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(54) **METHOD AND A SYSTEM FOR PREDICTING PROTEIN FUNCTIONAL SITE, A METHOD FOR IMPROVING PROTEIN FUNCTION, AND A FUNCTION-MODIFIED PROTEIN**

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(57) **ABSTRACT**

The present application provides a method for predicting the functional site of a protein using data of the entire proteins of an organism of which genome data or cDNA data is known. More specifically, the present application provides a method for predicting a protein functional site, comprising the steps of calculating the frequency of occurrence of an oligopeptide in the entire proteins, calculating the value of each amino-acid residue contributing to the frequency of occurrence as the representative value of the function, and predicting the protein functional site by using the representative value of function as an indicator. The present also provides a system for predicting a functional site for automatically performing said methods. Additionally, the present application provides a method for preparing a function-modified protein comprising subjecting the amino-acid residues composing the functional site identified by the method described above to artificial mutation, and a novel thermophilic DNA polymerase prepared by the method.

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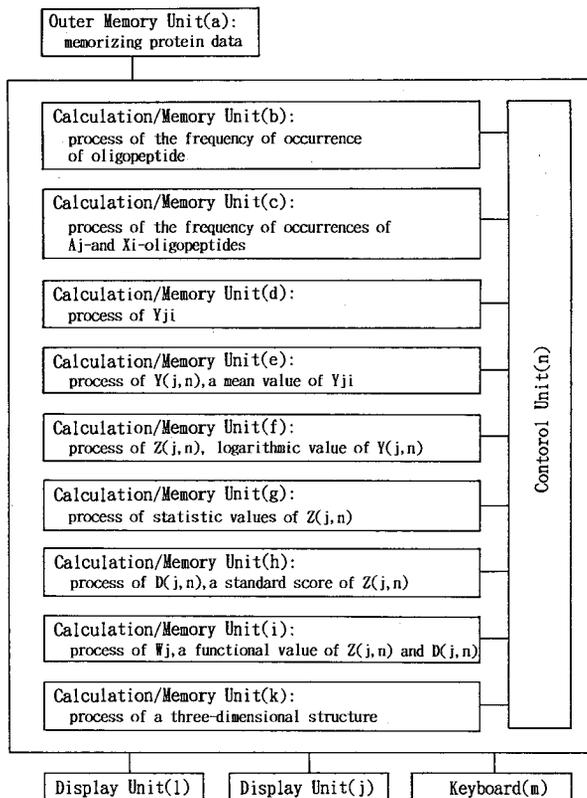


