAGCTGCATCTAGAGGCTTCCTTCATTTCCCTCC
GTCACCTCAGAGACATACACCTATGTCATTTTCAATTTCCCTATTTTTGGAAGAGGACTCCTTAAATTTGGGGGACTTACATG
ATTCATTTTAAACATCTGAGAAAAGCTTTGAACCCTGGGACGTGGCTAGTCATAACCTTACCAGATTTTACACATGTATC
TATGCATTTTCTGGACCCGTTCAACTTTTCTTTGAATCCTCTCTGTGTTACCCAGTAACTCATCTGTCACCAAGCCTT
GGGATTCTCCATCTGATTGTGATGTGAGTTGCACAGCTATGAAGGCTGTACACTGCACGAATGGAAGAGGCACCTGT
CCCAGAAAAGCATCATGGCTATCTGTGGGTAGTATGATGGGTGTTTTTAGCAGGTAGGAGGCAAATATCTTGAAAGG
GGTTGTGAAGAGGTGTTTTTCTAATTGGCATGAAGGTGTCATACAGATTTGCAAAGTTTAAATGGTGCCTTCATTTGGGA
TGCTACTCTAGTATTCCAGACCTGAAGAATCACAATAATTTTCTACCTGGTCTCTCCTTGTCTGATAATGAAAATTATG
ATAAGGATGATAAAAAGCACTTACTTCGTGTCCGACTCTTCTGAGCACCTACTTACATGCATTACTGCATGCACCTTCTTAC
AATAATTCTATGAGATAGGTACTATTATCCCATTTCTTTTTAAATGAAGAAAGTGAAGTAGGCCGGGCACGGTGGCT
CACGCCTGTAATCCCAGCACTTTGGGAGGCCAAAGCGGGTGGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAA
CATGGTGAACCCCATCTCTAATAAAAAATACAAAAAATTAGCTGGGCGTGGTGGCAGACGCCTGTAGTCCCAGCTACT
CGGAAGGCTGAGGCAGGAGAATGGCATGAACCCAGGAGGCAGAGCTTGCAGTGAGCCGAGTTTGCGCCACTGCACTC
CAGCCTAGGTGACAGAGTGAGACTCCATCTCAAAAAAATAAAAAATAAAAAATAAAAAAATGAAAAAAGAAAGTG
AAGTATAGAGTATCTCATAGTTTGTGAGTATAGAAAACAGGTTTCAAACCTCAGTCAATCTGACCGTTTGATACATCTCA
GACACCACTACATTCAGTAGTTTAGATGCCTAGAATAAATAGAGAAGGAAGGAGATGGCTCTTCTCTTGTCTCATTGTG
TTTCTTCTGAGTGAGCTTGAATCACATGAAGGGGAACAGCAGAAAACAACCAACTGATCCTCAGCTGTCATGTTTCCTT
TAAAAGTCCCTGAAGGAAGTCTTGAATGTGACTCCCTTGTCTCTTGTCTCTTTTGGCATTTCATTTCTTTGGACC
CTACGCAAGGACTGTAATTGGTGGGACAGCTAGTGGCCCTGCTGGGCTCACACACGGTGTCTCCTCCCTAGGCCAGTGC
CTCTGGAGTCAGAACTTGGTGGTATTCCCTCAATGAAGTGGAGTAAGCTCTCTCATTGAGATGGTATAATGGAAG
CCACCAAGTGGCTTAGAGGATGCCAGGTCCTTCATGGAGCCACTGGGGTTCGGGTGCACATTAATAAAAAAATCTA
ACCAGGACATTCAGGAATTGCTAGATTCTGGGAAATCAGTTCACCATGTTCAAAGAGTCTTTTTTTTTTTTTTGGAGACT

CTATTGCCAGGCTGGAGTGCAATGGCATGATCTCGGCTCACTGTAACCTCTGCCTCCCAGGTTCAAGCGATTCTCCTGT
 CTCAGCCTCCCAAGTAGCTGGGATTACAGGCGTGCACCACCATGCCCGGCTAATTTTTGTATTTTAGTAGAGACAGGG
 TTTCACCATGTTGGCCAGGCTGGTCTCGAACTCTCCTGACCTCGTGATCCGCCTGCCTCGGCCTCCCAAAGTGCTGAGAT
 TACAGGTGTGAGCCACCCTGCCAGCCGTCAAAAGAGTCTTAATATATATATCCAGATGGCATGTGTTTACTTTATGTT
 ACTACATGCACCTGGCTGCATAAATGTGGTACAAGCATTCTGTCTTGAAGGGCAGGTGCTTCAGGATACCATATACAGC
 TCAGAAAGTTTCTTTAGGCATTAATTTTAGCAAAGATATCTCATCTCTTCTTTTAAACCATTTTCTTTTTTTGTGGTT
 AGAAAAGTTATGTAGAAAAAAGTAAATGTGATTTACGCTCATTGTAGAAAAGCTATAAAATGAATACAATTAAGCTG
 TTATTTAATTAGCCAGTGAAAACTATTAACAACCTGTCTATTACCTGTTAGTATTATTGTTGCATTAATAAATGCATATA
 CTTTAATAAATGTATATTGTATTGTATACTGCATGATTTTATTGAAGTTCTTGTTCATCTTGTGTATATACTTAATCGCTT
 TGTCAATTTGGAGACATTTATTTGCTTCTAATTTCTTTACATTTTGTCTTACGGAATATTTTCATTCAACTGTGGTAGCC
 GAATTAATCGTGTCTTCTCACTCTAGGGACATTGTCGTCTAAGTTGTAAGACATTGGTTATTTTACCAGCAAACCATTT
 GAAAGCATATGACAAATTTTCTCTTAATATCTTACTATACTGAAAGCAGACTGCTATAAAGGCTTCACTTACTTCTC
 TACCTCATAAGGAATATGTTACAATTAATTTATTAGGTAAGCATTGTTTATATTGGTTTTATTTCACCTGGGCTGAGA
 TTTCAAGAAACACCCAGTCTTACAGTAACACATTTCACTAACACATTTACTAAACATCAGCAACTGTGGCCTGTTAA
 TTTTTTAATAGAAATTTAAGTCCTCATTTTCTTTCGGTGTTTTTTAAGCTTAATTTTCTGGCTTATTTCATAAATCTT
 AAGGTCAACTACATTTGAAAAATCAAAGACCTGCATTTTAAATTTCTTATTACCTCTGGCAAACCATTCACAAACCAT
 GGTAGTAAAGAGAAGGGTGACACCTGGTGGCCATAGGTAATGTACCACGGTGGTCCGGTGACCAGAGATGCAGCGCT
 GAGGGTTTTCTGAAGGTAAAGGAATAAAGAATGGGTGGAGGGCGTGCCTGGAAATCACTTGTAGAGAAAAGCCC
 CTGAAAATTTGAGAAAACAAACAAGAACTACTTACCAGCTATTTGAATTGCTGGAATCACAGGCCATTGCTGAGCTG
 CCTGAACTGGGAACACAACAGAAGGAAAACAAACACTCTGATAATCATTGAGTCAAGTACAGCAGGTGATTGAGGAC
 TGCTGAGAGGTACAGGCCAAAATTTCTTATGTTGTATTATAAATAATGTCATCTTATAAATACTGTCAGTATTTTATAAAACA
 TTCTTCACAACTCACACACATTTAAAAACAAACACTGTCTCTAAAATCCCCAAATTTTTTCATAAACTCAGTTTTAAAC
 TAACTTTTTTTCAAACCACAATCTGATTTAACAATGACTATCATTTAAATATTTCTGACTTTCAAATTAAGATTTTCAC
 ATGCAGGCTGATATTTGTAATTGTGATTCTCTCTGTAGGCTTTGGGTATAATGTGTTCTTTTCTTTTTGCATCAGCGAT
 TAACTTCTACACTCTAACATGTAGAATGTTACTACAATATTAAGTATTTGTATGACAATTTTATTTGAAAGCCTAGGA
 TGCGTTGACATCCTGCATGCATTTATTACTTGATATGCATGCATTCTGGTATCTCAAGCATTCTATTTCTGAGTAATTGTT
 TAAGGTGTAGAAGAGATAGATATGGTGGATTTGGAGTTGATACTTATATATTTTCTATTTCTTGGATGGATGAATTTGTA
 CATTAAGTTTTCCATGGCAGAAA

