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(54) Titre : SYSTÈME DE TRAITEMENT DE L'INFORMATION UTILISANT DE L'INFORMATION RELIÉE À DES
SEQUENCES NUCLEOTIDIQUES

(54) Title: INFORMATION PROCESSING SYSTEM USING NUCLEOTIDE SEQUENCE RELATED INFORMATION

(57) **Abrégé/Abstract:**

Information on a nucleotide sequence is processed by receiving request information of an object or service. A first processor searches and retrieves from a first memory area positional information corresponding to information on classification of the object or service. The positional information represents a position in a nucleotide sequence. The retrieved positional information is transmitted to a second processor which accesses a second memory area storing polymorphism information regarding the individual. Polymorphism information is received based on the transmitted positional information. The received polymorphism information is output to a user. The user includes a third processor which accesses a third memory area. The third memory area is searched to retrieve, semantic information based on the transmitted polymorphism information. The semantic information is output to a device that utilizes the semantic information or information corresponding to the semantic information for a provision of the object or service.



ABSTRACT

Information on a nucleotide sequence is processed by receiving request information of an object or service. A first processor searches and retrieves from a first
5 memory area positional information corresponding to information on classification of the object or service. The positional information represents a position in a nucleotide sequence. The retrieved positional information is transmitted to a second processor which accesses a second memory area storing polymorphism information regarding the individual. Polymorphism information is received based on the transmitted positional
10 information. The received polymorphism information is output to a user. The user includes a third processor which accesses a third memory area. The third memory area is searched to retrieve, semantic information based on the transmitted polymorphism information. The semantic information is output to a device that utilizes the semantic information or information corresponding to the semantic information for a provision of
15 the object or service.

INFORMATION PROCESSING SYSTEM USING NUCLEOTIDE SEQUENCE RELATED INFORMATION

This is a division of co-pending Canadian Patent Application No. 2,387,277 filed
5 May 23, 2002.

FIELD OF THE INVENTION

The present invention relates to an information processing system that provides
information through a communication network.

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BACKGROUND TECHNIQUE

Currently, genomic nucleotide sequences of various organisms including humans
are being rapidly determined and information on genomic nucleotide sequences is being
accumulated in various databases. For example, currently in progress is the construction
15 of a system which will enable various research institutes and researchers to utilize
information on genomic nucleotide sequences accumulated in databases through an
information network such as the Internet.

At the same time, research for the purpose of genomic drug discovery and
analysis of genetic information and the like have been actively conducted using
20 nucleotide sequences contained in such information on genomic nucleotide sequences,
and differences in nucleotide sequences among individual organisms represented by the
single nucleotide polymorphism are attracting attention. In general, differences in
nucleotide sequences among individual organisms refer to a polymorphism defined by
existence of a predetermined nucleotide difference at a frequency of 1% or more in an
25 individual species and a variation defined by a predetermined nucleotide difference of
less than 1% in an individual species. In particular, known polymorphisms are SNP
(single nucleotide polymorphism), in which there is one nucleotide difference among
individual organisms; an insertion/deletion polymorphism, in which one to several tens
of nucleotides (sometimes several thousands of nucleotides) have been deleted or
30 inserted; VNTR (variable number of tandem repeat), in which the number of repetitions
of a sequence comprising two to several tens of nucleotides as one unit varies; and a

microsatellite polymorphism (a repetition sequence having about two to four nucleotides).

Such polymorphisms sometimes affect, for example, differences in amino acid sequences of proteins among individual organisms or differences in expression efficiency concerning predetermined genes among individual organisms. Such influences cause, for example, differences in the morbidity rate of predetermined diseases among individual organisms or differences in sensitiveness to predetermined medicaments among individual organisms.

A system, however, which provides semantic information useful for each organism among a plurality of individual organisms through effective utilization of differences in nucleotide sequence-related information, such as a polymorphism, is not yet constructed.

Under the above circumstances, the present invention is directed to construction of a system for processing information for providing semantic information and/or information associated with the semantic information useful for each individual organism through effective utilization of differences in nucleotide sequence-related information among individual organisms.

SUMMARY OF THE INVENTION

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Certain exemplary embodiments may provide a computer-implemented method for processing information on a nucleotide sequence comprising the steps of: (a) receiving, via a communication network, request information of an object or service, wherein the object or service is provided for an individual based on one or more genetic differences between individuals, and wherein the request information does not include genetic information; (b) searching a first memory area for positional information, and wherein the positional information corresponds to information on classification of the object or service, and wherein the positional information represents a position in a nucleotide sequence, and retrieving the positional information, wherein the first memory area is permitted to be accessed by a first processor; (c) transmitting, via a communication network, the positional information retrieved in step (b) to a second processor which is permitted to access a second memory area storing polymorphism

information regarding the individual; (d) receiving, via a communication network, polymorphism information based on the positional information transmitted in step (c); and (e) outputting, via a communication network, the polymorphism information received in step (d) to a user, and wherein the user includes a third processor which is permitted to access a third memory area storing semantic information, and which searches the third memory area to retrieve, in response to said transmission of the polymorphism information, semantic information based on the transmitted polymorphism information, and outputs the semantic information to a device that utilizes the semantic information or information corresponding to the semantic information, which is used for a provision of the object or service substantially included in the request information, wherein steps (a)-(e) are conducted under the control of the first processor.

Certain exemplary embodiments may provide a computer-readable medium encoded with a program for processing information on a nucleotide sequence which allows a computer to execute processes including: (a) receiving, via a communication network, request information of an object or service, wherein the object or service is provided for an individual based on one or more genetic differences between individuals, and wherein the request information does not include genetic information; (b) searching a first memory area for positional information, and wherein the positional information corresponds to information on classification of the object or service, and wherein the positional information represents a position in a nucleotide sequence, and retrieving the positional information, wherein the first memory area is permitted to be accessed by a first processor; (c) transmitting, via a communication network, the positional information retrieved in process (b) to a second processor which is permitted to access a second memory area storing polymorphism information regarding the individual; (d) receiving, via a communication network, polymorphism information based on the positional information transmitted in process (c); and (e) outputting, via a communication network, the polymorphism information received in process (d) to a user, and wherein the user includes a third processor which is permitted to access a third memory area storing semantic information, and which searches the third memory area to retrieve, in response to said transmission of the polymorphism information, semantic information based on the transmitted polymorphism information, and outputs the

semantic information to a device that utilizes the semantic information or information corresponding to the semantic information, which is used for a provision of the object or service substantially included in the request information, wherein processes (a)-(e) are conducted under the control of the first processor.

5 Certain exemplary embodiments may provide an apparatus for processing information on a nucleotide sequence comprising: a transmitter/receive in communication with a communication network that receives request information of an object or service via the communication network, the object or service being provided for an individual based on one or more genetic differences between individuals, wherein
10 the request information does not include genetic information; a controller having a first processor that searches a first memory area for positional information that corresponds to information on classification of the object or service, the positional information representing a position in a nucleotide sequence, and retrieves the positional information, wherein the transmitter/receiver transmits, via the communication network,
15 the retrieved positional information to a second processor which is permitted to access a second memory area storing polymorphism information regarding the individual, and the controller, followed by receiving, via the communication network, the polymorphism information based on the transmitted positional information, outputs, via the communication network, the received polymorphism information to a user, and
20 wherein the user includes a third processor which is permitted to access a third memory area storing semantic information, and which searches the third memory area to retrieve, in response to said transmission of the polymorphism information, semantic information based on the transmitted polymorphism information, and outputs the semantic information to a device that utilizes the semantic information or information
25 corresponding to the semantic information, which is used for a provision of the object or service substantially included in the request information.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic view showing a construction of a system for processing information to which the present invention has been applied.

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Fig. 2 is a schematic view showing a construction of a shared computer.

Fig. 3 is a diagram showing construction of an embodiment of data that is recorded in a main database (“database” is hereinafter abbreviated to “DB”).

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Fig. 4 is a schematic view showing a construction of a personal computer.

Fig. 5 is a diagram showing construction of an embodiment of data recorded on a genome-related information recording medium.

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Fig. 6 is a flow chart showing processing in a shared computer and a personal computer in a system for providing a morbidity rate of a predetermined disease.

Fig. 7 is a flow chart, which is a continuation of Fig. 6, showing processing in a shared computer and a personal computer in a system for providing a morbidity rate of a predetermined disease.

Fig. 8 is a flow chart showing other processing in a shared computer and a personal computer in a system for providing a morbidity rate of a predetermined disease.

Fig. 9 is a flow chart showing further processing in a shared computer and a personal computer in a system for providing a morbidity rate of a predetermined disease.

Fig. 10 is a diagram showing construction of an embodiment of data recorded in a main DB that is utilized in a system for providing a medicament that is appropriate for the diathesis.

Fig. 11 is a flow chart showing processing in a shared computer and a personal computer in a system for providing a medicament that is appropriate for the diathesis.

Fig. 12 is a flow chart, which is a continuation of Fig. 11, showing processing in a shared computer and a personal computer in a system for providing a medicament that is appropriate for the diathesis.

Fig. 13 shows a screen image indicated as an example of a portal screen for a tailor-made non-prescription pharmaceuticals.

Fig. 14 shows a screen image indicated as an example of a medicament category menu screen.

Fig. 15 shows a screen image shown as an example of an input screen for individual IC confirmation.

Fig. 16 shows a screen image indicated as an example of a confirmation screen.

Fig. 17 is a diagram showing construction of data stored in memory section A in a system for providing a medicament that is appropriate for the diathesis.

Fig. 18 is a diagram showing construction of data stored in memory section B in a system for providing a medicament that is appropriate for the diathesis.

Fig. 19 is a diagram showing construction of information that is transmitted to a provider of a medicament in a system for providing a medicament that is appropriate for the diathesis.

Fig. 20 is a diagram showing construction of an embodiment of data recorded in a main DB used in a system for providing medical examination items depending on diathesis.

Fig. 21 is a diagram showing construction of an embodiment of data recorded in a link DB used in a system for providing medical examination items depending on the diathesis.

Fig. 22 is a diagram showing construction of an embodiment of data recorded on a genome-related information recording medium used in a system for providing medical examination items depending on the diathesis.

Fig. 23 is a flow chart showing processing in a shared computer and a personal computer in a system for providing medical examination items depending on the diathesis.

Fig. 24 is a flow chart, which is a continuation of Fig. 23, showing processing in a shared computer and a personal computer in a system for providing medical examination items depending on the diathesis.

Fig. 25 shows a screen image indicated as an example of a portal screen for medical examination.

Fig. 26 shows a screen image indicated as an example of a medical examination course menu screen.

Fig. 27 is a diagram showing construction of an embodiment of data recorded in a medical examination table used in a system for providing medical examination items depending on the diathesis.

Fig. 28 shows a screen image indicated as an example of an input screen for individual IC confirmation.

Fig. 29 is a diagram showing construction of data stored in memory section B in a system for providing medical examination items depending on the diathesis.

Fig. 30 is a diagram showing construction of data stored in memory section A in a system for providing medical examination items depending on the diathesis.

Fig. 31 shows a screen image representing an example of an output format in a system for providing medical examination items depending on the diathesis.

Fig. 32 is a diagram showing construction of an embodiment of data recorded in a main DB used in a system for providing information on other individual organisms having properties compatible with a "given property" of the requestor.

Fig. 33 is a diagram showing construction of an embodiment of data recorded on a genome-related information recording medium used in a system for providing information on other individual organisms having properties compatible with a "given property" of the requestor.

Fig. 34 is a flow chart showing processing in a shared computer and a personal computer in a system for donor registration for registering an individual's own polymorphism pattern.

Fig. 35 is a diagram showing construction of a donor DB constructed in a system for donor registration for registering an individual's own polymorphism pattern.

Fig. 36 is a flow chart showing processing in a shared computer and a personal computer in a system for providing information on other individual organisms having properties compatible with a "given property" of the requestor.

Fig. 37 is a flow chart, which is a continuation of Fig. 36, showing processing in a shared computer and a personal computer in a system for providing information on other individual organisms having properties compatible with a "given property" of the requestor.

DESCRIPTION OF REFERENCE NUMERALS

- 1: communication network
- 2: shared computer
- 3: personal computer

EMBODIMENT FOR CARRYING OUT THE INVENTION

The present invention is described in detail below with reference to the drawings.

1. First embodiment

First, a system for processing information which provides the morbidity rate of predetermined diseases to a user is described as a first embodiment to which the present invention has been applied. The present embodiment is directed to explanation of a system for providing the morbidity rate in accordance with the request information requested by a user and, thus, is explained as a simple model for the convenience of explanation.

As shown in Fig. 1, the system for processing information comprises a communication network 1, such as the Internet, a shared computer 2 connected to communication network 1, and a plurality of personal computers 3 connected to communication network 1, and enables data communication between shared computer 2 and personal computers 3 through communication network 1.

As shown in Fig. 2, shared computer 2 is constituted by a CPU 4 that totally controls the operation of the shared computer 2; an input device 5, such as a keyboard and a mouse, with which information, instructions for executing a program and the like can be input; a display device 6 such as a display apparatus; a storage 7 in which temporary information, unrewritable information and the like are recorded; a database 8 for storing various data; a recorder 9 for writing predetermined information in storage 7 and database 8; and a

transmitter/receiver 17 for transmission and reception of information to and from personal computers 3 through communication network 1.

Storage 7 in shared computer 2 is constituted by a memory section A10 and a memory section B11 which respectively record different types of information; a screen memory 12 having recorded therein screen data displayed, for example, on personal computer 3 or display device 6; and a processing program 13 for operating the system. Shared computer 2 may have screen memory 12, processing program 13 and the like in an external recording apparatus (not shown) connected to shared computer 2 through communication network 1 instead of containing those in storage 7 inside shared computer 2.

Database 8 (described as “memory” in Claim) in shared computer 2 is constituted by a main DB 14 in which a polymorphism address, a polymorphism pattern, and semantic information are recorded; a storage DB-A15 for saving information recorded in memory section A10; and a storage DB-B16 for saving information recorded in memory section B11.

As shown in Fig. 3, polymorphism addresses, a plurality of possible polymorphism patterns in the polymorphism address respectively, and semantic information implied by each of the plurality of polymorphism patterns respectively are stored in association with one another in main DB 14. Main DB 14 may also have recorded therein semantic information implied by a combination of polymorphism patterns in a plurality of polymorphism addresses (such as haplotype).

The “polymorphism address (positional information)” refers to, at least, a position in a nucleotide sequence where a polymorphism is present. In general, the term “polymorphism” includes, for example, a so-called SNP (single nucleotide polymorphism), RFLP (restriction fragment length of polymorphism), VNTR (variable number of tandem repeat), and microsatellite. However, the term “polymorphism” used herein is not limited to these and also includes a variation in nucleotides and nucleotide sequences existing only at a frequency of less than 1% in an individual species. Therefore, “polymorphism address” also includes a position in a nucleotide sequence which indicates a variation of a nucleotide and nucleotide sequences existing only at a frequency of less than 1% in an individual species. Specifically, the “polymorphism address” indicates a position representing a polymorphism or the like by a combination of numerical values, letters, symbols, and the like. The polymorphism address is not particularly limited, for example, may be represented by a combination of a chromosome number, a symbol indicating a gene having a polymorphism therein, and a numerical value indicating a position of a polymorphism in the gene.

Alternatively, it may be a combination of a symbol indicating a gene having polymorphism therein and a numerical value indicating a position of polymorphism in the gene.

Further, a “polymorphism address” may be a notation peculiar to a polymorphism imparted to each polymorphism. When the notation peculiar to a polymorphism is used as a polymorphism address, the polymorphism address does not directly indicate the position in the nucleotide sequence, instead, the position can be indirectly found by the notation peculiar to the polymorphism. Therefore, the “polymorphism address” includes the notation peculiar to the polymorphism.

A “polymorphism pattern (nucleotide sequence-related information)” is information on nucleotide sequences which differ among individual organisms, and contains, at least, a pattern of nucleotides or nucleotide sequences in a polymorphism. In addition, the “polymorphism pattern” includes a pattern of nucleotide and nucleotide sequences existing only at a frequency of less than 1% in an individual species and is not limited to a polymorphism. For example, in a polymorphism address known to have A or G, the “polymorphism pattern” is represented either by “A” or “G”.

The “polymorphism pattern” may represent a heterozygote or homozygote in a homologous chromosome. For example, the “polymorphism pattern” can be represented by “AA”, “GG”, or “AG” in the polymorphism address known to have A or G.

Further, the “polymorphism pattern” may indirectly represent a possible pattern in the predetermined polymorphism address instead of direct representation of patterns. For example, in the polymorphism address known to have A or G the “polymorphism pattern” may be represented by “allele 1” when the polymorphism address has “A” or “allele 2” when the polymorphism address has “G”. As described above, when the “polymorphism pattern” can be expressed as “AA”, “GG”, or “AG”, the “polymorphism pattern” may be represented by “ α ” when expressed as “AA”, it may be represented by “ β ” when expressed as “GG”, and it may be represented by “ γ ” when expressed as “AG”.

When the polymorphism is the microsatellite type the “polymorphism pattern” may be represented, for example, by numerical values indicating “the number of repetitions” and when the polymorphism is the insertion/deletion type the “polymorphism pattern” may be represented, for example, by symbols indicating “presence/absence”.