Fig. 1

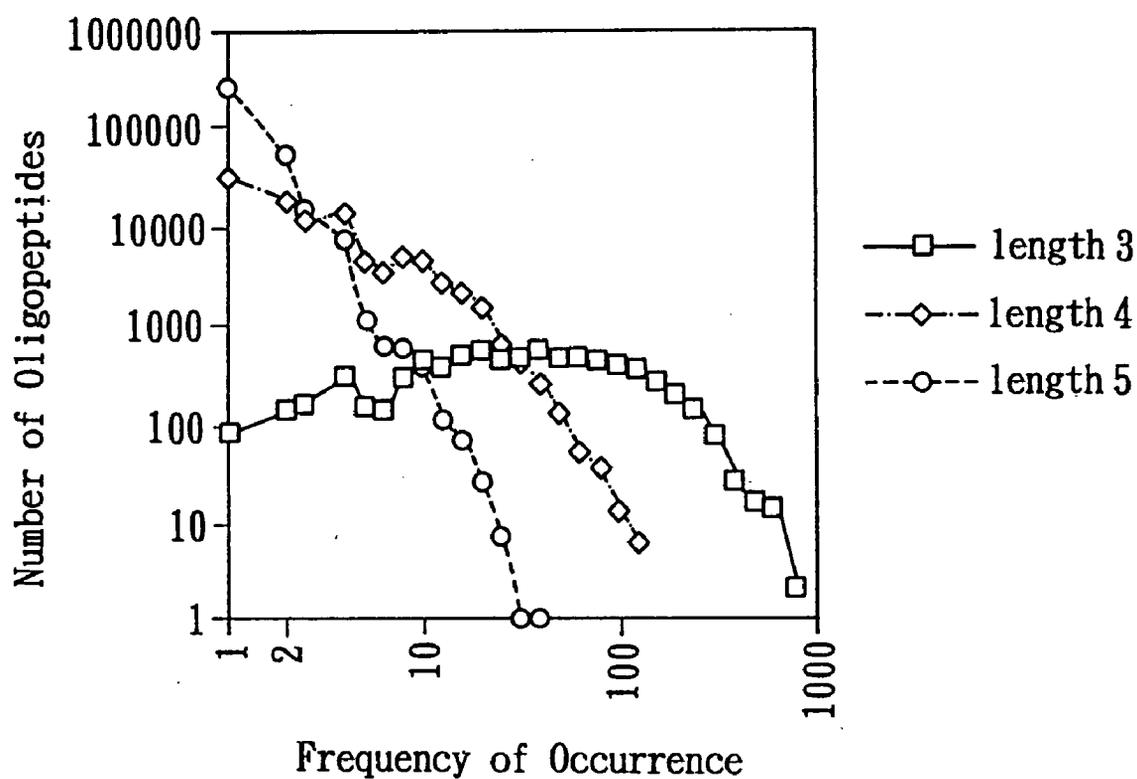


Fig. 3

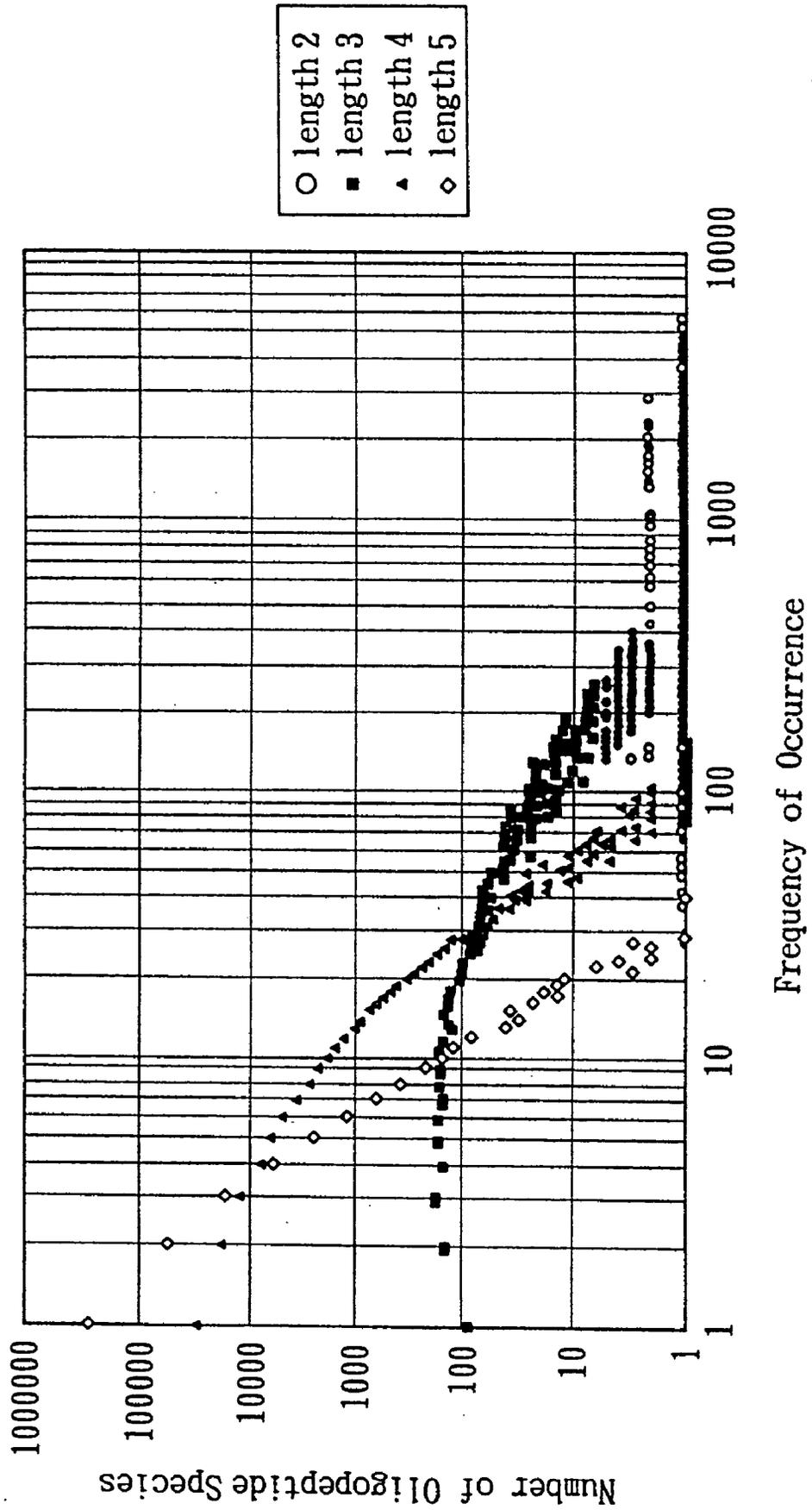


Fig. 4

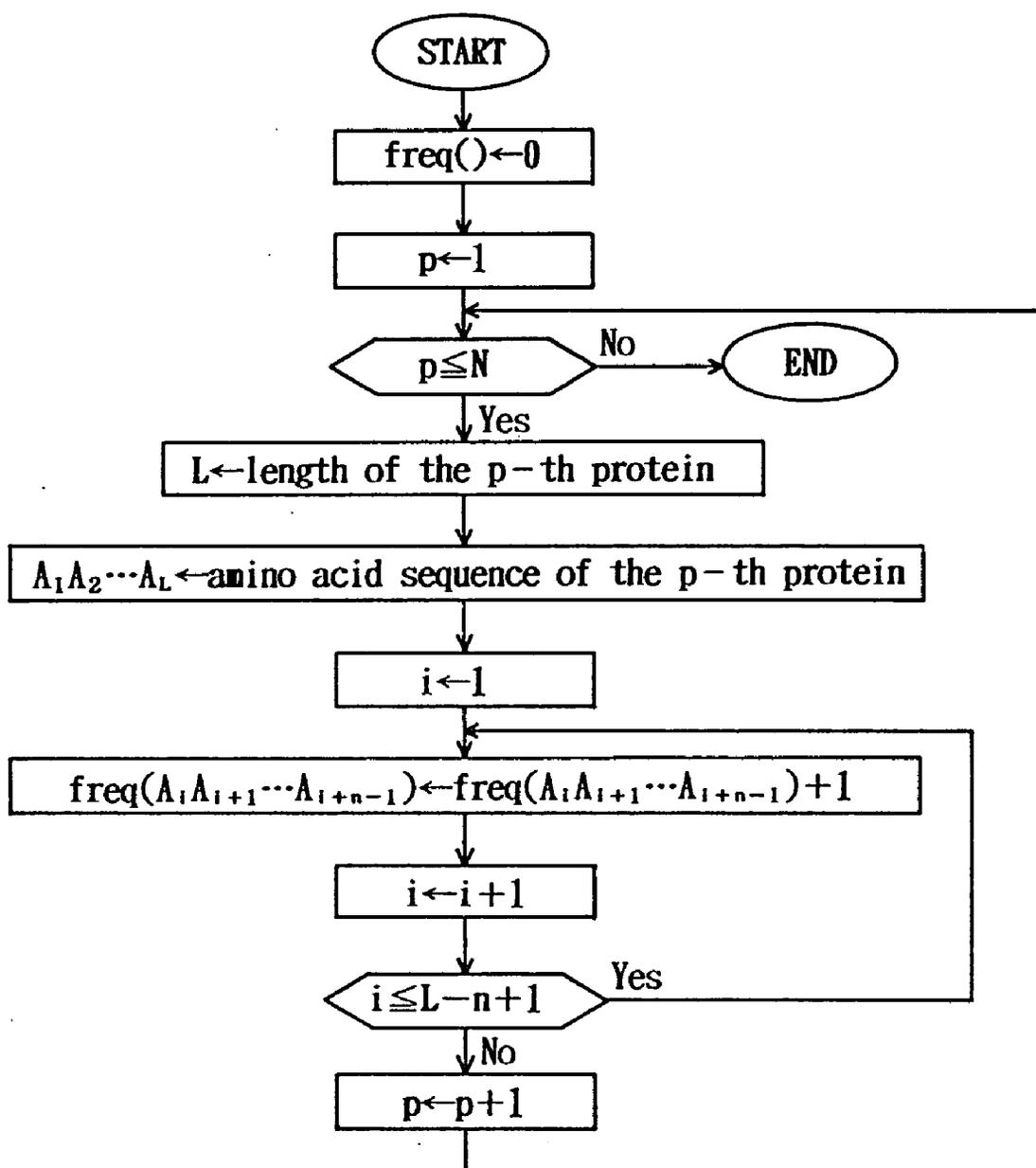


Fig. 5

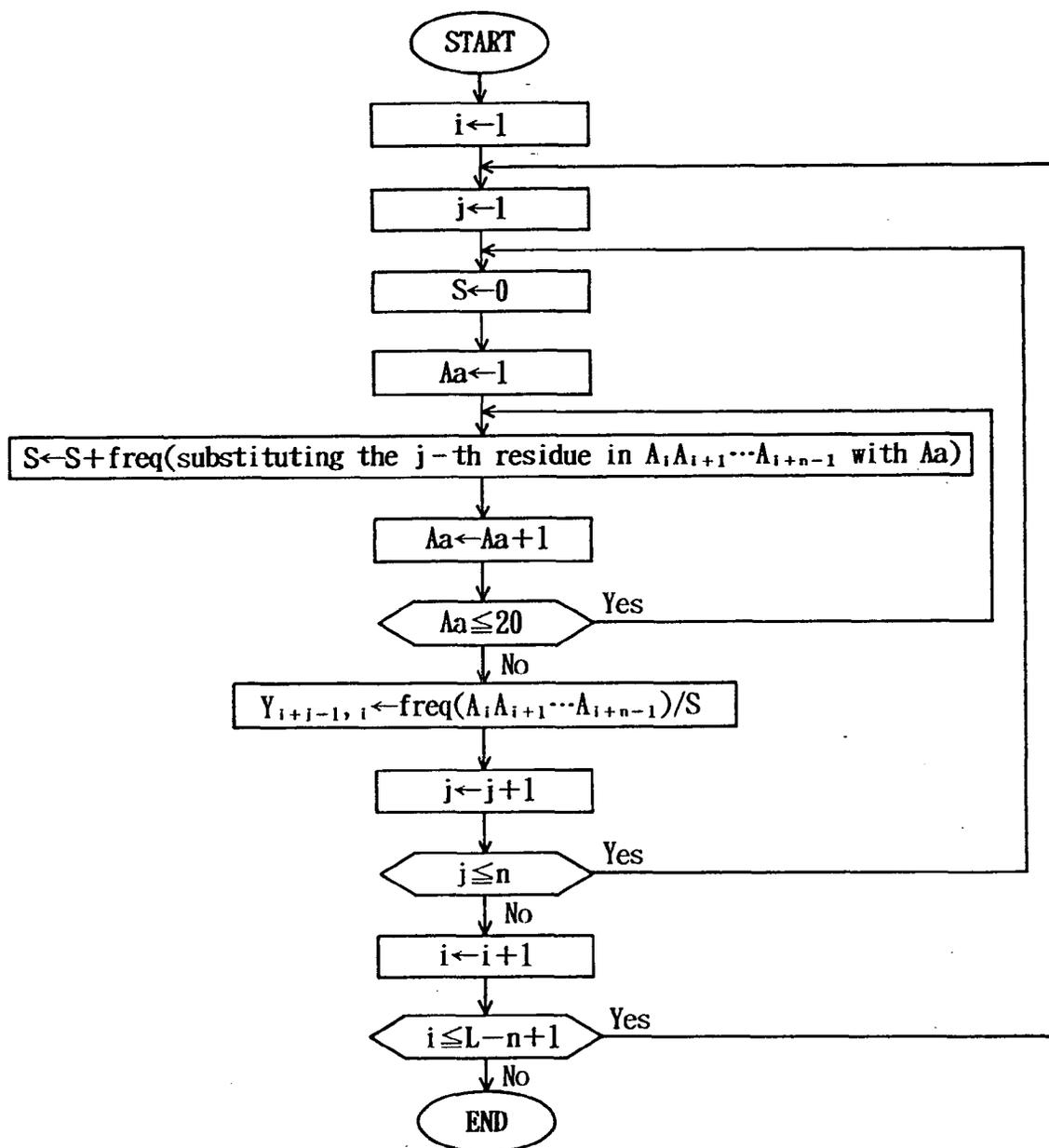


Fig. 6

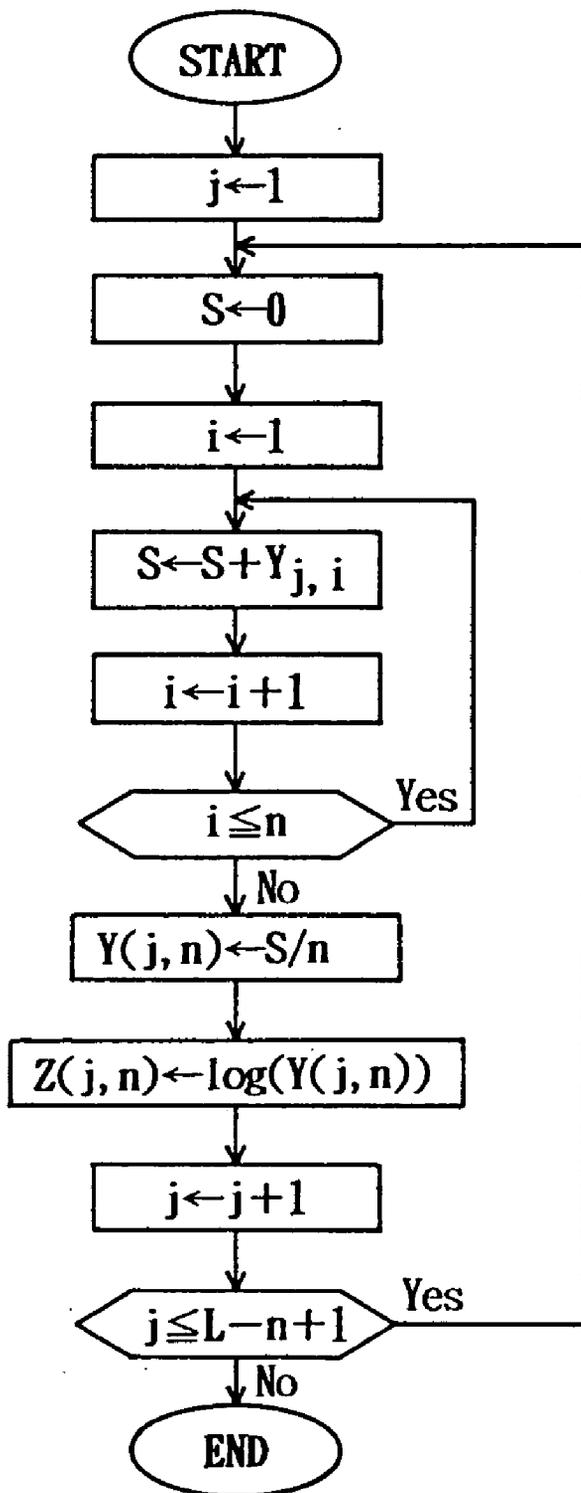


Fig. 7

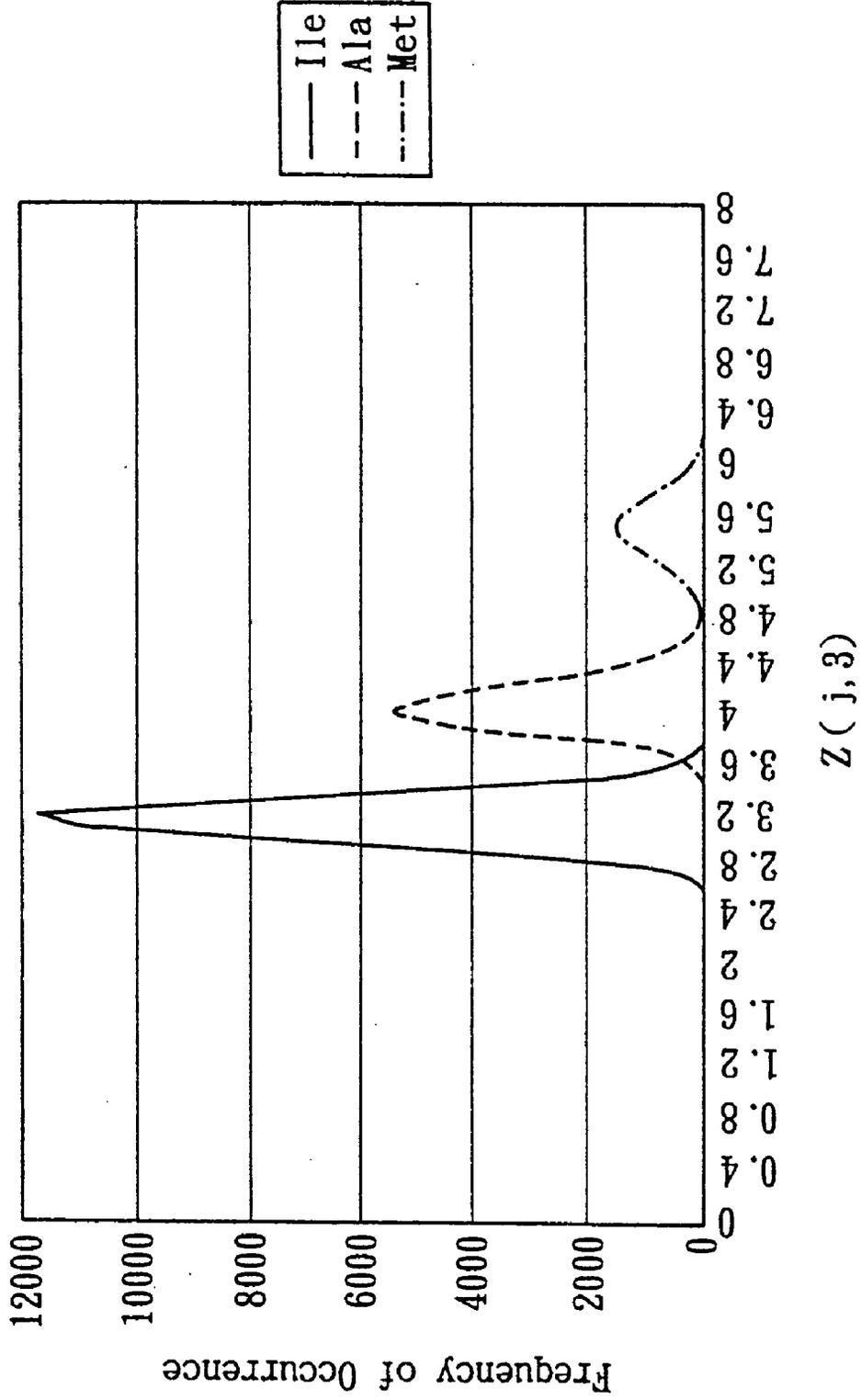


Fig. 8

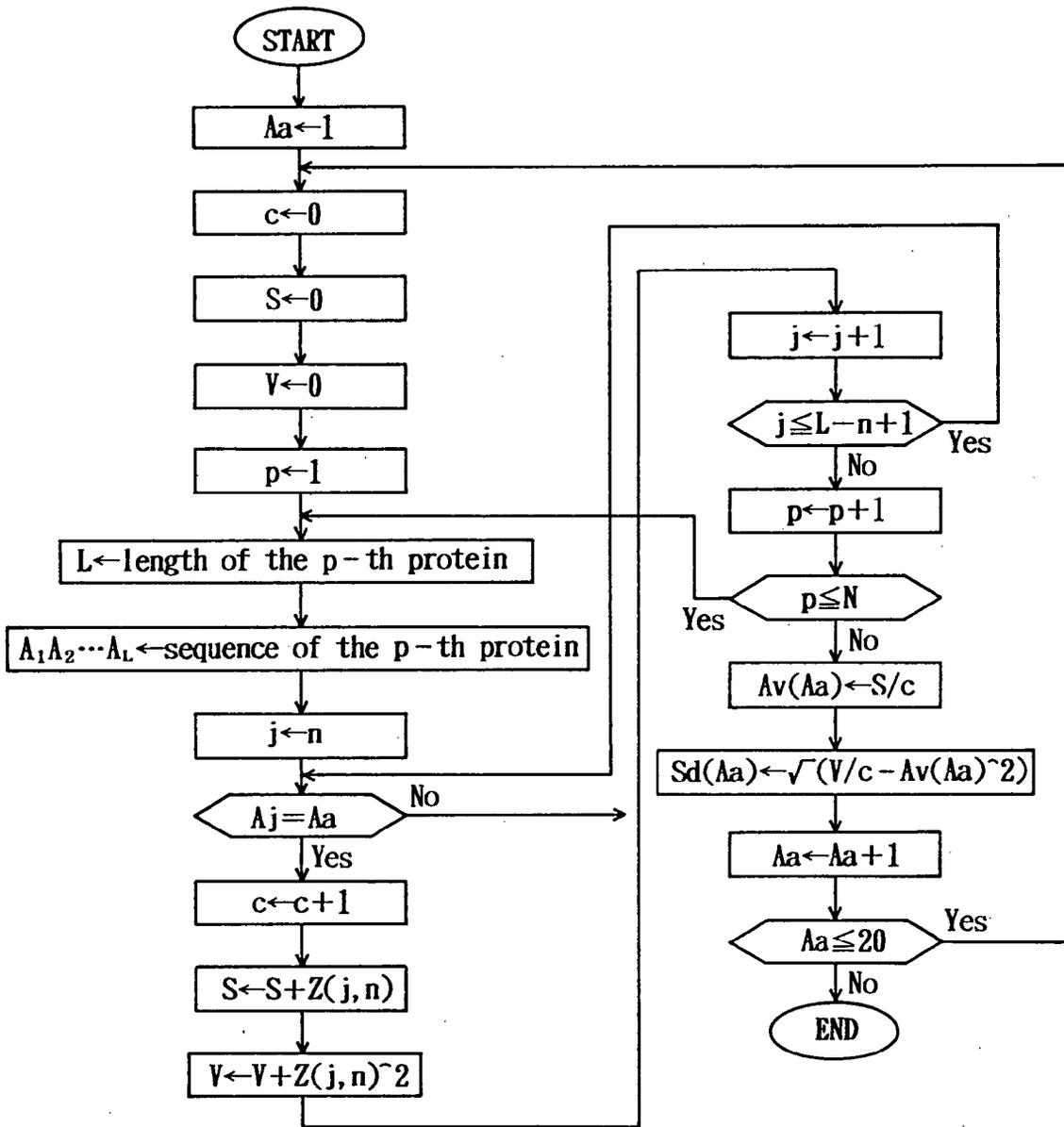


Fig. 9

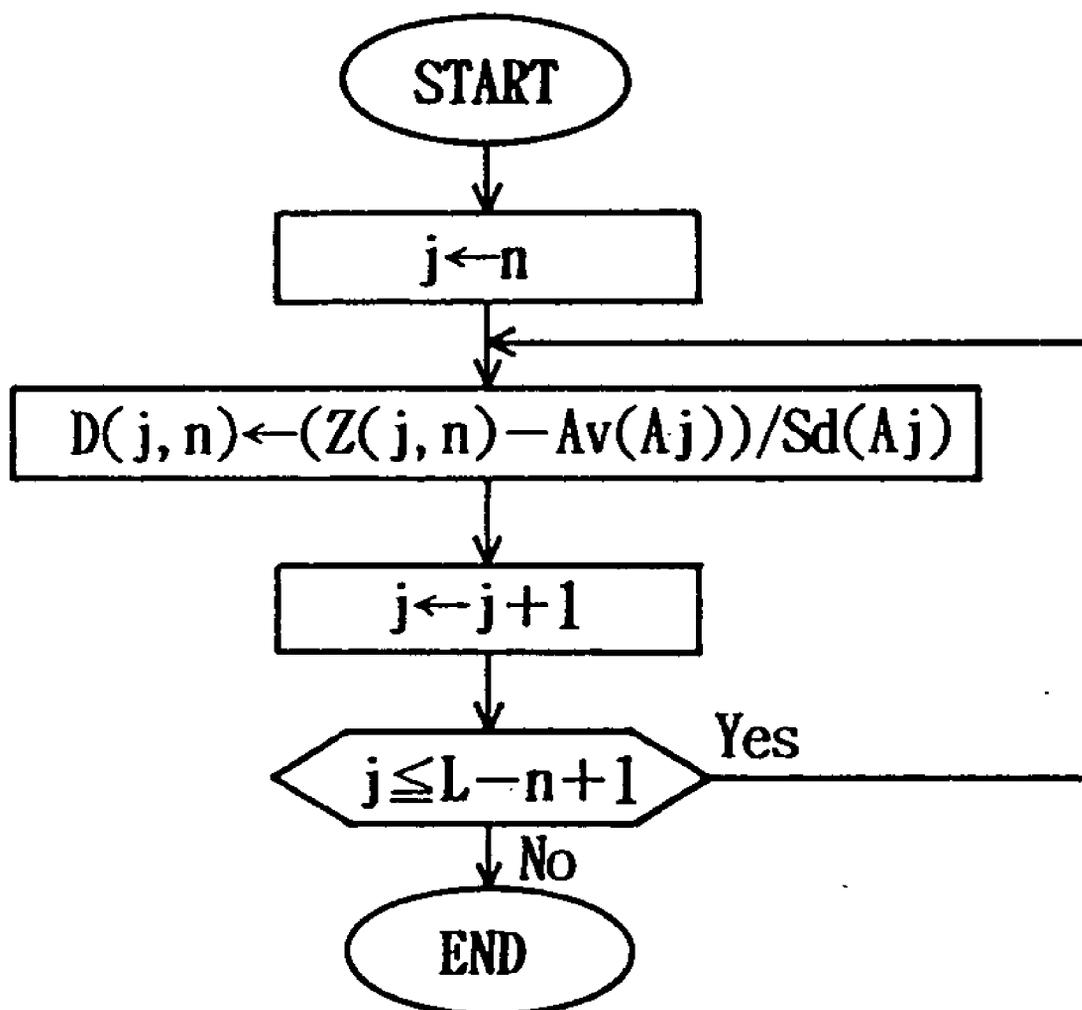


Fig. 10

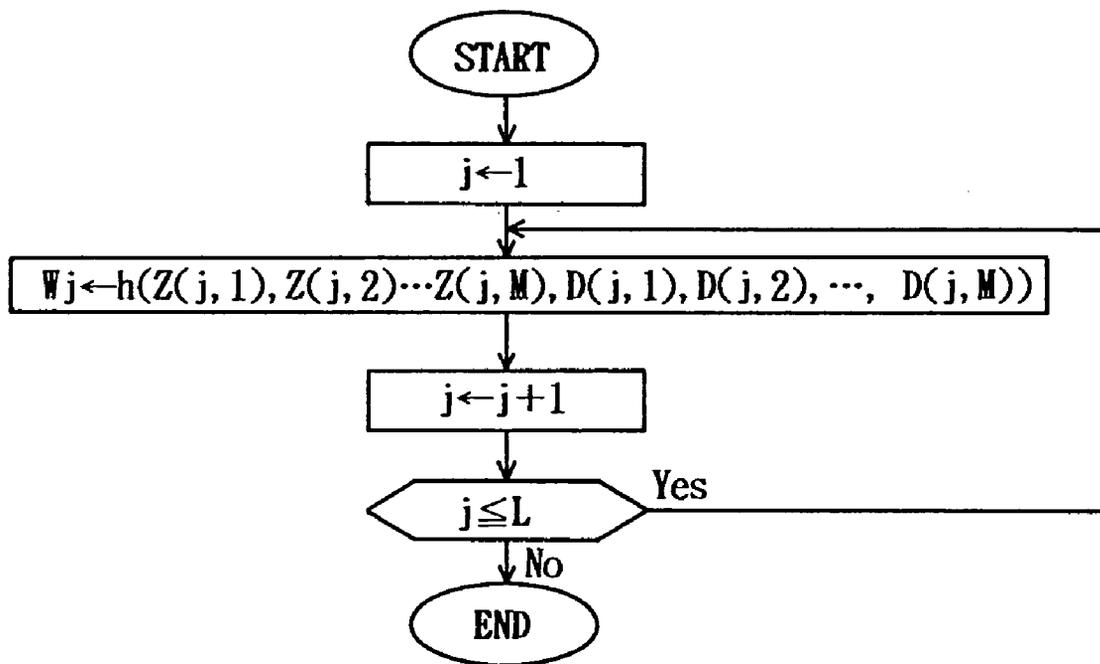


Fig. 11

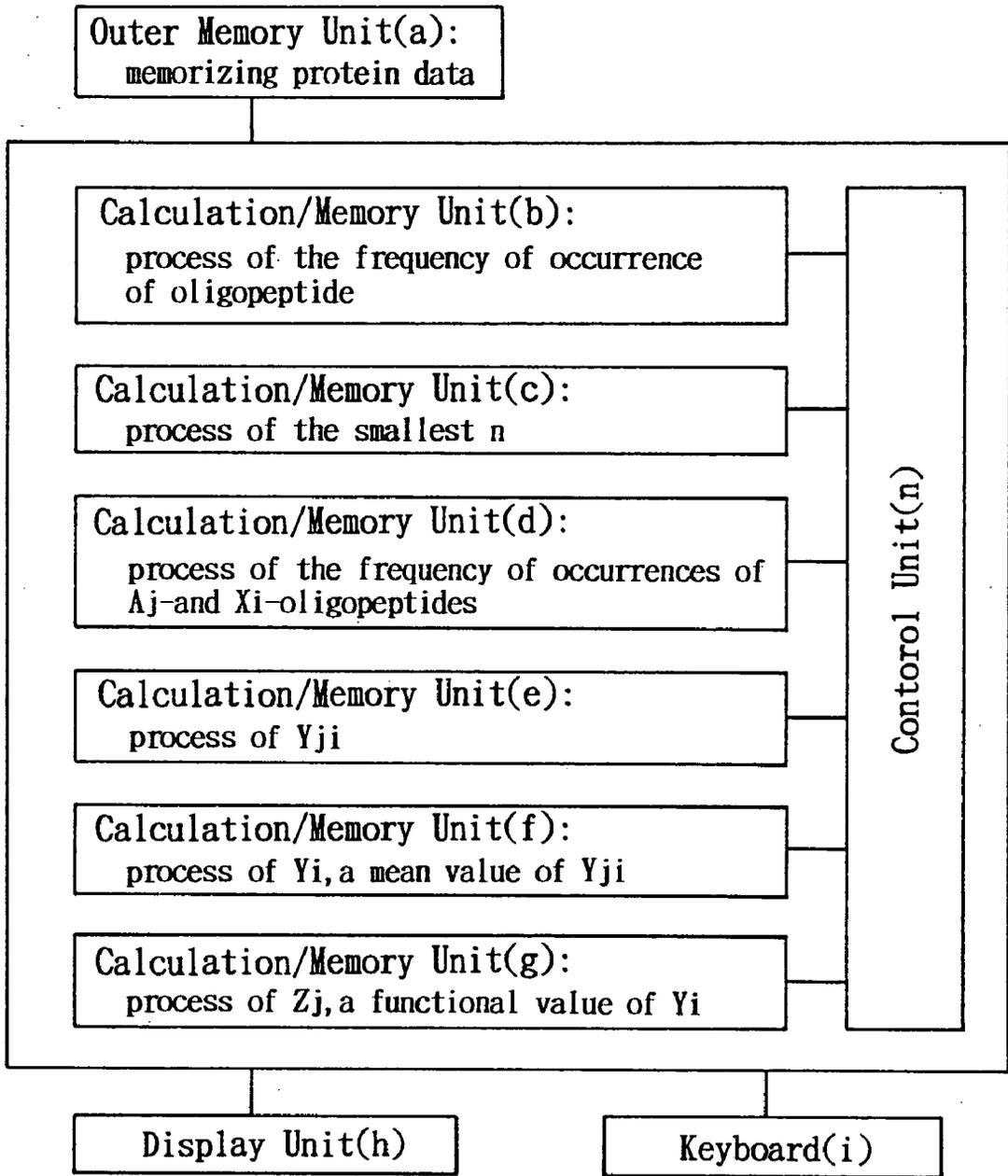


Fig. 12

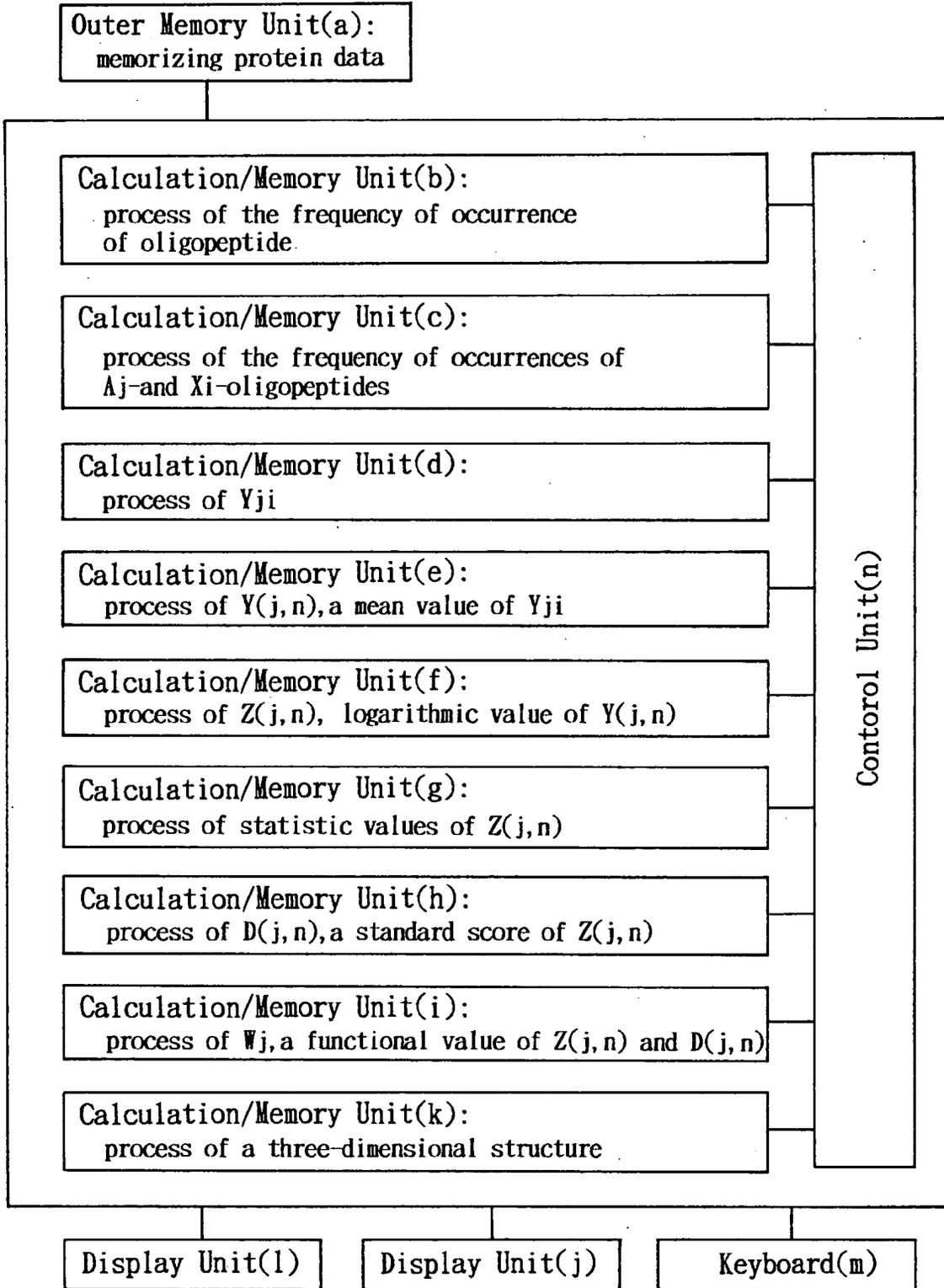


Fig. 13

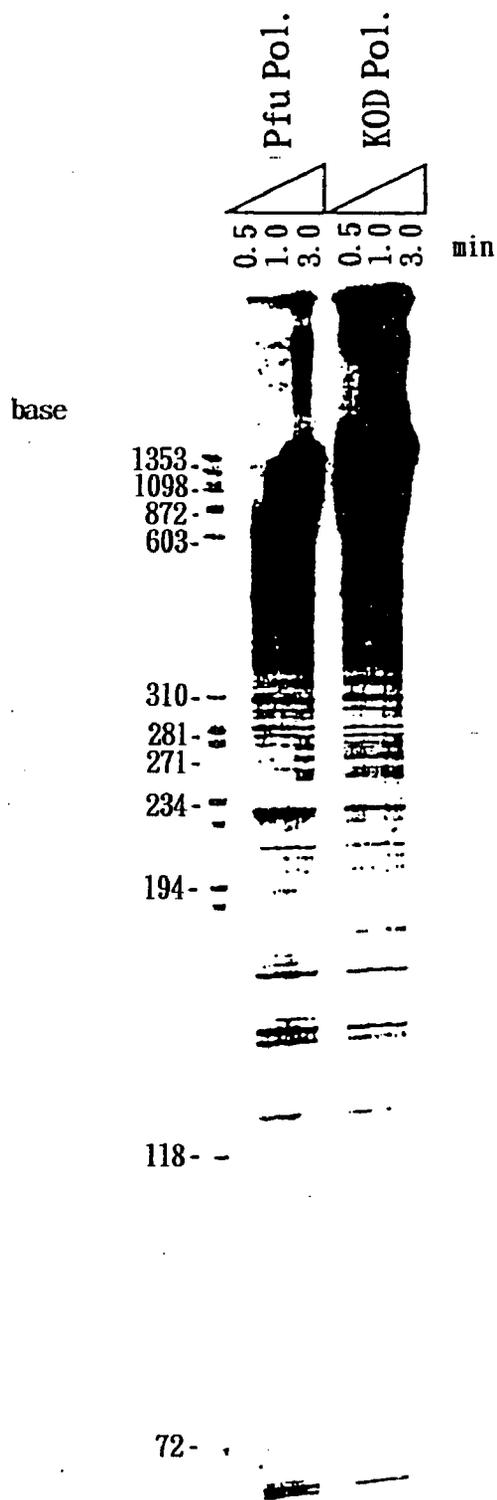


Fig. 14

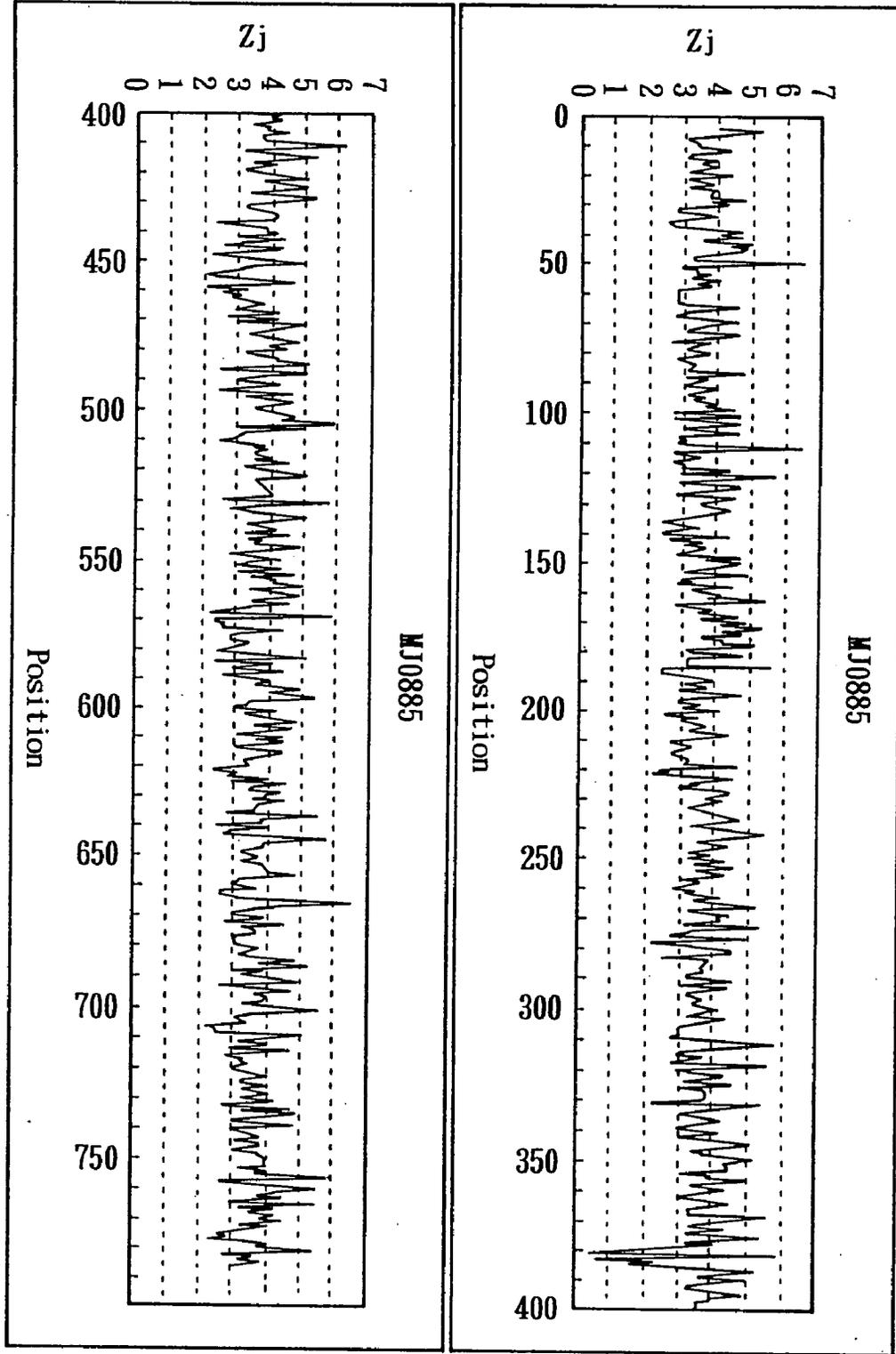
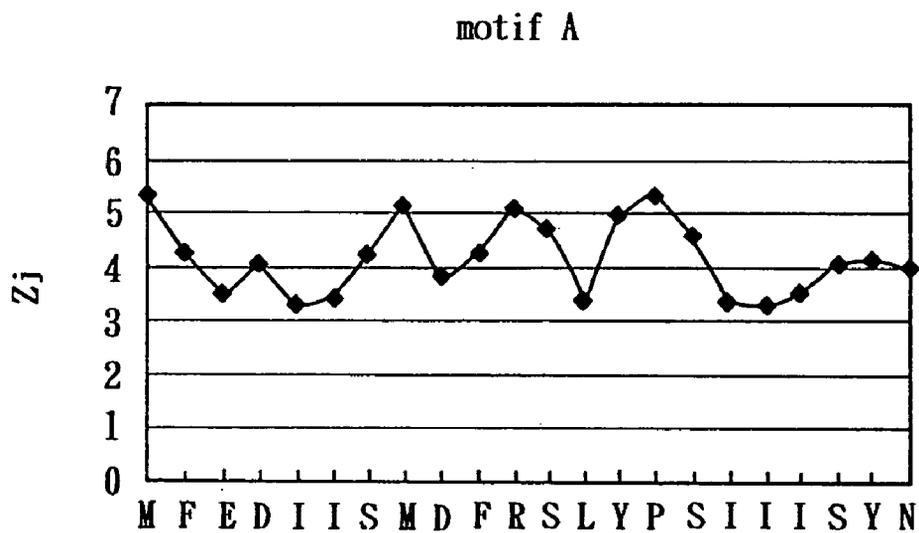
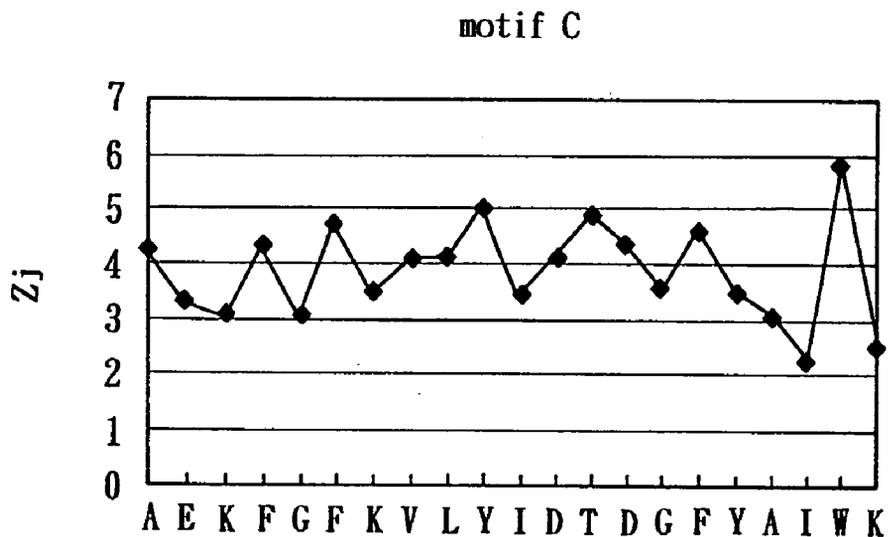