II. Pairwise Alignment result : Homo sapiens v/s Pan troglodytes

>>Pan Troglodytes HFE 3211 bp (3211 nt)
 Waterman-Eggert score: 9829; 857.5 bits; E(1) < 0
 99.3% identity (99.3% similar) in 1991 nt overlap (351-2341:1221-3211)

	360	370	380	390	400	410
Homo	GGAGTCCCACACCCTGCAGGTCATCCTGGGCTGTGAAATGCAAGAAGACAACAGTACCGA					
	: :::					
Pan	GCAGTCCCACACCCTGCAGGTCATCCTGGGCTGTGAAATGCAAGAAGACAACAGTACCGA					
	1230	1240	1250	1260	1270	1280

	420	430	440	450	460	470
Homo	GGGCTACTGGAAGTACGGGTATGATGGGCAGGACCACCTTGAATTCTGCCCTGACACACT					
	: :::					
Pan	GGGCTACTGGAAGTACGGGTATGATGGGCAGGACCACCTTGAATTCTGCCCTGACACACT					
	1290	1300	1310	1320	1330	1340

	480	490	500	510	520	530
Homo	GGATTGGAGAGCAGCAGAACCCAGGGCCTGGCCCACCAAGCTGGAGTGGGAAAGGCACAA					
	: :::					
Pan	GGATTGGAGAGCAGCAGAACCCAGGGCCTGGCCCACCAAGCTGGAGTGGGAAAGGCACAA					
	1350	1360	1370	1380	1390	1400

	540	550	560	570	580	590
Homo	GATTCGGGCCAGGCAGAACAGGGCCTACCTGGAGAGGGACTGCCCTGCACAGCTGCAGCA					
	: :::					
Pan	GATTCGGGCCAGGCAGAACAGGGCCTACCTGGAGAGGGACTGCCCTGCACAGCTGCAGCA					
	1410	1420	1430	1440	1450	1460

	600	610	620	630	640	650
--	-----	-----	-----	-----	-----	-----

Homo GTTGCTGGAGCTGGGGAGAGGTGTTTTGGACCAACAAGTGCCTCCTTTGGTGAAGGTGAC
 ::
 Pan GTTGCTGGAGCTGGGGAGAGGTGTTTTGGACCAACAAGTGCCTCCTTTGGTGAAGGTGAC
 1470 1480 1490 1500 1510 1520

660 670 680 690 700 710
 Homo ACATCATGTGACCTCTTCAGTGACCACTCTACGGTGTCTGGGCCTTGA ACTACTACCCCCA
 ::
 Pan ACATCATGTGACCTCTTCAGTGACCACTCTACGGTGTCTGGGCCTTGA ACTACTACCCCCA
 1530 1540 1550 1560 1570 1580

720 730 740 750 760 770
 Homo GAACATCACCATGAAGTGGCTGAAGGATAAGCAGCCAATGGATGCCAAGGAGTTCGAACC
 ::
 Pan GAACATCACCATGAAGTGGCTGAAGGATAAGCAGCCAATGGATGCCAAGGAGTTCGAACC
 1590 1600 1610 1620 1630 1640

780 790 800 810 820 830
 Homo TAAAGACGTATTGCCCAATGGGGATGGGACCTACCAGGGCTGGATAACCTTGGCTGTACC
 ::
 Pan TAAAGACGTATTGCCCAATGGGGATGGGACCTACCAGGGCTGGATAACCTTGGCTGTATC
 1650 1660 1670 1680 1690 1700

840 850 860 870 880 890
 Homo CCCTGGGGAAGAGCAGAGATATACGTGCCAGGTGGAGCACCCAGGCCTGGATCAGCCCCT
 ::
 Pan CCCTGGGGAAGAGCAGAGATATACGTGCCAGGTGGAGCACCCAGGCCTGGATCAGCCCCT
 1710 1720 1730 1740 1750 1760

900 910 920 930 940 950
 Homo CATTGTGATCTGGGAGCCCTCACCGTCTGGCACCCCTAGTCATTGGAGTCATCAGTGGAAT
 ::
 Pan CATTGTGATCTGGGAGCCCTCACCGTCTGGCACCCCTAGTCATTGGAGTCATCAGTGGAAT
 1770 1780 1790 1800 1810 1820

960 970 980 990 1000 1010
 Homo TGCTGTTTTGTCTCGTCATCTTGTTTATTGGAATTTTGTTCATAATATTAAGGAAGAGGCA
 ::
 Pan TGCTGTTTTGTCTCGTCATCTTGTTTATTGGAATTTTGTTCATAATATTAAGGAAGAGGCA
 1830 1840 1850 1860 1870 1880

1020 1030 1040 1050 1060 1070
 Homo GGGTTCAAGAGGAGCCATGGGGCACTACGTCTTAGCTGAACGTGAGTGACACGCAGCCTG
 ::
 Pan GGGTTCAAGAGGAGCCATGGGGCACTACGTCTTAGCTGAACGTGAGTGACACGCAGCCTG
 1890 1900 1910 1920 1930 1940