The term “semantic information” used herein refers to information associated with the “polymorphism pattern,” for example, information including responsiveness to medicaments, side-effect caused by medicaments, a risk against diseases and disorders,

diatheses and properties, interaction among proteins, and various phenotypes caused by differences in polymorphism patterns. “Semantic information” is a type of information which is corrected and increases in the numbers of types accompanied by progress in research on genome and genetics, and constant updating is preferred. In other words, “semantic information” becomes more accurate through increases and decreases in the amount of information accumulated by updating a database using the results of research on genome and genetics.

Information that is further induced from “semantic information” is “information associated with the semantic information” although it is not directly associated with the “polymorphism pattern.” When “semantic information” is a risk against diseases, when the relevant risk exceeds a given standard, for example, specific “medical examination items” are derived. These specific “medical examination items” are “information associated with the semantic information.”

In the present embodiment, semantic information is recorded in main DB 14 as “annotative information on the polymorphism pattern” associated with at least the predetermined “polymorphism address” and “polymorphism pattern” as shown in Fig. 3. Also, semantic information is associated with, for example, “polymorphism classification,” “classification (name of diseases)” and the like corresponding to the predetermined “polymorphism address.” Consequently, when a predetermined “polymorphism address” is a predetermined “polymorphism pattern,” types of diseases and annotative information (semantic information) on the morbidity rates of diseases can be obtained. For example, semantic information can be associated with a combination of respective polymorphism patterns corresponding to a plurality of polymorphism addresses (such as haplotype). In other words, each combination of polymorphism patterns in a plurality of polymorphism addresses can be respectively associated with annotative information (semantic information) representing different morbidity rates for predetermined diseases. In this case, when a plurality of polymorphism addresses are a combination of predetermined polymorphism patterns, annotative information (semantic information) indicating the morbidity rate of a predetermined disease can be obtained.

Semantic information can be further associated with a “level of disclosure” which is set in accordance with a predetermined standard. For example, a standard in setting a “level of disclosure” can be determined by taking into consideration unpredictable disbenefits and the like for individuals that would be caused by disclosure of semantic information, i.e., the morbidity rate of “classification (name of disease)”. In particular, in shared computer 2, a

“level of disclosure” can be set such that semantic information, the disclosure of which is inappropriate from the view point of, for example, law, regulations, the behavioral norms of an organization having the shared computer 2 or a contract with the user, is not disclosed. In this case, with this system, annotative information representing a morbidity rate associated with a “level of disclosure” at which disclosure is not possible is not disclosed to users. This can prevent the provision of semantic information which could result in unpredictable disbenefit for users or the disclosure of semantic information to parties other than the contract party.

The system may disclose semantic information having a predetermined “level of disclosure” associated therewith to the user through approval by the user of disclosure of semantic information having a predetermined “level of disclosure” associated therewith through, for example, informed consent.

The “level of disclosure” can be set as a plurality of levels of three or more, for example, “1, 2, 3 ...” or “a, b, c ...”. In this case, the level can be set on the shared computer 2 side according to the type of user, such as the user’s age, the user’s qualification, and whether or not a contract exists with the user. The user can select the level of disclosure such that only annotative information is provided which represents the morbidity rate associated with the level of disclosure that is above (or below) the predetermined level of disclosure determined in accordance with the informed consent or the like.

In database 8, for example, data such as nucleotide sequence-related information that is the genetic information of the individual requester utilizing the system can be recorded in storage DB-B16. In storage DB-A15, for example, data such as information distinguishing the requester from others utilizing the system can be recorded. In this way, the separate recording of the genetic information of individuals and the information for specifying individuals in storage DB-A15 and storage DB-B16, respectively, makes it difficult to associate a user’s genetic information with data that specifies the user.

Shared computer 2 is not limited to one having database 8 therein, and it may have an external database (not shown) connected to shared computer 2 through communication network 1. Shared computer 2 may have a plurality of databases 8 therein or may have an internal database 8 and an external database connected to shared computer 2 through communication network 1.

As shown in Fig. 4, personal computer 3 is constituted by CPU 20 that totally controls operation of personal computer 3, input device 21 such as a keyboard and a mouse with which information and instructions for executing a program are input, display device 22

such as a display apparatus, storage 23 having temporary information, rewritable information and the like recorded therein, reading apparatus 25 for reading data from genome-related information recording medium 24, and transmitter/receiver 29 for transmitting and receiving information to and from shared computer 2 through communication network 1. Personal computer 3 is not limited to a commonly used computer. For example, it may be any form of cellular phone, personal digital assistance, or other mobile communication tool.

Storage 23 in personal computer 3 has a memory section 26 for recording information provided from genome-related information recording medium 24 and the like, and is recorded a processing program 27 for operating the system for processing information.

Genome-related information recording medium 24 has genome-related information 28 of an individual recorded thereon. Genome-related information recording medium 24 includes, for example, a magnetic recording medium such as a magnetic disk or a magnetic card, an optical recording medium employing such as a magneto-optic recording system or a phase-change recording system, and a semiconductor memory. This genome-related information recording medium 24 may be in any form such as, for example, card, disk, stick, tape, or drum. Further, this genome-related information recording medium 24 may comprise genome-related information 28 of a single individual (an individual organism) recorded thereon. Alternatively, it may comprise a plurality of pieces of genome-related information 28 on a plurality of individuals (individual organisms) recorded thereon.

Genome-related information 28 contained in genome-related information recording medium 24 refers to, at least, a "polymorphism address" and a "polymorphism pattern" in the predetermined polymorphism address obtained as a result of analysis of an individual's (individual organism's) nucleotide sequences. Genome-related information 28 may contain various information, such as information concerning anamnesis, characteristics, an individual's clinical record, or a result of medical examination.

On genome-related information recording medium 24, recorded as genome-related information 28 is, for example, as shown in Fig. 5, the individual's number "Gno." (G number) peculiar to genome-related information 28 as well as the individual's information, such as date of birth, as data I; polymorphism addresses and polymorphism patterns as data II; anamnesis information as data III; characteristics as data IV; and information concerning the individual's clinical record and the like as data V. In other words, genome-related information 28 is constituted by data I, data II, data III, data IV, and data V. Data I and data II contain essential information and data III, data IV, and data V are respectively constituted by additional information.

In genome-related information 28, the “polymorphism address” corresponding to the position on the nucleotide sequence is linked with the “polymorphism pattern” in the polymorphism address and recorded. Additional information in a predetermined polymorphism address may be recorded in data II as a “comment” linked with a
 5 “polymorphism address”. All the nucleotide sequences of a predetermined individual organism may be recorded in data II. Even when all the nucleotide sequences are recorded in data II, “polymorphism addresses” and “polymorphism patterns” are contained within data II.

According to the present invention, personal computer 3 and genome-related information recording medium 24 are not limited to the construction as shown in Figs. 4 and
 10 5 respectively. For example, a genome-related information recording medium may be equipped with a memory section having a processing program and a personal computer may have the genome-related information recording medium mounted thereon to operate the processing program. In this case, a personal computer can be operated in accordance with a processing program recorded in a memory section on a genome-related information recording
 15 medium.

In a system for processing information having the above construction, processing program 13 recorded in storage 7 in shared computer 2 and processing program 27 recorded in storage 23 in personal computer 3 process information in accordance with, for example, flow charts as shown in Figs. 6 and 7. In the flow charts as shown in Figs. 6 and 7, a step
 20 described as “(shared)” refers to processing in shared computer 2 and a step described as “(personal)” refers to processing in personal computer 3.

The system for processing information is a system in which an individual possessing genome-related information recording medium 24 accesses shared computer 2 using personal computer 3 through communication network 1 and utilizes semantic information recorded in
 25 main DB 14 in shared computer 2. The system for processing information may be a system comprising the genome-related information recording medium 24, having genome-related information 28 on a plurality of individuals recorded thereon, to which individuals respectively access.

In this case, when utilizing the system, the requester first starts processing program
 30 27, which is recorded in storage 23, in step A1 (SA1). Processing program 27 drives reading apparatus 25 in personal computer 3 to access genome-related information recording medium 24. Thus, “Gno.” recorded as data I on genome-related information recording medium 24 is read out and the read-out “Gno.” is stored in memory section 26.

In step A2 (SA2), based on a screen image displayed by processing program 27 on display device 22, information, the provision of which is desired by the requester wishes to receive, for example, the “morbidity rate of large-bowel cancer” (requested information), is input to personal computer 3. At the same time, the “morbidity rate of large-bowel cancer” and “Gno.” are transmitted to shared computer 2 from personal computer 3 through communication network 1. Alternatively, the requester writes the “morbidity rate of large-bowel cancer” and “Gno.” in shared computer 2 from personal computer 3 through communication network 1.

In step A3 (SA3), shared computer 2 receives the “morbidity rate of large-bowel cancer” and “Gno.” The received “morbidity rate of large-bowel cancer” and “Gno.” are stored in memory section A10 as request information.

In step A4 (SA4), upon the reception of request information, processing program 13 recorded in storage 7 is started to access main DB 14. This processing program 13 performs processing in shared computer 2.

In step A5 (SA5), in accordance with processing program 13, “classification (name of disease)” recorded in main DB 14 is searched and information matching with the requested “morbidity rate of large-bowel cancer” (large-bowel cancer) is extracted.

In step A6 (SA6), from among data recorded in main DB 14, a “polymorphism address” associated with “classification (name of disease)” (large-bowel cancer) that matches with the “morbidity rate of large-bowel cancer” is read out. The read-out “polymorphism address” is stored as positional information associated with request information in memory section A10. Specifically, the “morbidity rate of large-bowel cancer” and “polymorphism address” are recorded in memory section A10 in association with a predetermined “Gno.”

In step A7 (SA7), “Gno.” and “polymorphism address” recorded in memory section A10 are transmitted to personal computer 3 and instruction information instructing submission of a “polymorphism pattern” corresponding to the transmitted “polymorphism address” is transmitted to personal computer 3. At this time, the submission of additional information such as that concerning anamnesis and characteristics may be optionally instructed depending on the types of request information.

In step A8 (SA8), “Gno.,” “polymorphism address,” and instruction information transmitted from shared computer 2, are received. The received “Gno.” and “polymorphism address” are recorded in memory section 26.

In step A9 (SA9), data II recorded on genome-related information recording medium 24 is accessed in accordance with the received instruction information. In step A10 (SA10),

in accordance with processing program 27, data II recorded on genome-related information recording medium 24 is searched, a polymorphism pattern in the instructed polymorphism address is read out, and the polymorphism pattern is then recorded in memory section 26 in association with the polymorphism address. In this case, whether the “Gno.” received in step 5 A8 is correct or not is preferably confirmed by accessing data I. In step A10, additional information recorded in data III, data IV, and data V is read out simultaneously with the polymorphism pattern and may be optionally recorded in memory section 26.

In step A11 (SA11), the temporarily-recorded polymorphism pattern associated with the polymorphism address and the optionally-recorded additional information in memory 10 section 26 are output to shared computer 2 together with “Gno.” through communication network 1. In step A12 (SA12), shared computer 2 receives the polymorphism pattern associated with the polymorphism address and the optionally-recorded additional information, and the received polymorphism pattern is recorded in memory section A10 in association with the polymorphism address.

15 In this embodiment, in step A7 shared computer 2 transmits instruction information instructing submission of the “polymorphism pattern”, and in step A10 personal computer 3 reads out the polymorphism patterns from genome-related information recording medium 24 in accordance with instruction information. The system, however, may not transmit the instruction information in step A7. In this case, personal computer 3 searches data II in step 20 A10 based on the polymorphism address received in step A8 and reads out polymorphism patterns of the received polymorphism address in accordance with processing program 27. Then, in step A11, personal computer 3 outputs polymorphism patterns and the like to shared computer 2. Also in this case, in step A12 shared computer 2 can obtain the polymorphism pattern of the “polymorphism address” associated with “classification (name of disease)” that 25 matches the “morbidity rate of large-bowel cancer”.

In step A13 (SA13), main DB 14 is accessed to search information matching with the received polymorphism address and polymorphism patterns. More specifically, a plurality of polymorphism patterns are recorded in main DB 14 for one polymorphism address. Thus, which polymorphism pattern in main DB 14 matches with the received 30 polymorphism address and the polymorphism pattern thereof is searched.

In step A14 (SA14), the morbidity rate of large-bowel cancer which is associated with the polymorphism pattern matching the received polymorphism pattern is read out in accordance with processing program 13. Specifically, in step A14, the morbidity rate of large-bowel cancer of a requester can be read out in accordance with the polymorphism

address and polymorphism pattern submitted by the requester. The read-out morbidity rate is stored in memory section A10 in association with the requester's "Gno." At this time, the morbidity rate of large-bowel cancer may be corrected in accordance with additional information and then stored. Alternatively, other information obtained from additional information may be stored in association with the requester's "Gno."

Subsequently, in step A15 (SA15), the requester's "Gno." and morbidity rate, which are stored in memory section A10, are transmitted as semantic information to personal computer 3 through communication network 1. Personal computer 3 receives the requester's "Gno." and morbidity rate (semantic information) in step A16 (SA16). The received semantic information is recorded in memory section 26.

In step A17 (SA17), the morbidity rate of large-bowel cancer is displayed on display device 22 based on semantic information recorded in memory section 26 in accordance with processing program 27. Instead of steps A15 to A17, shared computer 2 can read out (prepare) a screen that displays semantic information in accordance with processing program 13, and display it on display device 22 of personal computer 3 through communication network 1. Also in this case, semantic information is considered to be transmitted from shared computer 2 to personal computer 3. As a result, the requester can obtain the morbidity rate of large-bowel cancer using genome-related information 28 recorded on genome-related information recording medium 24.

As described above, in this system, utilization of genome-related information recording medium 24, which has individuals' polymorphism patterns in association with polymorphism addresses recorded thereon, enables individuals to use semantic information recorded in main DB 14 through the polymorphism addresses. In other words, an individual utilizing this system does not have to record semantic information on a genome-related information recording medium. Instead, the individual can obtain various semantic information simply by possessing genome-related information 28 having the polymorphism pattern associated with the polymorphism address.

More particularly, as described above, semantic information is corrected and increases in the number of types. Thus, by updating main DB 14 the semantic information becomes more accurate and includes a wider range of information. According to this system, individuals can utilize the newest semantic information by updating main DB 14 in line with such increase, correction or the like of semantic information.

Further, utilization of genome-related information recording medium 24 having genome-related information 28 recorded thereon eliminates the need for the user to undergo

an examination to obtain genome-related information every time he/she utilizes this system. More specifically, once genome-related information recording medium 24 is produced, thereafter the user can obtain the newest semantic information utilizing this system.

Retention by the user of genome-related information recording medium 24 on which the user's genome-related information 28 has been recorded can prevent apprehension in consigning an external organization to store the user's genome-related information 28 and the risk of leakage of genome-related information 28 through unauthorized access to the organization. On the other hand, when storage of genome-related information recording medium 24 having a plurality of pieces of genome-related information 28 of a plurality of individuals recorded thereon is consigned to an external organization, inefficient handling of genome-related information recording medium 24 or loss of genome-related information recording medium 24 can be prevented, unlike the case where each individual retains their own genome-related information recording medium 24.

In particular, in accordance with the flow charts shown in Figs. 6 and 7, all the genome-related information 28 recorded on genome-related information recording medium 24 is not necessarily output through communication network 1, and only the part of genome-related information 28 for which an instruction for the submission was received may be output. According to this system, leakage of highly confidential polymorphism addresses and polymorphism patterns peculiar to individuals can be prevented.

Also, in accordance with the flow charts shown in Figs. 6 and 7, personal computer 3 does not need to handle information recorded in main DB 14 since shared computer 2 obtains semantic information to provide to the requester. Thus, in accordance with the flow charts shown in Figs. 6 and 7, desired semantic information can be adequately obtained even if the capacity of personal computer 3 to process information is relatively low. Further, since personal computer 3 does not need to handle information recorded in main DB 14, standardization of processing program 27 in personal computer 3 in compliance with a card drive and the like having genome-related information recording medium 24 mounted thereon is facilitated.

In the system for processing information, processing program 13 recorded in storage 7 of shared computer 2 and processing program 27 recorded in storage 23 of personal computer 3 may process information in accordance with, for example, the flow chart shown in Fig. 8. Also in the flow chart shown in Fig. 8, a step described as "(shared)" refers to processing in shared computer 2 and a step described as "(personal)" refers to processing in personal computer 3.

When utilizing the system, the requester first starts processing program 27 recorded in storage 23 in step B1 (SB1). Processing program 27 drives reading apparatus 25 in personal computer 3 and accesses genome-related information recording medium 24, thereby reading out a "Gno." recorded in genome-related information recording medium 24 as data I.
5 The read-out "Gno." is stored in memory section 26.

In step B2 (SB2), based on a screen image displayed by processing program 27 on display device 22, information which the requester wishes to receive, for example, the "morbidity rate of large-bowel cancer" (request information), is input to personal computer 3. At the same time, the "morbidity rate of large-bowel cancer" and "Gno." are transmitted to
10 shared computer 2 from personal computer 3 through communication network 1. Concurrently requested is the submission of "polymorphism address" in which "classification (name of disease)" in main DB 14 is large-bowel cancer, all the "polymorphism patterns" associated with the "polymorphism address," and the "morbidity rate" implied by all the "polymorphism patterns" respectively. More specifically, in step B2, the requester requests
15 information comprising the "polymorphism address" in which "classification (name of disease)" in main DB 14 is large-bowel cancer, all the "polymorphism patterns" associated with the "polymorphism address," and the "morbidity rate" implied by all the "polymorphism patterns" respectively.