Fig. 15



MJ0558 Position 325..346



MJ0558 Position 550..570

Fig. 16

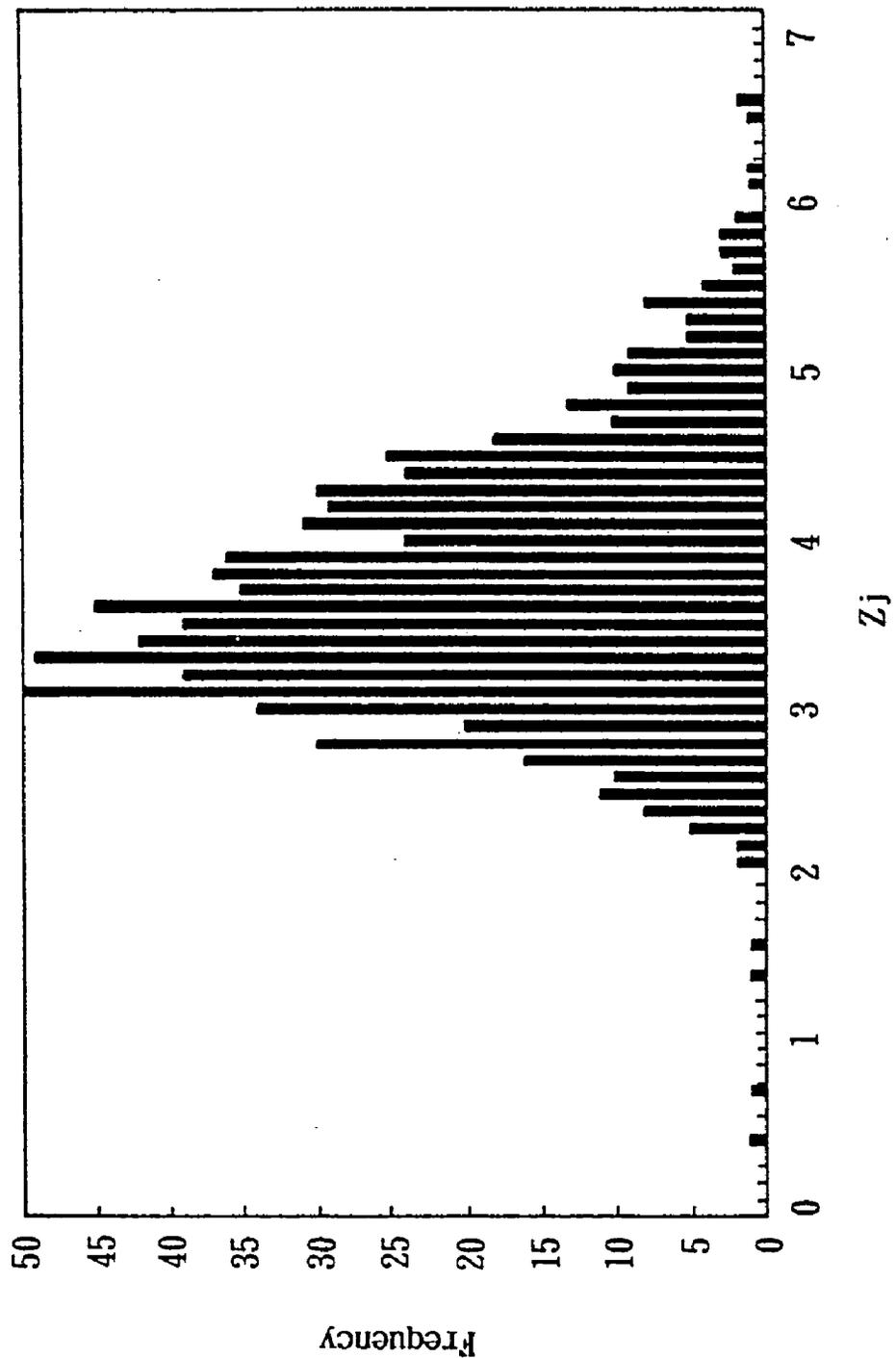


Fig. 17

motif C

	550		570
MJ	AEKFGFKVLYIDTDGFYAIWK		
KOD	EEKYGFKVIYSDDGFFATIP		
Pfu	EEKFGFKVLYIDTDGLYATIP		

Fig. 18

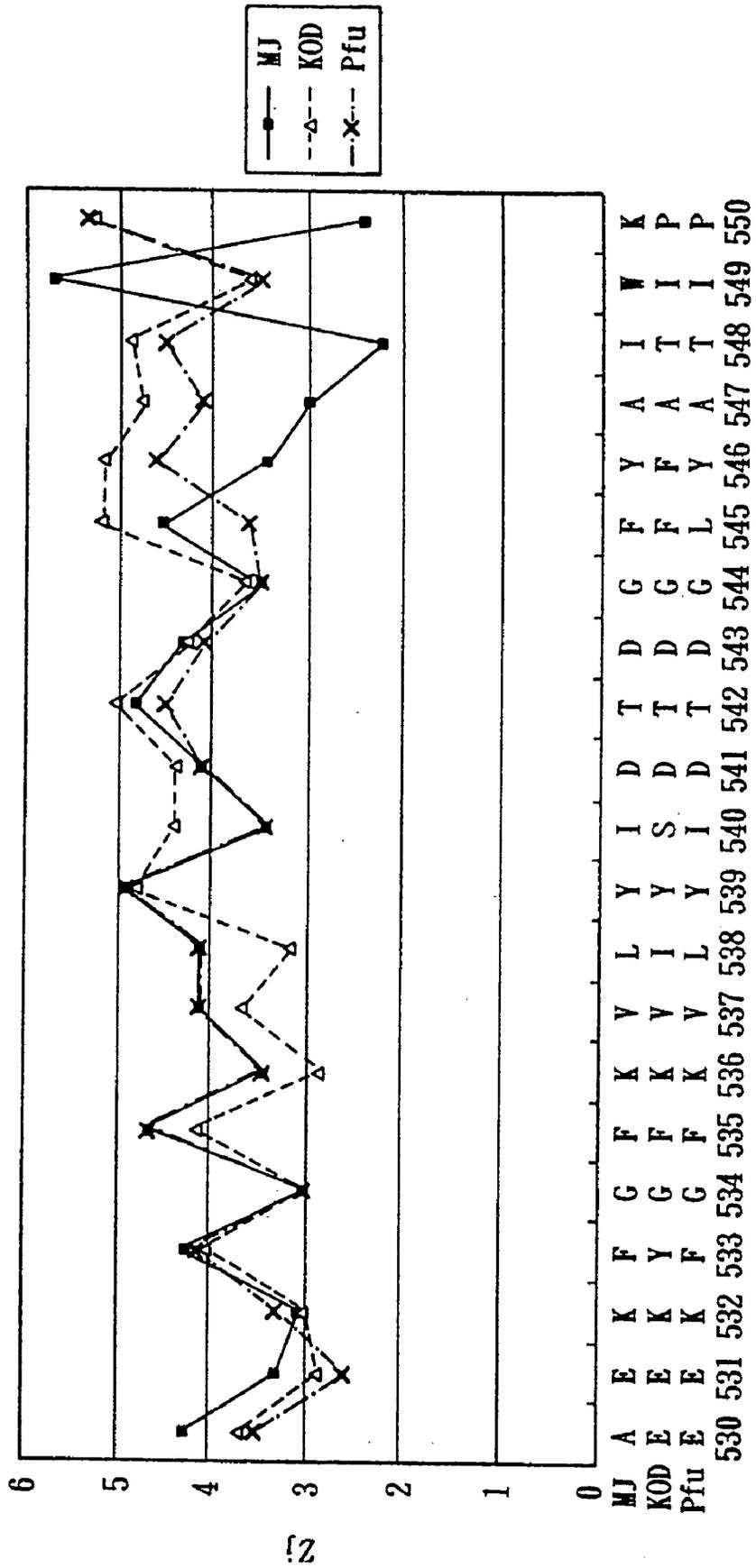


Fig. 19

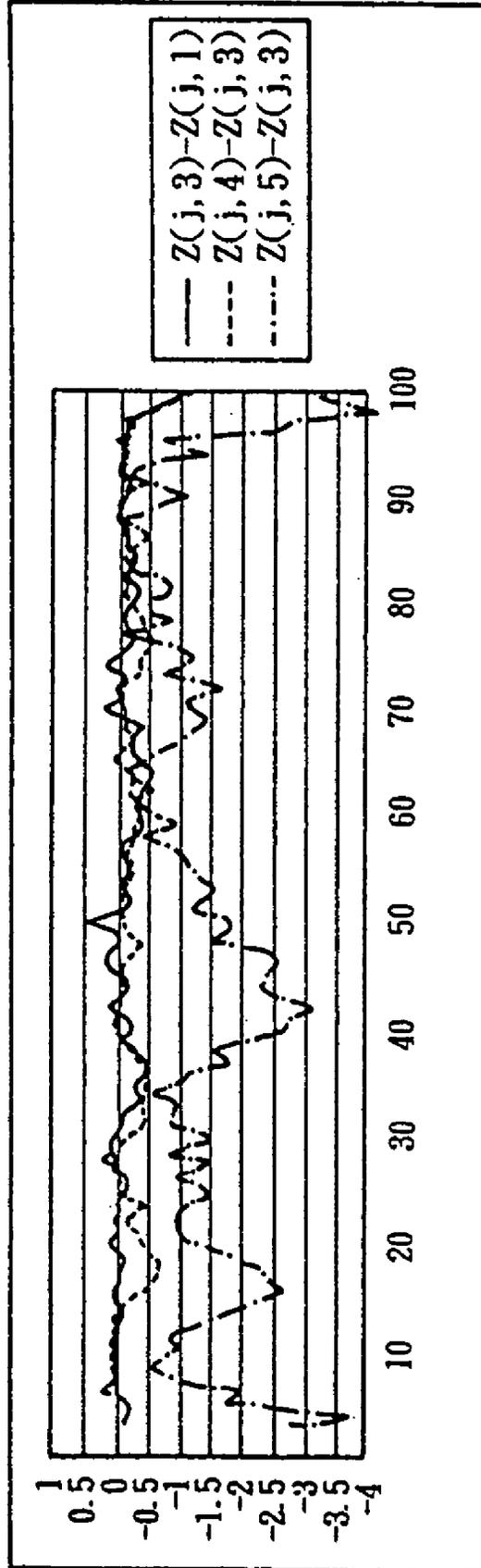


Fig. 20

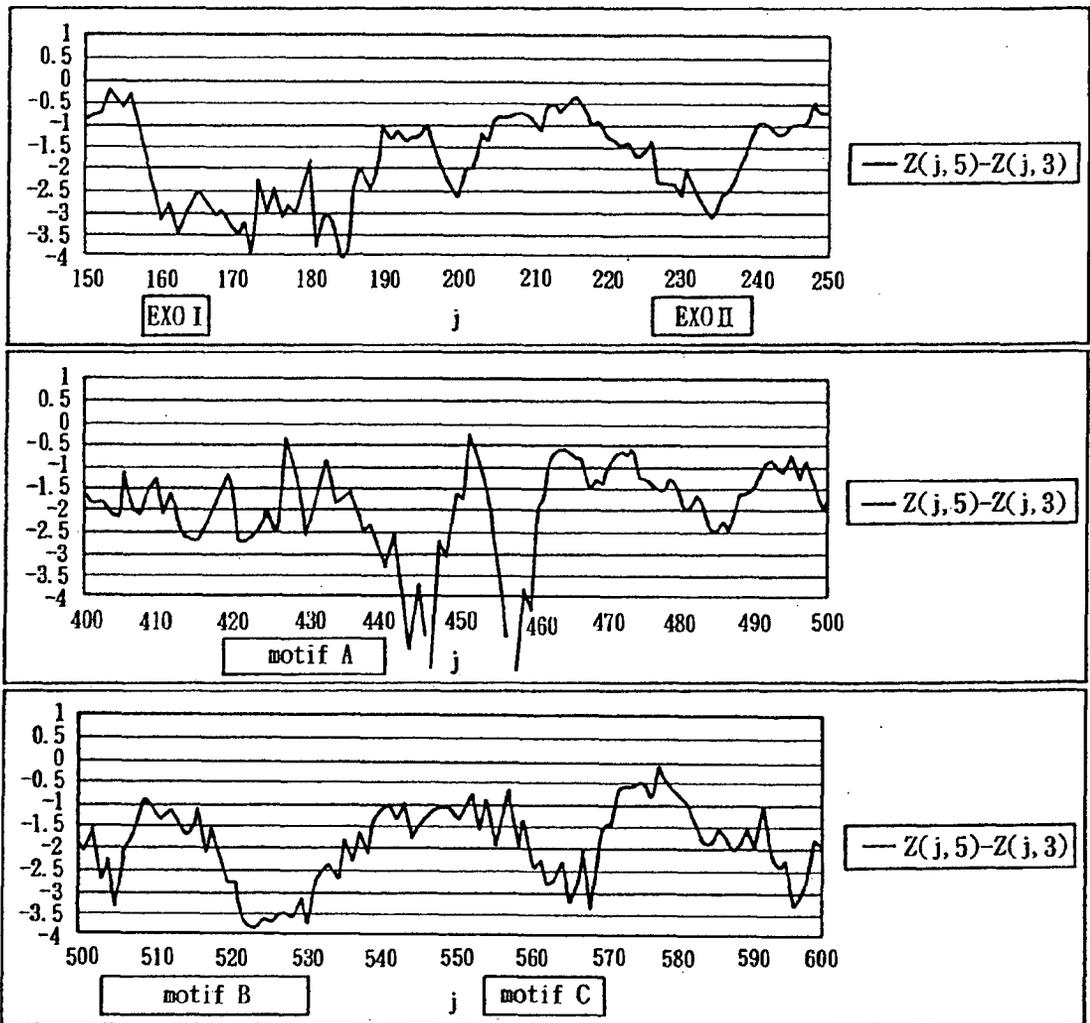


Fig. 21

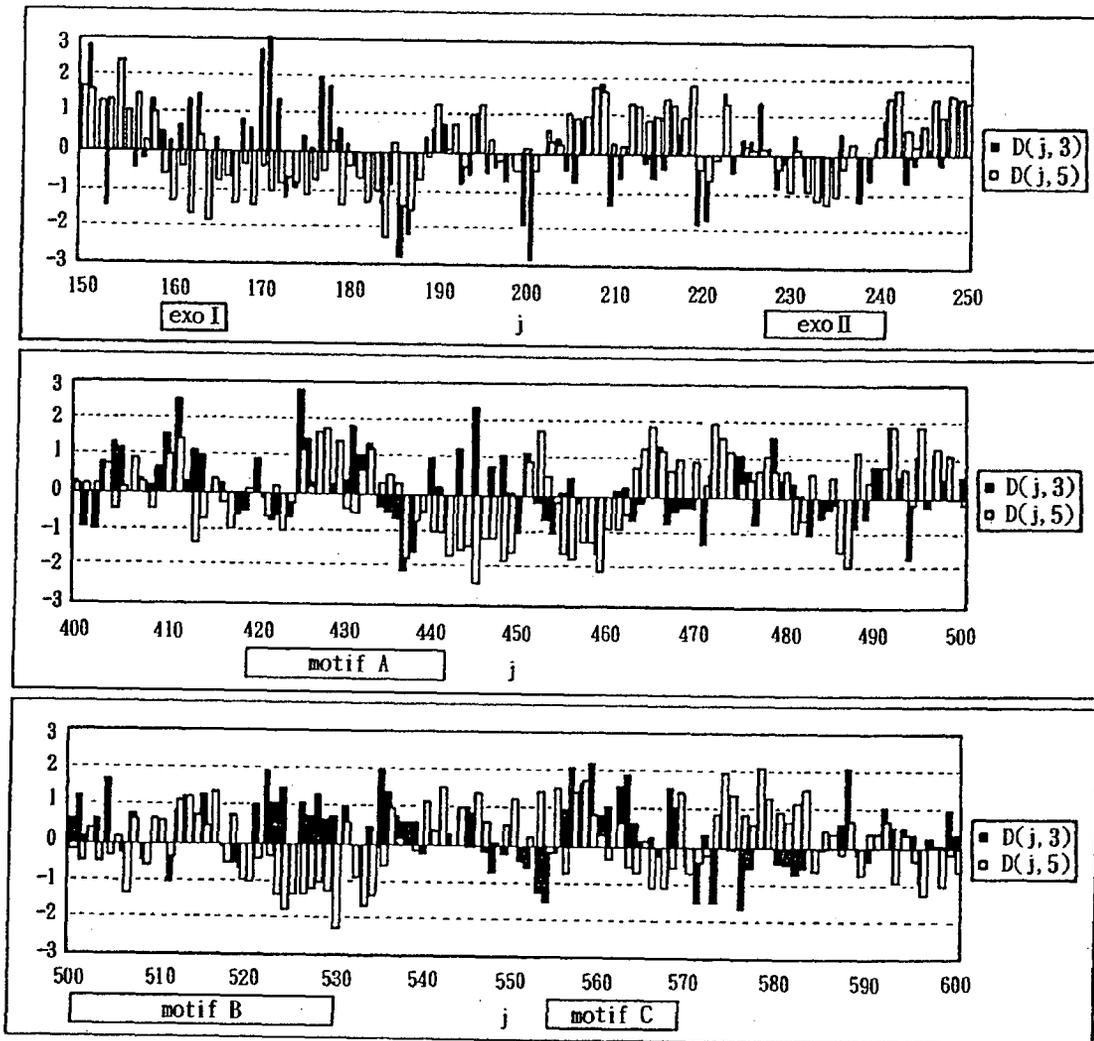


Fig. 22

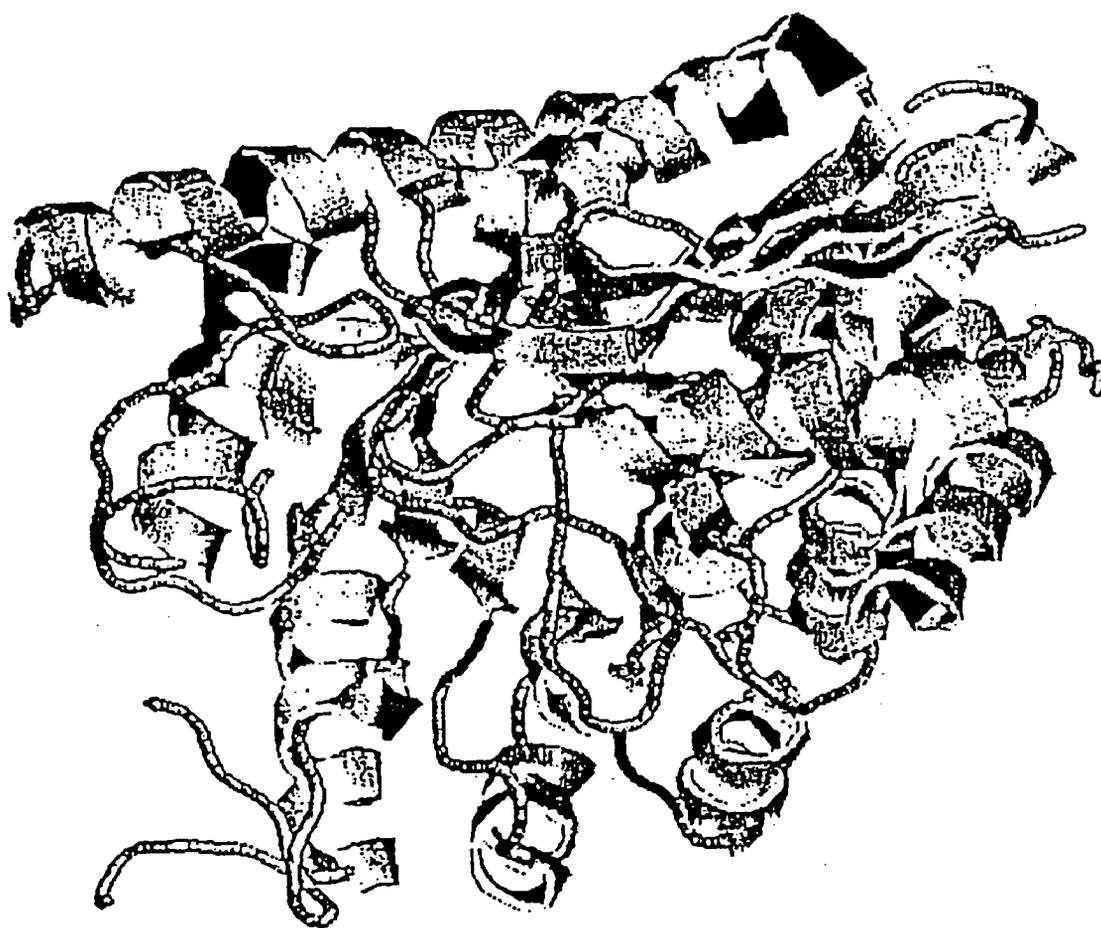


Fig. 23

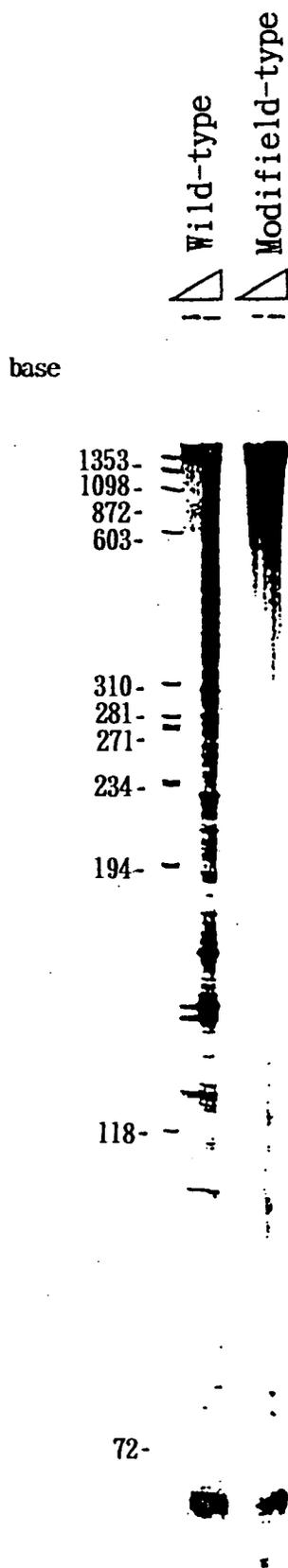
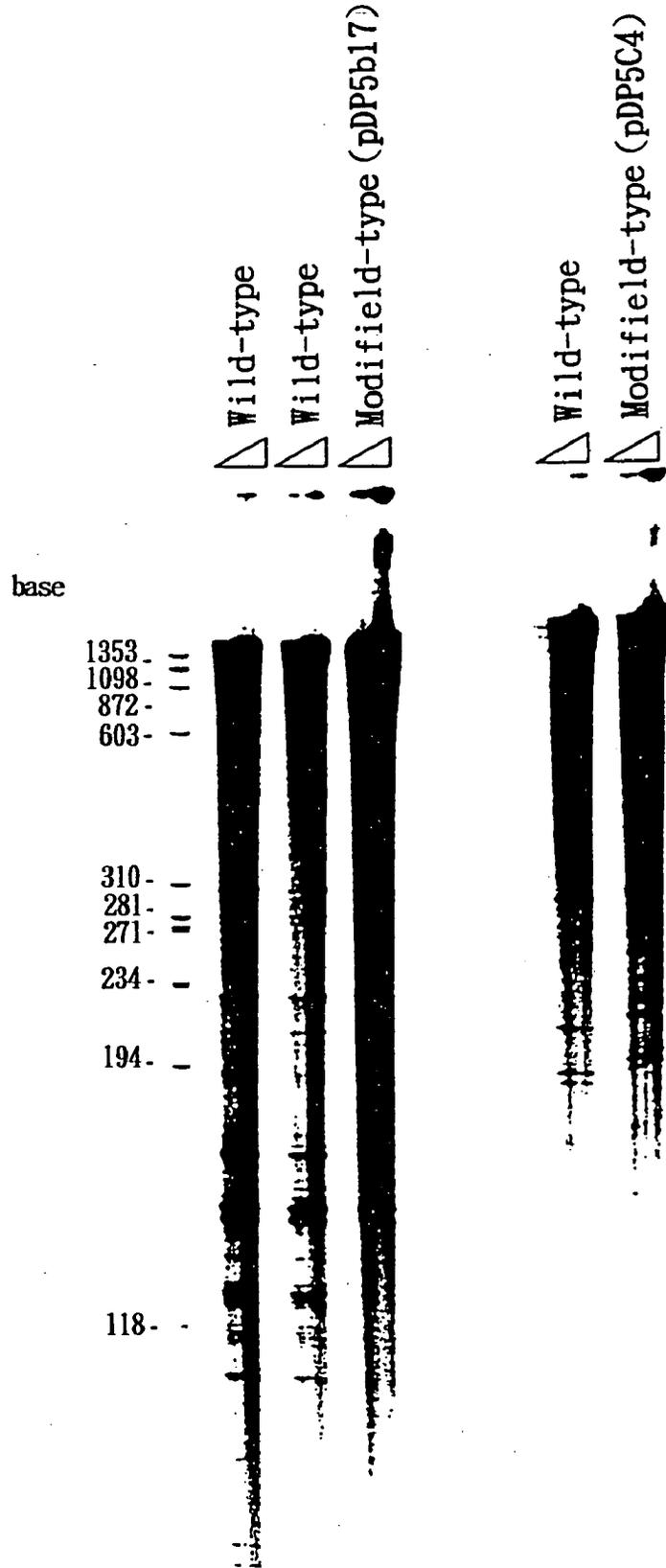