1080 1090 1100 1110 1120 1130
 Homo CAGACTCACTGTGGGAAGGAGACAAA ACTAGAGACTCAAAGAGGGAGTGCATTTATGAGC
 ::
 Pan CAGACTCATTGTGGGAAGGAGACAAA ACTAGAGACTCAAAGAGGGAGTGCATTTATGAGC
 1950 1960 1970 1980 1990 2000

1140 1150 1160 1170 1180 1190
 Homo TCTTCATGTTTCAGGAGAGAGTTGAACCTAAACATAGAAATTGCCTGACGAACTCCTTGA
 ::
 Pan TCTTCATGTTTCAGGAGAGAGTTGAACCTAAACATAGAAATTGCCTGACGAACTCCTTGA
 2010 2020 2030 2040 2050 2060

1200 1210 1220 1230 1240 1250
 Homo TTTTAGCCTTCTCTGTTCATTTCCTCAAAAAGATTTCCCCATTTAGGTTTCTGAGTTCT


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.....
Pan  TTTTAGCCTTCTCTGTTTCATTTCTCTCAAAAAGATTTCCCCATTTAGGTTTCTGAGTTCTCT
    2070  2080  2090  2100  2110  2120

    1260  1270  1280  1290  1300  1310
Homo  GCATGCCGGTGATCCCTAGCTGTGACCTCTCCCCTGGAACCTGTCTCTCATGAACCTCAAG
.....
Pan  GCATGCCAGTGATCCCTAGCTGTGACCTCTCCCCTGGAACCTGTCTCTCATGAACCTCAAG
    2130  2140  2150  2160  2170  2180

    1320  1330  1340  1350  1360  1370
Homo  CTGCATCTAGAGGCTTCCTTCATTTCTCCGTCACCTCAGAGACATACACCTATGTCATT
.....
Pan  CTGCATCTAGAGGCTTCCTTCATTTCTCCGTCACCTCAGAGACATACACCTATGTCATT
    2190  2200  2210  2220  2230  2240

    1380  1390  1400  1410  1420  1430
Homo  TCATTTCTATTTTTGGAAGAGGACTCCTTAAATTTGGGGGACTTACATGATTCATTTTA
.....
Pan  TCATTTCTATTTTTGGAAGAGGACTCCTTAAATTTGGGGGACTCACATGATTCATTTTA
    2250  2260  2270  2280  2290  2300

    1440  1450  1460  1470  1480  1490
Homo  ACATCTGAGAAAAGCTTTGAACCCCTGGGACGTGGCTAGTCATAACCTTACCAGATTTTAA
.....
Pan  ACATCTGAGAAAAGCTTTGAACCCCTGGGACATGGCTAGTCATAACCTTACCAGATTTTAA
    2310  2320  2330  2340  2350  2360

    1500  1510  1520  1530  1540  1550
Homo  CACATGTATCTATGCATTTTCTGGACCCGTTCAACTTTTCCTTTGAATCCTCTCTCTGTG
.....
Pan  CACATGTATCTATGCATTTTCTGGACCCGTTCAACTTTTCCTTTGAATCCTCTCTCTGTG
    2370  2380  2390  2400  2410  2420

    1560  1570  1580  1590  1600  1610
Homo  TTACCCAGTAACTCATCTGTACCAAGCCTGGGGATTCTTCCATCTGATTGTGATGTGA
.....
Pan  TTACCCAGTAACTCATCTGTACCAAGCCTGGGGATTCTTCCATCTGATTGTGATGTGA
    2430  2440  2450  2460  2470  2480

    1620  1630  1640  1650  1660  1670
Homo  GTTGACAGCTATGAAGGCTGTACACTGCACGAATGGAAGAGGCACCTGTCCCAGAAAAA
.....
Pan  GTTGACAGCTATGAAGGCTGTACACTGCACGAATGGAAGAGGCACCTGTCCCAGAAAAA
    2490  2500  2510  2520  2530  2540

    1680  1690  1700  1710  1720  1730
Homo  GCATCATGGCTATCTGTGGGTAGTATGATGGGTGTTTTTAGCAGGTAGGAGGCAAATATC
.....
Pan  GCATCATGGCTATCTGTGGGTAGTATGATGGGTGTTTTTAGCAGGTAGGAGGCAAATATC
    2550  2560  2570  2580  2590  2600

    1740  1750  1760  1770  1780  1790
Homo  TTGAAAGGGGTTGTGAAGAGGTGTTTTTCTAATTGGCATGAAGGTGTCATACAGATTTG
.....
Pan  TTGAAAGGGGTTGTGAAGAGGTGTTTTTCTAATTGGCATGAAGGTGTCATACAGATTTG
    2610  2620  2630  2640  2650  2660

    1800  1810  1820  1830  1840  1850
Homo  CAAAGTTTAATGGTGCCTTCATTTGGGATGCTACTCTAGTATTCCAGACCTGAAGAATCA
.....

```

Pan CAAAGTTTAATGGTGCCTTCATTTGGGATGCTACTCTAGTATTCCAGACCTGAAGAATCA
2670 2680 2690 2700 2710 2720

1860 1870 1880 1890 1900 1910

Homo CAATAATTTTCTACCTGGTCTCTCCTTGTTCTGATAATGAAAATTATGATAAGGATGATA
::: :::

Pan CAATAATTTTCTACCTGGTCTCTCCTTGTTCTGATAATGAAAATTATGATAAGGATGATA
2730 2740 2750 2760 2770 2780

1920 1930 1940 1950 1960 1970

Homo AAAGCACTTACTTCGTGTCCGACTCTTCTGAGCACCTACTTACATGCATTACTGCATGCA
::: :::

Pan AAAGCACTTACTTCGTGTCCGACTCTTCTGAGCACCTACTTACATGCATTACTGCATGCA
2790 2800 2810 2820 2830 2840

1980 1990 2000 2010 2020 2030

Homo CTTCTTACAATAATTCTATGAGATAGGTAATAATCCCATTTCTTTTTTAAATGAAGA
::: :::

Pan ATTCTTACAATAATTCTATGAGATAGGTAATAATCCCATTTCTTTTTTAAATGAAGA
2850 2860 2870 2880 2890 2900

2040 2050 2060 2070 2080 2090

Homo AAGTGAAGTAGGCCGGGCACGGTGGCTCACGCCTGTAATCCAGCACTTTGGGAGGCCAA
::: :::

Pan AAGTGAAGTAGGCCGGGCACGGTGGCTCACGCCTGTAATCCAGCACTTTGGGAGGCCAA
2910 2920 2930 2940 2950 2960

2100 2110 2120 2130 2140 2150

Homo AGCGGGTGGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAACATGGTGAACCCCA
::: :::

Pan AGTGGGTGGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAACATGGTGAACCCCA
2970 2980 2990 3000 3010 3020

2160 2170 2180 2190 2200 2210

Homo TCTCTAATAAAAAATACAAAAAATTAGCTGGGCGTGGTGGCAGACGCCTGTAGTCCCAGCT
::: :::

Pan TCTCTAATAAAAAATACAAAAAATTAGCTGGGCGTGGTGGCAGACGCCTGTAGTCCCAGCT
3030 3040 3050 3060 3070 3080

2220 2230 2240 2250 2260 2270

Homo ACTCGGAAGGCTGAGGCAGGAGAATGGCATGAACCCAGGAGGCAGAGCTTGCAGTGAGCC
::: :::

Pan ACTCGGAAGGCTGAGGCAGGAGAATGGCATGAACCCAGGAGGCAGAGCTTGCAGTGAGCC
3090 3100 3110 3120 3130 3140

2280 2290 2300 2310 2320 2330

Homo GAGTTTGCGCCACTGCACTCCAGCCTAGGTGACAGAGTGAGACTCCATCTCAAAAAATA
::: :::

Pan GAGTTTGCGCCACTGCACTCCAGCCTAGGTGACAGAGTGAGACTCCATCTCAAAAAATA
3150 3160 3170 3180 3190 3200

2340

Homo AAAATAAAAAAT
::: :::

Pan AAAATTAAAAAT
3210

III. Pairwise Alignment result 2: Homo sapiens v/s Gorilla gorilla

>>Gorilla gorilla HFE 5015 bp (5015 nt)

Waterman-Eggert score: 23206; 1309.2 bits; E(1) < 0

93.0% identity (93.0% similar) in 5190 nt overlap (1-5176:111-5015)