In step B3 (SB3), shared computer 2 receives the request information. Shared
20 computer 2 starts processing program 13 upon reception of the request information and, in step B4 (SB4), accesses main DB 14 in accordance with processing program 13.

In step B5 (SB5), in accordance with processing program 13, "classification (name of disease)" recorded in main DB 14 is searched and information matching with the requested "morbidity rate of large-bowel cancer" (large-bowel cancer) is extracted. In step B6 (SB6),
25 main DB 14 is accessed according to processing program 13 to read out the "polymorphism address" associated with "classification (name of disease)" (large-bowel cancer) that matches the "morbidity rate of large-bowel cancer," all the "polymorphism patterns" associated with the polymorphism address, and the "morbidity rate" in all the polymorphism patterns. The read-out "polymorphism address," "polymorphism pattern," and "morbidity rate" are stored
30 in memory section A10 in association with the request information. Specifically, the "polymorphism address," the "polymorphism pattern," and the "morbidity rate" are recorded in memory section A10 for a predetermined "Gno."

In step B7 (SB7), the "Gno." "polymorphism address," "polymorphism pattern," and "morbidity rate" recorded in memory section A10 are transmitted to personal computer 3

through communication network 1. In step B8 (SB8), the “Gno.” “polymorphism address,” “polymorphism pattern,” and “morbidity rate” transmitted from the shared computer 2 are received. The received “Gno.” “polymorphism address,” “polymorphism pattern,” and “morbidity rate” are recorded in memory section 26.

5 In step B9 (SB9), data II, recorded on genome-related information recording medium 24, is accessed according to processing program 27. In this case, data I, recorded on genome-related information recording medium 24, is also accessed and whether the received “Gno.” is correct or not is preferably confirmed.

10 In step B10 (SB10), the polymorphism pattern in the polymorphism address matching with the received “polymorphism address” is extracted from genome-related information 28 according to processing program 27. In step B10 (SB10), polymorphism pattern matching with the extracted polymorphism pattern is searched from all the received “polymorphism patterns” associated with the received polymorphism address.

15 In step B11 (SB11), the “morbidity rate” associated with the matching polymorphism pattern among all the received “polymorphism patterns” associated with the received polymorphism address is extracted and the extracted “morbidity rate” is output. This enables the requester to obtain the morbidity rate of large-bowel cancer (semantic information). In step B11, additional information recorded in data III, data IV, and data V is simultaneously read out and the morbidity rate of large-bowel cancer may be corrected by the
20 additional information and then output.

More specifically, in accordance with the flow chart shown in Fig. 8, genome-related information 28 recorded on genome-related information recording medium 24 is output only to personal computer 3 and other than this is not output at all to the outside. That is, genome-related information 28 is transmitted/received only between genome-related information
25 recording medium 24 and personal computer 3. Thus, the system can more accurately prevent leakage of highly confidential genome-related information 28 peculiar to the individual.

30 In the system for processing information, processing program 13 recorded in storage 7 of shared computer 2 and processing program 27 recorded in storage 23 of personal computer 3 may process information, for example, in accordance with the flow chart shown in Fig. 9. In the flow chart shown in Fig. 9, a step described as “(shared)” refers to processing in shared computer 2 and a step described as “(personal)” refers to processing in personal computer 3.

When utilizing the system, the requester first starts processing program 27 recorded in storage 23 in step C1 (SC1). Processing program 27 drives reading apparatus 25 in personal computer 3 and accesses genome-related information recording medium 24 to read out a "Gno." recorded therein as data I and all "polymorphism addresses" and
 5 "polymorphism patterns" recorded therein as data II. The read-out "Gno.", "polymorphism address", and "polymorphism pattern" are stored in memory section 26.

In step C2 (SC2), based on a screen image displayed by processing program 27 on display device 22, information which the requester wishes to receive, for example the "morbidity rate of large-bowel cancer" (request information), is input to personal computer 3.
 10 At the same time, the "morbidity rate of large-bowel cancer" and "Gno.", "polymorphism address," and "polymorphism pattern" recorded in memory section 26 are transmitted to shared computer 2 from personal computer 3 through communication network 1.

In step C3 (SC3), shared computer 2 receives "morbidity rate of large-bowel cancer," "Gno.", "polymorphism address," and "polymorphism pattern." The received
 15 "morbidity rate of large-bowel cancer" is recorded as request information in memory section A10 and "Gno.", "polymorphism address" and "polymorphism pattern" are also stored in memory section A10. Shared computer 2 starts processing program 13 upon reception of the request information and, in step C4 (SC4), accesses main DB 14 in accordance with processing program 13.

20 In step C5 (SC5), in accordance with processing program 13, "classification (name of disease)" recorded in main DB 14 is searched and classification matching with the requested "morbidity rate of large-bowel cancer" (large-bowel cancer) is extracted.

In step C6 (SC6), main DB 14 is accessed in accordance with processing program 13 to read out from main DB 14 the "polymorphism address" classified into "large-bowel
 25 cancer", all the "polymorphism patterns" associated with the polymorphism address, and the "morbidity rate" in all the polymorphism patterns. The read-out "polymorphism address," "polymorphism pattern," and "morbidity rate" are stored in memory section A10.

In step C7 (SC7), the data stored in memory section A10 in step C6 is searched based on the "polymorphism address" and the "polymorphism pattern" received in step C3,
 30 and a morbidity rate associated with polymorphism pattern matching with the received "polymorphism pattern" is extracted from memory section A10.

In step C8 (SC8), the result of step C7, that is, the morbidity rate extracted according to which polymorphism pattern in main DB 14 matches with the polymorphism pattern contained in the received information in step C3, is transmitted to personal computer 3

through communication network 1. In this case, shared computer 2 transmits the extracted morbidity rate together with the requester's "Gno."

In step C9 (SC9), the "Gno." and "morbidity rate (semantic information)" transmitted from shared computer 2 is received. The received "Gno." and "morbidity rate" are recorded in memory section 26. At this time, data I recorded on genome-related information recording medium 24 is accessed and whether the received "Gno." is correct or not can be confirmed.

In step C10 (SC10), in accordance with processing program 27, the morbidity rate of large-bowel cancer is displayed on display device 22 based on semantic information recorded in memory section 26. Instead of steps C8 to C10, shared computer 2 can read out (prepare) a screen that displays semantic information in accordance with processing program 13, and display it on display device 22 of personal computer 3 through communication network 1. Also in this case, semantic information is considered to be transmitted from shared computer 2 to personal computer 3. This enables the requester to obtain the morbidity rate of large-bowel cancer using genome-related information 28 recorded on genome-related information recording medium 24.

More particularly, in accordance with the flow chart shown in Fig. 9, all the genome-related information 28 recorded on genome-related information recording medium 24 is output to shared computer 2 and semantic information to provide to the requester is obtained in shared computer 2. In accordance with the flow chart shown in Fig. 9, even with a relatively small number of times of reception/transmission of information between personal computer 3 and shared computer 2, the requester can obtain semantic information. Therefore, in accordance with the flow chart shown in Fig. 9, even if the information processing capacity of personal computer 3 is relatively low, the desired semantic information can be adequately provided. In addition, the requester can obtain semantic information in a very simple manner.

As described above, according to the system, on genome-related information recording medium 24 and in main DB 14, standardization of only "polymorphism addresses" and the "polymorphism patterns" thereof eliminates the need for standardization of other specific data. Thus, the system can be utilized in a wide range of industries. That is, when providing information using genome-related information recording medium 24, the provider of objects or services can provide information in various manners without the need to standardize semantic information to correspond to the polymorphism pattern or a unified standard such as a method for transmitting/receiving data.

Furthermore, according to the system, a third party or third organization can easily monitor and control shared computer 2 by examining main DB 14. Accordingly, as the system can, for example, execute administrative control over the provider of semantic information, adequate and ethical control over the provider of semantic information can be executed .

Meanwhile, in the system for processing information, a recording medium in which information contained in data II is removed from a genome-related information recording medium, that is, a recording medium having only data I and additionally data III to V, may be used. In this case, information contained in data II is recorded in an external database (genome-related information recording medium) connected to personal computer 3 through communication network 1. In such a system, for example, in above-described step A10, the external database is accessed through communication network 1 and a polymorphism pattern in the instructed polymorphism address is read out, and the polymorphism pattern can be recorded in association with the polymorphism address in memory section 26. Thus, in this system, as with the flow charts shown in Figs. 6 and 7, 8, and 9, the requester can obtain semantic information.

In the system for processing information, the requester may have a genome-related information recording medium 24 connected to personal computer 3 through communication network 1 instead of the requester him/herself has genome-related information recording medium 24 or a recording medium in which information contained in data II is removed from the genome-related information recording medium. In such a system, the requester can access genome-related information recording medium 24 through communication network 1 to download information such as “polymorphism addresses” and “polymorphism patterns” recorded on genome-related information recording medium 24 into personal computer 3. In this case, genome-related information recording medium 24 may have genome-related information of a plurality of individuals for each individual (each “Gno.”) recorded thereon.

In addition, the present invention is not limited to the above-described construction in which shared computer 2 comprises main DB 14, and, for example, is applicable to a system for processing information equipped with main DB 14 connected to shared computer 2 through communication network 1. In this case, shared computer 2 accesses main DB 14 through communication network 1 as shown in the flow charts in Figs. 6 and 7, 8, or 9. In this case also, according to the system for processing information, the requester can also obtain desired semantic information in accordance with the flow charts shown in Figs. 6 and 7, 8, or 9.

More specifically, shared computer 2 can access a plurality of main DBs 14 owned by different organizations or groups through communication network 1 and can utilize semantic information contained in these plurality of main DBs 14, thereby providing information to the requester. That is, in the system for processing information, in step A5 in the flow charts shown in Figs. 6 and 7, in step B5 in the flow chart shown in Fig. 8, or in step C5 in the flow chart shown in Fig. 9, shared computer 2 accesses various main DBs 14 having information on the morbidity rate of large-bowel cancer as semantic information. According to the system for processing information, therefore, the requester can obtain information on the morbidity rate of large-bowel cancer from information contained in various main DBs 14.

In this system, as shown in the flow charts shown in Figs. 6 and 7, 8, or 9, shared computer 2 may transmit, at least, the request information received from personal computer 3 to a so-called agent and obtain semantic information (“morbidity rate of large-bowel cancer” in this embodiment) through the agent.

2. Second Embodiment

Next, a system for processing information for providing a medicament appropriate for the diathesis, such as regarding responsiveness to a medicament or immunity to substances, is explained as a second embodiment to which the present invention has been applied. Regarding the system for processing information explained as the present embodiment, explanation of the constitution, operation, and terms thereof is omitted by employing the same appellations, symbols, and definitions for the same constitution and terms as with the system for processing information according to the above described first embodiment. The present embodiment is directed to explanation of a system for providing an object (a medicament) in accordance with request information from the user, and thus, is explained as a simple model for the convenience of the explanation. More specifically, semantic information (drug type) and the like, as shown in Fig. 10, is recorded in main DB 14 in the present embodiment.

The system for processing information for providing a medicament can provide the most suitable medicament to the user, for example, as according to the flow charts shown in Figs. 11 and 12. In the flow charts shown in Figs. 11 and 12, a step described as “(shared)” refers to processing in shared computer 2 and a step described as “(personal)” refers to processing in personal computer 3.

In step II-1 (SII-1), the requester first accesses, for example, shared computer 2 of a pharmaceutical company from personal computer 3 through communication network 1.

Shared computer 2 of a pharmaceutical company may be, for example, an exclusive computer, a computer for controlling an order for a medicament and information on ordering and shipping, or a computer for controlling production of a desired medicament by controlling a pharmaceutical plant. Shared computer 2 is not limited to those belonging to a pharmaceutical company and it may belong to an agency or institution for providing information, such as a type of drug or number of medicaments to be provided, to a provider of a medicament such as a pharmaceutical company.

Examples of requesters in the present embodiment include individuals possessing genome-related information recording medium 24, medical institutions possessing genome-related information recording medium 24, and medical institutions utilizing genome-related information recording medium 24 possessed by individuals.

In step II-2 (SII-2), in shared computer 2, processing program 13 is started in response to the requester's access and a "portal screen for tailor-made non-prescription pharmaceuticals" is read from screen memory 12 to display on display device 22 of personal computer 3. Examples of the "portal screen for tailor-made non-prescription pharmaceuticals" include a screen image as shown in Fig. 13.

In step II-3 (SII-3), the requester equips genome-related information recording medium 24 into reading apparatus 25 in accordance with an instruction on the "portal screen for tailor-made non-prescription pharmaceuticals" so as to enable access to genome-related information recording medium 24 using personal computer 3. In this case, whether or not the requester is an authorized user of genome-related information recording medium 24 is preferably authenticated. In step II-4 (SII-4), the requester then clicks an "OK" button displayed on the "portal screen for tailor-made non-prescription pharmaceuticals" to transmit one's intent to utilize the system to shared computer 2.

In step II-5 (SII-5), a "medicament category menu" screen is read from screen memory 12 in shared computer 2 in accordance with processing program 13 and displayed on display device 22 of personal computer 3. Examples of a "medicament category menu" screen include a screen image as shown in Fig. 14.

In step II-6 (SII-6), the requester selects the desired medicament category (medicament) in accordance with the "medicament category menu" screen that has been accessed with personal computer 3.

In step II-7 (SII-7), reading apparatus 25 of personal computer 3 accesses genome-related information recording medium 24 and reads out “Gno.” from data I. In step II-8 (SII-8), “Gno.” read out in step II-7 and the “medicament category” (request information) selected in step II-6 are transmitted to shared computer 2 through communication network 1.

In step II-9 (SII-9), shared computer 2 receives “Gno.” and “medicament category” and the “Gno.” and the “medicament category” are stored in memory section A10. In the present embodiment, the “level of disclosure” is associated with semantic information that is recorded in main DB 14. The “level of disclosure” in this embodiment indicates whether or not semantic information and information associated with the semantic information should be disclosed to the user of the information. More specifically, recorded, in association with semantic information, are: “level of disclosure A” designating a level that may be disclosed only when the requester has been authenticated; “level of disclosure B” designating a level that may be disclosed to any requester irrespective of whether the requester has been authenticated or not; and “prohibition against disclosure” designating a level that should not be disclosed to any requester.

Shared computer 2 comprises, for example, a database having “information regarding the authenticated requester such as ‘requester No.’ ” (not shown, hereinafter referred to as “requester information database”) previously recorded in database 8.

The requester transmits the “requester No.” to shared computer 2 together with “Gno.” and request information. When services using semantic information associated with “level of disclosure A” and information associated with the semantic information are to be provided to the requester, shared computer 2 accesses the “requester information database” to judge, based on the received “requester No.”, whether or not the requester has been authenticated.

When the requester is judged to be authenticated, shared computer 2 registers “level of disclosure A” in association with the “Gno.” received from personal computer 3 in storage 7. In the subsequent step, shared computer 2 searches the “level of disclosure” in main DB 14 based on the registered “level of disclosure A” and provides services using semantic information associated with “level of disclosure A” and information associated with the semantic information to the requester. On the other hand, when the requester is judged to be unauthenticated, shared computer 2 immediately cancels processing or registers “level of disclosure B” in association with the received “Gno.” in storage 7. In the subsequent step, shared computer 2 searches the “level of disclosure” in main DB 14 based on the registered “level of disclosure B” and provides services using semantic information associated with

“level of disclosure B” and information associated with the semantic information to the requester. Thus, use of the “level of disclosure” enables the determination whether or not the disclosure of semantic information and information associated with the semantic information is approved.

In the present embodiment, because the system is one for providing non-prescription medicaments, the requesters are an unspecified large number of people without the need of previous authentication. This system, therefore, eliminates the need to judge whether or not the requester has been authenticated. For this reason, in step II-9, upon the receipt of request information (medicament category), “level of disclosure B” is automatically selected and registered in storage 7 in association with the “Gno.”

Next, in step II-10 (SII-10), main DB 14 is accessed in accordance with processing program 13. Then, in step II-11 (SII-11), from the polymorphism addresses contained in main DB 14, processing program 13 selects the polymorphism address which matches with the received “medicament category” and for which disclosure is approved by the “level of disclosure,” that is, one for which “level of disclosure B” has been set. In this case, as shown in Fig. 10, “major classification,” “middle classification,” and “minor classification” are recorded in association with one another based on predetermined polymorphism addresses in main DB 14. Among them, the middle classification categorizes the medicament category. Thus, a polymorphism address which matches with the received “medicament category” can be selected by searching “middle classification” based on the received “medicament category.” For example, in the “medicament category menu” in step II-5, when, in addition to a non-prescription medicament, there is a prescription medicament to be provided as an option only to a user with authentication for “level of disclosure A,” if a requester without the authentication for “level of disclosure A” requests both the non-prescription medicament and the prescription medicament, the request for the prescription medicament will be denied based on the level of disclosure in main DB 14, and in step II-11, only the polymorphism address associated with the non-prescription medicament is selected.