Fig. 24



METHOD AND A SYSTEM FOR PREDICTING PROTEIN FUNCTIONAL SITE, A METHOD FOR IMPROVING PROTEIN FUNCTION, AND A FUNCTION-MODIFIED PROTEIN

TECHNICAL FIELD

[0001] The present invention relates to a method for predicting a functional site of a protein, a system for predicting the function thereof, and a method for modifying the function of a protein and a function-modified protein. More specifically, the present invention relates to prediction of a functional site of a functionally unknown protein prepared by genome analysis or cDNA analysis, prediction of a novel function and a novel functional site of a protein with a known function, and prediction of a site on a protein to be modified for improving the function of the protein, and a protein with a modified function based on the prediction.

BACKGROUND ART

[0002] Following the progress of genome analysis and cDNA analysis of various organisms including pathogenic microorganisms, the number of novel genes whose functions are unknown is rapidly increasing, together with the number of proteins encoded by the genes. So far, the analysis of the nucleotide sequence of the whole genome of a microorganism, for example *Mycoplasma genitalium* (Fraser et al., Science 270, 397-403, 1995), *Haemophilus influenzae* (Fleischman et al., Science 269, 496-512, 1995), and *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996), has been completed, so that numerous novel proteins predicted from the genome sequence have been discovered. For humans and mice, the cDNA analysis is under way in combination with the genome analysis, which brings about the discovery of a great number of novel proteins.

[0003] In such circumstance, the prediction of the function of a functionally unknown protein or a functional site thereof has been a significant issue. If not only a novel protein but also a novel function or a novel functional site of a protein with a known function is discovered, whether or not these proteins are worth industrial or clinical application is possibly determined. Furthermore, such prediction of function possibly enables to prepare a modified protein with a further improved function.

[0004] Whether or not a protein encoded by a gene elucidated by genome analysis or cDNA analysis is novel or has a known function has been determined conventionally by searching the homology through protein databases such as Swiss-Prot. So as to predict a functional site, additionally, functionally identical proteins derived from various organisms are extracted from a protein database and are then subjected to alignment, to identify a region conserved in common to them and predict the conserved region as a functional site.

[0005] However, disadvantageously, such alignment method cannot be used if a protein obtained by genome analysis or cDNA analysis is an absolutely novel protein. Even if the protein has homology with known proteins in a protein database, the conserved region occupies most of the amino acid sequence of the protein in case that the protein is homologous to proteins derived from closely related organisms, so that it is impossible to predict the functional site. As to modification of protein, generally, the function of a protein is potentially deteriorated irrespective of the fact that the function is known or unknown once the conserved region is modified,

even if the functional site is predicted by alignment. Accordingly, the amino-acid residues outside the conserved region should be modified to improve the function. In other words, it is required to find a novel functional site in such protein to be modified. Using the conventional alignment method, disadvantageously, a novel functional site cannot be discovered or which amino-acid residue should be modified cannot be predicted.

[0006] Taking account of such circumstance, the present invention has been carried out. It is an object of the present invention to provide a novel method for predicting a functional site of a functionally unknown protein obtained by genome analysis or cDNA analysis.

[0007] In accordance with the present invention, furthermore, it is an object to provide a system for predicting the function.

[0008] In accordance with the present invention, still furthermore, it is an object to provide a method for predicting a novel functional site of a protein with an unknown function or with a known function and subjecting the functional site to mutation to prepare a modified protein.

[0009] Still furthermore, it is an object of the present invention to provide a protein with a function modified by the method described above.

DISCLOSURE OF INVENTION

[0010] A first aspect of the present invention proposed by the present application is a method for predicting a functional site of a protein derived from an organism "a" whose genome data, therefore the entire putative proteins, or cDNA data is known, which method comprises the steps of:

[0011] (1) determining in the amino acid sequences of the entire proteins of the organism "a", the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, and determining the smallest length (n) of oligopeptides satisfying the following criteria;

[0012] among oligopeptides of length (n), the number of oligopeptides which occur once in the entire proteins is smaller than the number of oligopeptides which occur twice in the entire proteins; among oligopeptides of length (n+1), the number of oligopeptides which occur once in the entire proteins is larger than the number of oligopeptides which occur twice in the entire proteins,

[0013] (2) determining in the entire proteins of the organism "a", the frequency of occurrence of the following Aji-oligopeptide of length (n+1), which is a part of the amino acid sequence of the protein as a subject for predicting a functional site and contains the j-th amino-acid residue Aj from the amino terminal end (N-terminus) of the amino acid sequence (length of L) of the protein, on condition that the j-th amino-acid residue Aj is defined as the i-th residue Aji from the N-terminus of the Aji-oligopeptide;

[0014] Aji-oligopeptide: aj1aj2 . . . Aji . . . ajnaj(n+1)

[0015] (wherein, $1 \leq i \leq n+1$; $n+1 \leq j \leq L-n$; $A_j = A_{ji}$; and Aji is the i-th residue of the oligopeptide; and moreover, $aj1 = A_{j-i+1}, \dots, aj(n+1) = A_{j-i+(n+1)}$),

and determining in the entire proteins of the organism "a", the frequency of occurrence of the following Xji-oligopeptide of length (n+1);

[0016] Xji-oligopeptide: aj1aj2 . . . Xji . . . ajnaj(n+1)

[0017] (wherein, $1 \leq i \leq n+1$; $n+1 \leq j \leq L-n$; and the i-th amino-acid residue Xji is any amino acid; and moreover, $aj1 = A_{j-i+1}, \dots, aj(n+1) = A_{j-i+(n+1)}$),

[0018] (3) calculating ratio Yji of the frequency of occurrence of the Aji-oligopeptide to that of the Xji-oligopeptide,

[0019] (4) determining mean Y_j of the Y_{ji} ;

$$Y_j = \sum_{i=1}^{n+1} Y_{ji}/(n+1),$$

[0020] (5) determining functional value Z_j of Y_j ;

$$Z_j = f(Y_j)$$

[0021] (Function f is a monotonously decreasing function or a monotonously increasing function), and defining the Z_j value as the representative value of the function of the j -th amino-acid residue A_j of the amino acid sequence (length of L), and

[0022] (6) repeating the steps (2) to (5) sequentially and determining the Z_j value of each A_j of all the amino-acid residues at positions between $n+1 \leq j \leq L-n$ in the amino acid sequence (length of L) of the protein, thereby predicting the degree of the involvement of each amino-acid residue in the function of the protein by using the dimension of the Z_j value as an indicator.

[0023] A second aspect of the present invention is a method for predicting a functional site of a protein derived from the entire putative proteins of an organism "a" of which genome data or cDNA data is known, which method comprises the steps of:

[0024] (1) determining the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, in the amino acid sequences of the entire proteins of the organism "a",

[0025] (2) as to an appropriate protein of the organism "a",

[0026] (2') determining in the entire proteins of the organism "a", the frequency of occurrence of the following A_{ji} -oligopeptide of given length of (n) ($1 \leq n \leq M$), provided that M is the smallest length of oligopeptides satisfying the criterion that all the oligopeptides of length M are at frequency 1 of the occurrence), which the A_{ji} -oligopeptide is a part of the amino acid sequence of the protein and contains the j -th amino-acid residue A_j , on condition that the j -th amino-acid residue A_j is defined as the i -th residue A_{ji} from the N-terminus of the A_{ji} -oligopeptide ($n \leq j \leq L-n+1$);

[0027] A_{ji} -oligopeptide: $a_{j1}a_{j2} \dots a_{ji} \dots a_{jn}$

[0028] (wherein, $1 \leq i \leq n$; $A_j = A_{ji}$, and A_{ji} is the i -th residue of the oligopeptide; and moreover, $a_{j1} = A_{j-i+1}, \dots, a_{jn} = A_{j-i+n}$),

and determining in the entire proteins of the organism "a", the frequency of occurrence of the following X_{ji} -oligopeptide of length (n) corresponding to the length of A_{ji} -oligopeptide;

[0029] X_{ji} -oligopeptide: $a_{j1}a_{j2} \dots X_{ji} \dots a_{jn}$

[0030] (wherein, $1 \leq i \leq n$; and the i -th residue X_{ji} is any amino acid; and moreover, $a_{j1} = A_{j-i+1}, \dots, a_{jn} = A_{j-i+n}$),

[0031] (3) calculating ratio Y_{ji} of the frequency of occurrence of the A_{ji} -oligopeptide to that of the X_{ji} -oligopeptide,

[0032] (4) determining mean $Y(j,n)$ of the Y_{ji} ;

$$Y(j,n) = \sum_{i=1}^n Y_{ji}/n,$$

[0033] (5) determining functional value $Z(j,n)$ of $Y(j,n)$;

$$Z(j,n) = -\log(Y(j,n)),$$

[0034] (6) repeating the steps (2') to (5) sequentially and determining the $Z(j,n)$ value of each amino-acid residue A_j at position j ($n \leq j \leq L-n+1$) in the amino acid sequence (length of L),

[0035] (7) sequentially repeating the steps (2) to (6) for the entire proteins of the organism "a", thereby determining the distribution of the $Z(j,n)$ value of each amino-acid residue in the entire proteins, and the $Z(j,n)$ values are classified into twenty according to the twenty amino acids, and then determining mean $Av(Aa)$ of the $Z(j,n)$ values for each amino acid Aa and the standard deviation $Sd(Aa)$ of the distribution thereof, on the basis of the distribution, to determine function g to the j -th amino-acid residue A_j of a protein for normalizing the difference in distribution due to the species of amino-acid residues;

$$g = g(Z(j,n), A_j) = [Z(j,n) - Av(Aa)] / Sd(Aa)$$

[0036] (provided that $A_j = Aa$; and $1 \leq n \leq M$),

[0037] (8) determining value $D(j,n)$ of the function g of each A_j of all the amino-acid residues at position j ($n \leq j \leq L-n+1$ and $1 \leq n \leq M$) of a protein in the entire proteins as recovered in the step (7);

$$D(j,n) = g(Z(j,n), A_j),$$

and

[0038] (9) defining the representative value of the function of the j -th amino-acid residue in the amino acid sequence (length of L) as functional value W_j of the $Z(j,n)$ and $D(j,n)$;

$$W_j = h(Z(j,1), Z(j,2), \dots, Z(j,M), D(j,1), D(j,2), \dots, D(j,M))$$

thereby predicting the degree of the involvement of each amino-acid residue in the function of the protein by using the dimension of the W_j value as an indicator.

[0039] A third aspect of the present invention is a system for automatically conducting the method according to claim 1, at least comprising the following units (a) to (g);

[0040] (a) an outer memory unit memorizing the amino acid sequence data of the entire putative proteins derived from organism "a" of which genome data or cDNA data is known, as well as an existing protein data base,

[0041] (b) a calculation/memory unit, composed of CPU calculating the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, in the amino acid sequences of the entire proteins from the organism "a", and a memory unit having the memory of the calculation results,

[0042] (c) a calculation/memory unit, composed of CPU calculating the smallest length (n) of oligopeptide satisfying the following criteria among the individual oligopeptides of which the frequencies of the occurrences being memorized in the unit (b);

[0043] among oligopeptides of length (n), the number of oligopeptides which occur once in the entire proteins is smaller than the number of oligopeptides which occur twice in the entire proteins; among oligopeptides of length ($n+1$), the number of oligopeptides which occur once in the entire proteins is larger than the number of oligopeptides which occur twice in the entire proteins, and a memory unit having the memory of the length (n),

[0044] (d) a calculation/memory unit, composed of CPU determining in the entire proteins of the organism "a", the frequency of occurrence of the following A_{ji} -oligopeptide of length ($n+1$), which is a part of the amino acid sequence

of the protein as a subject for predicting a functional site and contains the *j*-th amino-acid residue *A_j* from the amino terminal end (N-terminus) of the amino acid sequence (length of *L*) of the protein, on condition that the *j*-th amino-acid residue *A_j* is defined as the *i*-th residue *A_{ji}* from the N-terminus of the *A_{ji}*-oligopeptide;

[0045] *A_{ji}*-oligopeptide: *aj1aj2 . . . A_{ji} . . . ajn_{aj(n+1)}*
 [0046] (wherein, $1 \leq i \leq n+1$; $n+1 \leq j \leq L-n$; *A_j*=*A_{ji}*; and *A_{ji}* is the *i*-th residue of the oligopeptide; and moreover, $aj1=A_{j-i+1}, \dots, aj(n+1)=A_{j-i+(n+1)}$),

and determining in the entire proteins of the organism “a”, the frequency of occurrence of the following *X_{ji}*-oligopeptide of length (*n*+1);

[0047] *X_{ji}*-oligopeptide: *aj1aj2 . . . X_{ji} . . . ajn_{aj(n+1)}*
 (wherein, $1 \leq i \leq n+1$; $n+1 \leq j \leq L-n$; and the *i*-th amino-acid residue *X_{ji}* is any amino acid; and moreover, $aj1=A_{j-i+1}, \dots, aj(n+1)=A_{j-i+(n+1)}$),

and a memory unit having the memory of the calculation results,

[0048] (e) a calculation/memory unit, composed of CPU determining ratio *Y_{ji}* of the frequency of occurrence of the *A_{ji}*-oligopeptide to that of the *X_{ji}*-oligopeptide, and a memory unit having the memory of *Y_{ji}*,

[0049] (f) a calculation/memory unit, composed of CPU determining mean *Y_j* of the *Y_{ji}*;

$$Y_j = \sum_{i=1}^{n+1} Y_{ji}/(n+1),$$

and a memory unit having the memory of *Y_j*, and

[0050] (g) a calculation/memory unit, composed of CPU determining functional value *Z_j* of *Y_j*;

$$Z_j = f(Y_j)$$

[0051] (Function *f* is a monotonously decreasing function or a monotonously increasing function),

and a memory unit having the memory of *Z_j*.

[0052] A fourth aspect of the present invention is a system for automatically conducting the method according to claim 3, at least comprising the following units (a) to (i);

[0053] (a) an outer memory unit memorizing the amino acid sequence data of the entire putative proteins of the organism “a” of which genome data or cDNA data is known, as well as an existing protein data base,

[0054] (b) a calculation/memory unit, composed of CPU calculating in the amino acid sequences of the entire proteins of the organism “a”, the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, and a memory unit having the memory of the calculation results,

[0055] (c) a calculation/memory unit, composed of CPU determining in the entire proteins of the organism “a”, the frequency of occurrence of the following *A_{ji}*-oligopeptide of given length of (*n*) ($1 \leq n \leq M$, provided that *M* is the smallest length of oligopeptides satisfying the criterion that all the oligopeptides of length *M* are at frequency 1 of the occurrence), which the *A_{ji}*-oligopeptide is a part of the amino acid sequence of the protein and contains the *j*-th amino-acid residue *A_j*, on condition that the *j*-th amino-

acid residue *A_j* is defined as (or identical to?) the *i*-th residue *A_{ji}* from the N-terminus of the *A_{ji}*-oligopeptide ($n \leq j \leq L-n+1$);

[0056] *A_{ji}*-oligopeptide: *aj1aj2 . . . A_{ji} . . . ajn*
 [0057] (wherein, $1 \leq i \leq n$; *A_j*=*A_{ji}*, and *A_{ji}* is the *i*-th residue of the oligopeptide; and moreover, $aj1=A_{j-i+1}, \dots, ajn=A_{j-i+n}$),

and determining in the entire proteins of the organism “a”, the frequency of occurrence of the following *X_{ji}*-oligopeptide of length (*n*) corresponding to the length of *A_{ji}*-oligopeptide;

[0058] *X_{ji}*-oligopeptide: *aj1aj2 . . . X_{ji} . . . ajn*
 [0059] (wherein, $1 \leq i \leq n$; and the *i*-th residue *X_{ji}* is any amino acid; and moreover, $aj1=A_{j-i+1}, \dots, ajn=A_{j-i+n}$),

and a memory unit memorizing the calculation results,
 [0060] (d) a calculation/memory unit, composed of CPU calculating ratio *Y_{ji}* of the frequency of occurrence of the *A_{ji}*-oligopeptide to that of the *X_{ji}*-oligopeptide, and a memory unit having the memory of the *Y_{ji}*,

[0061] (e) a calculation/memory unit, composed of CPU determining mean *Y(j,n)* of the *Y_{ji}*;

$$Y(j, n) = \sum_{i=1}^n Y_{ji}/n,$$

and a memory unit having the memory of *Y(j,n)*,

[0062] (f) a calculation/memory unit, composed of CPU determining functional value *Z(j,n)* of *Y(j,n)*;

$$Z(j, n) = -\log(Y(j, n)),$$

and a memory unit having the memory of *Z(j,n)*,

[0063] (g) a calculation/memory unit, composed of CPU determining the distribution of the *Z(j,n)* value of each amino-acid residue in the entire proteins, and the *Z(j,n)* values are classified into twenty according to the twenty amino acids, and then determining mean *Av(Aa)* of the *Z(j,n)* values for each amino acid *Aa* and the standard deviation *Sd(Aa)* of the distribution thereof, on the basis of the distribution, to determine function *g* to the *j*-th amino-acid residue *A_j* of a protein for normalizing the difference in distribution due to the species of amino-acid residues;

$$g = g(Z(j, n), A_j) = [Z(j, n) - Av(Aa)] / Sd(Aa)$$

[0064] (wherein, *A_j*=*Aa*; and $1 \leq n \leq M$)

and a memory unit having the memory of *g*,
 [0065] (h) a calculation/memory unit, composed of CPU determining value *D(j,n)* of function *g* memorized in the unit (g) concerning each of all the amino-acid residues *A_j* at position *j* ($n \leq j \leq L-n+1$) in the amino acid sequence (length of *L*);

$$D(j, n) = g(Z(j, n), A_j)$$

and a memory unit having the memory of the *D(j, n)* value, and

[0066] (i) a calculation/memory unit, composed of a calculation unit determining appropriate functional value *W_j* of the *Z(j,n)* and *D(j,n)* of each amino-acid residue in the amino acid sequence;

$$W_j = h(Z(j, 1), Z(j, 2), \dots, Z(j, M), D(j, 1), D(j, 2), \dots, D(j, M))$$

and a memory unit having the memory of the *W_j* value.

[0067] A fifth aspect of the present invention is a method for modifying the known function of protein "A" derived from the entire proteins of organism "a" of which genome data or cDNA data has been known, which method comprises the steps of:

[0068] (1) extracting a protein closely related to the protein "A" from an existing protein data base and subjecting the proteins to alignment,

[0069] (2) determining in the amino acid sequences of the entire proteins of the organism "a", the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, and determining the smallest length (n) of oligopeptides satisfying the following criteria;

[0070] among oligopeptides of length (n), the number of oligopeptides which occur once in the entire proteins is smaller than the number of oligopeptides which occur twice in the entire proteins; among oligopeptides of length (n+1), the number of oligopeptides which occur once in the entire proteins is larger than the number of oligopeptides which occur twice in the entire proteins,

[0071] (3) determining in the entire proteins of the organism "a", the frequency of occurrence of the following Aji-oligopeptide of length (n+1), which is a part of the amino acid sequence of the protein as a subject for predicting a functional site and contains the j-th amino-acid residue Aj from the amino terminal end (N-terminus) of the amino acid sequence (length of L) of the protein, on condition that the j-th amino-acid residue Aj is defined as the i-th residue Aji from the N-terminus of the Aji-oligopeptide;

[0072] Aji-oligopeptide: aj1aj2 . . . Aji . . . ajnaj(n+1)

[0073] (wherein, $1 \leq n+1$; $n+1 \leq j \leq L-n$; $A_j = A_{ji}$; and Aji is the i-th residue of the oligopeptide; and moreover, $aj1 = A_{j-i+1}$, . . . , $aj(n+1) = A_{j-i+(n+1)}$),

and determining in the entire proteins of the organism "a", the frequency of occurrence of the following Xji-oligopeptide of length (n+1);

[0074] Xji-oligopeptide: aj1aj2 . . . Xji . . . ajnaj(n+1)

(wherein, $1 \leq n+1$; $n+1 \leq j \leq L-n$; and the i-th amino-acid residue Xji is any amino acid, and moreover, $aj1 = A_{j-i+1}$, . . . , $aj(n+1) = A_{j-i+(n+1)}$),

[0075] (4) calculating ratio Yji of the frequency of occurrence of the Aji-oligopeptide to that of the Xji-oligopeptide,

[0076] (5) determining mean Yj of the Yji;

$$Y_j = \sum_{i=1}^{n+1} Y_{ji}(n+1),$$

[0077] (6) determining functional value Zj of Yj;

$$Z_j = f(Y_j)$$

[0078] (Function f is a monotonously decreasing function or a monotonously increasing function),

and defining the Zj value as the representative value of the function of the j-th amino-acid residue Aj of the amino acid sequence (length of L) of the protein "A",

[0079] (7) sequentially repeating the steps (3) to (6) and determining the Zj value of each of all the amino-acid residues at positions between $n+1 \leq j \leq L-n$ in the amino acid sequence (length of L) of the protein "A",

[0080] (8) selecting at least one amino-acid residue to be subjected to mutation from the amino acid sequence (length of L) of the protein "A" on the basis of the alignment data carried out in the step (1), sequentially repeating the steps (3) to (6) for variant amino-acid residues in various mutated (or mutant?) amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residue, to determine the Zj value of the variant amino-acid residues,

[0081] (9) selecting a mutated amino acid sequence wherein the Zj value of the variant amino-acid residue as determined in the step (8) is larger or smaller than the Zj value of the wild type amino-acid residue as determined in the step (7), and

[0082] (10) preparing a modified gene encoding the modified amino acid sequence from the protein "A" gene, and producing the modified protein as the expression product of the gene.