```

      10   20   30   40   50   60
Homo  AGAGCTGGGGAAATGGGCCCCGCGAGCCAGGCCGGCGCTTCTCCTCCTGATGCTTTTGCAG
      .....
Gorill AGAGCTGGGGAAATGGGCCCCGCGAGCCAGGCCGGCGCTTCTCCTCCTGATGCTTTTGCAG
      120  130  140  150  160  170
    
```

```

      70   80   90  100  110  120
Homo  ACCGCGGTCCTGCAGGGGCGCTTGCTGCGTTCACACTCTCTGCACTACCTCTTCATGGGT
      .....
Gorill ACCGCGGTCCTGCAGGGGCGCTTGCTGCGTTCACACTCTCTGCACTACCTCTTCATGGGT
      180  190  200  210  220  230
    
```

```

      130  140  150  160  170  180
Homo  GCCTCAGAGCAGGACCTTGGTCTTTCCCTGTTTGAAGCTTTGGGCTACGTGGATGACCAG
      .....
Gorill GCCTCAGAGCAGGACCTTGGTCTTTCCCTGTTTGAAGCTTTGGGCTACGTGGATGACCAG
      240  250  260  270  280  290
    
```

```

      190  200  210  220  230  240
Homo  CTGTTCGTGTTCTATGATCATGAGAGTCGCCGTGTGGAGCCCCGAACTCCATGGGTTTCC
      .....
Gorill CTGTTCGTGTTCTATGATCATGAGAGTCGCCGTGTGGAGCCTCGAACTCCATGGGTTTCC
      300  310  320  330  340  350
    
```

```

      250  260  270  280  290  300
Homo  AGTAGAATTTCAAGCCAGATGTGGCTGCAGCTGAGTCAGAGTCTGAAAGGGTGGGATCAC
      .....
Gorill AGTAGAATTTCAAGCCAGGTGTGGCTGCAGCTGAGTCAGAGTCTGAAAGGGTGGGATCAC
      360  370  380  390  400  410
    
```

```

      310  320  330  340  350  360
Homo  ATGTTCACTGTTGACTTCTGGACTATTATGGAAAATCACAACCACAGCAAGGAGTCCCAC
      .....
Gorill ATGTTCACTGTTGACTTCTGGACTATTATGGAAAATCACAATCACAGCAAGG-----
      420  430  440  450  460
    