In step II-12 (SII-12), the selected “polymorphism address” “medicament category” and “Gno.” are recorded in memory section B11. At this time, in step II-13 (SII-13), in the system, an “anonymous polymorphism address” (secondary positional information) is set in compliance with a predetermined regulation and the like based on the “polymorphism address” (positional information) which is recorded in memory section B11, and linked to the “polymorphism address” in memory section B11 to record the “anonymous polymorphism address” in memory section B11. This “anonymous polymorphism address” is linked only to

the “polymorphism address” selected in step II-11 whereas the “polymorphism address” itself directly represents a position of the polymorphism pattern in a nucleotide sequence. That is, the “anonymous polymorphism address” does not directly represent a position of a polymorphism pattern in a nucleotide sequence. In other words, “the anonymous
 5 polymorphism address” is set by allocating, for example, serial numbers, alphabetical characters, or random numbers to the “polymorphism address” selected in step II-11.

In step II-14 (SII-14), in accordance with processing program 13, informed consent (hereinafter abbreviated to “IC”) concerning the submission of polymorphism patterns is asked of the requester. Specifically, in step II-14 (SII-14), an “input screen for individual IC
 10 confirmation” is read from screen memory 12 in accordance with processing program 13 and information on the received “medicament category” is incorporated therein. Then, the “input screen for individual IC confirmation” is displayed on display device 22 of personal computer 3. Examples of an “input screen for individual IC confirmation” include a screen image as shown in Fig. 15.

15 In step II-15 (SII-15), in accordance with the “input screen for individual IC confirmation” displayed on display device 22 of personal computer 3, the requester inputs their consent for submission of polymorphism patterns. This indicates that the requester has consented to submit polymorphism patterns.

In step II-16 (SII-16), data I on genome-related information recording medium 24 is
 20 accessed to read out “Gno.” In step II-17 (SII-17), the read-out “Gno.” and the “individual IC confirmation” input in step II-15 are transmitted to shared computer 2.

In step II-18 (SII-18), shared computer 2 receives the “Gno.” and “individual IC confirmation” transmitted from personal computer 3, and the received “individual IC confirmation” is stored in memory section A10. When storing the “individual IC
 25 confirmation,” the “individual IC confirmation” is associated with the “Gno.” in the data stored in memory section A10 that matches with the received “Gno.”

In step II-19 (SII-19), the “polymorphism address” and the “anonymous polymorphism address” stored in steps II-12 and II-13 are read from memory section B11. In step II-20 (SII-20), the read-out “polymorphism address” is transmitted to personal computer
 30 3 in association with the read-out “anonymous polymorphism address,” and instruction information that instructs the submission of the polymorphism pattern corresponding to the “polymorphism address” and the “anonymous polymorphism address” associated with the polymorphism pattern is output to personal computer 3.

In step II-21 (SII-21), the “polymorphism address” and “anonymous polymorphism address” transmitted from shared computer 2 as well as the instruction information are received, and genome-related information recording medium 24 is accessed in accordance with processing program 27. In step II-22 (SII-22), from data II in genome-related information 28, the processing program 27 extracts and reads out the polymorphism address as well as the polymorphism pattern corresponding to the polymorphism address requested by the instruction information, and reads out “Gno.” from data I.

In step II-23 (SII-23), the read-out polymorphism pattern is associated with the anonymous polymorphism address set for the polymorphism address of the polymorphism pattern, and these are transmitted together with “Gno.” from personal computer 3 to shared computer 2. Specifically, in step II-23, the anonymous polymorphism address, the polymorphism pattern, and the Gno. are transmitted from personal computer 3 to shared computer 2 through communication network 1. In this case, the polymorphism address is not transmitted in association with the polymorphism pattern from personal computer 3. Also, the anonymous polymorphism address does not directly represent a position of the polymorphism pattern in a nucleotide sequence. Thus, even if the content transmitted in step II-23 is leaked externally due to unforeseen circumstances, the position of the polymorphism pattern in the nucleotide sequence cannot be determined.

In step II-24 (SII-24), the anonymous polymorphism address, the polymorphism pattern associated with the anonymous polymorphism address, and “Gno.” transmitted from personal computer 3 are received, and the received polymorphism pattern is stored in memory section B11. When storing the “polymorphism pattern,” the “polymorphism pattern” is associated with the “Gno.” and the “anonymous polymorphism address” in the data stored in memory section B11 that match with the received “Gno.” and “anonymous polymorphism address.” According to the procedure in step II-24, the “Gno.”, the “polymorphism address,” the “anonymous polymorphism address” set for each polymorphism address and “polymorphism pattern” are recorded in association with one another in memory section B11.

In step II-25 (SII-25), main DB 14 is accessed in accordance with processing program 13. In step II-26 (SII-26), main DB 14 is searched based on the “middle classification (medicament category),” the “polymorphism address,” and the “polymorphism pattern” thereof stored in memory section B11, and the “drug type” associated with the matching polymorphism pattern is read out (see Fig. 10). “Drug type” herein is, in the medicament category classified by the “middle classification” as shown in Fig. 10, semantic

information implying information such as a component, a dose, and the like suitable for individuals having predetermined polymorphism patterns. When a plurality of pieces of semantic information is associated with one polymorphism pattern, each piece of semantic information is read out.

5 In step II-27 (SII-27), the obtained “drug type” is stored in memory section B11 in accordance with processing program 13. In step II-28 (SII-28), the “medicament category” and “drug type” are read from memory section B11 in accordance with processing program 13. In step II-29 (SII-29), a “confirmation screen” corresponding to the read-out “medicament category” and “drug type” is read from screen memory 12 in accordance with
10 processing program 13 and displayed on display device 22 of personal computer 3. Examples of a “confirmation screen” include a screen image as shown in Fig. 16.

In step II-30 (SII-30), the requester clicks an “order screen” button on the “confirmation screen” of personal computer 3 upon confirmation of “name of medicament,” “drug type” and the like on the “confirmation screen” to thereby read out the “order screen”
15 (not shown) from screen memory 12 and input order information. Order information includes information for identifying the requester, the quantity with respect to a medicament name falling within the drug type displayed on the order screen, the methods for reception and payment, and the like.

In step II-31 (SII-31), a “Gno.” is read from data I of genome-related information 28
20 by accessing genome-related information recording medium 24 in accordance with processing program 27. In step II-32 (SII-32), the read-out “Gno.” and order information input in step II-30 are transmitted to shared computer 2 in accordance with processing program 27.

In step II-33 (SII-33), shared computer 2 receives “Gno.” and “order information”
25 transmitted from personal computer 3 and stores the order information in memory section A10. When storing the order information, among the data stored in memory section A10, the “order information” is associated with a “Gno.” matching the received “Gno.”

In step II-34 (SII-34), a “confirmation screen for acceptance of order” (not shown) is read from screen memory 12 in accordance with processing program 13 to display the
30 “confirmation screen for acceptance of order” on display device 22 of personal computer 3. In step II-35 (SII-35), the requester then confirms the order contents in accordance with the instructions on the “confirmation screen for acceptance of order”. The requester, by the above processes, thus completes ordering of a medicament suitable for the individual possessing genome-related information recording medium 24.

Meanwhile, in step II-36 (SII-36), essential information such as “medicament name (information associated with semantic information),” “drug type (semantic information)” and “quantity” is extracted from order information stored in memory section A10 and the extracted information is then transmitted to a provider of medicaments (the user of semantic information and information associated with the semantic information) such as a pharmaceutical plant or distribution center. Essential information refers to information which is required in the production or delivery of medicaments requested by the requester. A provider of medicaments includes a producer of medicaments suitable for the requester through the adjustment of medicament ingredients, component ratio and the like in accordance with the “medicament name” and the “drug type,” or a provider of medicaments that provides medicaments by selecting from plural types of medicaments that were previously produced with varying medicament ingredients, component ratios and the like based on the “medicament name” and “drug type”. According to the above procedures, the provider of medicaments produces or delivers medicaments suitable for individuals possessing genome-related information recording medium 24.

In shared computer 2, the procedure finally advances to step II-37 (SII-37). In step II-37, the contents stored in memory section A10 and memory section B11 are separately stored in storage DB-A15 and storage DB-B16 respectively. The contents stored in memory section A10 and memory section B11 are then deleted. At this time, the “anonymous polymorphism address” in the content stored in memory section B11 is preferably deleted.

Steps II-34 and II-35 are directed to a final confirmation of the order by the requester, and thus, are not essential in the system. Specifically, the system may proceed in such a manner that step II-33 is followed by step II-36. In this case, the requester completes ordering of medicaments suitable for an individual possessing genome-related information recording medium 24 in step II-32.

According to the flow charts shown in Figs. 11 and 12, information associated with the semantic information recorded in main DB14 as the “drug type” can be utilized using genome-related information 28 having individuals’ polymorphism patterns in association with polymorphism addresses recorded therein. According to the system, therefore, a medicament suitable for an individual possessing genome-related information recording medium 24 can be obtained based on the diathesis, including responsiveness to a medicament or immunity against substances, of the individual.

More particularly, semantic information is corrected and increases the number of types thereof as described above. Thus by updating main DB14 the semantic information

becomes more accurate and includes a wider range of information. According to this system, individuals can utilize the newest semantic information by updating main DB14 in line with such increase, correction or the like of semantic information.

In accordance with flow charts such as shown in Figs. 11 and 12, using the level of disclosure which has been previously recorded in main DB 14, the allowable range of disclosure of semantic information and information associated with the semantic information can be limited depending on the requester. According to the system, therefore, the disclosure of highly confidential semantic information and information associated with the semantic information in response to a request made by an unapproved requester can be prevented.

Thus, use of the level of disclosure enables the observation of laws, regulations, behavioral standards of an organization having the shared computer 2 or the like.

Further, in the flow charts as shown in Figs. 11 and 12, in step II-23 the anonymous polymorphism addresses and polymorphism patterns are transmitted from personal computer 3 to shared computer 2 through communication network 1. In the system, therefore, even if information transmitted in step II-23 was leaked externally due to unforeseen circumstances, the position of the polymorphism pattern in a nucleotide sequence contained in the information cannot be determined. In other words, use of the anonymous polymorphism address in the system can prevent leakage of an individual's information without the use of a highly developed encryption technique. Therefore, the concealment of an individual's information is enhanced in the system since information transmitted in step II-23 is unavailable to others. Furthermore, in the transmission of the information in step II-23, if the information is encrypted in accordance with a commonly performed system, concealment can be further enhanced.

In accordance with the flow charts shown in Figs. 11 and 12, memory section A10 has the "Gno.", "requested medicament category," "IC confirmation," and "order information" stored therein, as shown in Fig. 17, and memory section B11 has the "Gno.", "requested medicament category," "polymorphism address," "anonymous polymorphism address," "polymorphism pattern," and "drug type" stored therein, as shown in Fig. 18. Linking of the information for identifying the requester and the genetic information of the requester is made complicated by separating memory section A10, which stores the "order information" containing information to identify the requester from memory section B11, which stores the "polymorphism pattern" as genetic information of the requester oneself. This can conceal the correlation between a specific "polymorphism pattern" and information identifying the individual described in the "order information." The specific "polymorphism

pattern” is linked to the “order information” through “Gno.” and thus they can be associated with each other if necessary.

In the system, information stored in memory section A10 and information stored in memory section B11 are separately stored in storage DB-A15 and storage DB-B16 respectively. Thus, even when the “order information”, “polymorphism address” and “polymorphism pattern” of the requester is stored for a long period of time, the correlation between information for identifying the individual described in the “order information” and the “polymorphism address” and “polymorphism pattern” can be concealed according to the system. In this case, therefore, security against information leakage can be further enhanced.

In the system, when transmitting information to the provider of medicaments such as a pharmaceutical plant or a distribution center in step II-36, “Gno.” is not contained in the information. Instead, the information preferably comprises, for example, the “order No.”, the “medicament name,” the “drug type,” the “number of tablets,” the “number of boxes,” and the like, as shown in Fig. 19. In this case, information transmitted to the provider of medicaments contains no “Gno.”, and thus the “Gno.” cannot be identity from the information. This makes it impossible to specify the individual who possesses the genome-related information recording medium 24 corresponding to the “Gno.” In this case, therefore, the “polymorphism pattern” and “order information” of an individual can be surely concealed and the security against information leakage can be further enhanced.

In the above embodiment, the system for processing information for providing medicaments suitable for individuals using polymorphism addresses and polymorphism patterns contained in genome-related information 28 was described. The system, however, is not limited to medicaments but is also applicable to the provision of objects such as foods, cosmetics, or organism-related materials. In this case, for example, as semantic information implied by the polymorphism pattern and information associated with semantic information, a type of substance contained in an object such as food, cosmetics, or organism-related materials is selected using, for example, immunity against substances or responsiveness to medicaments as an index. A suitable object for the individual can be provided by producing or selecting an object in accordance with the type of selected substances and the like.

As described above, according to the system, on genome-related information recording medium 24 and in main DB 14, standardization of only “polymorphism addresses” and the “polymorphism patterns” thereof eliminates the need for standardization of other specific data. Thus, the system can be utilized in a wide range of industries. That is, when providing information using genome-related information recording medium 24, the provider

of objects or services can provide information in various manners without the need to standardize semantic information to correspond to the polymorphism pattern or a unified standard such as a method for transmitting/receiving data.

Furthermore, according to the system, a third party or third organization can easily monitor and control shared computer 2 by examining main DB 14. Accordingly, as the system can, for example, execute administrative control over the provider of semantic information, adequate and ethical control over the provider of semantic information can be executed.

3. Third Embodiment

Next, a system for processing information is explained as a third embodiment to which the present invention has been applied, wherein suitable medical examination items are provided to the individual based on the morbidity rate of a predetermined disease.

Regarding the system for processing information explained as the present embodiment, the explanation of its constitution, operation, and terms thereof is omitted by employing the same appellations, symbols, and definitions for the same constitution and terms as with the system for processing information according to the above described the first and the second embodiments. The present embodiment is directed to explanation of a system for providing medical examination items based on request information from the user and, thus, is explained as a simple model for the convenience of explanation.

More specifically, semantic information and the like such as is shown in Fig. 20 is recorded in main DB 14 in the present embodiment. Main DB 14 according to the present embodiment has semantic information that is implied by a polymorphism pattern in association with the polymorphism pattern of a predetermined polymorphism address as “annotative information on the polymorphism pattern.” In this “annotative information on the polymorphism pattern,” five value levels are set from the highest level “5,” which represents a high morbidity rate with respect to a disease classified in the “minor classification,” to “1,” which represents a low morbidity rate. When indicating the morbidity rate of the predetermined disease by a combination of the polymorphism pattern of the predetermined polymorphism address with a polymorphism pattern of other polymorphism address, a value “Z” is set in “annotative information on the polymorphism pattern” corresponding to the predetermined polymorphism address and a value “Y” is set in “annotative information on the polymorphism pattern.” corresponding to the other polymorphism address.

In the present embodiment, link DB 30 as shown in Fig. 21 is used, which represents a correspondence between a polymorphism address with the value “Z” being set in “annotative information on the polymorphism pattern” and a polymorphism address where the value “Y,” which is to be combined with the polymorphism address with the value “Z,” is set. In link DB 30, a possible polymorphism pattern (described as “pattern Z” in Fig. 21) of the polymorphism address with the value “Z” being set (described as “polymorphism address Z” in Fig. 21) is associated with a possible polymorphism pattern (described as “pattern Y” in Fig. 21) of the polymorphism address with the value “Y” being set (described as “polymorphism address Y” in Fig. 21) which is to be combined with the “polymorphism address Z”. Further, in link DB 30, the morbidity rate is associated with a combination of “pattern Z” and “pattern Y”. Link DB 30 may be stored in database 8 of shared computer 2 or may be provided externally to shared computer 2.

Main DB 14 described in the present embodiment may be controlled by, for example, a public organization or group. In this case, information that is not directly required when selecting medical examination items, which have been classified in a major classification according to, for example, “responsiveness to drug” or “drug side-effect,” is also included.

Meanwhile, in the present embodiment genome-related information recording medium 24 on which polymorphism addresses and polymorphism patterns as shown in Fig. 22 are recorded is exemplified and explained. Exemplified in Fig. 22 are a polymorphism address, which corresponds to the polymorphism address exemplified in main DB 14 shown in Fig. 20, and a polymorphism pattern thereof.

According to the system for processing information that provides medical examination items, the most suitable medical examination can be provided to the user in accordance with, for example, the flow charts shown in Figs. 23 and 24. In the flow charts shown in Figs. 23 and 24, a step described as “(shared)” refers to processing in shared computer 2 and a step described as “(personal)” refers to processing in personal computer 3.

In step III-1 (SIII-1), the requester first accesses shared computer 2 belonging to, for example, a medical examination center (hereinafter referred to as a “medical center”), from personal computer 3 through communication network 1. Examples of the requester include individuals who possess genome-related information recording medium 24, medical institutions that possess genome-related information recording medium 24, and medical institutions that utilize genome-related information recording medium 24 possessed by individuals.

When individuals possess genome-related information recording medium 24, a card-type recording medium can be exemplified as genome-related information recording medium 24 and the individuals, the medical institutions using genome-related information recording medium 24 of the individuals, or the like utilize the system as the requester. In contrast, when medical institutions possess genome-related information recording medium 24, examples of genome-related information recording medium 24 include a database-type recording medium having a plurality of pieces of genome-related information 28 of a plurality of individuals recorded thereon and the medical institutions utilize the system as the requester.

In the present embodiment, an individual possesses genome-related information recording medium 24 and the individual (the examinee) is the subject of a medical examination, while the medical institution that carries out the medical examination on the examinee is the requester. Thus, personal computer 3 belongs to a medical institution in the following description.