[0083] A sixth aspect of the present invention is a method for modifying the function of protein "B" derived from an organism "b" of which genome data or cDNA data has been unknown, which method comprises the steps of:

[0084] (1) extracting protein "A" most closely related to protein "B" from the entire proteins of organism "a" of which genome data or cDNA data being known and subjecting the protein to alignment, or extracting a protein closely related to protein "B" from an existing protein data base to subject the protein to alignment,

[0085] (2) determining in the amino acid sequences of the entire proteins of the organism "a", the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, and determining the smallest length (n) of oligopeptides satisfying the following criteria;

[0086] among oligopeptides of length (n), the number of oligopeptides which occur once in the entire proteins is smaller than the number of oligopeptides which occur twice in the entire proteins; among oligopeptides of length (n+1), the number of oligopeptides which occur once in the entire proteins is larger than the number of oligopeptides which occur twice in the entire proteins,

[0087] (3) determining in the entire proteins of the organism "a", the frequency of occurrence of the following Aji-oligopeptide of length (n+1), which is a part of the amino acid sequence of the protein as a subject for predicting a functional site and contains the j-th amino-acid residue Aj from the amino terminal end (N-terminus) of the amino acid sequence (length of L) of the protein, on condition that the j-th amino-acid residue Aj is defined as the i-th residue Aji from the N-terminus of the Aji-oligopeptide;

[0088] Aji-oligopeptide: aj1aj2 . . . Aji . . . ajnaj(n+1)

[0089] (wherein, $1 \leq i \leq n+1$; $n+1 \leq j \leq L-n$; $A_j = A_{ji}$; and Aji is the i-th residue of the oligopeptide; and moreover, $aj1 = A_{j-i+1}$, . . . , $aj(n+1) = A_{j-i+(n+1)}$),

and determining in the entire proteins of the organism "a", the frequency of occurrence of the following Xji-oligopeptide of length (n+1);

[0090] Xji-oligopeptide: aj1aj2 . . . Xji . . . ajnaj(n+1)

[0091] (wherein, $1 \leq i \leq n+1$; $n+1 \leq j \leq L-n$; and the i-th amino-acid residue Xji is any amino acid; and moreover, $aj1 = A_{j-i+1}$, . . . , $aj(n+1) = A_{j-i+(n+1)}$),

[0092] (4) calculating ratio Yji of the frequency of occurrence of the Aji-oligopeptide to that of the Xji-oligopeptide,

[0093] (6) determining mean Y_j of the Y_{ji} ;

$$Y_j = \sum_{i=1}^{n+1} Y_{ji}/(n+1),$$

[0094] (6) determining functional value Z_j of Y_j ;

$$Z_j = f(Y_j)$$

[0095] (Function f is a monotonously decreasing function or a monotonously increasing function),

and defining the Z_j value as the representative value of the function of the j -th amino-acid residue A_j of the amino acid sequence (length of L) of the protein "A",

[0096] (7) sequentially repeating the steps (3) to (6) and determining the Z_j value of each of all the amino-acid residues at positions between $n+1 \leq j \leq L-n$ in the amino acid sequence (length of L),

[0097] (8) selecting at least one amino-acid residue to be subjected to mutation from the amino acid sequence (length of L) of the protein "A" on the basis of the alignment data carried out in the step (1), sequentially repeating the steps (3) to (6) for variant amino-acid residues in various mutated amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residues, to determine the Z_j value of the variant amino-acid residues,

[0098] (9) selecting the mutation position and the mutated amino-acid residue wherein the Z_j value of the variant amino-acid residue as determined in the step (8) is larger or smaller than the Z_j value of the wild type amino-acid residue as determined in the step (7), and

[0099] (10) preparing a modified gene encoding the modified amino acid sequence having the mutated amino-acid residue at the position from the protein "B" gene, and producing the modified protein as the expression product of the gene.

[0100] A seventh aspect of the present invention is a method for modifying the known function of protein "A" derived from the entire proteins of organism "a" of which genome data or cDNA data has been known, which method comprises the steps of:

[0101] (1) extracting proteins closely related to the protein "A" from an existing protein data base and subjecting the proteins to alignment,

[0102] (2) determining the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, in the amino acid sequences of the entire proteins of the organism "a",

[0103] (3) as to an appropriate protein of the organism "a",

[0104] (3') determining in the entire proteins of the organism "a", the frequency of occurrence of the following A_{ji} -oligopeptide of given length of (n) ($1 \leq n \leq M$, provided that M is the smallest length (n) of oligopeptides satisfying the criterion that all the oligopeptides of length M are at frequency 1 of the occurrence), which the A_{ji} -oligopeptide is a part of the amino acid sequence of the protein and contains the j -th amino-acid residue A_j , on condition that the j -th amino-acid residue A_j is defined as the i -th residue A_{ji} from the N-terminus of the A_{ji} -oligopeptide ($n \leq j \leq L-n+1$);

[0105] A_{ji} -oligopeptide: $a_j1a_j2 \dots a_j \dots a_{jn}$

[0106] (wherein, $1 \leq i \leq n$; $A_j = A_{ji}$, and A_{ji} is the i -th residue of the oligopeptide; and moreover, $a_j1 = A_{j-i+1}, \dots$, $a_{jn} = A_{j-i+n}$),

and determining in the entire proteins of the organism "a", the frequency of occurrence of the following X_{ji} -oligopeptide of length (n) corresponding to the length of A_{ji} -oligopeptide;

[0107] X_{ji} -oligopeptide: $a_j1a_j2 \dots X_{ji} \dots a_{jn}$

[0108] (wherein, $1 \leq i \leq n$; and the i -th residue X_{ji} is any amino acid; and moreover, $a_j1 = A_{j-i+1}, \dots, a_{jn} = A_{j-i+n}$),

[0109] (4) calculating ratio Y_{ji} of the frequency of occurrence of the A_{ji} -oligopeptide to that of the X_{ji} -oligopeptide,

[0110] (5) determining mean $Y(j,n)$ of the Y_{ji} ;

$$Y(j, n) = \sum_{i=1}^n Y_{ji}/n,$$

[0111] (6) determining functional value $Z(j,n)$ of $Y(j,n)$;

$$Z(j,n) = -\log(Y(j,n)),$$

[0112] (7) repeating the steps (3') to (6) sequentially and determining the $Z(j,n)$ value of each amino-acid residue A_j at position j ($n \leq j \leq L-n+1$) in the amino acid sequence (length of L),

[0113] (8) sequentially repeating the steps (3) to (7) for the entire proteins of the organism "a", thereby determining the distribution of the $Z(j,n)$ value of each amino-acid residue in the entire proteins, and the $Z(j,n)$ values are classified into twenty according to the twenty amino acids, and then determining mean $Av(A_a)$ of the $Z(j,n)$ values for each amino acid A_a and the standard deviation $Sd(A_a)$ of the distribution thereof, on the basis of the distribution, to determine function g to the j -th amino-acid residue A_j of a protein for normalizing the difference in distribution due to the species of amino-acid residues;

$$g = g(Z(j,n), A_j) = [Z(j,n) - Av(A_a)] / Sd(A_a)$$

[0114] (provided that $A_j = A_a$; and $1 \leq n \leq M$),

[0115] (9) determining value $D(j,n)$ of the function g of each A_j of all the amino-acid residues at position j ($n \leq j \leq L-n+1$ and $1 \leq n \leq M$) of a protein in the entire proteins as recovered in the step (8);

$$D(j,n) = g(Z(j,n), A_j),$$

and

[0116] (10) defining the representative value of the function of the j -th amino-acid residue in the amino acid sequence (length of L) as functional value W_j of the $Z(j,n)$ and $D(j,n)$;

$$W_j = h(Z(j,1), Z(j,2), \dots, Z(j,M), D(j,1), D(j,2), \dots, D(j,M))$$

[0117] (11) sequentially repeating the steps (3) to (10), to determine the individual W_j values of all the amino-acid residues at position $n \leq j \leq L-n+1$ in the amino acid sequence (length of L),

[0118] (12) selecting at least one amino-acid residue to be subjected to mutation on the basis of the alignment data carried out in the step (1) from the amino acid sequence (length of L) of the protein "A", and sequentially repeating the steps (3) to (10) for variant amino-acid residues in

various mutated amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residue, to determine the W_j value of the variant amino-acid residue,

[0119] (13) selecting a mutated amino acid sequence wherein the W_j value of the variant amino-acid residue as determined in the step (12) is larger or smaller than the W_j value of the wild type amino-acid residue as determined in the step (10), and

[0120] (14) preparing a modified gene encoding the modified amino acid sequence from the protein "A" gene, and producing the modified protein as the expression product of the gene.

[0121] An eighth aspect of the present invention is a protein of which function is artificially modified by the methods for modifying the function of protein set forth above.

[0122] One embodiment of the function-modified protein is a thermophilic DNA polymerase, prepared by artificially modifying the amino acid sequence of Pfu DNA polymerase so that the elongation of synthesized DNA chain might not be terminated intermediately during the catalysis for the synthesis of a DNA chain complimentary to a single-stranded DNA, and more specifically, the thermophilic DNA polymerase is one comprising the amino acid sequence of SQ ID No.1. In association with this thermophilic DNA polymerase, the present application provides a DNA sequence encoding the amino acid sequence of SQ ID No.1 and a recombinant vector carrying the DNA sequence. Such recombinant vector includes recombinant plasmid pDP320 carried on *Escherichia coli* HMS174 (DE3) /pDP320 (FERM P-16052). Still furthermore, in accordance with the present invention, it is provided a method for preparing this thermophilic DNA polymerase, comprising culturing a cell transformed with an expression vector carrying the DNA sequence and isolating and purifying the objective enzyme generated in a culture medium.

[0123] Another embodiment of the function-modified protein is a thermophilic DNA polymerase, prepared by artificially modifying the amino acid sequence of Pfu DNA polymerase so that the synthesized DNA chain might be more elongated during the catalysis for the synthesis of a DNA chain complimentary to a single-stranded DNA, and more specifically, the thermophilic DNA polymerase is one comprising the amino acid sequence of SQ ID No.6 or a DNA polymerase comprising the amino acid sequence of SQ ID No.7. In association with this thermophilic DNA polymerase, the present application provides a DNA sequence encoding the amino acid sequence of SQ ID No.6 or 7, and a recombinant vectors carrying such DNA sequences, respectively. As such vectors, there are provided recombinant plasmid pDP5b17 carried on *Escherichia coli* HMS174 (DE3)/pDP5b17 (FERM BP-6189) (vector carrying the DNA sequence encoding the amino acid sequence of SQ ID No.1), and recombinant plasmid pDP5C4 carried on *Escherichia coli* HMS174 (DE3)/pDP5C4 (FERM BP-6190) (vector carrying the DNA sequence encoding the amino acid sequence of SQ ID No.1). Still furthermore, it is provided a method for producing the DNA polymerase, comprising culturing a cell transformed with an expression vector carrying the DNA sequence and isolating and purifying the objective enzyme produced in a culture medium.

[0124] The method for predicting a protein functional site in accordance with the first aspect of the present invention has been established on what will be described below. More spe-

cifically, protein is composed of a sequence of twenty amino acids, but the sequence is not random. Hence, the frequency of occurrence of a specific oligopeptide as a partial amino acid sequence in the entire proteins encoded by genome derived from an appropriate organism species is not constant, but some oligopeptides occur at high frequencies in various proteins while other oligopeptides rarely occur therein. It is recognized that among them, oligopeptides highly frequently occurring in common to various proteins do not have any potency to determine the uniqueness (specificity) of individual proteins, namely any potency to determine the functions, while oligopeptides occurring at low frequencies adversely determine the uniqueness and functions of individual proteins.

[0125] It is suggested that the functional site of protein is composed of oligopeptides occurring at low frequencies. Additionally, longer oligopeptides, more rarely occurring, increase in number. In other words, oligopeptide of length $(n+1)$ as shown in the step (3) according to the method of the first aspect of the present invention is mostly the shortest oligopeptide occurring at a low frequency, and the calculated functional value Z_j of amino-acid residue A_j at an appropriate position j in the oligopeptide is the coefficient of the occurrence (namely, the representative value of the function) of the amino-acid residue A_j at the position.

[0126] According to the method for predicting the protein functional site in accordance with the second aspect of the present invention, the contribution degree of the amino-acid residue A_j to the frequency of the occurrence of A_j -oligopeptide can be evaluated on the basis of the ratio Y_{ji} of the frequency of the occurrence of the A_j -oligopeptide to that of the X_{ji} -oligopeptide as shown in the step (3), and thus, the calculated functional value $Z(j, n)$ of the amino-acid residue A_j at an appropriate position of a protein serves as the coefficient of the occurrence of the amino-acid residue A_j at the position (namely, the representative value of the function).

[0127] Furthermore, the value $Z(j, n)$ varies, depending on the species of amino-acid residue A_j . In the step (7) according to the inventive method, the distribution of the $Z(j, n)$ value of each of twenty amino acids is determined in the entire proteins of organism "a", to determine $D(j, n)$ value by normalizing the $Z(j, n)$ value on the basis of the mean and standard deviation of $Z(j, n)$ value of each amino acid, as determined on the basis of the distribution, which serves as the representative value of the function, after correction of the bias due to each amino-acid residue species.

[0128] Furthermore, longer oligopeptides, more rarely occurring, increase in number. Because the $Z(j, n)$ and $D(j, n)$ values generally vary, depending on the length (n) , accordingly, the functional value W_j of the $Z(j, n)$ and $D(j, n)$ as determined on a variety of length (n) is defined as the representative value of the function.

[0129] The systems for predicting a protein functional site in accordance with the third and fourth aspects of the present invention are individually systems for automatically carrying out the methods in accordance with the first and second aspects of the present invention; the methods for modifying protein in accordance with the fifth and sixth aspects of the present invention are methods for preparing mutant proteins by substituting the amino-acid residue at the functional site predicted by the method of the first aspect of the present invention with another amino-acid residue. Still further, the method for modifying a protein in accordance with the seventh aspect of the present invention is a method for preparing

a mutant protein by substituting the amino-acid residue at the functional site predicted by the method in accordance with the second aspect of the present invention with another amino-acid residue. In accordance with the eighth aspect of the present invention, a function-modified protein is provided. Furthermore, thermophilic DNA polymerase is provided as an embodiment of such protein.

[0130] The term "DNA polymerase" is the generic name of enzymes catalyzing the synthesis of a DNA chain complementary to a single-stranded DNA. DNA polymerase is an essential enzyme for DNA sequencing and in vitro DNA amplification, and "thermophilic DNA polymerase" is inevitable for PCR (polymerase chain reaction) in terms of the automation of a series of the reaction cycles.

[0131] Such thermophilic DNA polymerase includes known ones, for example Taq, Pfu, KOD, which are separately used, depending on the characteristic performance. Pfu DNA polymerase in particular has been known as an enzyme with an extremely low frequency of erroneous reading during the synthesis of DNA strands (at a high fidelity). However, the Pfu DNA polymerase is inappropriate for the amplification of polymeric DNAs such as genome DNA, because the synthetic DNA yielded by the Pfu DNA polymerase is low and the activity thereof to elongate a synthetic chain is insufficient. Thus, the present application provides a novel Pfu DNA polymerase prepared according to the method of the fifth aspect of the present invention.

BRIEF DESCRIPTION OF DRAWINGS

[0132] FIG. 1 depicts graphically the individual frequencies of the occurrences of oligopeptides of lengths 3, 4 and 5, and the distributions of the individual frequencies thereof according to the method of the first aspect of the present invention;

[0133] FIG. 2 depicts an example of an amino acid sequence of a length of 20, and examples of Aji-oligopeptide of a length of 4, containing the amino-acid residue Met at position 5 in the sequence and examples of Xji-oligopeptides;

[0134] FIG. 3 depicts graphically the individual frequencies of the occurrences of oligopeptides of lengths of 2, 3, 4 and 5, and the distributions of the individual frequencies thereof according to the method of the second aspect of the present invention;

[0135] FIG. 4 depicts an example of the flow chart for conducting the step (1) of the method of the second aspect of the present invention;

[0136] FIG. 5 depicts an example of the flow chart for conducting the steps (2') to (3) of the method of the second aspect of the present invention;

[0137] FIG. 6 depicts an example of the flow chart for conducting the steps (4) to (5) of the method of the second aspect of the present invention;

[0138] FIG. 7 depicts a distribution of the frequency of $Z(j, 3)$ of each of three amino acids, see below, according to the method of the second aspect of the present invention, wherein the solid line expresses the distribution of that of isoleucine (Ile); the dotted line expresses the distribution of that of alanine (Ala); and the alternate long and short dash line expresses the distribution of that of methionine (Met);

[0139] FIG. 8 depicts an example of the flow chart for conducting the step (7) of the method of the second aspect of the present invention;

[0140] FIG. 9 depicts an example of the flow chart for conducting the step (8) of the method of the second aspect of the present invention;

[0141] FIG. 10 depicts an example of the flow chart for conducting the step (9) of the method of the second aspect of the present invention;

[0142] FIG. 11 depicts a block diagram illustrating the system of the third aspect of the present invention;

[0143] FIG. 12 depicts a block diagram illustrating the system of the fourth aspect of the present invention;

[0144] FIG. 13 depicts the electrophoresis results of conventional Pfu DNA polymerase and KOD DNA polymerase, indicating the primer elongating activities thereof;

[0145] FIG. 14 depicts a distribution chart of the plotted $Z_j = -\log Y_j$ value for the whole amino acid sequence of the α -type DNA polymerase MJ encoded by MJ0885 (gene name from the genome of *Methanococcus jannaschii*), which value is calculated by the first aspect of the present invention;

[0146] FIG. 15 depicts a distribution chart of the plotted $Z_j = -\log Y_j$ value for partial sequences (motif A and motif C) of the amino acid sequence of which the distribution chart is shown in FIG. 14;

[0147] FIG. 16 depicts a frequency distribution chart of the $Z_j = -\log Y_j$ value calculated on the basis of the amino acid sequence of the α -type DNA polymerase MJ encoded by MJ0885;

[0148] FIG. 17 depicts alignment charts of the amino acid sequences of the individual motif Cs from the α -type DNA polymerases Pfu, KOD and MJ derived from *Pyrococcus furiosus*, *Pyrococcus* sp. KOD, and *Methanococcus jannaschii*, respectively;

[0149] FIG. 18 depicts distribution charts of the plotted $Z_j = -\log Y_j$ values for the individual motif Cs of the α -type DNA polymerases Pfu, KOD and MJ;

[0150] FIG. 19 depicts distribution charts of the plotted values of $W_j = Z(j, 3) - Z(j, 1)$ (in solid line), $W_j = Z(j, 4) - Z(j, 3)$ (in dotted line) and $W_j = Z(j, 5) - Z(j, 3)$ (in alternate long and short dash line) of the 100 residues from the N-terminus of the whole amino acid sequence of the α -type DNA polymerase MJ encoded by MJ0885, and these values are calculated by the method of the second aspect of the present invention;

[0151] FIG. 20 depicts distribution charts of the plotted value $W_j = Z(j, 5) - Z(j, 3)$ for partial sequences (regions comprising exoI, exoII, motif A, motif B and motif C) of the amino acid sequence of the α -type DNA polymerase MJ encoded by MJ0885;

[0152] FIG. 21 depicts distribution charts of the plotted values of $W_j = D(j, 3)$ (in dark color) and $W_j = D(j, 5)$ (in pale color) for partial sequences (regions comprising exoI, exoII, motif A, motif B and motif C) of the amino acid sequence of the α -type DNA polymerase MJ encoded by MJ0885;

[0153] FIG. 22 depicts distribution charts in dark color of the positions of amino-acid residues with $W_j = D(j, 3)$ of 2 or more or of 2 or less in the three-dimensional structure of the amino acid sequence of enolase encoded by MJ0232 (gene name from the genome of *Methanococcus jannaschii*) on a three-dimensional structure model;

[0154] FIG. 23 depicts the results of electrophoresis, indicating the primer elongating activities of the conventional Pfu DNA polymerase (wild type) and the modified Pfu DNA polymerase I of the present invention; and

[0155] FIG. 24 depicts the results of electrophoresis, indicating the primer elongating activities of the conventional Pfu

DNA polymerase (wild type) and the modified Pfu DNA polymerases II and III of the present invention.

BEST MODE FOR CARRYING OUT THE INVENTION

[0156] The method for predicting a functional site of a protein in accordance with the first aspect of the present invention is a method for predicting the functional site of an appropriate protein of organism "a" with a known genome data or cDNA analysis data, in the entire putative proteins of the organism "a", essentially comprising the following steps (1) to (6).

Step (1)

[0157] By determining the frequency of the occurrence of each amino acid and the frequencies of the occurrences of individual oligopeptides produced by permutations of twenty amino acids in the amino acid sequences of the entire proteins of the organism "a", the oligopeptide length (n) is determined.

[0158] The length n is determined, then, as the smallest integer satisfying the following criteria.

[0159] "Among oligopeptides of length n, the number of oligopeptides that occur once in the entire proteins is smaller than the number of oligopeptides that occur twice in the entire proteins; among oligopeptides of length (n+1), the number of oligopeptides that occur once in the entire proteins is larger than the number of oligopeptides that occur twice in the entire proteins."

[0160] For example, FIG. 1 depicts distribution charts of the frequencies of the occurrences of oligopeptides of a length of 3, oligopeptides of a length of 4 and oligopeptides of a length of 5, in the entire proteins encoded by the genome of a microorganism *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996). In the case of the three types of length of the oligopeptides shown in FIG. 1, the smallest n in the step (1) is 3.

Step (2)

[0161] Given that the j-th amino-acid residue from the N-terminus of the amino acid sequence (length of L) of the protein as a subject for predicting a functional site is described here as A_j ($n+1 \leq j \leq L-n$), the frequency of occurrence of a partial sequence of the amino acid sequence of the protein, which sequence corresponds to the following Aji-oligopeptide of length (n+1), containing the j-th amino-acid residue A_j ;

[0162] Aji-oligopeptide: $a_j1a_j2 \dots A_ji \dots a_jn a_j(n+1)$

[0163] ($1 \leq i \leq n+1$; $A_j = A_ji$; and A_j is the i-th residue of the oligopeptide; and moreover, $a_j1 = A_j-i+1, \dots, a_j(n+1) = A_j-i+(n+1)$),

and the frequency of the occurrence of the following Xji-oligopeptide of length (n+1);

[0164] Xji-oligopeptide: $a_j1a_j2 \dots X_ji \dots a_jn a_j(n+1)$

[0165] (the i-th residue X_ji is any amino acid; and moreover, $a_j1 = A_j-i+1, \dots, a_j(n+1) = A_j-i+(n+1)$)

should be determined in the entire proteins of the organism "a".