```

```

      370  380  390  400  410  420
Homo  ACCCTGCAGGTCATCCTGGGCTGTGAAATGCAAGAAGACAACAGTACCGAGGGCTACTGG
Gorill -----
    
```

```

      430  440  450  460  470  480
Homo  AAGTACGGGTATGATGGGCAGGACCACCTTGAATTCTGCCCTGACACACTGGATTGGAGA
Gorill -----
    
```

```

      490  500  510  520  530  540
Homo  GCAGCAGAACCCAGGGCCTGGCCCACCAAGCTGGAGTGGGAAAGGCACAAGATTCGGGCC
Gorill -----
    
```

```

      550  560  570  580  590  600
    
```

Homo AGGCAGAACAGGGCCTACCTGGAGAGGGACTGCCCTGCACAGCTGCAGCAGTTGCTGGAG

Gorill -----

610 620 630 640 650 660
Homo CTGGGGAGAGGTGTTTTGGACCAACAAGTGCCTCCTTTGGTGAAGGTGACACATCATGTG

.....
Gorill -----TG CCTCCTTTGGTGAAGGTGACACATCATGTG
470 480 490

670 680 690 700 710 720
Homo ACCTCTTCAGTGACCACTCTACGGTGTGCGGCCTTGA ACTACTACCCCCAGAACATCACC

.....
Gorill ACCTCTTCAGTGACCACTCTACGGTGTGCGGCCTTGA ACTACTACCCCCAGAACATCACC
500 510 520 530 540 550

730 740 750 760 770 780
Homo ATGAAGTGGCTGAAGGATAAGCAGCCAATGGATGCCAAGGAGTTCGAACCTAAAGACGTA

.....
Gorill ATGAAGTGGCTGAAGGATAAGCAGCCAATGGATGCCAAGGAGTTCGAACCTAAAGACGTA
560 570 580 590 600 610

790 800 810 820 830 840
Homo TTGCCCAATGGGGATGGGACCTACCAGGGCTGGATAACCTTGGCTGTACCCCCTGGGGAA

.....
Gorill TTGCCCAATGGGGATGGGACCTACCAGGGCTGGATAACCTTGGCTGTACCCCCTGGGGAA
620 630 640 650 660 670

850 860 870 880 890 900
Homo GAGCAGAGATATACGTGCCAGGTGGAGCACCCAGGCCTGGATCAGCCCCTCATTGTGATC

.....
Gorill GAGCAGAGATATACGTGCCAGGTGGAGCACCCAGGCCTGGATCAGCCCCTCATTGTGATC
680 690 700 710 720 730

910 920 930 940 950 960
Homo TGGGAGCCCTCACCGTCTGGCACCCCTAGTCATTGGAGTCATCAGTGGAATTGCTGTTTT

.....
Gorill TGGGAGCCCTCACCGTCTGGCACCCCTAGTCATTGGAGTCATCAGTGGAATTGCTGTTTT
740 750 760 770 780 790

970 980 990 1000 1010 1020
Homo GTCGTCATCTTGTTTCATTGGAATTTTGTTTCATAATATTAAGGAAGAGGCAGGGTTCAAGA

.....
Gorill TTCGTAATCTTGTTTCATTGGAATTTTGTTTCATAATATTAAGGAAGAGGCAGGGTTCAAGA
800 810 820 830 840 850

1030 1040 1050 1060 1070 1080
Homo GGAGCCATGGGGCACTACGTCTTAGCTGAACGTGAGTGACACGCAGCCTGCAGACTCACT

.....
Gorill GGAGCCATGGGGCACTACGTCTTAGCTGAACGTGAGTGACACGCAGCCTGCAGACTCATT
860 870 880 890 900 910

1090 1100 1110 1120 1130 1140
Homo GTGGGAAGGAGACAAA ACTAGAGACTCAAAGAGGGAGTGCATTTATGAGCTCTTCATGTT

.....
Gorill GTGGGAAGGAGACAAA ACTAGAGACTCAA AAAGGGAGTGCATTTATGAGTTCTTCATGTT
920 930 940 950 960 970

1150 1160 1170 1180 1190 1200
Homo TCAGGAGAGAGTTGAACCTAAACATAGAAATTGCCTGACGAACTCCTTGATTTTAGCCTT

.....
Gorill TCAGGACAGAGTTGAACCTAAACATAGAAATTGCCTGAAGAACTCCCTGATTTTAGCCTT
 980 990 1000 1010 1020 1030

1210 1220 1230 1240 1250 1260
Homo CTCTGTTCAATTCCTCAAAAAGATTTCCCCATTTAGGTTTCTGAGTTCCTGCATGCCGGT

Gorill CTCTGTTCAATTTCTCAGAAAAGATTTCCCCATTTAGGTTTCTGAGTTCCTGCATGCCAGT
 1040 1050 1060 1070 1080 1090

1270 1280 1290 1300 1310 1320
Homo GATCCCTAGCTGTGACCTCTCCCCTGGAAGTGTCTCTCATGAACCTCAAGCTGCATCTAG

Gorill GATCCCTAGCTGTGACCTCTCCCCTGGAAGTGTCTCTCATGAACCTCAAGCTGCATCTAG
 1100 1110 1120 1130 1140 1150

1330 1340 1350 1360 1370 1380
Homo AGGCTTCCTTCATTTCCCTCCGTACCTCAGAGACATACACCTATGTCATTTTCATTTTCCTA

Gorill AGGCTTCCTTCATTTCCCTCCATCACCTCAGAGACATACACCTATGTCATTTTCATTTTCCTA
 1160 1170 1180 1190 1200 1210

1390 1400 1410 1420 1430 1440
Homo TTTTGGGAAGAGGACTCCTTAAATTTGGGGGACTTACATGATTCATTTTAACATCTGAGA

Gorill TTTTGTGAAGAGGACTCCTTAAATTTGGGGGACTTACATGATTCATTTTAACATCTGAGA
 1220 1230 1240 1250 1260 1270

1450 1460 1470 1480 1490 1500
Homo AAAGCTTTGAACCCTGGGACGTGGCTAGTCATAACCTTACCAGATTTTACACATGTATC

Gorill AAAGCTTTGAACCCTGGGACATGGCTAGTCATAACCTTACCAGATTTTACACATGTATC
 1280 1290 1300 1310 1320 1330

1510 1520 1530 1540 1550 1560
Homo TATGCATTTTCTGGACCCGTTCAACTTTTCCTTTGAATCCTCTCTCTGTGTTACCCAGTA

Gorill TATGCATTTTCTGGACCCGTTCAACTTTTCCTTTGAATCCTCTCTCTGTGTTACCCAGTA
 1340 1350 1360 1370 1380 1390

1570 1580 1590 1600 1610 1620
Homo ACTCATCTGTCACCAAGCCTTGGGGATTCTTCCATCTGATTGTGATGTGAGTTGCACAGC

Gorill ACTCATCTGTCACCAAGCCCTGGGGATTCTTCCATCTGATTGTGATGTGAGTTGCACAGC
 1400 1410 1420 1430 1440 1450

1630 1640 1650 1660 1670 1680
Homo TATGAAGGCTGTACTGACACTGCACGAATGGAAGAGGCACCTGTCCCAGAAAAAGCATCATGGC

Gorill TATGAAGGCTGTACTGACACTGCACGAATGGAAGAGGCACCTGTCCCAGAAAAAGCATCATGGC
 1460 1470 1480 1490 1500 1510

1690 1700 1710 1720 1730 1740
Homo TATCTGTGGGTAGTATGATGGGTGTTTTTAGCAGGTAGGAGGCAAATATCTTGAAAGGGG

Gorill TATCTGTGGGTAGTATGATGGGTGTTTTTAGCAGGTAGGAGGCAAATATCTTGAAAGGGG
 1520 1530 1540 1550 1560 1570

1750 1760 1770 1780 1790