Shared computer 2 may be that of an agent or group for providing desired information on medical examination items and information associated therewith to a requester of information on medical examination items as well as that of a medical center.

In step III-2 (SIII-2), processing program 13 is started upon access by the requester and a “portal screen for medical examination” is read from screen memory 12 in shared computer 2 to display on display device 22 of personal computer 3. Examples of “a portal screen for medical examination” include a screen image as shown in Fig. 25.

In step III-3 (SIII-3), the requester loads genome-related information recording medium 24 into reading apparatus 25 in accordance with instructions on the “portal screen for medical examination,” thereby enabling access to genome-related information recording medium 24 using personal computer 3. In this case, whether or not the requester is an authorized user of genome-related information recording medium 24 is preferably confirmed. Thereafter, in step III-4 (SIII-4), the requester clicks an “OK” button displayed on the “portal screen for medical examination” to transmit their intent to utilize the system to shared computer 2.

In step III-5 (SIII-5), a “medical examination course menu” screen is read from screen memory 12 in accordance with processing program 13 in shared computer 2 and displayed on display device 22 of personal computer 3. Examples of the “medical examination course menu” screen include a screen image as shown in Fig. 26.

In step III-6 (SIII-6), in accordance with the “medical examination course menu” screen to which the requester has accessed with personal computer 3, the requester selects a desired medical examination course for the examinee based on the intent of the examinee. The present embodiment describes the case where an “entrusted check-up course” was
 5 selected from among the medical examination courses. The “entrusted check-up course” refers to a course that conducts a medical examination using a combination of examination items that are common for all courses and specific examination items selected according to the medical examinee’s diathesis and the like, i.e., examination items to be selected when the morbidity rate of a predetermined disease of the examinee is high.

10 Shared computer 2 has a “medical examination table” as a database, which has a series of examination items that were used in medical examinations previously accumulated therein. In this “medical examination table,” for example, as shown in Fig. 27, “major classification,” “middle classification,” and “minor classification” are associated with information on the examination method, examinee preparation and the like. “Major
 15 classification” categorizes examination items which are common among all medical examination courses as “basic” regardless of the level of morbidity rate, and examination items which are selected when the morbidity rate is high are categorized as “morbidity rate.” However, even though an examination item is classified as “basic,” it is also classified in the “morbidity rate” category when the morbidity rate is preferably judged prior to the medical
 20 examination from the viewpoint of guidance for improvement of living conditions after the medical examination. The “morbidity rate” in “major classification” in the medical examination table corresponds to the “morbidity rate” in “major classification” in main DB 14. Also, “middle classification” and “minor classification” in the medical examination table correspond to “middle classification” and “minor classification,” respectively, in main
 25 DB 14.

In step III-7 (SIII-7), reading apparatus 25 of personal computer 3 accesses genome-related information recording medium 24 to read out “Gno.” from data I. In step III-8 (SIII-8), “Gno.” read out in step III-7 and “medical examination course (entrusted check-up course)” (request information), which was selected in step III-6, are transmitted to shared
 30 computer 2 through communication network 1.

In step III-9 (SIII-9), shared computer 2 receives the “Gno.” and the “medical examination course,” and the “Gno.” and the “medical examination course” are stored in memory section A10. In the present embodiment, the “level of disclosure” is associated with semantic information that is recorded in main DB 14. The “level of disclosure” in this

embodiment indicates whether or not semantic information and information associated with the semantic information should be disclosed to the user of the information. More specifically, recorded, in association with semantic information, are: “level of disclosure A” designating a level that may be disclosed only when the requester has been authenticated;
 5 “level of disclosure B” designating a level that may be disclosed to any requester irrespective of whether the requester has been authenticated or not; and “prohibition against disclosure” (not shown in Fig. 20) designating a level that should not be disclosed to any requester.

Shared computer 2 comprises, for example, a database having “information regarding the authenticated requester such as ‘requester No.’ ” (not shown, hereinafter
 10 referred to as “requester information database”) previously recorded in database 8.

The requester transmits the “requester No.” to shared computer 2 together with “Gno.” and request information. When services using semantic information associated with “level of disclosure A” and information associated with the semantic information are to be provided to the requester, shared computer 2 accesses the “requester information database” to
 15 judge, based on the received “requester No.”, whether or not the requester has been authenticated.

When the requester is judged to be authenticated, shared computer 2 registers “level of disclosure A” in association with the “Gno.” received from personal computer 3 in storage 7. In the subsequent step, shared computer 2 searches the “level of disclosure” in
 20 main DB 14 based on the registered “level of disclosure A” and provides services using semantic information associated with “level of disclosure A” and information associated with the semantic information to the requester. On the other hand, when the requester is judged to be unauthenticated, shared computer 2 immediately cancels processing or registers “level of disclosure B” in association with the received “Gno.” in storage 7. In the subsequent step,
 25 shared computer 2 searches the “level of disclosure” in main DB 14 based on the registered “level of disclosure B” and provides services using semantic information associated with “level of disclosure B” and information associated with the semantic information to the requester. Thus, use of the “level of disclosure” enables the determination whether or not the disclosure of semantic information and information associated with the semantic information
 30 is approved.

In the present embodiment, because the system is one for providing medical examination items and examinee is an unspecified large number of people, the requesters need not to be authenticated previously. This system, therefore, eliminates the need to judge whether or not the requester has been authenticated. For this reason, in step III-9, upon the

receipt of request information (medical examination course), “level of disclosure B” is automatically selected and registered in storage 7 in association with the “Gno.” Examples of a case where “level of disclosure A” is selected include the case in which the requester is an authenticated medical institution or the like that intends to obtain all the information
 5 necessary for treating a patient (e.g., morbidity rate and responsiveness to medicaments).

Next, in step III-10 (SIII-10), main DB 14 is accessed in accordance with processing program 13. In step III-11 (SIII-11), from main DB 14, processing program 13 selects polymorphism address that is classified as “morbidity rate” in “major classification” and for which “level of disclosure B” has been set, and reads out the “polymorphism address,”
 10 “middle classification,” and “minor classification.” More specifically, since “entrusted check-up course” was received as request information in step III-9, examination items should be selected by judging the level of morbidity rate of the predetermined disease of the examinee in advance. In order to comply with the request information, “morbidity rate” classified in the major classification of main DB 14 (i.e. having “morbidity rate” as semantic
 15 information) is selected and the “polymorphism address” corresponding thereto is read out. That is to say, the “polymorphism address” corresponding to the request information is read out.

In step III-12 (SIII-12), the “polymorphism address,” “middle classification,” “minor classification,” and “Gno.”, which were read-out in accordance with the request information,
 20 are recorded in memory section B11. At this time, according to the present system, in step III-13 (SIII-13), an “anonymous polymorphism address” is set for the “polymorphism address” recorded in memory section B11 and linked with the “polymorphism address” in memory section B11, thereby recording the “anonymous polymorphism address” in memory section B11. This “anonymous polymorphism address” is of the same concept as the
 25 “anonymous polymorphism address” described in the second embodiment.

In step III-14 (SIII-14), in accordance with processing program 13, informed consent (hereinafter abbreviated to “IC”) concerning the submission of genome-related information
 28 and the content of semantic information and information associated with the semantic information is asked of the requester. Specifically, in step III-14, an “input screen for
 30 individual IC confirmation” is read from screen memory 12 in accordance with processing program 13 and information on the received “medical examination course” is incorporated therein. Then, the “input screen for individual IC confirmation” is displayed on display device 22 of personal computer 3. Examples of the “input screen for individual IC confirmation” include a screen image as shown in Fig. 28.

In step III-15 (SIII-15), in accordance with the “input screen for individual IC confirmation” displayed on display device 22 of personal computer 3, the requester obtains the consent of the examinee regarding the submission of the polymorphism patterns and the content of semantic information and information associated with the semantic information, and inputs the fact that the examinee has consented. This indicates that the examinee has consented to the submission of the polymorphism pattern and the content of semantic information and information associated with the semantic information. The consent to the submission of polymorphism patterns and the content of semantic information and information associated with the semantic information in step III-15 may be carried out by using the “medical examination portal” screen in step III-2 or the “medical examination course menu” screen in step III-5.

In step III-16 (SIII-16), the requester accesses data I of genome-related information recording medium 24 to read out “Gno.” In step III-17 (SIII-17), the read-out “Gno.” and the “individual IC confirmation” input in step III-15 are transmitted to shared computer 2.

In step III-18 (SIII-18), shared computer 2 receives the “Gno.” and “individual IC confirmation” transmitted from personal computer 3 and stores the received “individual IC confirmation” in memory section A10. When storing the “individual IC confirmation,” among the data stored in memory section A10, “individual IC confirmation” is associated with a “Gno.” matching with the received “Gno.”

In step III-19 (SIII-19), the “polymorphism address” and the “anonymous polymorphism address,” stored in steps III-12 and III-13, are read out from memory section B11. In step III-20 (SIII-20), the read-out “polymorphism address” is transmitted to personal computer 3 in association with the read-out “anonymous polymorphism address” and instruction information that instructs the submission of the polymorphism pattern corresponding to the “polymorphism address” and the “anonymous polymorphism address” associated with the polymorphism pattern is output to personal computer 3.

In step III-21 (SIII-21), the “polymorphism address” and “anonymous polymorphism address” transmitted from shared computer 2 as well as instruction information are received. At the same time, genome-related information recording medium 24 is accessed in accordance with the instruction information. At this time, the received “polymorphism address” and “anonymous polymorphism address” are stored in memory section 26. In step III-22 (SIII-22), in accordance with the instruction information, processing program 27 extracts and reads out the requested polymorphism address and the polymorphism pattern

corresponding to the polymorphism address from data II of genome-related information 28,
and reads out "Gno." from data I.

In step III-23 (SIII-23), the polymorphism pattern read-out in step III-22 is
transmitted in association with the anonymous polymorphism address corresponding to the
polymorphism address corresponding to the polymorphism pattern, together with the "Gno."
from personal computer 3 to shared computer 2. At this time, personal computer 3 transmits
the polymorphism pattern and the anonymous polymorphism address associated with the
polymorphism pattern, although personal computer 3 does not transmit any polymorphism
address associated with the polymorphism pattern. Since the anonymous polymorphism
address does not directly represent a position of the polymorphism pattern in the nucleotide
sequence, the position of the polymorphism pattern in the nucleotide sequence cannot be
determined even if the contents transmitted in step III-23 are leaked externally due to
unforeseen circumstances.

In step III-24 (SIII-24), the anonymous polymorphism address, the polymorphism
pattern associated with the anonymous polymorphism address and "Gno." transmitted from
personal computer 3 are received and the received polymorphism pattern is stored in memory
section B11. When storing the "polymorphism pattern," the "polymorphism pattern" is
associated with the "Gno." and "anonymous polymorphism address" in the data stored in
memory section B11 that match with the received "Gno." and "anonymous polymorphism
address".

In step III-25 (SIII-25), main DB 14 is accessed in accordance with processing
program 13. In step III-26 (SIII-26), main DB 14 is searched based on "middle
classification," "minor classification," "polymorphism address" and the "polymorphism
pattern" thereof stored in memory section B11, and the "annotative information on
polymorphism pattern" associated with the matching polymorphism pattern is read out (see
Fig. 20). The "annotative information on the polymorphism pattern" herein is semantic
information implying the morbidity rate of the disease classified in the "middle
classification" and the "minor classification" as shown in Fig. 20. When a plurality of pieces
of semantic information is associated with one polymorphism pattern, each piece of semantic
information is read out.

The "annotative information on the polymorphism pattern" in the present
embodiment includes five levels of values that indicate morbidity rate, a value "Z" that
indicates the morbidity rate of a predetermined disease by combining a polymorphism pattern

of a predetermined polymorphism address with the polymorphism pattern of an other polymorphism address, and a value “Y” that indicates the other polymorphism address.

In step III-26, when the read-out “annotative information on the polymorphism pattern” is value “Z,” processing program 13 accesses link DB 30 shown in Fig. 21.

5 Processing program 13 first searches link DB 30 based on the “polymorphism address” and the “polymorphism pattern” associated with value “Z,” and selects the other polymorphism address (polymorphism address Y) to be combined with the “polymorphism address Z.” Subsequently, based on the selected polymorphism address Y, the polymorphism pattern (pattern Y) of the polymorphism address Y is read from memory section B11. Link DB 30 is
10 then searched based on the read-out pattern Y of the polymorphism address Y to select the combination corresponded to the pattern Z of the polymorphism address Z and the pattern Y of the polymorphism address Y. The morbidity rate associated with the selected combination is read out from link DB 30.

In step III-27 (SIII-27), in accordance with processing program 13, the obtained
15 “annotative information on the polymorphism pattern,” that is, a morbidity rate, is stored in memory section B11 in association with the “polymorphism pattern.” Thus, as shown in Fig. 29, “annotative information on the polymorphism pattern” is associated with the “middle classification” and “minor classification” in memory section B11. Thus, for example, a morbidity rate of the disease indicated by “middle classification” and “minor classification”
20 is associated. In memory section A10, “Gno.,” “name of medical examination course,” and “IC confirmation” are stored in association with one another as shown in Fig. 30.

In step III-28 (SIII-28), in accordance with processing program 13, among the items in “middle classification,” “minor classification,” and “annotative information on the polymorphism pattern” stored in memory section B11, those items in which morbidity rate
25 exceeds a given level are read out. For example, “middle classification” and “minor classification” items for which the value for morbidity rate in the “annotative information on polymorphism pattern” is at least “4” with respect to the five levels, are read out.

In step III-29 (SIII-29), a medical examination table (Fig. 27) is searched based on the read-out “middle classification” and the “minor classification,” and those items
30 corresponded to the read-out “middle classification” and the “minor classification” are selected.

In step III-30 (SIII-30), essential items are extracted from the medical examination table. More specifically, examination methods, examination items, examinee preparation, inquiry items and the like, for the “middle classification” and the “minor classification”

selected in step III-29, are read out. Further, examination methods, examination items, examinee preparation, inquiry items, and the like, which are items classified into “basic” by the “major classification” in the medical examination table, are also read out.

In step III-31 (SIII-31), an output format as shown in, for example, Fig. 31 is prepared based on the examination methods, examination items, examinee preparation, inquiry items and the like, which were extracted in step III-30, and the output format data is transmitted to personal computer 3. The output format includes examination items that are common among all medical examination courses (classified into “basic” by the major classification in the medical examination table) and specific examination items concerning a disease for which the morbidity rate exceeds a given level (classified into “morbidity rate” by the major classification in the medical examination table). The examination items concerning diseases for which the morbidity rate exceeds a given level are “information associated with semantic information” induced from semantic information (morbidity rate).

This output format is used in a medical institution conducting the medical examination on the examinee. On the other hand, an output format for an examinee receiving a medical examination on which the examination classification and the examinee preparation are displayed, may be output. In this case, information such as the method and purpose of the examination from which the examinee can ascertain his or her own morbidity rate is not disclosed to the examinee. Thus, the imparting of anxiety to the examinee can be prevented. Also in this case, a medical institution conducting the medical examination can suitably select examining methods for the disease in such a manner that, for example, the examinee would not know what disease is subject of examination.

In step III-31, the output format may be directly transmitted to the examinee instead of being transmitted to a medical institution conducting the medical examination. In this case, the output format transmitted to the examinee does not preferably contain any information that would impart anxiety to the examinee, such as a morbidity rate of a predetermined disease. However, information that may impart anxiety to the examinee can be disclosed in accordance with IC.

In step III-32 (SIII-32), personal computer 3 receives the output format data that was output by shared computer 2 and stores it in memory section 26. Then, in step III-33 (SIII-33), by outputting the output format, the requester can obtain information on medical examination items including specific examination items that are appropriate for the diathesis of the individual possessing genome-related information recording medium 24.

In shared computer 2, in step III-34 (SIII-34) subsequent to step III-31, the contents stored in memory section A10 and memory section B11 are separately stored in storage DB-A15 and storage DB-B16, respectively, and the contents stored in memory section A10 and memory section B11 may then be deleted. At this time, the “anonymous polymorphism address” among the contents stored in memory section B11 is preferably deleted.

As described above, with this system, by following the flow charts shown in Figs. 23 and 24, the requester can obtain information on medical examination items based on the individual’s (the examinee’s) diathesis and the like. The column for “category” on the output format shown in Fig. 31 includes “basic,” which implies examination items that are common among predetermined medical examination courses, and “specific,” which implies examination items for diseases having a high morbidity rate that were obtained in step III-29. Through utilization of this system, medical examination services including “specific” examination items that are appropriate for the individual can be provided.

Further, in the system, even when semantic information is associated with a combination of polymorphism patterns in a plurality of polymorphism addresses, using link DB 30, semantic information which is specific to the individual possessing genome-related information recording medium 24 can be obtained.

In the system, the output format may be prepared by constructing a database in which additional information regarding the medical examinees, such as the results of past medical examination, has been accumulated as a history, reading the history of the results of past medical examination of the examinee from the database, and associating the history with the “annotative information on the polymorphism pattern” and information associated therewith. This database may be provided in shared computer 2 or may be provided in personal computer 3 or genome-related information recording medium 24. According to the system, the output format shows the history of the results of medical examination of the examinee, and thus, comparison with the results of medical examination thereafter enables the provision of enhanced medical examination services. The results of medical examination, for example, include the presence or absence of oxidative damage in genes or gene expression found by genetic testing in addition to the results of conventional medical examination. Further, in the system, in step III-9, as request information in addition to the desired “medical examination course,” for example, a request for “functional food that prevents diseases in respect of which the morbidity rate of the examinee is high” is received, and the requested functional food can be provided together with information on medical examination items based on the examinee’s diathesis.