[0166] Such Aji-oligopeptide and Xji-oligopeptide can be illustrated for example in FIG. 2. The upper row (1) in FIG. 2 expresses in single letter code the partial sequence from the N-terminus to the 20-th amino-acid residue of the putative amino acid sequence on the basis of the gene MJ0885, which is believed to encode the α -type DNA polymerase of *Metha-*

nococcus jannaschii (Bult et al., Science 273, 1058-1073, 1996); the middle row (2) expresses examples of Aji-oligopeptide of a length of 4, containing the 5-th amino-acid residue Met(M) in the amino acid sequence; and the rows (3) to (6) further below express examples of Xji-oligopeptide containing the 5-th amino-acid residue M.

Step (3)

[0167] Calculating ratio Y_{ji} of the frequency of the occurrence of the Aji-oligopeptide to that of the Xji-oligopeptide.

Step (4)

[0168] The mean Y_j of the Y_{ji} is determined as follows.

$$Y_j = \sum_{i=1}^{n+1} Y_{ji} / (n+1),$$

Step (5)

[0169] Monotonously decreasing functional value or monotonously increasing functional value Z_j of Y_j is determined as follows;

$$Z_j = f(Y_j).$$

[0170] The Z_j value is defined as the representative value of the function of the j-th amino-acid residue of the amino acid sequence (length of L).

Step (6)

[0171] By subsequently repeating the steps (2) to (5) sequentially and determining the Z_j value of each of all the amino-acid residues at position $n+1 \leq j \leq L-n$, the degree of the involvement of each amino-acid residue in the function of the protein is predicted by using the dimension of the Z_j value as an indicator. More specifically, because the manner of occurring of each amino-acid residue in the context is expressed as the functional value Z_j of Y_j , a larger Z_j value indicates a lower frequency of occurrence of the amino-acid residue if Z_j is a monotonously decreasing functional value, which suggests that the amino-acid residue has higher responsibility over the fulfillment of the function. If Z_j is a monotonously increasing function, additionally, it is suggested that an amino-acid residue with a smaller Z_j value has greater responsibility over the function.

[0172] By expressing the Z_j value of each amino-acid residue for example in a distribution chart wherein the Z_j value is plotted on the vertical axis while the amino acid sequence is shown on the horizontal axis, furthermore, the functional site can be confirmed at a glance, which is preferable as an embodiment for carrying out the present invention.

[0173] The method for predicting a protein functional site in accordance with the second aspect of the present invention is a method for predicting a functional site of an appropriate protein in the entire putative proteins of the organism "a" with a known genome data or cDNA analysis data, essentially comprising the following steps (1) to (9).

Step (1)

[0174] The frequency of the occurrence of each amino acid and the frequencies of the occurrences of individual oligopep-

tides produced by permutations of twenty amino acids, in the amino acid sequences of the entire proteins of the organism "a", are determined.

[0175] For example, FIG. 3 shows a distribution chart of the frequencies of the occurrences of oligopeptides of a length of 3, oligopeptides of a length of 4 and oligopeptides of a length of 5, which are determined in the entire proteins encoded by the genome of a microorganism *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996) on the basis of the genome data of the microorganism.

[0176] FIG. 4 depicts an example of the flow chart for carrying out the step (1).

Step (2)

[0177] As to an appropriate protein of organism "a",

Step (2')

[0178] Given that the j-th amino-acid residue from the N-terminus of the amino acid sequence (length of L) of the protein is described here as A_j , the frequency of the occurrence of a partial sequence of the amino acid sequence of the protein, which sequence corresponds to the following A_{ji} -oligopeptide of an appropriate length n ($1 \leq n \leq M$), provided that "M" is the smallest length of oligopeptides satisfying the following criterion; all the oligopeptides of length M are at frequency 1 of the occurrence), containing the j-th amino-acid residue A_j ($n \leq j \leq L-n+1$);

[0179] A_{ji} -oligopeptide: $aj_1 aj_2 \dots A_j \dots aj_n$

[0180] ($1 \leq i \leq n$; $A_j = A_j$ and A_{ji} is the i-th residue of the oligopeptide; and moreover, $aj_1 = A_{j-i+1}, \dots, aj_{(n+1)} = A_{j-i+(n+1)}$),

and the frequency of the occurrence of the following X_{ji} -oligopeptide of the length n corresponding to the length of the A_{ji} -oligopeptide;

[0181] X_{ji} -oligopeptide: $aj_1 aj_2 \dots X_{ji} \dots aj_n$

[0182] (the i-th residue X_{ji} is any amino acid; and moreover, $aj_1 = A_{j-i+1}, \dots, aj_{(n+1)} = A_{j-i+(n+1)}$),

are determined in the entire proteins of the organism "a".

[0183] In the same manner as by the method of the first aspect of the present invention, such A_{ji} -oligopeptide and X_{ji} -oligopeptide are for example illustrated as in FIG. 2.

Step (3)

[0184] The ratio Y_{ji} of the frequency of the occurrence of the A_{ji} -oligopeptide to that of the X_{ji} -oligopeptide is determined.

[0185] FIG. 5 depicts an example of the flow chart for carrying out the aforementioned steps (2') to (3).

Step (4)

[0186] The mean $Y(j, n)$ of the Y_{ji} is determined as described below.

$$Y(j, n) = \sum_{i=1}^n Y_{ji}/n.$$

Step (5)

[0187] The logarithmic value $Z(j, n)$ of $Y(j, n)$ is determined as follows.

$$Z(j, n) = -\log(Y(j, n))$$

[0188] FIG. 6 depicts an example of the flow chart for carrying out the aforementioned steps (4) to (5).

Step (6)

[0189] By subsequently repeating the steps (2') to (5) sequentially, the $Z(j, n)$ value of each of all the amino-acid residues at position $n \leq j \leq L-n+1$ in the amino acid sequence (length of L) is determined.

Step (7)

[0190] By sequentially repeating the steps (2) to (6) over the entire proteins of the organism "a", thereby determining the distribution of the $Z(j, n)$ value of each amino-acid residue in the entire proteins, and the $Z(j, n)$ values are classified into twenty according to the twenty amino acids, and then determining mean $Av(Aa)$ of the $Z(j, n)$ values for each amino acid Aa and the standard deviation $Sd(Aa)$ of the distribution thereof, on the basis of the distribution, to determine function g to the j-th amino-acid residue A_j of a protein for normalizing the difference in distribution due to (or among?) the species of amino-acid residues is determined;

$$g = g(Z(j, n), A_j) = [Z(j, n) - Ad(Aa)] / Sd(Aa)$$

[0191] (provided that $A_j = Aa$; and $1 \leq n \leq M$).

[0192] For example, FIG. 7 depicts a distribution of the frequency of $Z(j, n)$ for three species of amino acids, namely isoleucine (Ile), alanine (Ala) and methionine (Met), in the entire proteins encoded by the genome of *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996). Based on the distribution, the mean and standard deviation of the $Z(j, n)$ values for an amino acid isoleucine (Ile), namely $Ad(Ile)$ and $Sd(Ile)$, respectively, are determined as $Ad(Ile) = 3.16$ and $Sd(Ile) = 0.17$, and the function g for $A_j = Ile$ is determined as follows.

$$g = g(Z(j, n), A_j) = (Z(j, n) - 3.16) / 0.17$$

[0193] FIG. 8 depicts an example of the flow chart for carrying out the step (7).

Step (8)

[0194] The value $D(j, n)$ of the function g of each of all the amino-acid residues A_j at position $n \leq j \leq L-n+1$ in the amino acid sequence (length of L) as recovered in the step (7) is determined;

$$D(j, n) = g(Z(j, n), A_j).$$

[0195] FIG. 9 depicts an example of the flow chart for carrying out the step (8).

Step (9)

[0196] The functional value W_j of the $Z(j, n)$ value and the $D(j, n)$ value is determined as follows.

$$W_j = h(Z(j, 1), Z(j, 2), \dots, Z(j, M), D(j, 1), D(j, 2), \dots, D(j, M))$$

[0197] By defining the value of the W_j as the representative value of the function of the j-th amino-acid residue in the amino acid sequence (length of L), the degree of the responsibility of each amino-acid residue over the function of the protein is estimated by using the dimension of the W_j value as an indicator.

[0198] FIG. 10 depicts an example of the flow chart for carrying out the step (9).

[0199] By expressing the W_j value of each amino-acid residue for example in a distribution chart wherein the W_j value is plotted on the vertical axis while the amino acid sequence is shown on the horizontal axis, furthermore, the functional site can be confirmed at a glance, which is preferable as an embodiment for carrying out the present invention.

[0200] If the three-dimensional structure of the protein as a subject for predicting the functional site is known or if a three-dimensional structure model thereof can be prepared by known methods (for example, homology modeling method, Peitsch, Proceedings of the Fifth International Conference on Intelligent Systems for Molecular Biology, 1997, 5, 234-236), the distribution is expressed on the three-dimensional structure, whereby a spatial arrangement of an amino-acid residue as a candidate of a novel functional site can be confirmed, which is preferable as an embodiment for carrying out the invention.

[0201] The system for predicting a functional site of a protein in accordance with the third aspect of the present invention is a system for automatically carrying out the method for predicting a functional site in accordance with the first aspect of the present invention, at least comprising the following units (a) to (g) for conducting the steps (1) to (6) according to the method of the first aspect of the present invention, as shown for example in the composition example in FIG. 11.

Outer Memory Unit (a):

[0202] Unit memorizing the amino acid sequence data of a protein or an existing protein data base for use in the step (1).

Calculation/Memory Unit (b):

[0203] Unit, composed of CPU calculating the frequencies of the occurrences of individual oligopeptides as determined in the step (1), and a memory unit having the memory of the calculation results.

Calculation/Memory Unit (c):

[0204] Unit, composed of CPU calculating the smallest length (n) of oligopeptides as determined in the step (1) and a memory unit having the memory of the length n.

Calculation/Memory Unit (d):

[0205] Unit, composed of CPU calculating the frequencies of occurrence of each amino acid and the frequencies of occurrence of A_{ji} -oligopeptide and X_{ji} -oligopeptide in the entire proteins as determined in the step (2) and a memory unit having the memory of the calculation results.

Calculation/Memory Unit (e):

[0206] Unit, composed of CPU calculating the Y_{ji} value as determined in the step (3) and a memory unit having the memory of the Y_{ji} value.

Calculation/Memory Unit (f):

[0207] Unit, composed of CPU calculating the Y_j value as determined in the step (4) and a memory unit having the memory of the Y_j value.

Calculation/Memory Unit (g):

[0208] Unit, composed of CPU calculating the Z_j value as determined in the step (5) and a memory unit having the memory of Z_j .

[0209] Additionally, the system for predicting a functional site is provided with the following display unit (h) in a preferable embodiment.

Display Unit (h):

[0210] Unit displaying the Z_j value of each amino-acid residue recovered in the calculation/memory unit (g) in a distribution chart.

[0211] The system of the present invention may be equipped with keyboard (i) and control unit (j) and the like as illustrated in FIG. 11, in addition to these units (a) to (h).

[0212] According to the fourth aspect of the present invention, the system for predicting a protein functional site is a system for automatically conducting the method of the second aspect of the present invention, at least comprising the following units (a) to (i) for carrying out the steps (1) to (9) according to the method of the second aspect of the present invention, as shown in the composition example in FIG. 12.

Outer Memory Unit (a):

[0213] Unit memorizing the amino acid sequence data and an existing protein data base for use in the step (1).

Calculation/Memory Unit (b):

[0214] Unit, composed of CPU calculating the frequencies of the occurrences of individual oligopeptides as determined in the step (1) and a memory unit having the memory of the calculation results.

Calculation/Memory Unit (c):

[0215] Unit, composed of CPU calculating the frequencies of occurrence of each amino acid and the individual frequencies of the occurrences of A_{ji} -oligopeptide and X_{ji} -oligopeptide in the entire proteins as determined in the step (2') and a memory unit having the memory of the calculation results.

Calculation/Memory Unit (d):

[0216] Unit, composed of CPU calculating Y_{ji} as determined in the step (3) and a memory unit having the memory of the Y_{ji} value.

Calculation/Memory Unit (e):

[0217] Unit, composed of CPU calculating the $Y(j, n)$ value as determined in the step (4) and a memory unit having the memory of the $Y(j, n)$ value.

Calculation/Memory Unit (f):

[0218] Unit, composed of CPU calculating the $Z(j, n)$ value as determined in the steps (5) and (6) and a memory unit having the memory of the $Z(j, n)$ value.

Calculation/Memory Unit (g):

[0219] Unit, composed of CPU calculating the g value as determined in the step (7) and a memory unit having the memory of the g value.

Calculation/Memory Unit (h):

[0220] Unit, composed of CPU calculating the $D(j, n)$ value as determined in the step (8) and a memory unit having the memory of the $D(j, n)$ value.

Calculation/Memory Unit (i):

[0221] Unit, composed of CPU calculating the W_j value as determined in the step (9) and a memory unit having the memory of the W_j value.

[0222] Additionally, the system for predicting a functional site in accordance with the fourth aspect of the present invention may be equipped with an appropriate combination of the following units (j) to (l).

Display Unit (j):

[0223] Unit displaying the W_j value of each amino-acid residue as recovered with the unit (i) in a distribution chart.

Calculation/Memory Unit (k):

[0224] Unit memorizing an existing database of protein three-dimensional structures or unit preparing a three-dimensional structure model based on an amino acid sequence according to a known method and memorizing the three-dimensional structure.

Display Unit (l):

[0225] Unit displaying the W_j value of each amino-acid residue in a distribution chart on the three-dimensional structure stored in the database or three-dimensional structure model recorded on the unit (k).

[0226] The system of the present invention may satisfactorily be equipped with keyboard (m) and control unit (n) and the like as illustrated in FIG. 12, in addition to these units (a) to (l).

[0227] The method for modifying the function of a protein in accordance with the fifth aspect of the present invention will be described below. The method is a method for modifying the function of protein "A", which function has been known, derived from the entire putative proteins of organism "a" with a known genome data or cDNA analysis data, essentially comprising the following steps (1) to (10).

Step (1):

[0228] Extracting proteins closely related to the protein "A" from an existing protein data base and subjecting the proteins to alignment.

Steps (2) to (7):

[0229] Subjecting the amino acid sequences of the entire proteins of the organism "a" to the steps (1) to (6) according to the method of the first aspect of the present invention.

Step (8):

[0230] Selecting at least one amino-acid residue to be subjected to mutation from the amino acid sequence (length of L) of the protein "A" on the basis of the alignment data recovered in the step (1), sequentially repeating the steps (3) to (6) for a variant amino-acid residue of various mutated amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residue, to determine the Z_j value of the variant amino-acid residue.

Step (9):

[0231] Selecting a mutated amino acid sequence wherein the Z_j value of the variant amino-acid residue as determined in the step (8) is larger or smaller than the Z_j value of the intact amino-acid residue as determined in the step (7).

Step (10):

[0232] Preparing a modified gene of the protein "A", which gene encodes the mutant amino acid sequence selected in the step (9), and expressing the modified gene in an appropriate host-vector system to prepare modified protein "A".

[0233] In accordance with the sixth aspect of the present invention, the method for modifying a protein function is a method for modifying the function of protein "B" derived from organism "b" with an unknown genome data or cDNA analysis data, essentially comprising the following steps (1) to (10).

Step (1):

[0234] Extracting protein "A" most closely related to protein "B" from the entire putative proteins of organism "a" with a known genome data or cDNA analysis data and subjecting the protein "A" to alignment, or extracting proteins closely related to protein "B" from an existing protein data base to subject the proteins to alignment.

Steps (2) to (8):

[0235] Conducting the steps (2) to (8) of the method of the third aspect of the present invention over the amino acid sequences of the entire proteins of the organism "a".

Step (9):

[0236] Selecting a position that should be mutated and an amino acid residue for which should be substituted wherein the Z_j value of the substituted amino-acid residue as determined in the step (8) is larger or smaller than the Z_j value of the intact amino-acid residue as determined in the step (7).

Step (10):

[0237] Preparing a modified gene of the protein B according to a known method, which gene encodes the amino acid sequence mutated at the position to another amino acid residue as selected in the step (9), and expressing the modified gene in an appropriate host-vector system to prepare modified protein B.

[0238] As has been described above, the methods for modifying the protein function in the fifth and sixth aspects of the present invention, comprising the method for predicting a functional site of the first aspect of the present invention, are characterized in that an unknown functional site of protein is newly found and is subjected to mutation.

[0239] Furthermore, the method for modifying a protein function according to the seventh aspect of the present invention also utilizes the method for predicting the function in accordance with the second aspect of the present invention, whereby the method can be carried out in the same manner as in the case of the fifth aspect of the present invention.

[0240] The eighth aspect of the present invention is a protein of which function is artificially modified by the method of the fifth to seventh aspect of the present invention. An embodiment of such protein is thermophilic DNA polymerase, which is more specifically an enzyme prepared by modifying, in accordance with the sixth aspect of the present invention, a thermophilic Pfu DNA polymerase derived from *Pyrococcus furiosus* in a genetic engineering manner by the known method for preparing mutant gene (Strategies, Vol. 9, p. 3-4, 1996) (the thermophilic DNA polymerase of the present invention is sometimes referred to as "modified Pfu

DNA polymerase"). The enzyme can be prepared as follows. More specifically, because the nucleotide sequence of the gene of Pfu DNA polymerase is known (Nucleic Acids Research, Vol. 21, p. 259-265, 1993), the gene of Pfu DNA polymerase is prepared by PCR comprising synthetically preparing an oligopeptide complementary to both the ends by using the genome DNA of the archaeobacterium as template and using the oligopeptide as primer. The DNA fragment of the gene is cloned into a vector, and the gene is subsequently subjected to mutation by the method described in the reference mentioned above. In accordance with the present invention, in particular, the mutation of the gene was executed by nucleotide substitution, so that a part of the amino acid sequence of Pfu DNA polymerase might be substituted with the amino-acid residues from KOD DNA polymerase. In terms of amino acid sequence, Pfu DNA polymerase has about 80% homology with KOD DNA polymerase, and therefore, similar synthetic termination occurs during PCR (FIG. 13), but the elongation rate with KOD DNA polymerase is about 6-fold the rate with Pfu DNA polymerase. By substituting some amino-acid residues of Pfu DNA polymerase with some amino-acid residues of KOD DNA polymerase, the synthetic termination of the chain elongation might be improved or the elongation rate turns rapid, which possibly enables the recovery of an enzyme capable of elongating a DNA chain under way of synthesis more longer. By expressing in *Escherichia coli* the mutant gene that was mutated in such a manner and recovering and purifying the expression product, the modified Pfu DNA polymerase of the present invention was recovered.

[0241] The thermophilic DNA polymerase (modified Pfu DNA polymerase I) of the present invention is more specifically an enzyme of the amino acid sequence of SQ ID No.1. The amino acid sequence is a novel sequence prepared by identifying potentially function-modifiable amino-acid residues of the amino acid sequence of the conventionally known Pfu DNA polymerase, according to the inventive method for predicting a functional site, and substituting the amino-acid residues as shown in Table 1. By using the novel enzyme then for DNA synthesis by PCR, for example, the synthetic termination occurring when using the conventional DNA polymerases is almost totally overcome, as shown in the following examples. It is needless to say that template DNA chains to be highly efficiently amplified with the conventional polymerase can be amplified at high efficiency in the same manner.

[0242] Furthermore, the thermophilic DNA polymerases (modified Pfu DNA polymerases II and III) of the present invention are more specifically enzymes of amino acid sequences of SQ ID Nos. 6 and 7, which are novel sequences prepared by identifying potentially function-modifiable amino-acid residues of the amino acid sequence of the Pfu DNA polymerase, according to the inventive method for predicting a functional site, and substituting the amino-acid residues as shown in Table 1. By using the novel enzymes then for DNA synthesis by PCR, for example, synthetic polymeric products can be recovered at large scales, as shown in the following examples.

TABLE 1

Modified DNA polymerases	Positions	Wild-type amino acid	Modified amino acid
I	2	Ile	Val
	533	Phe	Tyr
	538	Leu	Ile

TABLE 1-continued

Modified DNA polymerases	Positions	Wild-type amino acid	Modified amino acid
II	540	Ile	Ser
	545	Leu	Phe
	546	Tyr	Phe
	2	Ile	Val
	710	Pro	Arg
	712	Ser	Arg
	713	Asn	Asp
III	717	Leu	Pro
	2	Ile	Val
	717	Leu	Pro

[0243] The DNA sequences encoding these modified Pfu DNA polymerases include for example the mutant genes of the Pfu DNA polymerase gene, as recovered during the process of enzyme preparation. As to these mutant genes, the DNA sequences encoding the amino acid sequences of SQ ID Nos.1, 6 and 7 for example have been cloned in recombinant plasmids p320, pDP5b17 and pDP5C4, respectively, and these recombinant plasmids have been integrated in *Escherichia coli* HMS174 (DE3) and deposited at the Life Engineering and Industrial Technology Research Institute, the Agency of Industrial Science and Technology, Japan (Accession Nos. FERM P-16052, FERM BP-6189 and FERM BP-6190, respectively).

[0244] Additionally, the DNA sequences of the present invention may appropriately be designed as DNA sequences with conjugated nucleotide codons corresponding to the individual amino-acid residues of SQ ID No.1, 6 or 7.

[0245] The thermophilic DNA polymerases of the present invention may be expressed in microorganisms such as *Escherichia coli*, which may thereafter be recovered. By inserting and integrating the DNA sequence into an expression vector with an origin of replication in a microorganism, a promoter, a ribosome-binding site, a cDNA cloning site, and a terminator and the like to prepare an expression vector, transforming a host cell with the expression vector and thereafter culturing the resulting transformant, an enzyme encoded by the DNA sequence can be generated in the microorganism at a large scale.

EXAMPLES

[0246] The present invention will now be described more specifically in more detail with reference to examples, but the invention is not limited to the following examples.

Example 1

[0247] According to the method of the first aspect of the present invention and based on the genome data of *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996), $Z_j = -\log Y_j$ ($f = -\log$) was calculated, concerning each amino-acid residue in the amino acid sequence (from the N-terminus to the C terminus) of a DNA polymerase speculated from the microbial gene MJ0885 believed to encode the α -type DNA polymerase. The results are plotted in a distribution chart in FIG. 14.

[0248] Among the motifs known as the functional sites of the α -type DNA polymerase, furthermore, motif A and motif C were extracted, and the Z_j values of the individual amino-acid residues were plotted in FIG. 15. FIG. 15 and FIG. 16 below suggest that the $Z_j = -\log Y_j$ values of amino-acid resi-

dues responsible for the function are larger than those of the remaining amino-acid residues:

[0249] FIG. 16 depicts a distribution chart of the frequency of the value $Z_j = -\log Y_j$ for the amino acid sequence of the α -type DNA polymerase encoded by MJ0885. It is confirmed in the figure that amino-acid residues with value $Z_j = -\log Y_j$ of 4.8 or more are highly possibly amino-acid residues responsible for the protein function.

Example 2

[0250] Following the chart on FIG. 15 in Example 1, the characteristic properties of α -type DNA polymerase Pfu (DDBJ Accession No. D12983) derived from *Pyrococcus furiosus* were modified, on the basis of the amino acid sequence of an α -type DNA polymerase KOD (DDBJ Accession No. D29671) derived from *Pyrococcus* sp., and the genome data of *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996), and the amino acid sequence of the α -type DNA polymerase (named here as MJ) encoded by MJ0885.

[0251] FIG. 17 depicts alignment charts of the amino acid sequences of the motif Cs from Pfu, KOD and MJ, with no difference between the region 531 to 544 from Pfu and the region 551 to 564 from MJ (the numbers 550 and 570 indicate the N-terminus and C-terminus amino acid residues of the motif C shown here from MJ).

[0252] FIG. 18 depicts the results of the prediction of functional sites in the amino acid sequences of the motif Cs of Pfu, KOD and MJ according to the inventive method, on the basis of the genome data of *Methanococcus jannaschii*. The results indicate that mutations Ile540Ser, Leu545Phe, Tyr546Phe, and Ile548Thr increase the $Z_j = -\log Y_j$ values of the amino-acid residues. Furthermore, the values $Z_j = -\log Y_j$ of Asp541 and Ala547 are increased. By subjecting the α -type DNA polymerase MJ of *Methanococcus jannaschii* to such mutation, the resulting sequence turns more unique (specific) which possibly brings about the improvement of some function.