Homo TTGTGAAGAGGTGT-TTTTTCTAATTGGCATGAAGGTGTCATACAGATTTGCAAAGTTTA

Gorill TTGTGAAGAGGTGTCTTTTTCTAATTGGCATGAAGTTGTCATACAGATTGCAAAGTTA
1580 1590 1600 1610 1620 1630

1800 1810 1820 1830 1840 1850
Homo ATGGTGCCTTCATTTGGGATGCTACTCTAGTATTCCAGACCTGAAGAATCACAATAATTT
:: ::

Gorill ATGATGCCTTCATTTGGGATGCTACTCTAGTATTCCAGACCTGAAGAATCACAATAATTT
1640 1650 1660 1670 1680 1690

1860 1870 1880 1890 1900
Homo TCTACCTGGTCTCTCCTTGTCTGATAATGAAAATTATGATAAGGATGAT-----
:: ::

Gorill TCTACCTGGTCTCTCCTTGTCTGATAATGAAAATTATGATAAGGATGATGATAATGATG
1700 1710 1720 1730 1740 1750

1910 1920 1930 1940 1950 1960
Homo --AAAAGCACTTACTTCGTGTCCGACTCTTCTGAGCACCTACTTACATGCATTACTGCAT
:: ::

Gorill AATAAAGCACTTACTTCGTGTCCGACTCTTCTGAGCACCTACTTACATGCATTACTGCAT
1760 1770 1780 1790 1800 1810

1970 1980 1990 2000 2010 2020
Homo GCACTTCTTACAATAATTCTATGAGATAGGTACTATTATCCCCATTTCTTTTTTAAATGA
:: ::::::::::::::::::::::::::::::: ::::::::::::::::::::::

Gorill GCAATTCTTACAATAATTCTATGAGATAGGTACTATCATCCCCATTTCTTTTTTAAATGA
1820 1830 1840 1850 1860 1870

2030 2040 2050 2060 2070 2080
Homo AGAAAGTGAAGTAGGCCGGGCACGGTGGCTCACGCCTGTAATCCCAGCACTTTGGGAGGC
:: ::

Gorill AGAAAGTGAAGTAGGCCGGGCACGGTGGCTCACGCCTGTAATCCCAGCACTTTGGGAGGC
1880 1890 1900 1910 1920 1930

2090 2100 2110 2120 2130 2140
Homo CAAAGCGGGTGGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAACATGGTCAAACC
:: ::::::::::::::::::::::::::::::: ::::::::::::::::::::::

Gorill CAAAGCGAGTGGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAACATGGTCAAACC
1940 1950 1960 1970 1980 1990

2150 2160 2170 2180 2190 2200
Homo CCATCTCTAATAAAAAATACAAAAAATTAGCTGGGCGTGGTGGCAGACGCCTGTAGTCCCA
:: ::

Gorill CCATCTCTAATAAAAAATACAAAAAATTAGCTGGGCGTGGTGGCAGACGCCTGTAGTCCCA
2000 2010 2020 2030 2040 2050

2210 2220 2230 2240 2250 2260
Homo GCTACTCGGAAGGCTGAGGCAGGAGAATGGCATGAACCCAGGAGGCAGAGCTTGCAGTGA
:: ::::::::::::::::::::::::::::::: ::::::::::::::::::::::

Gorill GCTACTCGGAAGGCTGAGGCAGGAGAATGGCATGAACCCAGGAGGCAGAACTTGCAGTGA
2060 2070 2080 2090 2100 2110

2270 2280 2290 2300 2310 2320
Homo GCCGAGTTTGCGCCACTGCACTCCAGCCTAGGTGACAGAGTGAGACTCCATCTCAAAAAA
:: ::

Gorill GCCGAGTTTGCGCCACTGCACTCCAGCCTAGGTGACAGAGTGAGACTCCATCTCAAAAAA
2120 2130 2140 2150 2160 2170

2330 2340 2350 2360 2370 2380
Homo ATAAAAATAAAAAATAAAAAAATGAAAAAAAAAAGAAAGTGAAGTATAGAGTATCTCATAG
:: ::::::::::::::::::::::::::::::: ::::::::::::::::::::::

Gorill ATAAAAATAAAAAATAAAAAAATGAAAAAAAAAAGAAAGTGAAGTATAGAGTATCTCATAG

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    2180    2190    2200    2210    2220    2230
Homo TTTGTCAGTGATAGAAACAGGTTTCAAACCTCAGTCAATCTGACCGTTTGATACATCTCAG
:
:
:
Gorill CTTGTCAGTGATAG----AGGTTTCAAACCTCAGTCAATCTGACCGTTTGATACATCTCAG
    2240    2250    2260    2270    2280    2290

    2450    2460    2470    2480    2490    2500
Homo ACACCACTACATTACAGTAGTTTAGATGCCTAGAATAAATAGAGAAGGAAGGAGATGGCTC
:
:
:
Gorill ACACCACTACATTACAGTAGTTTAGATGCCTAGAGTAAATAGAGAAGGAAGTAGATGGCTC
    2300    2310    2320    2330    2340    2350

    2510    2520    2530    2540    2550    2560
Homo TTCTCTTGCTCATTGTGTTTCTTCTGAGTGAGCTTGAATCACATGAAGGGGAACAGCAG
:
:
:
Gorill TTCTCTTGCTCATTGTGTTTCTTCTGAGTGAGCTTGAATCACATGAAGGGGAACAGCAG
    2360    2370    2380    2390    2400    2410

    2570    2580    2590    2600    2610    2620
Homo AAAACAACCAACTGATCCTCAGCTGTCATGTTTCCTTTAAAAGTCCCTGAAGGAAGGTCC
:
:
:
Gorill AAAACAACCAACTGATCCTCAGCTGTCATGTTTCCTTTAAAAGTCCCTGAAGGAAGGTCC
    2420    2430    2440    2450    2460    2470

    2630    2640    2650    2660    2670    2680
Homo TGGAAATGTGACTCCCTTGCTCCTCTGTTGCTCTCTTTGGCATTTCATTTCTTTGGACCCTA
:
:
:
Gorill TGGAAATGTGACTCCCTTGCTCCTCTGTTGCTCTCTTTGGCATTTCATTTCTTTGGACCCTA
    2480    2490    2500    2510    2520    2530

    2690    2700    2710    2720    2730    2740
Homo CGCAAGGACTGTAATTGGTGGGGACAGCTAGTGGCCCTGCTGGGCTTCACACACGGGTGTC
:
:
:
Gorill CGCAAGGACTGTAATTGGTGGGGACAGCTAGTGGCCCTGCTGGGCTTCACACACAGTGTC
    2540    2550    2560    2570    2580    2590

    2750    2760    2770    2780    2790    2800
Homo CTCCCTAGGCCAGTGCCTCTGGAGTCAGAACTCTGGTGGTATTTCCCTCAATGAAGTGGA
:
:
:
Gorill CTCCCTAGGCCAGTGCCTCTGGAGTCAGAACTCTGGTGGTATTTCCCTCGATGAAGTGGA
    2600    2610    2620    2630    2640    2650

    2810    2820    2830    2840    2850    2860
Homo GTAAGCTCTCTCATTTTGAGATGGTATAATGGAAGCCACCAAGTGGCTTAGAGGATGCCC
:
:
:
Gorill GTAAGCTCTCTCATTTTGAGATGGTATAATGGAAGCCACCAAGTGGCTTAGAGGATGCCC