In the present embodiment, the system for providing medical examination services including examination items that are “specific” for the examinee in which the requester utilizes a medical center having main DB 14 was described although the system is not limited thereto. The system may comprise personal computer 3 belonging to the examinee who
5 handles genome-related information 28, a medical center having a database in which polymorphism addresses are recorded in a manner capable of corresponding to the request information, and organizations or groups other than the medical center having main DB 14. The examinee may be the requester in the system for processing information. In this case, the examinee accesses the medical center to request the medical examination services, and the
10 medical center accesses main DB 14 of the other organization, to provide medical examination services containing “specific” examination items appropriate for the examinee (requester). More particularly, the other organization having main DB 14 provides semantic information and information associated with semantic information without handling genome-related information 28 and the medical center handles genome-related information 28 without
15 possessing main DB 14. In this case, main DB 14 of the other organization can be utilized in a plurality of medical centers. Further, the other organization performs control, updating, or the like of main DB 14 and the medical center handles genome-related information 28 of the requester. This enables the requester to receive the medical services through the medical center. In this case, medical examination items, which contain the “specific” examination
20 items appropriate for the examinee are transmitted to the examinee through the medical center. However, it may not be limited to this and the medical examination items may be directly transmitted from the other organization to the examinee. Alternatively, the medical examination items may be transmitted to the third organization actually conducting the medical examination from the other organization.

As described above, according to the system, on genome-related information recording medium 24 and in main DB 14, standardization of only “polymorphism addresses” and the “polymorphism patterns” thereof eliminates the need for standardization of other specific data. Thus, the system can be utilized in a wide range of industries. That is, when providing information using genome-related information recording medium 24, the provider
30 of objects or services can provide information in various manners without the need to standardize semantic information to correspond to the polymorphism pattern or a unified standard such as a method for transmitting/receiving data.

Furthermore, according to the system, a third party or third organization can easily monitor and control shared computer 2 by examining main DB 14. Accordingly, as the

system can, for example, execute administrative control over the provider of semantic information, adequate and ethical control over the provider of semantic information can be executed.

4. Fourth embodiment

Next, a system for donor registration for registering an individuals' own polymorphism pattern based on solicitation for a predetermined purpose, and a system for processing information that provides information on other individual organisms having properties compatible with a "given property" of the requester in relation to the predetermined purpose are described as the fourth embodiment to which the present invention has been applied. For the convenience of explanation, it is described as a simple model.

Regarding the system for processing information explained as the present embodiment, the explanation of its constitution, operation, and terms thereof is omitted by employing the same appellations, symbols, and definitions for the same constitution and terms as with the system for processing information according to the above described the first, the second and the third embodiments. More particularly, semantic information (an individual property type, a compatible type) or the like as shown in Fig. 32 is recorded in main DB 14 according to the present embodiment. In main DB 14, the "polymorphism address" and the "polymorphism pattern" in the polymorphism address are recorded in association with each other depending on the "classification" for categorizing a predetermined property. In main DB 14 is also recorded an "individual property type" implied by a predetermined "polymorphism pattern" (information on a predetermined individual organism) and a "compatible type" indicating a compatible type with the "individual property type" (information on an other individual organism) as semantic information. The "level of disclosure" defining whether information on the "individual property type" and the "compatible type" in the predetermined polymorphism address can be disclosed or not may be further recorded in main DB 14 in association with this information.

More specifically, polymorphism addresses "1000" and "2000" are associated with the "given property," and the individual property type implied by a possible combination of polymorphism patterns in these polymorphism addresses is recorded as "a" and "b." The compatible types of these individual property types "a" and "b" are recorded as "b" and "a," respectively. That is to say, Fig. 32 demonstrates that, for example, when polymorphism patterns in polymorphism addresses "1000" and "2000" are respectively "A" and "C," the

individual property type regarding the “given property” is “a,” and the compatible type that is compatible to individual property type “a” is “b.”

The present embodiment is explained by using genome-related information recording medium 24 having polymorphism addresses and polymorphism patterns as shown in Fig. 33 recorded thereon as an example. Exemplified in Fig. 33 is genome-related information recording medium 24 containing polymorphism addresses and polymorphism patterns thereof corresponding to the polymorphism addresses (“1000” and “2000”) exemplified in main DB 14 in Fig. 32.

First, a system for donor registration for registering an individual’s own polymorphism pattern based on solicitation for a predetermined purpose is explained. In the system for donor registration, for example, as in accordance with the flow chart shown in Fig. 34, a donor database (hereinafter referred to as a “donor DB”) for a predetermined purpose such as shown in Fig. 35 can be constructed. In the flow chart shown in Fig. 34, a step described as “(shared)” refers to processing in shared computer 2 and a step described as “(personal)” refers to processing in personal computer 3.

In the system, processing program 27, which is recorded in storage 23 of personal computer 3, is first started based on solicitation for donors in step D-1 (SD-1). In step D-1, processing program 27 drives reading apparatus 25 of personal computer 3 and accesses genome-related information recording medium 24 to read out a “Gno.” recorded thereon as data I. The read-out “Gno.” is stored in memory section 26.

In step D-2 (SD-2), the read-out “Gno.” is transmitted together with a “given property,” which is registration-request information, through communication network 1 to shared computer 2 belonging to the party soliciting donors. This enables declaration of intention (i.e., request for services) to register as a donor for the “given property,” using the system. At this time, strict enforcement of obtainment of informed consent (hereinafter abbreviated to “IC”) is preferably conducted regarding the purpose of donor registration and its content and the method and range of utilization thereof.

In step D-3 (SD-3), shared computer 2 receives the “Gno.” and “given property” transmitted from personal computer 3. After reception of the “Gno.” and “given property,” shared computer 2 starts processing program 13. The received “Gno.” and “given property” are stored in memory section A10.

In step D-4 (SD-4), main DB 14 is accessed in accordance with processing program 13. In step D-5 (SD-5), main DB 14 is searched based on the “given property” as request

information in accordance with processing program 13 to extract the classification corresponded to the “given property.”

In step D-6 (SD-6), the “polymorphism address” associated with the extracted “given property” is read from main DB 14 in accordance with processing program 13. The read-out “polymorphism address” is stored in memory section A10.

In step D-7 (SD-7), “Gno.” and “polymorphism address” stored in memory section A10 are read out in accordance with processing program 13, “Gno.” and “polymorphism address” are transmitted to personal computer 3, and instruction information instructing submission of the “polymorphism pattern” corresponding to the transmitted “polymorphism address” is transmitted to personal computer 3. In this case, the submission of additional information such as anamnesis and characteristics may be optionally instructed within a range regulated by the IC.

In step D-8 (SD-8), the “Gno.” and “polymorphism address” transmitted from shared computer 2 are received. The received “Gno.” and “polymorphism address” are recorded in memory section 26. In step D-9 (SD-9), data II which is recorded on genome-related information recording medium 24 is accessed in accordance with the received instruction information (“Gno.” and “polymorphism address”).

In step D-10 (SD-10), data II on genome-related information recording medium 24 is searched to extract the instructed polymorphism address. In step D-11 (SD-11), data II on genome-related information recording medium 24 is searched in accordance with the extracted polymorphism address to read out the corresponding polymorphism pattern. The read-out polymorphism pattern is recorded in association with the polymorphism address in memory section 26. At this time, data I is preferably accessed to confirm whether the “Gno.” contained in instruction information is correct or not. In step D-10, additional information recorded in data III, data IV, and data V may be simultaneously read out in addition to the polymorphism pattern to record in memory section 26 if necessary.

In step D-12 (SD-12), the temporarily-recorded polymorphism pattern associated with the polymorphism address and the optionally-recorded additional information in memory section 26 are output to shared computer 2 through communication network 1 together with the “Gno.” In step D-13 (SD-13), shared computer 2 receives the polymorphism pattern associated with the polymorphism address, the Gno., and the optionally-recorded additional information. The received polymorphism pattern is recorded in memory section A10 in association with the polymorphism address.

In step D-14 (SD-14), the polymorphism pattern and the Gno., which are recorded in memory section A10, and additional information are associated with the “given property” and then registered in the donor DB as shown in Fig. 35. In this donor DB, the polymorphism address associated with a given property and the polymorphism pattern of the polymorphism address are recorded for each “Gno.”

A donor DB is provided in, for example, database 8 (memory) of shared computer 2 although it is not limited thereto. For example, a donor DB may be provided in a memory external to shared computer 2.

As described above, donor registration can be very easily carried out according to the system without the necessity to conduct any examination, investigation, or test when soliciting for the predetermined purpose. In particular, in the system, when performing donor registration in accordance with the flow chart shown in Fig. 34, all of genome-related information 28 which has been recorded on genome-related information recording medium 24 is not necessarily output through communication network 1. Instead, only the part of genome-related information 28 for which submission has been instructed may be output. According to the system, therefore, donor registration can be carried out while preventing leakage of highly confidential polymorphism addresses and polymorphism patterns that are peculiar to individuals. Also in the system, use of the “anonymous polymorphism address” can further prevent leakage of polymorphism addresses and polymorphism patterns.

Meanwhile, in the system, a donor DB may comprise a polymorphism address and a polymorphism pattern associated with the “given property” as well as semantic information implied by the polymorphism pattern recorded therein. More specifically, the “individual property type” and the “compatible type,” which are recorded in main DB 14, may be recorded in the donor DB in association with the “Gno.” or “polymorphism pattern.”

Next, a system for processing information that provides information on other individual organisms having properties compatible with the “given property” of the requester (referred to as a “recipient” in the present embodiment) for a predetermined purpose is described. This system for processing information utilizes main DB 14 shown in Fig. 32 and the donor DB shown in Fig. 35 and can provide information on other individual organisms in accordance with, for example, the flow charts shown in Figs. 36 and 37. In the flow charts shown in Figs. 36 and 37, a step described as “(shared)” refers to processing in shared computer 2 and a step described as “(personal)” refers to processing in personal computer 3.

In the system, a recipient first starts processing program 27 recorded in storage 23 of personal computer 3 in step R-1 (SR-1) in order to obtain information on other individual

organisms that is compatible with the recipient's "given property." In step R-1, processing program 27 drives reading apparatus 25 of personal computer 3 and accesses genome-related information recording medium 24 to read out a "Gno." recorded as data I on genome-related information recording medium 24. The read-out "Gno." is stored in memory section 26.

5 In step R-2 (SR-2), the read-out "Gno." and the "given property" that is the request information are transmitted through communication network 1 to shared computer 2 of the provider of information. This enables declaration of intention to request information on other individual organisms which are compatible with respect to the "given property," utilizing the system. At this time, strict enforcement of obtainment of IC is preferably conducted
10 regarding acceptable use regulations for information on the polymorphism pattern and the compatible other individual organisms.

In step R-3 (SR-3), shared computer 2 receives the "Gno." and "given property" transmitted from personal computer 3. After reception of the "Gno." and "given property," shared computer 2 starts processing program 13. The received "Gno." and "given property"
15 are stored in memory section A10.

In step R-4 (SR-4), main DB 14 is accessed in accordance with processing program 13. In step R-5 (SR-5), main DB 14 is searched in accordance with processing program 13 and based on the "given property," and the classification matching with the "given property" is extracted.

20 In step R-6 (SR-6), the "polymorphism address" associated with the extracted classification "given property" is read from main DB 14 in accordance with processing program 13. The read-out "polymorphism address" is stored in memory section A10. More specifically, in the case of main DB 14 shown in Fig. 32, "1000" and "2000" are extracted and stored as polymorphism addresses associated with the "given property."

25 In step R-7 (SR-7), "Gno." and "polymorphism address" stored in memory section A10 are read out in accordance with processing program 13, the "Gno." and the "polymorphism address" are transmitted to personal computer 3, and instruction information instructing the submission of the "polymorphism pattern" corresponding to the transmitted "polymorphism address" is transmitted to personal computer 3. At this time, the submission
30 of additional information such as anamnesis and characteristics may be optionally instructed within a range regulated by IC.

In step R-8 (SR-8), personal computer 3 receives the "Gno." and the "polymorphism address" transmitted from shared computer 2. The received "Gno." and "polymorphism address" are recorded in memory section 26. In step R-9 (SR-9), data II recorded on genome-

related information recording medium 24 is accessed in accordance with the received instruction information (“Gno.” and “polymorphism address”).

In step R-10 (SR-10), data II on genome-related information recording medium 24 is searched and the instructed polymorphism address is extracted. In step R-11 (SR-11), data II
5 on genome-related information recording medium 24 is searched based on the extracted polymorphism address to read out the corresponding polymorphism pattern. The read-out polymorphism pattern is recorded in memory section 26 in association with the polymorphism address. In this case, data I is preferably accessed to confirm whether the “Gno.” contained in the instruction information is correct or not. Also, in step R-10,
10 additional information recorded in data III, data IV and data V is simultaneously read out in addition to the polymorphism pattern and it may be recorded in memory section 26 if necessary. More specifically, in the case of genome-related information recording medium 24 shown in Fig. 33, polymorphism patterns “A (corresponding to polymorphism address 1000)” and “C (corresponding to polymorphism address 2000)” are read out in compliance
15 with the instructed polymorphism address.

In step R-12 (SR-12), the temporarily-recorded polymorphism pattern associated with the polymorphism address and the optionally-recorded additional information in memory section 26 are output to shared computer 2 through communication network 1 together with the “Gno.” In step R-13 (SR-13), shared computer 2 receives the
20 polymorphism pattern associated with the polymorphism address, the Gno., and the optionally-recorded additional information. The received polymorphism pattern is recorded in memory section A10 in association with the polymorphism address. More specifically, polymorphism pattern “A” corresponding to polymorphism address “1000” and polymorphism pattern “C” corresponding to polymorphism address “2000” are recorded in
25 memory section A10.

In step R-14 (SR-14), main DB 14 is accessed in accordance with processing program 13. In step R-15 (SR-15), main DB 14 is searched based on the received polymorphism address and the polymorphism pattern thereof to extract the “individual property type” associated with the matched polymorphism pattern (information on the
30 predetermined individual organisms). Specifically, in the case of Fig. 32, the individual property type “a” is extracted for the combination in which the polymorphism patterns of polymorphism addresses “1000” and “2000” are “A” and “C” respectively.

In step R-16 (SR-16), main DB 14 is searched based on the “individual property type” extracted in step R-15, and the “compatible type” being compatible to the “individual

property type" (information on other individual organisms) is extracted. More specifically, in the case of Fig. 32, since the individual property type is "a," "b" is extracted as the compatible type for the individual property type "a."

In step R-17 (SR-17), main DB 14 is searched based on the extracted compatible type, and an individual property type which is the same as the compatible type is searched. The polymorphism pattern, associated with the searched individual property type, and the polymorphism address thereof are read out. Specifically, in the case of Fig. 32, since the extracted compatible type is "b," the polymorphism pattern having "b" as the individual property type and the polymorphism address thereof are read out. That is, in the case of Fig. 32, a combination in which the polymorphism pattern of polymorphism address "1000" is "G" and the polymorphism pattern of polymorphism address "2000" is "C" and a combination in which the polymorphism pattern of polymorphism address "1000" is "A" and the polymorphism pattern of polymorphism address "2000" is "T" are read out.

In step R-18 (SR-18), a donor DB is accessed in accordance with processing program 13. The donor DB (for example, as shown in Fig. 35) is constructed as described above and polymorphism addresses and the polymorphism pattern thereof associated with a "given property" for a plurality of individual organisms are recorded.

In step R-19 (SR-19), the donor DB is searched in accordance with processing program 13 based on the polymorphism address and the polymorphism pattern thereof that were read out in step R-17, and a "Gno." (information for discriminating an individual organism) having a combination of the polymorphism address and the polymorphism pattern matching with the combination of the polymorphism address and the polymorphism pattern thereof, which were read out in step R-17, is extracted. More specifically, in the case of Fig. 35, when searching for a "Gno." having combination in which "1000" is "G" and "2000" is "C" and a "Gno." having combination in which "1000" is "A" and "2000" is "T," "Gno." 0003 and "Gno." 0004 can be extracted. The extracted "Gno." is information associated with semantic information, that was induced from semantic information.

In step R-20 (SR-20), the extracted "Gno." is transmitted to personal computer 3. At this time, the "Gno." and other information may be transmitted to personal computer 3 within the range regulated by the IC made by the donor registrant. Examples of other information include information that was recorded when constructing the donor DB shown in Fig. 35, for example, additional information recorded in data III, data IV, and data V on genome-related information recording medium 24. More specifically, in the case of Fig. 35, "Gno." 0003 and "Gno." 0004 are transmitted to personal computer 3.

In step R-21 (SR-21), personal computer 3 receives the “Gno.” and other information transmitted from shared computer 2. In step R-22 (SR-22), the received “Gno.” and other information are output, thereby obtaining information on other individual organisms having properties compatible with the “given property.” That is, the recipient can obtain information on other individual organisms which is compatible with respect to the recipient’s “given property.” In the case of Fig. 35, the recipient can obtain information associated with semantic information, i.e., that other individual organisms which are compatible with respect to the “given property” are “Gno.” 0003 and “Gno.” 0004. In step R-20, the results of extraction may be simultaneously transmitted to an other organization providing objects or services to the requester using the result of extraction.