Example 3

[0253] Based on the genome data of *Methanococcus jannaschii* (Bult et al., Science 273, 1058-1073, 1996), the values $Z(j, 1) = -\log Y(j, 1)$, $Z(j, 3) = -\log Y(j, 3)$, $Z(j, 4) = -\log Y(j, 4)$, and $Z(j, 5) = -\log Y(j, 5)$ were calculated, concerning the individual amino-acid residues of the amino acid sequence (from the N to C termini) of a DNA polymerase speculated on the basis of the microbial gene MJ0885 believed to encode the α -type DNA polymerase, according to the method of the second aspect of the present invention, so that $W_j = Z(j, 3) - Z(j, 1)$ ($h = Z(j, 3) - Z(j, 1)$) was calculated. Similarly, $W_j = Z(j, 4) - Z(j, 3)$ ($h = Z(j, 4) - Z(j, 3)$) and $W_j = Z(j, 5) - Z(j, 3)$ ($h = Z(j, 5) - Z(j, 3)$) were also calculated.

[0254] FIG. 19 depicts the results of the 100 residues from the N-terminus in a plotted distribution chart. Given $h = Z(j, 5) - Z(j, 3)$, regions with significantly different distributions from those of the remaining two cases are present in the region from the 35-th to 60-th residues and the like. The distributions indicate that smaller $W_j = Z(j, 5) - Z(j, 3)$ more specifically characterizes the amino acid sequence.

[0255] Among the motifs known as the functional sites of the α -type DNA polymerases, furthermore, regions containing exoI, exoII, motif A, motif B and motif C were extracted, and subsequently, the W_j values of the individual amino-acid

residues are plotted in FIG. 20. As shown in FIG. 20, the regions with the characteristic reduction of W_j are consistent with the functional sites.

Example 4

[0256] FIG. 21 depicts the values $W_j = D(j, 3)$ and $W_j = D(j, 5)$ of individual amino-acid residues in the regions containing the exoI, exoII, motif A, motif B and motif C extracted among the motifs known as the functional sites in the α -type DNA polymerases ($h = D(j, 3)$ and $h = D(j, 5)$). Amino-acid residues with $W_j = D(j, n)$ of 2 or more or of 2 or less are present outside the motifs, and these amino-acid residues are candidates of new functional sites.

Example 5

[0257] FIG. 22 depicts in a dark color the positions of amino-acid residues having $W_j = D(j, 3)$ of 2 or more or of 2 or less in the amino acid sequence of MJ0232, which is speculated as enolase of *Methanococcus jannaschii*, on a three-dimensional structure model prepared on the basis of the enolase of budding yeast. It is indicated that residues positioned apart on the amino acid sequence are closely positioned on the three-dimensional structure.

Example 6

[0258] Modified Pfu DNA polymerase I was prepared, by applying the mutation of putative amino-acid residues for improving the function of the DNA polymerase MJ in Example 2 to the Pfu DNA polymerase.

(1) Preparation of Modified Pfu DNA Polymerase Gene

Cloning of Pfu DNA Polymerase Gene:

[0259] Following the nucleotide sequence of Pfu DNA polymerase gene (Nucleic Acids Research, Vol. 21, p. 259-265, 1993), a PCR primer was prepared, for amplifying the objective gene by PCR by using the genome DNA of *P. furiosus* as template, which was then cloned in an expression vector for *Escherichia coli*. The detail is described below.

[0260] *P. furiosus* DSM3638 was cultured according to the method described in the reference described above. First, the culture medium described in the reference was prepared, followed by sterilization at a high temperature under pressure, and subsequently, nitrogen gas was purged into the resulting culture medium. The bacterium was inoculated into the culture medium, for stationary culturing at 95° C. for 15 hours. From 200 ml of the culture broth were recovered the bacteria of about 0.5 mg by centrifugation. The collected bacteria were suspended in buffer A (10 mM Tris/HCl (pH 8.0) 1 mM EDTA, 100 mM NaCl), followed by addition of 1 ml of 10% SDS and subsequent agitation, and to the resulting suspension was added 0.5 mg of proteinase K for reaction at 55° C. for 60 minutes. The reaction solution was extracted sequentially in phenol, phenol/chloroform, and chloroform, and to the extract was added ethanol to make the DNA insoluble, which was then recovered. The resulting DNA was dissolved in 1 ml of TE buffer (10 mM Tris/HCl (pH 8.0), 1 mM EDTA), followed by addition of 0.5 mg RNase A for reaction at 37° C. for 60 minutes and re-extraction sequentially in phenol, phenol/chloroform, and chloroform, and subsequent ethanol precipitation, to recover the DNA, which was then dissolved in the TE buffer, to recover the DNA at about 0.3 mg.

[0261] For PCR amplification of the objective DNA polymerase gene, then, two primer DNAs of SQ ID Nos.2 and 3 were synthesized on the basis of the known sequence data. More specifically, it was designed that the initiation codon ATG of the objective gene and a restriction nuclease NcoI sequence (5'-CCATGG-3') might be introduced in the forward primer sequence, while the reverse primer might be conjugated at an appropriate position downstream the termination codon. PCR was conducted in a reaction system of 50 μ l, by using 2 μ g of *P. furiosus* DNA and 10 pmol each of the primers under conditions for LA Taq (manufactured by TaKaRa Brewery) and attached buffers. The cycle conditions were as follows; 93° C./3 minutes prior to the addition of the enzyme, and 30 cycles of each cycle composed of 94° C./0.5 minute, 55° C./0.5 minute and 72° C./1.0 minute. The amplified DNA fragment was purified, followed by treatment with NcoI, and the resulting DNA fragment was similarly cleaved with NcoI and subsequently blunt ended, and the resulting fragment was then integrated downstream the T7 promoter of an NcoI-treated expression vector pET-15b. The expression vector was defined as pDPWT100, to confirm the nucleotide sequence of the inserted gene.

Modification of Pfu DNA Polymerase Gene:

[0262] According to the known method (Strategies, Vol. 9, p. 3-4, 1996) and for the expression vector pDPWT100 with the cloned Pfu DNA polymerase gene integrated therein, a modified Pfu DNA polymerase gene was prepared on the expression vector pDPWT100 by using oligopeptides containing desired mutations (SQ ID Nos.4 and 5) and the mutation induction kit manufactured by Promega Corporation, whereby an expression vector pDP320 was constructed. By determining the nucleotide sequence of the modified gene, furthermore, the amino acid sequence of the modified Pfu DNA polymerase (SQ ID No.1) was verified.

(2) Expression and Purification of the Modified Pfu DNA Polymerase in *Escherichia coli*

[0263] The gene of the modified Pfu DNA polymerase I was expressed in *Escherichia coli* as follows, which was then purified.

[0264] The expression vector pDP320 with the modified Pfu DNA polymerase gene was inserted in a strain *Escherichia coli* HMS174 (DE3) and cultured in an LB culture medium supplemented with IPTG to a final concentration of 0.1 mM for 14 hours, to induce the expression of the enzyme in the bacteria of the strain *Escherichia coli*. After harvesting the bacteria by centrifugation, a modified Pfu DNA polymerase was extracted under ultrasonic treatment in a buffer containing 150 mM Tris/HCl (pH 7.5), 2 mM EDTA, 0.24 mM APMSF and 0.2% Tween 20. The crude extract solution was thermally treated at 80° C. for 15 minutes, to inactivate the DNA polymerase derived from *Escherichia coli* and partially purify the DNA polymerase of the present invention. The partially purified fraction was dialyzed against a buffer composed of 50 mM Tris/HCl (pH 7.5), 1 mM EDTA, 0.2% Tween 20, 7 mM 2-mercaptoethanol and 10% glycerol. At the stage was detected a DNA polymerizing activity specific to the modified Pfu DNA polymerase I.

Example 7

[0265] By using the modified Pfu DNA polymerase I partially purified in Example 6, the primer elongation reaction of a DNA chain complimentary to the template DNA was tested.

[0266] One μ g of the partially purified enzyme fraction described above was placed in 20 μ l of a reaction solution containing 20 mM Tris/HCl (pH 8.0), 2 mM MgCl₂, 50 μ g/ml BSA, 0.1% Triton X-100, 1 mM each of cold dNTPs (0.1 mM for dCTP), 0.63 μ g of pBLUESCRIPT plasmid prepared by annealing together 10 μ Ci of [α -³²P]dCTP and a primer of M13(-21), for reaction at 75° C. for one minute and 3 minutes. The elongated DNA chain was separated by electrophoresis on a polyacrylamide gel containing 8M urea, and the resulting pattern was analyzed with an image analyzer. As a control, additionally, the conventional wild-type Pfu DNA polymerase was used for the same DNA synthesis.

[0267] The results are shown in FIG. 23. When the conventional wild-type Pfu DNA polymerase was used, at least 10 bands indicating the presence of incomplete DNA chains due to synthetic termination were observed. However, these bands disappeared during the DNA synthesis with the modified Pfu DNA polymerase I of the present invention. Alternatively, no difference in the accumulation of highly elongated DNA chains around 1000 bases was observed.

Example 8

[0268] Pfu DNA polymerases II and III were prepared.

(1) Preparation of Modified Pfu DNA Polymerase Gene

[0269] In the same manner as in Example 6(1), the Pfu DNA polymerase gene was cloned, to prepare modified genes II and III as follows.

Preparation of Modified Pfu DNA Polymerase II:

[0270] According to the known method (Strategies, Vol. 9, p. 3-4, 1996) and for the expression vector pDPWT100 with the cloned Pfu DNA polymerase gene integrated therein, the gene of modified Pfu DNA polymerase II was prepared on the expression vector pDPWT100 by using oligopeptides containing desired mutations (SQ ID Nos.8 and 9) and the mutation induction kit manufactured by Promega Corporation, whereby an expression vector pDP5b17 was constructed. By determining the nucleotide sequence of the modified gene, furthermore, the amino acid sequence of the modified Pfu DNA polymerase II (SQ ID No.6) was confirmed.

Preparation of the Gene of Modified Pfu DNA Polymerase III:

[0271] By the same method as described above except for the use of the oligonucleotides of SQ ID Nos.10 and 11, the gene of modified Pfu DNA polymerase III was prepared, to construct an expression vector pDP5C4. By determining the nucleotide sequence of the modified gene, the amino acid sequence (SQ ID No.7) of the modified Pfu DNA polymerase III was confirmed.

(2) Expression in *Escherichia coli* and Purification of the Modified Pfu DNA Polymerases II and III

[0272] The genes of the modified Pfu DNA polymerases II and III, thus prepared, were expressed in *Escherichia coli* as follows, which were then purified.

[0273] The expression vectors pDP5b17 and pDP5C4 were independently inserted in a strain *Escherichia coli* HMS174 (DE3) and cultured in an LB culture medium supplemented with IPTG to a final concentration of 0.1 mM for 14 hours, to induce the expression of the enzymes in the bacteria of the strain *Escherichia coli*. After harvesting the bacteria by centrifugation, modified Pfu DNA polymerases II and III were

extracted, with ultrasonic treatment, in a buffer containing 150 mM Tris/HCl(pH 7.5), 2 mM EDTA, 0.24 mM APMSF and 0.2% Tween 20. The crude extract solution was thermally treated at 80° C. for 15 minutes, to inactivate the DNA polymerases derived from *Escherichia coli* and partially purify the modified DNA polymerases II and III. The partially purified fractions were dialyzed against a buffer composed of 50 mM Tris/HCl (pH 7.5), 1 mM EDTA, 0.2% Tween 20, 7 mM 2-mercaptoethanol and 10% glycerol. At the stage were detected DNA polymerizing activities specific to the modified Pfu DNA polymerases II and III.

Example 9

[0274] By using the modified Pfu DNA polymerases II and III partially purified in Example 8, the primer elongation reaction of a DNA chain complementary to the template DNA was tested.

[0275] One µg of each of the partially purified enzyme fractions described above was placed in 20 µl of a reaction solution containing 20 mM Tris/HCl(pH 8.0), 2 mM MgCl₂, 50 µg/ml BSA, 0.1% Triton X-100, 1 mM each of cold dNTPs (0.1 mM for dCTP), 0.63 µg of pBLUESCRIPT plasmid prepared by annealing together 10 µCi of [α-³²P]dCTP and a primer M13(-21), for reaction at 75° C. for one minute and 3 minutes. The elongated DNA chain was separated by electrophoresis on a polyacrylamide gel containing 8M urea, and the resulting pattern was analyzed with an image analyzer. As a control, additionally, the conventional wild-type Pfu DNA polymerase was used for the same DNA synthesis.

[0276] The results are shown in FIG. 24. When the conventional wild-type Pfu DNA polymerase was used, bands indicating the presence of incomplete DNA chains were present under observation, because of the presence of a large region at about 1000 bases where synthetic termination occurred. However, the yield of synthesized products including those of bands of about 1000 bases was elevated during the DNA synthesis with the modified Pfu DNA polymerases II and III of the present invention, together with bands indicating the presence of more polymeric (more elongated) PCR products under observation.

[0277] The results described above indicate that the DNA polymerases of the present invention can more markedly elongate DNA chains in the course of synthesis during the DNA synthesis by PCR.

INDUSTRIAL APPLICABILITY

[0278] In accordance with the present invention, the functional site of a functionally unknown protein recovered by genome analysis or cDNA analysis can be predicted. A novel functional site of a protein with a known function can also be predicted.

[0279] The thermophilic DNA polymerases provided by the present invention can highly efficiently synthesize and amplify the whole length of a polymeric DNA by PCR, whereby the in vitro synthesis and amplification of a DNA chain and the nucleotide sequencing thereof can be attained at a high precision in a simple manner.

SEQUENCE LISTING

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<212> TYPE: PRT
<213> ORGANISM: Unknown
<220> FEATURE:
<223> OTHER INFORMATION: Description of Unknown Organism: DNA polymerase

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Arg Leu Phe Lys Lys Glu Asn Gly Lys Phe Lys Ile Glu His Asp Arg
 20            25            30

Thr Phe Arg Pro Tyr Ile Tyr Ala Leu Leu Arg Asp Asp Ser Lys Ile
 35            40            45

Glu Glu Val Lys Lys Ile Thr Gly Glu Arg His Gly Lys Ile Val Arg
 50            55            60

Ile Val Asp Val Glu Lys Val Glu Lys Lys Phe Leu Gly Lys Pro Ile
 65            70            75            80

Thr Val Trp Lys Leu Tyr Leu Glu His Pro Gln Asp Val Pro Thr Ile
 85            90            95

Arg Glu Lys Val Arg Glu His Pro Ala Val Val Asp Ile Phe Glu Tyr
 100           105           110

Asp Ile Pro Phe Ala Lys Arg Tyr Leu Ile Asp Lys Gly Leu Ile Pro
 115           120           125
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Leu	Tyr	His	Glu	Gly	Glu	Glu	Phe	Gly	Lys	Gly	Pro	Ile	Ile	Met	Ile
	145				150					155					160
Ser	Tyr	Ala	Asp	Glu	Asn	Glu	Ala	Lys	Val	Ile	Thr	Trp	Lys	Asn	Ile
			165						170					175	
Asp	Leu	Pro	Tyr	Val	Glu	Val	Val	Ser	Ser	Glu	Arg	Glu	Met	Ile	Lys
			180					185					190		
Arg	Phe	Leu	Arg	Ile	Ile	Arg	Glu	Lys	Asp	Pro	Asp	Ile	Ile	Val	Thr
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Tyr	Asn	Gly	Asp	Ser	Phe	Asp	Phe	Pro	Tyr	Leu	Ala	Lys	Arg	Ala	Glu
	210					215					220				
Lys	Leu	Gly	Ile	Lys	Leu	Thr	Ile	Gly	Arg	Asp	Gly	Ser	Glu	Pro	Lys
	225				230					235					240
Met	Gln	Arg	Ile	Gly	Asp	Met	Thr	Ala	Val	Glu	Val	Lys	Gly	Arg	Ile
				245					250					255	
His	Phe	Asp	Leu	Tyr	His	Val	Ile	Thr	Arg	Thr	Ile	Asn	Leu	Pro	Thr
		260						265					270		
Tyr	Thr	Leu	Glu	Ala	Val	Tyr	Glu	Ala	Ile	Phe	Gly	Lys	Pro	Lys	Glu
		275					280					285			
Lys	Val	Tyr	Ala	Asp	Glu	Ile	Ala	Lys	Ala	Trp	Glu	Ser	Gly	Glu	Asn
	290					295					300				
Leu	Glu	Arg	Val	Ala	Lys	Tyr	Ser	Met	Glu	Asp	Ala	Lys	Ala	Thr	Tyr
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Glu	Leu	Gly	Lys	Glu	Phe	Leu	Pro	Met	Glu	Ile	Gln	Leu	Ser	Arg	Leu
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Val	Gly	Gln	Pro	Leu	Trp	Asp	Val	Ser	Arg	Ser	Ser	Thr	Gly	Asn	Leu
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Val	Glu	Trp	Phe	Leu	Leu	Arg	Lys	Ala	Tyr	Glu	Arg	Asn	Glu	Val	Ala
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Pro	Asn	Lys	Pro	Ser	Glu	Glu	Glu	Tyr	Gln	Arg	Arg	Leu	Arg	Glu	Ser
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Tyr	Thr	Gly	Gly	Phe	Val	Lys	Glu	Pro	Glu	Lys	Gly	Leu	Trp	Glu	Asn
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Ile	Val	Tyr	Leu	Asp	Phe	Arg	Ala	Leu	Tyr	Pro	Ser	Ile	Ile	Ile	Thr
			405						410					415	
His	Asn	Val	Ser	Pro	Asp	Thr	Leu	Asn	Leu	Glu	Gly	Cys	Lys	Asn	Tyr
		420						425					430		
Asp	Ile	Ala	Pro	Gln	Val	Gly	His	Lys	Phe	Cys	Lys	Asp	Ile	Pro	Gly
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Phe	Ile	Pro	Ser	Leu	Leu	Gly	His	Leu	Leu	Glu	Glu	Arg	Gln	Lys	Ile
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Lys	Thr	Lys	Met	Lys	Glu	Thr	Gln	Asp	Pro	Ile	Glu	Lys	Ile	Leu	Leu
	465				470					475					480
Asp	Tyr	Arg	Gln	Lys	Ala	Ile	Lys	Leu	Leu	Ala	Asn	Ser	Phe	Tyr	Gly
			485					490						495	
Tyr	Tyr	Gly	Tyr	Ala	Lys	Ala	Arg	Trp	Tyr	Cys	Lys	Glu	Cys	Ala	Glu
			500					505					510		
Ser	Val	Thr	Ala	Trp	Gly	Arg	Lys	Tyr	Ile	Glu	Leu	Val	Trp	Lys	Glu
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<400> SEQUENCE: 5

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ttcgag                                           66

<210> SEQ ID NO 6
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Thr Phe Arg Pro Tyr Ile Tyr Ala Leu Leu Arg Asp Asp Ser Lys Ile
 35         40         45
Glu Glu Val Lys Lys Ile Thr Gly Glu Arg His Gly Lys Ile Val Arg
 50         55         60
Ile Val Asp Val Glu Lys Val Glu Lys Lys Phe Leu Gly Lys Pro Ile
 65         70         75         80
Thr Val Trp Lys Leu Tyr Leu Glu His Pro Gln Asp Val Pro Thr Ile
 85         90         95
Arg Glu Lys Val Arg Glu His Pro Ala Val Val Asp Ile Phe Glu Tyr
 100        105        110
Asp Ile Pro Phe Ala Lys Arg Tyr Leu Ile Asp Lys Gly Leu Ile Pro
 115        120        125
Met Glu Gly Glu Glu Glu Leu Lys Ile Leu Ala Phe Asp Ile Glu Thr
 130        135        140
Leu Tyr His Glu Gly Glu Glu Phe Gly Lys Gly Pro Ile Ile Met Ile
 145        150        155        160
Ser Tyr Ala Asp Glu Asn Glu Ala Lys Val Ile Thr Trp Lys Asn Ile
 165        170        175
Asp Leu Pro Tyr Val Glu Val Val Ser Ser Glu Arg Glu Met Ile Lys
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Arg Phe Leu Arg Ile Ile Arg Glu Lys Asp Pro Asp Ile Ile Val Thr
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Pro Asn Lys Pro Ser Glu Glu Glu Tyr Gln Arg Arg Leu Arg Glu Ser
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385 390 395 400
Ile Val Tyr Leu Asp Phe Arg Ala Leu Tyr Pro Ser Ile Ile Ile Thr
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500 505 510
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Glu Leu Glu Tyr Glu Gly Phe Tyr Lys Arg Gly Phe Phe Val Thr Lys
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595 600 605
Leu Glu Ile Val Arg Arg Asp Trp Ser Glu Ile Ala Lys Glu Thr Gln

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Tyr	Asn	Gly	Asp	Ser	Phe	Asp	Phe	Pro	Tyr	Leu	Ala	Lys	Arg	Ala	Glu
	210					215					220				
Lys	Leu	Gly	Ile	Lys	Leu	Thr	Ile	Gly	Arg	Asp	Gly	Ser	Glu	Pro	Lys
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His	Phe	Asp	Leu	Tyr	His	Val	Ile	Thr	Arg	Thr	Ile	Asn	Leu	Pro	Thr
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Val	Glu	Trp	Phe	Leu	Leu	Arg	Lys	Ala	Tyr	Glu	Arg	Asn	Glu	Val	Ala
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Pro	Asn	Lys	Pro	Ser	Glu	Glu	Glu	Tyr	Gln	Arg	Arg	Leu	Arg	Glu	Ser
	370					375					380				
Tyr	Thr	Gly	Gly	Phe	Val	Lys	Glu	Pro	Glu	Lys	Gly	Leu	Trp	Glu	Asn
385					390					395					400
Ile	Val	Tyr	Leu	Asp	Phe	Arg	Ala	Leu	Tyr	Pro	Ser	Ile	Ile	Ile	Thr
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His	Asn	Val	Ser	Pro	Asp	Thr	Leu	Asn	Leu	Glu	Gly	Cys	Lys	Asn	Tyr
			420						425				430		
Asp	Ile	Ala	Pro	Gln	Val	Gly	His	Lys	Phe	Cys	Lys	Asp	Ile	Pro	Gly
		435					440						445		
Phe	Ile	Pro	Ser	Leu	Leu	Gly	His	Leu	Leu	Glu	Glu	Arg	Gln	Lys	Ile
	450					455					460				
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465					470					475					480
Asp	Tyr	Arg	Gln	Lys	Ala	Ile	Lys	Leu	Leu	Ala	Asn	Ser	Phe	Tyr	Gly
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Tyr	Tyr	Gly	Tyr	Ala	Lys	Ala	Arg	Trp	Tyr	Cys	Lys	Glu	Cys	Ala	Glu
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Ser	Val	Thr	Ala	Trp	Gly	Arg	Lys	Tyr	Ile	Glu	Leu	Val	Trp	Lys	Glu
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Leu	Glu	Glu	Lys	Phe	Gly	Phe	Lys	Val	Leu	Tyr	Ile	Asp	Thr	Asp	Gly
	530					535					540				
Leu	Tyr	Ala	Thr	Ile	Pro	Gly	Gly	Glu	Ser	Glu	Glu	Ile	Lys	Lys	Lys
545					550					555					560
Ala	Leu	Glu	Phe	Val	Lys	Tyr	Ile	Asn	Ser	Lys	Leu	Pro	Gly	Leu	Leu
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Glu	Leu	Glu	Tyr	Glu	Gly	Phe	Tyr	Lys	Arg	Gly	Phe	Phe	Val	Thr	Lys
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595 600 605