    2660    2670    2680    2690    2700    2710

    2870    2880    2890    2900    2910    2920
Homo AGGTCCTTCCATGGAGCCACTGGGGTTCCGGTGCACATTAAAAAAAAAATCTAACCAGGA
:
:
:
Gorill AGGTCCTTCCATGGGGCCACTGGGATTCGGTGCACATTAAAAAAAAAATCTAACCAGGA
    2720    2730    2740    2750    2760    2770

    2930    2940    2950    2960    2970    2980
Homo CATTACAGGAATTGCTAGATTCTGGGAAATCAGTTCACCATGTTCAAAAAGAGTCTTTTTTT
:
:
:
Gorill CACTCAGGAATTGCTAGATTCTGGGAAATCAGTTCACCATGTTCAAAAAGAGTCTTTTTTT
    2780    2790    2800    2810    2820    2830

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2990 3000 3010 3020 3030 3040
Homo TTTTTTTGAGACTCTATTGCCAGGCTGGAGTGCAATGGCATGATCTCGGCTCACTGTAA
: ::
Gorill T----TTGAGACTCTATTGCCAGGCTGGAGTGCAATGGCATGATCTCGGCTCACTGTAA
2840 2850 2860 2870 2880

3050 3060 3070 3080 3090 3100
Homo CCTCTGCCTCCCAGGTTCAAGCGATTCTCCTGTCTCAGCCTCCCAAGTAGCTGGGATTAC
:::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill CCTCTGCCTCCCAGGTTCAAGCGATTCTCCTGCCTCAGCCTCCCGAGTAGCTGGGATTAC
2890 2900 2910 2920 2930 2940

3110 3120 3130 3140 3150 3160
Homo AGGCGTGCACCACCATGCCCGGCTAATTTTTGTATTTTTAGTAGAGACAGGGTTTCACCA
:::::::::::::::::::::::::::::::::::: :: ::::::::::::::
Gorill AGGCGTGCACCACCATGCCCGGCTAATTTTTGTATTTTTAGTAGATACGGGGTTTCACCA
2950 2960 2970 2980 2990 3000

3170 3180 3190 3200 3210 3220
Homo TGTTGGCCAGGCTGGTCTCGAACTCTCCTGACCTCGTGATCCGCCTGCCTCGGCCTCCCA
:::::::::::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill TGTTGGCCAGGCTGGTCTCGAACTCTCCTGACCTCGTGATCCGCCTGCCTCGGCCTCCCA
3010 3020 3030 3040 3050 3060

3230 3240 3250 3260 3270 3280
Homo AAGTGCTGAGATTACAGGTGTGAGCCACCCTGCCAGCCGTCAAAAAGAGTCTTAATATAT
:::::::::::::::::::::::::::: :: ::::::::::::::::::::
Gorill AAGTGCTGAGATTACAGGTGTGAGCCACCAGCTCAGCCGTCAAAAAGAGTCTTAATATAT
3070 3080 3090 3100 3110 3120

3290 3300 3310 3320 3330 3340
Homo ATATCCAGATGGCATGTGTTTACTTTATGTTACTACATGCACTTGGCTGCATAAATGTGG
:::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill ATATCCAGATGGCATGTGTTTACTTTATGTTACTACATGCATTTGGCTGCATAAATGTGG
3130 3140 3150 3160 3170 3180

3350 3360 3370 3380 3390 3400
Homo TACAAGCATTCTGTCTTGAAGGGCAGGTGCTTCAGGATACCATATACAGCTCAGAAGTTT
:::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill TACAAGCATTCTGTCTTGAAGGGCAGGTGCTTCAGGATACCATATACAGCTCAGAAGTTT
3190 3200 3210 3220 3230 3240

3410 3420 3430 3440 3450 3460
Homo CTTCTTTAGGCATTAAATTTTAGCAAAGATATCTCATCTCTTCTTTTAAACCATTTTCTT
:::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill CTTCTTTAGGCATTAAATTTTAGCAAAGATATCTCATCTCTTCTTTTAAACCATTTTCTT
3250 3260 3270 3280 3290 3300

3470 3480 3490 3500 3510 3520
Homo TTTTGTGGTTAGAAAAGTTATGTAGAAAAAAGTAAATGTGATTTACGCTCATTGTAGAA
:::: :::::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill TTTTCTGGTTAGAAAAGTTATGTAGAAAAAAGTAAATGTGATTTACGCTTATTGTAGAA
3310 3320 3330 3340 3350 3360

3530 3540 3550 3560 3570 3580
Homo AAGCTATAAAAATGAATACAATTAAGCTGTTATTTAATTAGCCAGTGAAAACTATTAAC
:::::::::::::::::::::::::::: ::::::::::::::::::::
Gorill AAGCTATAAAAATGAATACAATTAAGCTGCTATTTAATTAGCCGGTGAAAACTATTAAC
3370 3380 3390 3400 3410 3420

3590 3600 3610 3620 3630 3640
Homo AACTTGTCTATTACCTGTTAGTATTATTGTTGCATTAAAAATGCATATACTTTAATAAAAT
: : : : : :

Gorill AACTTGTCTATTACCTGTTAGTATTATTGTTGCATTAAAAATGCATATACTTTAATAAAAT
3430 3440 3450 3460 3470 3480

3650 3660 3670 3680 3690 3700
Homo GTATATTGTATTGTATACTGCATGATTTTATTGAAGTTCTTGTTTCATCTTGTGTATATAC
: : : : : :

Gorill GTACATTGTATTGTATACTGCGTGATTTTATTGATGTTCTTGTTTCATCTTGTGTATATAC
3490 3500 3510 3520 3530 3540

3710 3720 3730 3740 3750 3760
Homo TTAATCGCTTTGTCATTTTGGAGACATTTATTTTGCTTCTAATTTCTTTACATTTTGTCT
: : : : : :

Gorill TTAATCGCTTTGTCATTTTGGAGACATTTATTTTGCTTCTAATTTCTTTACATTTTGTCT
3550 3560 3570 3580 3590 3600

3770 3780 3790 3800 3810 3820
Homo TACGGAATATTTTCATTCAACTGTGGTAGCCGAATTAATCGTGTTCCTTCACTCTAGGGA
: : : : : :

Gorill TACGGAATATTTTCATTCAACTGTGGTAGCCGAATTAATCATGTTCCTTCACTCTAGGGA
3610 3620 3630 3640 3650 3660

3830 3840 3850 3860 3870 3880
Homo CATTGTCGTCTAAGTTGTAAGACATTGGTTATTTTACCAGCAAACCATTCTGAAAGCATA
: : : : : :

Gorill CATTGTCGTCTAAGTTGTAAGACATTGGTTATTTTACCAACAAACCATTCTGAAAGCATA
3670 3680 3690 3700 3710 3720

3890 3900 3910 3920 3930 3940
Homo TGACAAATATTTCTCTCTTAATATCTTACTATACTGAAAGCAGACTGCTATAAGGCTTC
: : : : : :

Gorill TGACAAATCTTTCTCTCTTAATATCGTACTATACTGAAAGCAGACTGCTATAAGGCTTC