As described above, information on an other individual organism which has a property compatible with the recipient’s “given property” for a predetermined purpose can be obtained in accordance with the flow charts shown in Figs. 36 and 37 according to the system. More particularly, in this case, all of genome-related information 28 recorded on genome-related information recording medium 24 is not necessarily output through communication network 1, and instead, only the part of genome-related information 28 for which submission has been instructed may be output. According to the system, therefore, leakage of highly confidential polymorphism addresses and polymorphism patterns peculiar to the individuals can be prevented. Also in the system, use of the “anonymous polymorphism address” can further prevent leakage of the polymorphism addresses and polymorphism patterns.

In the system, since a classification such as “given property” and “compatible type,” which is compatible with the individual property type, are corrected and increases in the number of its type, a more accurate and wider range of information is included through updating of main DB 14. In addition, if many individual organisms are registered as donors in accordance with the flow charts shown in Fig. 34, a large amount of donor information is accumulated in the donor DB. According to the system, updating of main DB 14 in compliance with such increase, correction and the like of information and increase in the amount of information in the donor DB enable the recipient to utilize the newest information.

Furthermore, according to the system, a third party or third organization can easily monitor and control shared computer 2 by examining main DB 14 and donor DB. Accordingly, as the system can, for example, execute administrative control over the provider of information on other individual organisms, adequate and ethical control over the provider of information on other individual organisms can be executed.

When the donor DB has semantic information implied by the polymorphism pattern recorded therein, shared computer 2 searches main DB 14 based on, for example, the polymorphism pattern and the Gno. received in step R-13 (SR-13) to extract the “individual property type” and the “compatible type” of the recipient (step R-15). Shared computer 2 searches the “individual property type” of other individual organisms recorded in the donor DB based on the “individual property type” and “compatible type” of the recipient without conducting steps R-16 to R-19, and extracts other individual organisms having the “individual property type” that matches with the recipient’s “compatible type.” Subsequently, conducting steps R-20 to R-22, the recipient can obtain information on other individual organisms that are compatible with the recipient regarding the “given property.” In this case, in step R-20, the results of extraction may be simultaneously transmitted to another organization for providing objects or services to the requester using the results of extraction.

Further in the system, a donor registrant utilizing the system for donor registration can obtain information on other individual organisms as a recipient. In this case, donor registration is carried out through steps D-1 to D-14 shown in Fig. 34 and then advanced to step R-14, and information on other individual organisms can be obtained for the “given property” in the same manner as described. In this case, the user performs donor registration for a “given property” and at the same time can obtain information on other individual organisms for the “given property.” When an individual organism which has conducted donor registration afterwards becomes a recipient, shared computer 2 accesses the donor DB after steps R-1 and R-2 and the “polymorphism address” matching with the “Gno.” of the recipient and the “polymorphism pattern” thereof are read out, and the process then advances to step R-14. Information on other individual organisms for the “given property” can be obtained in the same manner as described. Also in this case, the user performs donor registration for a “given property” and at the same time can obtain information on other individual organisms for the “given property.”

As described above, according to the system, on genome-related information recording medium 24 and in main DB 14, standardization of only “polymorphism addresses” and the “polymorphism patterns” thereof eliminates the need for standardization of other specific data. Thus, the system can be utilized in a wide range of industries. That is, when providing information using genome-related information recording medium 24, the provider of objects or services can provide information in various manners without the need to

standardize semantic information to correspond to the polymorphism pattern or a unified standard such as a method for transmitting/receiving data.

In the above fourth embodiment, the system for processing information for obtaining information on other individual organisms having a “compatible type” compatible with an “individual property type,” as the “individual property type,” was explained. The system for processing information, however, is not limited to such embodiment. For example, the system may be directed to obtaining information on other individual organisms having a “compatible type” that is incompatible with an “individual property type” as the “individual property type.” In this case, an “individual property type” implied by a predetermined “polymorphism pattern” (information on a predetermined individual organism) and an “incompatible type” (information on an other individual organism) indicating a type incompatible with the “individual property type” are recorded as semantic information in main DB 14.

EFFECT OF THE INVENTION

As is apparent from the foregoing detailed description, a system for processing information can be constructed according to the present invention in which differences in information on nucleotide sequences among individual organisms are effectively utilized in order to provide semantic information and/or information associated with the semantic information that is useful for each individual. More particularly, the present invention enables use of the newest semantic information through updating, correction or the like of semantic information. Therefore, the present invention can provide high-quality objects and/or services relating to request information.

What is claimed is:

1. A computer-implemented method for processing information on a nucleotide sequence comprising the steps of:
 - 5 (a) receiving, via a communication network, request information of an object or service, wherein the object or service is provided for an individual based on one or more genetic differences between individuals, and wherein the request information does not include genetic information;
 - (b) searching a first memory area for positional information, and wherein the
 - 10 positional information corresponds to information on classification of the object or service, and wherein the positional information represents a position in a nucleotide sequence, and retrieving the positional information, wherein the first memory area is permitted to be accessed by a first processor;
 - (c) transmitting, via a communication network, the positional information
 - 15 retrieved in step (b) to a second processor which is permitted to access a second memory area storing polymorphism information regarding the individual;
 - (d) receiving, via a communication network, polymorphism information based on the positional information transmitted in step (c); and
 - (e) outputting, via a communication network, the polymorphism information
 - 20 received in step (d) to a user, and wherein the user includes a third processor which is permitted to access a third memory area storing semantic information, and which searches the third memory area to retrieve, in response to said transmission of the polymorphism information, semantic information based on the transmitted polymorphism information, and outputs the semantic information to a device that
 - 25 utilizes the semantic information or information corresponding to the semantic information, which is used for a provision of the object or service substantially included in the request information, wherein steps (a)-(e) are conducted under the control of the first processor.
 - 30 2. The method for processing information on a nucleotide sequence according to claim 1, further comprising the steps of:
 - in advance of step (c), receiving, via a communication network, consent

information regarding the provision of polymorphism information from a provider of the request information; or in advance of step (e), receiving, via a communication network, consent information regarding the content of semantic information or information corresponding to the semantic information from the provider of the request
5 information.

3. The method for processing information on a nucleotide sequence according to claim 1, wherein in step (c), secondary positional information corresponding to the positional information retrieved in step (b) is set and the positional information retrieved
10 in step (b) is transmitted along with the secondary positional information, and in step (d), the polymorphism information is received along with the secondary positional information.

4. A computer-readable medium encoded with a program for processing
15 information on a nucleotide sequence which allows a computer to execute processes including:

(a) receiving, via a communication network, request information of an object or service, wherein the object or service is provided for an individual based on one or more genetic differences between individuals, and wherein the request information does not
20 include genetic information;

(b) searching a first memory area for positional information, and wherein the positional information corresponds to information on classification of the object or service, and wherein the positional information represents a position in a nucleotide sequence, and retrieving the positional information, wherein the first memory area is
25 permitted to be accessed by a first processor;

(c) transmitting, via a communication network, the positional information retrieved in process (b) to a second processor which is permitted to access a second memory area storing polymorphism information regarding the individual;

(d) receiving, via a communication network, polymorphism information based
30 on the positional information transmitted in process (c); and

(e) outputting, via a communication network, the polymorphism information received in process (d) to a user, and wherein the user includes a third processor which

is permitted to access a third memory area storing semantic information, and which searches the third memory area to retrieve, in response to said transmission of the polymorphism information, semantic information based on the transmitted polymorphism information, and outputs the semantic information to a device that
 5 utilizes the semantic information or information corresponding to the semantic information, which is used for a provision of the object or service substantially included in the request information, wherein processes (a)-(e) are conducted under the control of the first processor.

10 5. The computer-readable medium according to claim 4, wherein the processes further comprise:

in advance of process (c), receiving, via a communication network, consent information regarding the provision of polymorphism information from a provider of the request information; or in advance of process (e), receiving, via a communication
 15 network, consent information regarding the content of semantic information or information corresponding to the semantic information from the provider of the request information.

6. The computer-readable medium according to claim 4, wherein in process (c),
 20 secondary positional information corresponding to the positional information retrieved in process (b) is set and the positional information retrieved in process (b) is transmitted along with the secondary positional information, and in process (d), the polymorphism information is received along with the secondary positional information.

25 7. An apparatus for processing information on a nucleotide sequence comprising:
 a transmitter/receive in communication with a communication network that receives request information of an object or service via the communication network, the object or service being provided for an individual based on one or more genetic differences between individuals, wherein the request information does not include
 30 genetic information;

a controller having a first processor that searches a first memory area for positional information that corresponds to information on classification of the object or

service, the positional information representing a position in a nucleotide sequence, and retrieves the positional information, wherein the transmitter/receiver transmits, via the communication network, the retrieved positional information to a second processor which is permitted to access a second memory area storing polymorphism information
 5 regarding the individual, and the controller, followed by receiving, via the communication network, the polymorphism information based on the transmitted positional information, outputs, via the communication network, the received polymorphism information to a user, and

wherein the user includes a third processor which is permitted to access a third
 10 memory area storing semantic information, and which searches the third memory area to retrieve, in response to said transmission of the polymorphism information, semantic information based on the transmitted polymorphism information, and outputs the semantic information to a device that utilizes the semantic information or information corresponding to the semantic information, which is used for a provision of the object or
 15 service substantially included in the request information.

8. The apparatus for processing information on a nucleotide sequence according to claim 7 wherein the transmitter/receiver performs one of:

(i) prior to transmitting, receiving, via the communication network, consent
 20 information regarding provision of the polymorphism information from a provider of the request information and

(ii) prior to outputting, receiving, via the communication network, consent
 information regarding the content of semantic information or information corresponding to the semantic information from the provider of the request information.

25

9. The apparatus for processing information on a nucleotide sequence according to claim 1 wherein the controller sets secondary positional information corresponding to the retrieved positional information and the transmitter/receiver transmits, via the communication network, the retrieved positional information along with the secondary
 30 position information and the controller receives, via the communication network, the polymorphism information is along with the secondary positional information.

Fig. 1

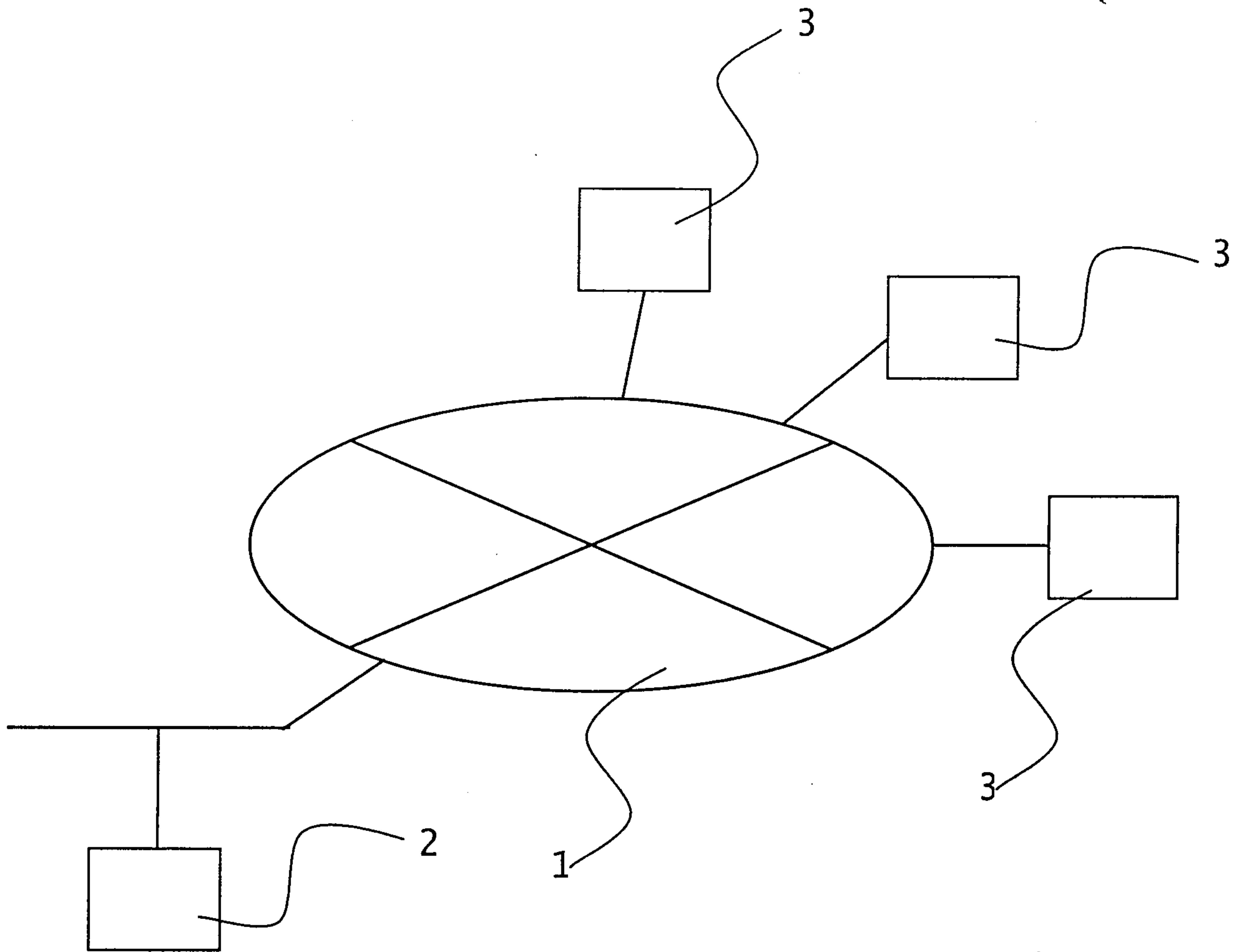


Fig. 2

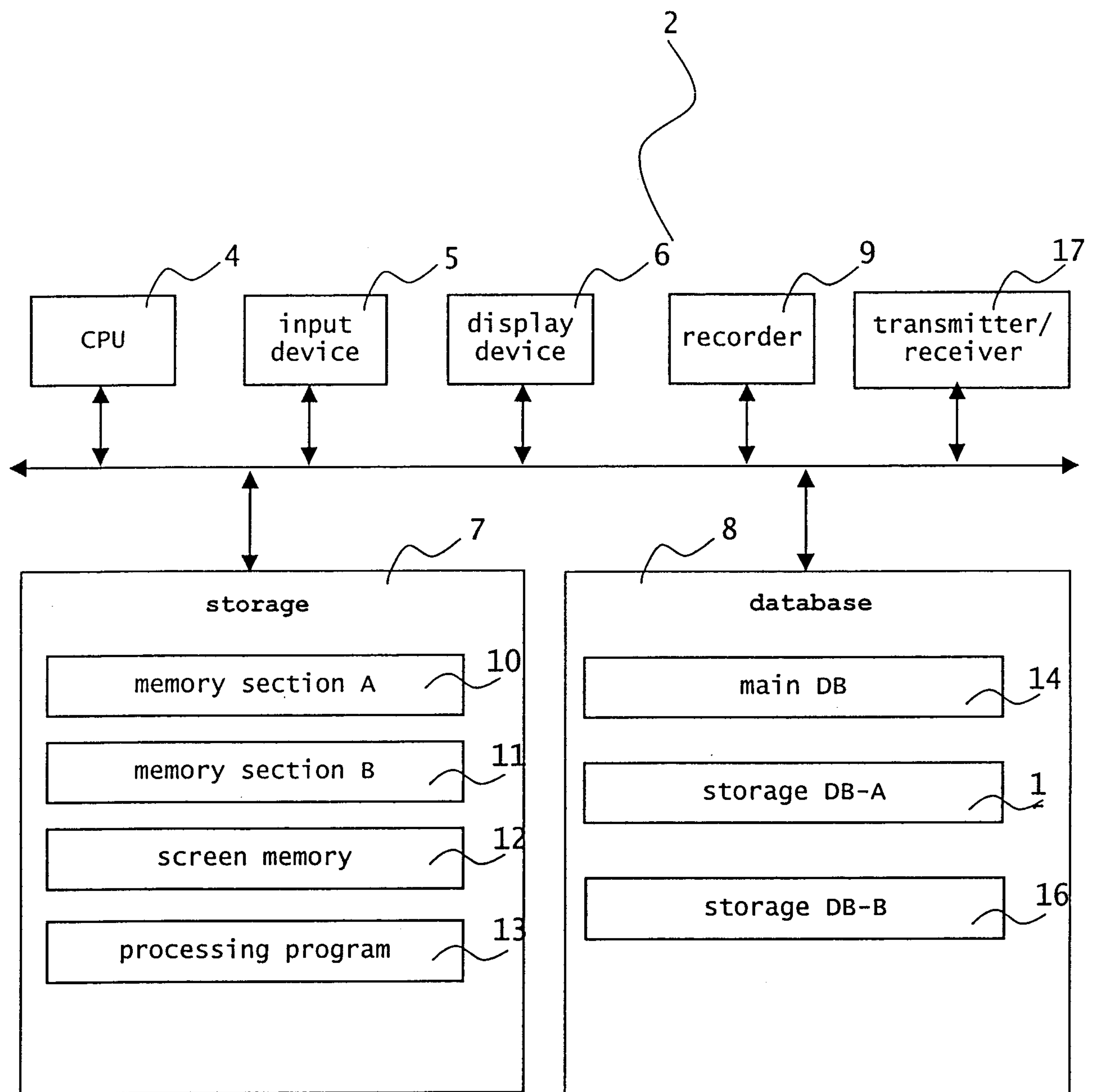


Fig. 3

Polymorphism address		Polymorphism type	Polymorphism pattern	Classification (name of disease)	Annotative information on polymorphism pattern (morbidity rate)		Level of disclosure (disclosability)
123456		SNP	A	hypertension	a		O
123456		SNP	G	hypertension	b		O
223456		SNP	G	large-boowel cancer	(i)		O
223456		SNP	A	large-bowel cancer	(ii)		O
234567		SNP	G	stomach cancer	c		O
234567		SNP	A	stomach cancer	d		O
334567		SNP	A	asthma	(iii)		O
334567		SNP	G	asthma	(iv)		O
345678		SNP	C	diabetes	e		O
345678		SNP	T	diabetes	f		O
445678		SNP	T	lung cancer	(v)		O
445678		SNP	C	lung cancer	(vi)		O
456789		SNP	T	pollinosis	g		O
456789		SNP	C	pollinosis	h		O
456789		SNP	T	hypertension	i		O
456789		SNP	C	hypertension	j		O
:	:	microsatellite	14 times	inmedicable disease	—	:	x
:	:	microsatellite	9 times	inmedicable disease	—	:	x
:	:	deletion	G	:	:	:	O
:	:	deletion	deletion	:	:	:	O