Leu Glu Ile Val Arg Arg Asp Trp Ser Glu Ile Ala Lys Glu Thr Gln
610 615 620

Ala Arg Val Leu Glu Thr Ile Leu Lys His Gly Asp Val Glu Glu Ala
625 630 635 640

Val Arg Ile Val Lys Glu Val Ile Gln Lys Leu Ala Asn Tyr Glu Ile
645 650 655

Pro Pro Glu Lys Leu Ala Ile Tyr Glu Gln Ile Thr Arg Pro Leu His
660 665 670

Glu Tyr Lys Ala Ile Gly Pro His Val Ala Val Ala Lys Lys Leu Ala
675 680 685

Ala Lys Gly Val Lys Ile Lys Pro Gly Met Val Ile Gly Tyr Ile Val
690 695 700

Leu Arg Gly Asp Gly Pro Ile Ser Asn Arg Ala Ile Pro Ala Glu Glu
705 710 715 720

Tyr Asp Pro Lys Lys His Lys Tyr Asp Ala Glu Tyr Tyr Ile Glu Asn
725 730 735

Gln Val Leu Pro Ala Val Leu Arg Ile Leu Glu Gly Phe Gly Tyr Arg
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Trp Leu Asn Ile Lys Lys Ser
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 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<400> SEQUENCE: 8

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<210> SEQ ID NO 9
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 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 DNA

<400> SEQUENCE: 9

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<210> SEQ ID NO 10
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 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 DNA

<400> SEQUENCE: 10

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      DNA

<400> SEQUENCE: 11

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40

1-2. (canceled)

3. A method for predicting a functional site of a protein derived from the entire putative proteins of an organism "a" of which genome data or cDNA data is known, which method comprises:

- (1) determining the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, in the amino acid sequences of the entire proteins of the organism "a",
- (2) as to an appropriate protein of the organism "a",
- (2') determining in the entire proteins of the organism "a", the frequency of occurrence of the following Aji-oligo-peptide of given length of (n) ($1 \leq n \leq M$, provided that M is the smallest length of oligopeptides satisfying the criterion that all the oligopeptides of length M are at frequency 1 of the occurrence), which Aji-oligo-peptide is a part of the amino acid sequence of the protein and contains the j-th amino-acid residue Aj ($n \leq j \leq L-n+1$), on condition that the j-th amino-acid residue Aj is defined as the i-th residue Aji from the N-terminus of the Aji-oligo-peptide, wherein
 Aji-oligo-peptide: $a_1 a_2 \dots a_j \dots a_n$, wherein $1 \leq i \leq n+1$;
 Aj = aji, and
 Aj is the i-th residue of the oligopeptide;
 $a_j = A_{j-i+n}$, and
 determining in the entire proteins of the organism "a", the frequency of occurrence of the following Xji-oligo-peptide of length (n) corresponding to the length of Aji-oligo-peptide, wherein
 Xji-oligo-peptide: $a_1 a_2 \dots X_{ji} \dots a_n$, wherein the i-th residue Xji is any amino-acid residue, and $a_1 = A_{j-i+1}, \dots, a_n = A_{j-i+n}$;
- (3) calculating ratio Yji of the frequency of occurrence of the Aji-oligo-peptide to that of the Xji-oligo-peptide;
- (4) determining mean Y(j,n) of the Yji, wherein

$$Y(j, n) = \sum_{i=1}^n Y_{ji}/n,$$

- (5) determining functional value Z(j,n) of Y(j,n), wherein
 $Z(j,n) = -\log(Y(j,n))$;
- (6) repeating steps (2') to (5) sequentially and determining the Z(j,n) value of each amino-acid residue Aj at position j ($n \leq j \leq L-n+1$) in the amino acid sequence (length of L);

- (7) repeating steps (2) to (6) sequentially for the entire proteins of the organism "a", thereby determining the distribution of the Z(j,n) value of each amino-acid residue in the entire proteins, and the Z(j,n) values are classified into twenty according to the twenty amino acids, and then determining mean Av(Aa) of the Z(j,n) values for each amino acid Aa and the standard deviation Sd (Aa) of the distribution thereof, on the basis of the distribution, to determine function g to the j -th amino-acid residue Aj of a protein for normalizing the difference in distribution due to the species of amino-acid residues, wherein

$$g = g(Z(j,n), Aj) = [Z(j,n) - Av(Aa)] / Sd(Aa),$$

provided that Aj Aa; and $1 \leq n \leq M$;

- (8) determining value D(j,n) of the function g of each Aj of all the amino-acid residues at position j ($n \leq j \leq L-n+1$ and $1 \leq n \leq M$) of a protein in the entire proteins as recovered in the step (7), wherein

$$D(j,n) = g(Z(j,n), Aj);$$

and

- (9) defining the representative value of the function of the j-th amino-acid residue in the amino acid sequence (length of L) as functional value Wj of the Z(j,n) and D(j,n), wherein

$$W_j = h(Z(j,1), Z(j,2), \dots, Z(j,M), D(j,1), D(j,2), \dots, D(j,M)), \text{ and}$$

thereby predicting the degree of the involvement of each amino-acid residue in the function of the protein by using the dimension of the Wj value as an indicator.

4. The method according to claim 3, wherein the Wj value of each amino-acid residue is expressed in a two-dimensional distribution chart.
5. The method according to claim 3, wherein the Wj value of each amino-acid residue is expressed on a three-dimensional structure model of the protein.

6-7. (canceled)

8. A system for automatically conducting the method according to claim 3, comprising;

- (a) an outer memory unit for memorizing the amino acid sequence data of the entire putative proteins of organism "a" of which genome data or cDNA data is known, as well as an existing protein data base;
- (b) a first calculation/memory unit, for calculating in the amino acid sequences of the entire proteins of the organism "a", the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopep-

- tides produced by permutations of twenty amino acids, and a memory unit for storing the calculation results;
- (c) a second calculation/memory unit for determining, in the entire proteins of the organism "a", the frequency of occurrence of the following Aji-oligopeptide of given length of (n) ($1 \leq n \leq M$, provided that M is the smallest length of oligopeptides satisfying the criterion that all the oligopeptides of length M are at frequency 1 of the occurrence), which Aji-oligopeptide is a part of the amino acid sequence of a given protein of the organism "a" and contains the j-th amino-acid residue Aj, on condition that the j-th amino-acid residue Aj is defined as the i-th residue Aji from the N-terminus of the Aji-oligopeptide ($n \leq j \leq L-n+1$), wherein
Aji-oligopeptide: $aj_1aj_2 \dots a_{j-1} \dots a_j \dots a_{j+1} \dots a_{j+n}$, wherein
 $1 \leq n+1$;
 $A_j = a_{j_i}$,
Aj is the i-th residue of the oligopeptide,
 $aj_1 = A_j - i + 1, \dots, aj_n = A_j - i + n$, and
determining in the entire proteins of the organism "a", the frequency of occurrence of the following Xji-oligopeptide of length (n) corresponding to the length of Aji-oligopeptide, wherein
Xji-oligopeptide: $aj_1aj_2 \dots X_{j_i} \dots a_{j+n}$, wherein
the i-th residue Xji is any amino acid,
 $aj_1 = A_j - i + 1, \dots,$
 $aj_n = A_j - i + n$, and
a memory unit for storing the calculation results;
- (d) a third calculation/memory unit for calculating ratio Yji of the frequency of occurrence of the Aji-oligopeptide to that of the Xji-oligopeptide, and a memory unit for storing the calculation results for ratio Yji;
- (e) a fourth calculation/memory unit for determining mean Y(j,n) of the Yji, wherein

$$Y(j, n) = \sum_{i=1}^n Y_{j_i} / n,$$

and a memory unit for storing the calculation results for mean Y(j,n),

- (f) a fifth calculation/memory unit for determining functional value Z(j,n) of Y(j,n), wherein

$$Z(j, n) = -\log(Y(j, n)),$$

and a memory unit for storing the calculation results of Z(j,n);

- (g) a sixth calculation/memory unit for determining the Z(j,n) value of each amino-acid residue in the amino acid sequences of the entire proteins of the organism "a", determining the distribution of the Z(j,n) value of each amino-acid residue in the entire proteins, and the Z(j,n) values are classified into twenty according to the twenty amino acids, and then determining mean Av(Aa) of the Z(j,n) values for each amino acid Aa and the standard deviation Sd(Aa) of the distribution thereof, on the basis of the distribution, to determine function g for normalizing the difference in distribution due to the species of amino-acid residues, wherein

$$g = g(Z(j, n), A_j) = [Z(j, n) - Av(Aa)] / Sd(Aa), \text{ wherein}$$

$A_j = Aa$; and $1 \leq n \leq M$,

and a memory unit for storing g,

- (h) a seventh calculation/memory unit for determining value D(j,n) of function g stored in the memory unit in step (g) concerning each of all the amino-acid residues Aj at position j ($n \leq j \leq L-n+1$) in the amino acid sequence (length of L), wherein

$$D(j, n) = g(Z(j, n), A_j),$$

and a memory unit for storing the calculation of the D(j, n) value; and

- (i) an eighth calculation/memory unit for determining appropriate functional value Wj of the Z(j,n) and D(j,n) of each amino-acid residue in the amino acid sequence, wherein

$$W_j = h(Z(j, 1), Z(j, 2), \dots, Z(j, M), D(j, 1), D(j, 2), \dots, D(j, M))$$

and a memory unit for storing the calculation of the Wj value.

9. The system according to claim 8, the system being equipped with a display unit displaying the Wj value of each amino-acid residue in a two-dimensional distribution chart.

10. The system according to claim 8, wherein the system is equipped with a calculation/memory unit for memorizing an existing database of three-dimensional structure of proteins or preparing a three-dimensional structure model based on an amino acid sequence according to a known method and memorizing the three-dimensional structure model, and a display unit displaying the Wj value of each amino-acid residue in the amino acid sequence in a distribution chart on the three-dimensional structure stored in the data base or three-dimensional structure model memorized in the calculation/memory unit.

11. A method for modifying the known function of protein "A" derived from the entire proteins of organism "a" of which genome data or cDNA data has been known, which method comprises:

- (1) extracting a protein closely related to the protein "A" from an existing protein data base and subjecting the proteins to alignment;
- (2) determining, in the amino acid sequences of the entire proteins of the organism "a", the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, and determining the smallest length (n) of oligopeptides satisfying the following criteria:

among oligopeptides of length (n), the number of oligopeptides which occur in the entire proteins is smaller than the number of oligopeptides which occur twice in the entire proteins; among oligopeptides of length (n+1), the number of oligopeptides which occur once in the entire proteins is larger than the number of oligopeptides which occur twice in the entire proteins;

- (3) determining, in the entire proteins of the organism "a", the frequency of occurrence of the following Aji-oligopeptide of length (n+1), which is a part of the amino acid sequence of the protein as a subject for predicting a functional site and contains the j-th amino-acid residue Aj from the amino terminal end (N-terminus) of the amino acid sequence (length of L) of the protein, on condition that the j-th amino-acid residue Aj is defined as, or is identical to, the i-th residue Aji from the N-terminus of the Aji-oligopeptide ($n+1 \leq j \leq L-n$), wherein Aji-oligopeptide: $aj_1aj_2 \dots a_{j-1} \dots a_{j+n}$, wherein

- $1 \leq i \leq n+1$;
 $A_j = A_{ji}$,
 A_j is the i -th residue of the oligopeptide, and
 $a_{j1} = A_{j-i+1}, \dots, a_{j(n+1)} = A_{j-i+(n+1)}$,
 and determining in the entire proteins of the organism "a", the frequency of occurrence of the following X_{ji} -oligopeptide of length $(n+1)$, wherein
 X_{ji} -oligopeptide: $a_{j1}a_{j2} \dots X_{ji} \dots a_{jn}a_{j(n+1)}$, wherein
 $n+1 \leq j \leq L-n$; and
 the i -th residue X_{ji} is any amino acid, and
 $a_{j1} = A_{j-i+1}, \dots, a_{j(n+1)} = A_{j-i+(n+1)}$;
 (4) calculating ratio Y_{ji} of the frequency of occurrence of the A_{ji} -oligopeptide to that of the X_{ji} -oligopeptide;
 (5) determining mean Y_j of the Y_{ji} , wherein

$$Y_j = \sum_{i=1}^{n+1} Y_{ji} / (n+1),$$

- (6) determining functional value Z_j of Y_j , wherein

$$Z_j = f(Y_j), \text{ and}$$

Function f is a monotonously decreasing function or a monotonously increasing function, and defining the Z_j value as the representative value of the function of the j -th amino-acid residue of the amino acid sequence (length of L) of the protein "A";

- (7) sequentially repeating steps (3) to (6) and determining the Z_j value of each of all the amino-acid residues at position between $n+1 \leq j \leq L-n$ in the amino acid sequence (length of L);
 (8) selecting at least one amino-acid residue to be subjected to mutation on the basis of the alignment data carried out in step (1) from the amino acid sequence (length of L) of the protein "A", sequentially repeating steps (3) to (6) for variant amino-acid residues in various mutated amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residue, to determine the Z_j value of the variant amino-acid residues;
 (9) selecting a mutated amino acid sequence wherein the Z_j value of the variant amino-acid residue as determined in step (8) is larger or smaller than the Z_j value of the wild type amino-acid residue as determined in step (7); and
 (10) preparing a modified gene encoding the modified amino acid sequence from the protein "A" gene, and producing the modified protein as the expression product of the gene.

12. A method for modifying the function of protein "B" derived from an organism "b" of which genome data or cDNA data has been unknown, which method comprises:

- (1) extracting protein "A" which is most closely related to protein "B" from the entire proteins of organism "a" of which genome data or cDNA data being known and subjecting the protein to alignment, or extracting a protein closely related to protein "B" from an existing protein data base to subject the protein to alignment;
 (2) determining, in the amino acid sequences of the entire proteins of the organism "a", the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permuta-

tions of twenty amino acids, and determining the smallest length (n) of oligopeptides satisfying the following criteria

among oligopeptides of length (n), the number of oligopeptides which occur once in the entire proteins is smaller than the number of oligopeptides which occur twice in the entire proteins; among oligopeptides of length $(n+1)$, the number of oligopeptides which occur once in the entire proteins is larger than the number of oligopeptides which occur twice in the entire proteins;

- (3) determining, in the entire proteins of the organism "a", the frequency of occurrence of the following A_{ji} -oligopeptide of length $(n+1)$, which is a part of the amino acid sequence of the protein as a subject for predicting a functional site and contains the j -th amino-acid residue A_j from the amino terminal end (N-terminus) of the amino acid sequence (length of L) of the protein, on condition that the j -th amino-acid residue A_j is defined as the i -th residue A_{ji} from the N-terminus of the A_{ji} -oligopeptide $(n+1 \leq j \leq L-n)$, wherein
 A_{ji} -oligopeptide: $a_{j1}a_{j2} \dots A_{ji} \dots a_{jn}a_{j(n+1)}$, wherein
 $1 \leq i \leq n+1$,
 $A_j = A_{ji}$, and
 A_j is the i -th residue of the oligopeptide,
 $a_{j1} = A_{j-i+1}, \dots, a_{j(n+1)} = A_{j-i+(n+1)}$,
 and determining, in the entire proteins of the organism "a", the frequency of occurrence of the following X_{ji} -oligopeptide of length $(n+1)$, wherein
 X_{ji} -oligopeptide: $a_{j1}a_{j2} \dots X_{ji} \dots a_{jn}a_{j(n+1)}$, wherein
 $1 \leq i \leq n+1$,
 $n+1 \leq j \leq L-n$, and
 the i -th residue X_{ji} is any amino acid, and
 $a_{j1} = A_{j-i+1}, \dots, a_{j(n+1)} = A_{j-i+(n+1)}$;

- (4) calculating ratio Y_{ji} of the frequency of occurrence of the A_{ji} -oligopeptide to that of the X_{ji} -oligopeptide;
 (6) determining mean Y_j of the Y_{ji} , wherein

$$Y_j = \sum_{i=1}^{n+1} Y_{ji} / (n+1),$$

- (6) determining functional value Z_j of Y_j , wherein

$$Z_j = f(Y_j)$$

Function f is a monotonously decreasing function or a monotonously increasing function, and defining the Z_j value as the representative value of the function of the j -th amino-acid residue A_j of the amino acid sequence (length of L) of the protein "A";

- (7) sequentially repeating steps (3) to (6) and determining the Z_j value of each of all the amino-acid residues at position between $n+1 \leq j \leq L-n$ in the amino acid sequence (length of L);
 (8) selecting at least one amino-acid residue to be subjected to mutation on the basis of the alignment data carried out in step (1) from the amino acid sequence (length of L) of the protein "A", sequentially repeating steps (3) and (6) for variant amino-acid residues in various mutated amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residues, to determine the Z_j value of the variant amino-acid residues;

- (9) selecting the mutation position and the mutated amino-acid residue wherein the Z_j value of the variant amino-acid residue as determined in the step (8) is larger or smaller than the Z_j value of the wild type amino-acid residue as determined in the step (7); and
- (10) preparing a modified gene encoding the modified amino acid sequence having the mutated amino-acid residue at the position from the protein "B" gene, and producing the modified protein as the expression product of the gene.

13. A method for modifying the known function of protein "A" derived from the entire proteins of organism "a" of which genome data or cDNA data has been known, which method comprises:

- (1) extracting proteins closely related to the protein "A" from an existing protein data base and subjecting the proteins to alignment;
- (2) determining the frequency of occurrence of each amino acid and the frequency of occurrence of individual oligopeptides produced by permutations of twenty amino acids, in the amino acid sequences of the entire proteins of the organism "a";
- (3) as to an appropriate protein of the organism "a";
- (3') determining, in the entire proteins of the organism "a", the frequency of occurrence of the following A_{ji} -oligopeptide of given length of (n) ($1 \leq n \leq M$, provided that M is the smallest length of oligopeptides satisfying the criterion that all the oligopeptides of length M are at frequency 1 of the occurrence), which A_{ji} -oligopeptide is apart of the amino acid sequence of the protein and contains the j-th amino-acid residue A_j , on condition that the j-th amino-acid residue A_j is defined as the i-th residue A_{ji} from the N-terminus of the A_{ji} -oligopeptide ($n \leq j \leq L-n+1$), wherein

A_{ji} -oligopeptide: $a_{j1}a_{j2} \dots a_{ji} \dots a_{jn}$, wherein

$$1 \leq i \leq n,$$

$$A_j = a_{ji}, \text{ and}$$

A_j is the i-th residue of the oligopeptide,

$$a_{j1} = A_{j-i+1}, \dots,$$

$$a_{jn} = A_{j-i+n},$$

and determining in the entire proteins of the organism "a", the frequency of occurrence of the following X_{ji} -oligopeptide of length (n) corresponding to the length of A_{ji} -oligopeptide, wherein

X_{ji} -oligopeptide: $a_{j1}a_{j2} \dots X_{ji} \dots a_{jn}$, wherein

$$1 \leq i \leq n, \text{ and}$$

the i-th residue X_{ji} is any amino acid,

$$a_{j1} = A_{j-i+1}, \dots, \text{ and}$$

$$a_{jn} = A_{j-i+n};$$

- (4) calculating ratio Y_{ji} of the frequency of occurrence of the A_{ji} -oligopeptide to that of the X_{ji} -oligopeptide;
- (5) determining mean $Y(j,n)$ of the Y_{ji} , wherein

$$Y(j, n) = \sum_{i=1}^n Y_{ji} / n,$$

- (6) determining functional value $Z(j,n)$ of $Y(j,n)$, wherein

$$Z(j,n) = -\log(Y(j,n));$$

- (7) repeating steps (3') to (6) sequentially and determining the $Z(j,n)$ value of each amino-acid residue A_j at position j ($n \leq j \leq L-n+1$) in the amino acid sequence (length of L);

- (8) repeating steps (3) to (7) sequentially for the entire proteins of the organism "a", thereby determining the distribution of the $Z(j,n)$ value of each amino-acid residue in the entire proteins, and the $Z(j,n)$ values are classified into twenty according to the twenty amino acids, and then determining mean $Av(Aa)$ of the $Z(j,n)$ values for each amino acid Aa and the standard deviation $Sd(Aa)$ of the distribution thereof, on the basis of the distribution, to determine function g to the j-th amino-acid residue A_j of a protein for normalizing the difference in distribution due to the species of amino-acid residues, wherein

$$g = g(Z(j,n), A_j) = [Z(j,n) - Av(Aa)] / Sd(Aa)$$

provided that $A_j = Aa$, and $1 \leq n \leq M$;

- (9) determining value $D(j,n)$ of the function g of each A_j of all the amino-acid residues at position j ($n \leq j \leq L-n+1$ and $1 \leq n \leq M$) of a protein in the entire proteins as recovered in step (8), wherein

$$D(j,n) = g(Z(j,n), A_j);$$

- (10) defining the representative value of the function of the j-th amino-acid residue in the amino acid sequence (length of L) as functional value W_j of the $Z(j,n)$ and $D(j,n)$, wherein

$$W_j = h(Z(j,1), Z(j,2), \dots, Z(j,M), D(j,1), D(j,2), \dots, D(j,M));$$

- (11) repeating steps (3) to (10) sequentially, to determine the individual W_j values of all the amino-acid residues at position $n \leq j \leq L-n+1$ in the amino acid sequence (length of L); (12) selecting at least one amino-acid residue to be subjected to mutation on the basis of the alignment data carried out in step (1) from the amino acid sequence (length of L) of the protein "A", and repeating steps (3) and (10) sequentially for variant amino-acid residues in various mutated amino acid sequences where the selected amino-acid residue has been mutated into another amino-acid residue, to determine the W_j value of the variant amino-acid residue;

- (13) selecting a mutated amino acid sequence wherein the W_j value of the variant amino-acid residue as determined in step (12) is larger or smaller than the W_j value of the wild type amino-acid residue as determined in step (10); and

- (14) preparing a modified gene encoding the modified amino acid sequence from the protein "A" gene, and producing the modified protein as the expression product of the gene.

14. A protein of which function is artificially modified by the method according to claim 11, 12 or 13.

15. The protein according to claim 14, which is a thermophilic DNA polymerase prepared by artificially modifying the amino acid sequence of Pfu DNA polymerase so that the elongation of synthesized DNA chain might not be terminated intermediately during the catalysis for the synthesis of a DNA chain complementary to a single-stranded DNA.

16. The thermophilic DNA polymerase according to claim 15, comprising the amino acid sequence of SEQ ID No.1.

17. The protein according to claim 14, which is a thermophilic DNA polymerase prepared by artificially modifying

the amino acid sequence of Pfu DNA polymerase so that the synthesized DNA chain might be more elongated during the catalysis for the synthesis of a DNA chain complimentary to a single-stranded DNA.

18. The thermophilic DNA polymerase according to claim **17**, comprising the amino acid sequence of SEQ ID No.6.

19. The thermophilic DNA polymerase according to claim **17**, comprising the amino acid sequence of SEQ ID. No.7.

20. A DNA sequence encoding the amino acid sequence of SEQ ID No.1.

21. A DNA sequence encoding the amino acid sequence of SEQ ID No. 6.

22. A DNA sequence encoding the amino acid sequence of SEQ ID No.7.

23. A recombinant vector carrying the DNA sequence according to claim **20**.

24. The recombinant vector according to claim **23**, which is a recombinant plasmid pDP320 carried on *Escherichia coli* HMS 174(DE3)/pDP 320 (FERM P-16052).

25. A recombinant vector carrying the DNA sequence according to claim **21**.

26. The recombinant vector according to claim **25**, which is a recombinant plasmid pDP 5b17 carried on *Escherichia coli* HMS 174(DE3)/pDP 5b17 (FERM BP-6189).

27. A recombinant vector carrying the DNA sequence according to claim **22**.

28. The recombinant vector according to claim **27**, which is a recombinant plasmid pDP 5c4 carried on *Escherichia coli* HMS 174(DE3)/pDP 5c4(FERM BP-6190).

29. A method for preparing a thermophilic DNA polymerase, comprising culturing a cell transformed with an expression vector carrying the DNA sequence according to claim **20** and isolating and purifying the objective enzyme produced in the culture medium.

30. A method for preparing a DNA polymerase, comprising culturing a cell transformed with an expression vector carrying the DNA sequence according to claim **21** or **22** and isolating and purifying the objective enzyme produced in the culture medium.

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