3730 3740 3750 3760 3770 3780

3950 3960 3970 3980 3990 4000
Homo ACTTACTCTTCTACCTCATAAGGAATATGTTACAATTAATTTATTAGGTAAGCATTTGTT
: : : : : :

Gorill ATTTACTCTTCTACCTCATAAGGAATATGTTACAATTAATTCATTAGGTAAGCATTTGTT
3790 3800 3810 3820 3830 3840

4010 4020 4030 4040 4050 4060
Homo TTATATTGGTTTTATTTCACCTGGGCTGAGATTTCAAGAAACACCCCAGTCTTCACAGTA
: : : : : :

Gorill TTATATTGGTTTTATTTCACCTGGGCTGAGATTTCAAGAAACACCCCAGTCTTCACAGTA
3850 3860 3870 3880 3890 3900

4070 4080 4090 4100 4110 4120
Homo ACACATTTCACTAACACATTTACTAAACATCAGCAACTGTGGCCTGTTAATTTTTTTT-AA
: : : : : : : :

Gorill ACACATTTGCTAACACATTTACTAAACATCAGCAACTGTGGCCTGTTAATTTTTTTTTAA
3910 3920 3930 3940 3950 3960

4130 4140 4150 4160 4170 4180
Homo TAGAAATTTTAAGTCCTCATTTTCTTTCGGTGTTTTTTAAGCTTAATTTTTCTGGCTTTA
: : : : : :

Gorill TAGAAATCTTAAGTCCCATTTTCTTTCGGTGTTTTTTAAGCTTAATTTTTCTGGCTTTA
3970 3980 3990 4000 4010 4020

4190 4200 4210 4220 4230 4240

Homo TTCATAAATTCTTAAGGTCAACTACATTTGAAAAATCAAAGACCTGCATTTTAAATTCTT

 Gorill TTCATAAATTCTTAAGGTCAACTACATTTGAAAAATCAAAGACCTGCATTTTAAATTCTT
 4030 4040 4050 4060 4070 4080

4250 4260 4270 4280 4290 4300
 Homo ATTCACCTCTGGCAAACCATTACAAACCATGGTAGTAAAGAGAAGGGTGACACCTGGT

 Gorill ATTCACCTCTGGCAAACCATTACAAACCATGGTAGTAAAGAGAAGGGTGACACCTGGT
 4090 4100 4110 4120 4130 4140

4310 4320 4330 4340 4350 4360
 Homo GGCCATAGGTAAATGTACCACGGTGGTCCGGTGACCAGAGATGCAGCGCTGAGGGTTTTTC

 Gorill GGCCATAGGTAAACGTACCACGGTGGTCCGGTGACCAGAGATGCAGCGCTGAGGGTTTTTC
 4150 4160 4170 4180 4190 4200

4370 4380 4390 4400 4410 4420
 Homo CTGAAGGTAAAGGAATAAAGAATGGGTGGAGGGGCGTGCCTGAAATCACTTGTAGAGA

 Gorill CTGAAGGTAAAGGAATAAAGAATGGGTGGAGGGGCGTGCCTGAAATCACTTGTAGAGA
 4210 4220 4230 4240 4250 4260

4430 4440 4450 4460 4470 4480
 Homo AAAGCCCCTGAAAATTTGAGAAAACAAACAAGAACTACTTACCAGCTATTTGAATTGCT

 Gorill AAAGCCCCTGAAAATTTGAGAAAACAAACAAGAACTACTTACCAGCTATTTGAATTGCT
 4270 4280 4290 4300 4310 4320

4490 4500 4510 4520 4530 4540
 Homo GGAATCACAGGCCATTGCTGAGCTGCCTGAACTGGGAACACAACAGAAGGAAAAACAAACC

 Gorill GGAATCACAGGCCATTGCTGAGCTGCCTGAACTGGGAACACAACAGAAGGAAAAACAAACC
 4330 4340 4350 4360 4370 4380

4550 4560 4570 4580 4590 4600
 Homo ACTCTGATAATCATTGAGTCAAGTACAGCAGGTGATTGAGGACTGCTGAGAGGTACAGGC

 Gorill ACTCTGATAATCATTGAGTCAAGTACAGCAGGTGATTGAGGACTGCTGAGAGGTACAGGC
 4390 4400 4410 4420 4430 4440

4610 4620 4630 4640 4650 4660
 Homo CAAAATTCTTATGTTGTATTATAATAATGTCATCTTATAATACTGTCAGTATTTTATAAA

 Gorill CAAAATTCTTATGTTGTATTATAATAATGTCATCTTATAATACTGTCAGTATTTTATAAA
 4450 4460 4470 4480 4490 4500

4670 4680 4690 4700 4710 4720
 Homo ACATTCTTCACAACTCACACACATTTAAAAACAAAACACTGTCTCTAAAATCCCCAAAT

 Gorill ACATTCTTCACAACTCACACACATTTAAAAACAAAACACTGTCTCTAAAATCCCCAAAT
 4510 4520 4530 4540 4550 4560

4730 4740 4750 4760 4770 4780
 Homo TTTTCATAAACTCAGTTTTAACTAACTTTTTTCAAACCACAATCTGATTTAACAATGA

 Gorill TTTTCATAAACTCAGTTTTAACTAACTTTTTTCAAACCACAATCTGATTTAACAATGA
 4570 4580 4590 4600 4610 4620

4790 4800 4810 4820 4830 4840
 Homo CTATCATTTAAATATTTCTGACTTTCAAATTAAGATTTTTCACATGCAGGCTGATATTG

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Gorill CTATCATTTAAATATTTCTGACTTTCAAATTAAGATTTTCACATGCAGGCTGATATTTG
4630 4640 4650 4660 4670 4680

4850 4860 4870 4880 4890 4900
Homo TAATTGTGATTCTCTCTGTAGGCTTTGGGTATAATGTGTTCTTTTCCTTTTTTGCATCAG
.....
Gorill TAATTGTGATTCTCTCTGTAGGCTTTGGGTATAATGTGTTCTTTTCCTTTTTTGCATCAG
4690 4700 4710 4720 4730 4740

4910 4920 4930 4940 4950 4960
Homo CGATTAACTTCTACACTCTAACATGTAGAATGTTACTACAATATTAAGTATTTTGTATG
.....
Gorill CGATTAACTTCTACACTCTAACATGTAGAATGTTACTACAATATTAAGTATTTTGTATG
4750 4760 4770 4780 4790 4800

4970 4980 4990 5000 5010 5020
Homo ACAATTTTATTTGAAAGCCTAGGATGCGTTGACATCCTGCATGCATTTATTACTTGATAT
.....
Gorill ACAATTTTATTTGAAAGCCTAGGATGCGTTGACATCCTGCATACATTTATTACTTGATAT
4810 4820 4830 4840 4850 4860

5030 5040 5050 5060 5070 5080
Homo GCATGCATTCTGGTATCTCAAGCATTCTATTTCTGAGTAATTGTTTAAGGTGTAGAAGAG
.....
Gorill GCATGCATTCTGGTATCTCAAGCATTCTATTTCTGAGTAATTGTTTAAGGTGTAGAAGAG
4870 4880 4890 4900 4910 4920

5090 5100 5110 5120 5130 5140
Homo ATAGATATGGTGGATTTGGAGTTGATACTTATATATTTTCTATTTCTTGGATGGATGAAT
.....
Gorill ATAGATATGGTGGATTTGGAGTTGATACTTATATATTTTCTATTTCTTGGATGGATGAAT
4930 4940 4950 4960 4970 4980

5150 5160 5170
Homo TTGTACATTA AAAAGTTTTCCATGGCAGAAA
.....
Gorill TTGTACATTA AAAAGTTTTCCATGGCAGAAA
4990 5000 5010

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