Fig. 4

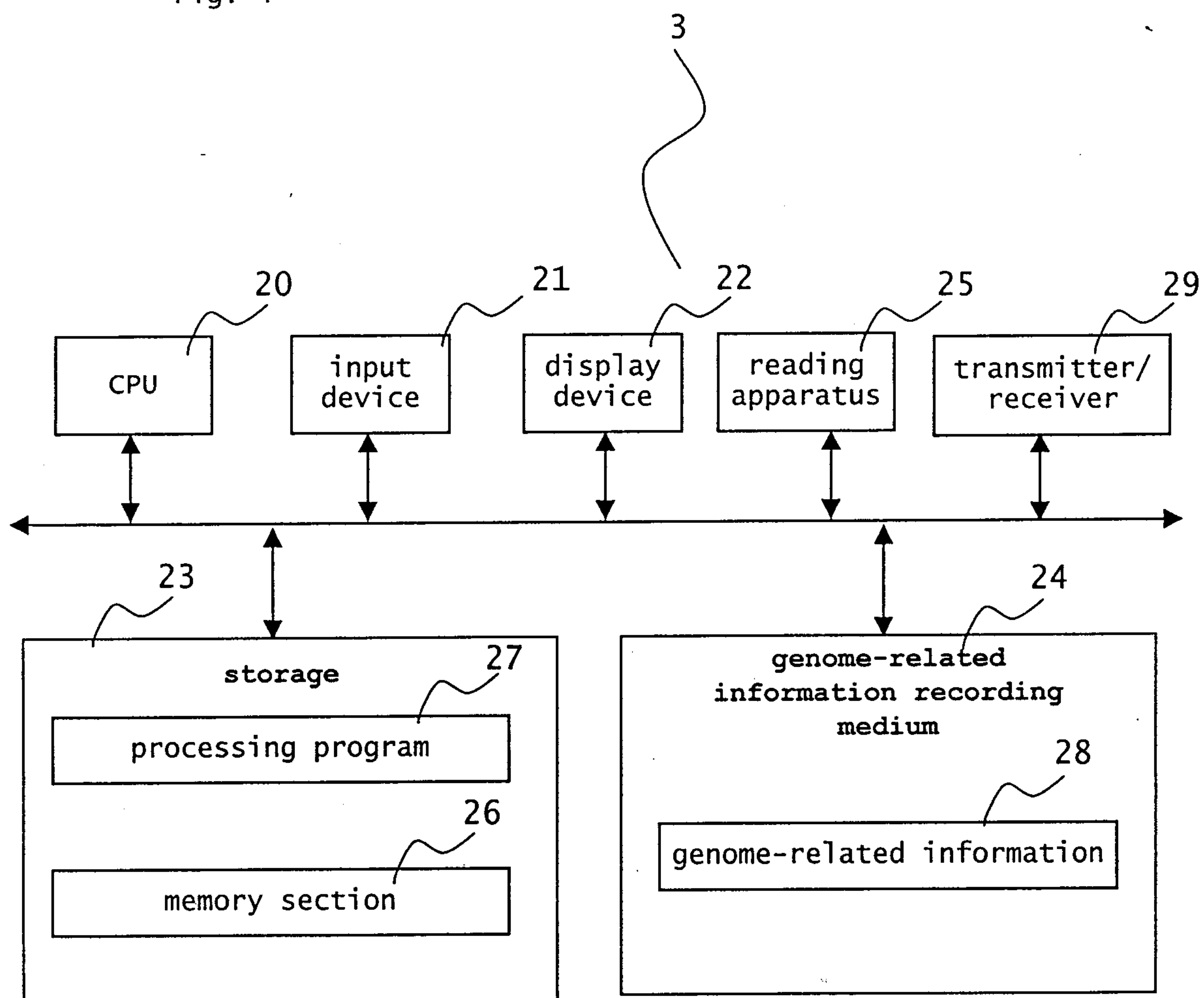


Fig. 5

I

Gno.	Date of birth
0001	**,**,****

II

Polymorphism address	Polymorphism pattern	Comment
000001	G	
000002	T	
:	:	:
123456	A	
:	:	:
223456	G	
:	:	:
234567	G	
:	:	:
334567	G	
:	:	:
345678	C	
:	:	:
445678	T	
:	:	:
456789	T	
456790	G	
456791	14 times	
456792	deletion	
:	:	:

III

Anamnesis
infantile asthma
gout
pollinosis
gastric ulcer
atopy
hypertension
diabetes

IV

Characteristics	Record
blood type	type A
body height	
weight	
vision	
running ability	
psychological test	
:	:
:	:
:	:
:	:
:	:

V.....

..... (clinical record, etc.)
.....
.....
.....
.....
.....
.....
.....
.....
.....

Fig. 6

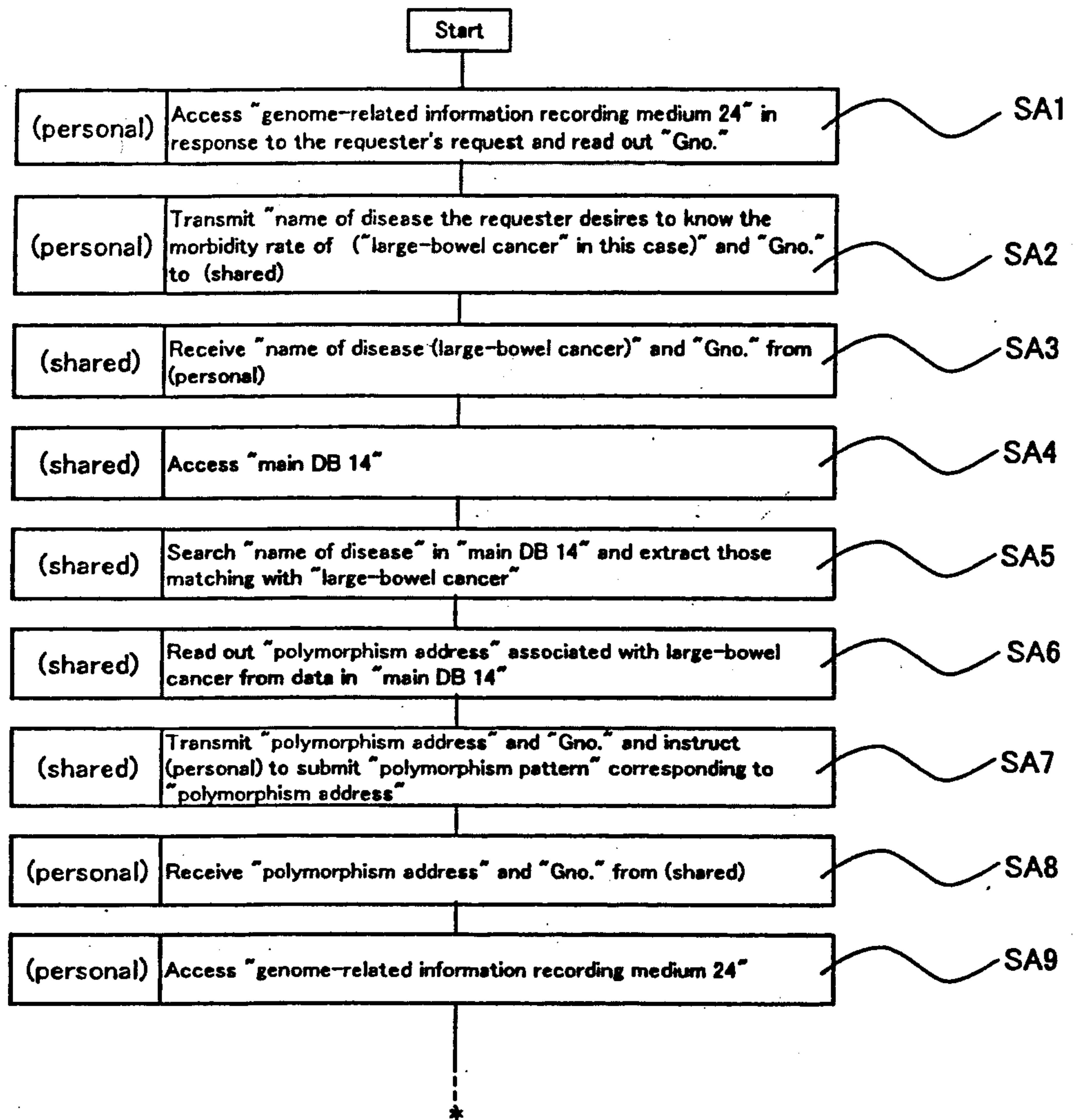


Fig. 7

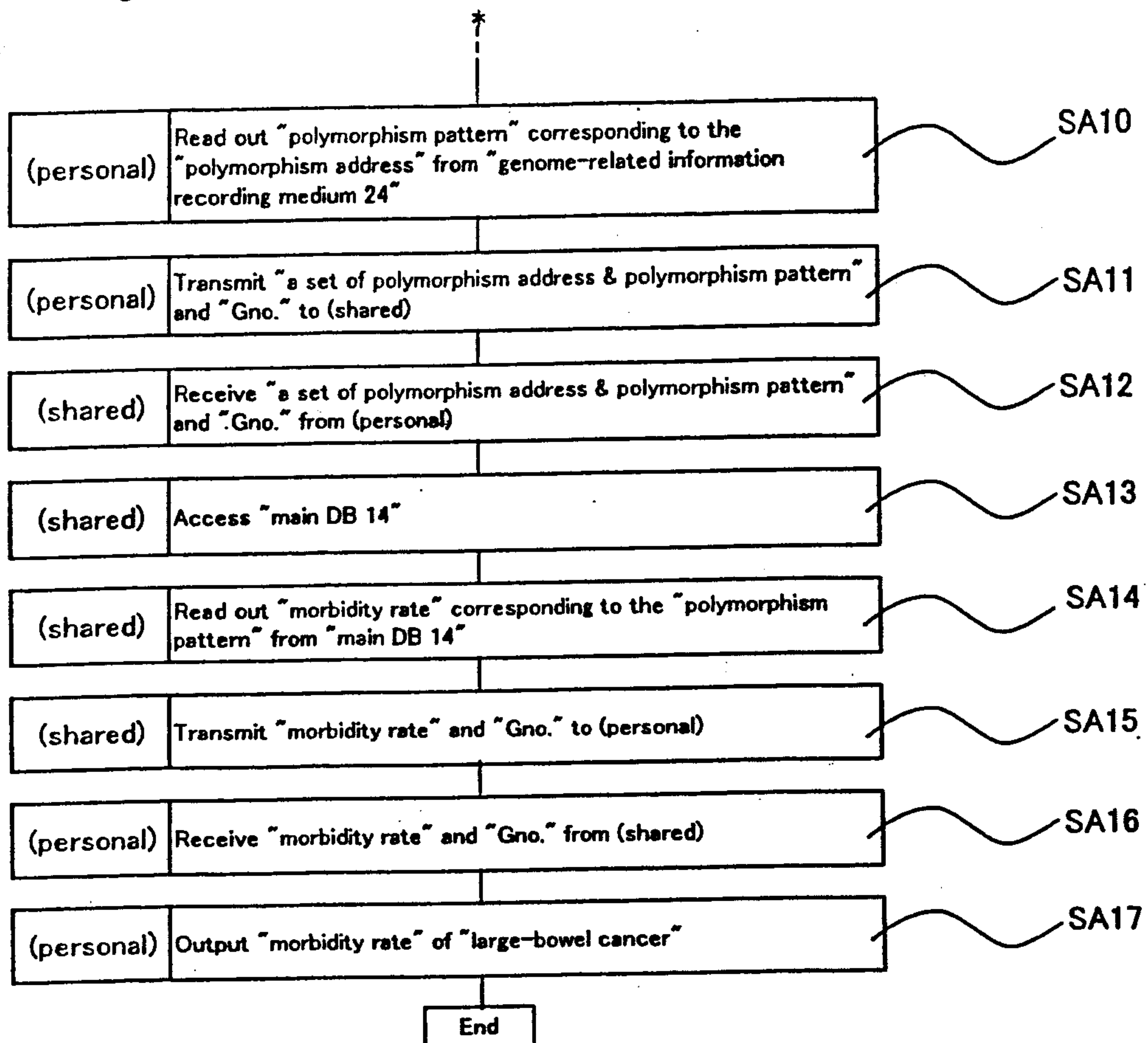


Fig. 8

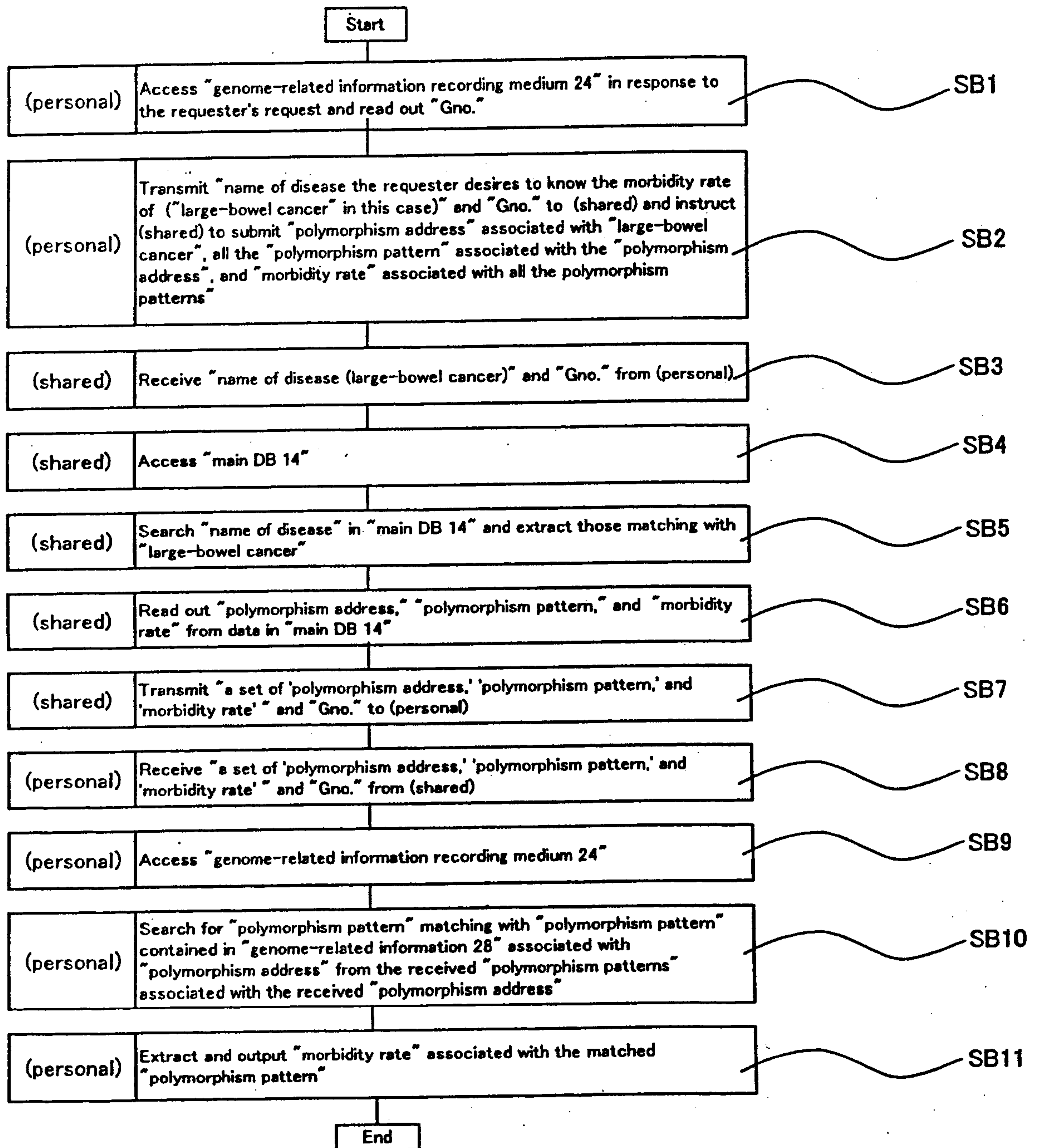


Fig. 9

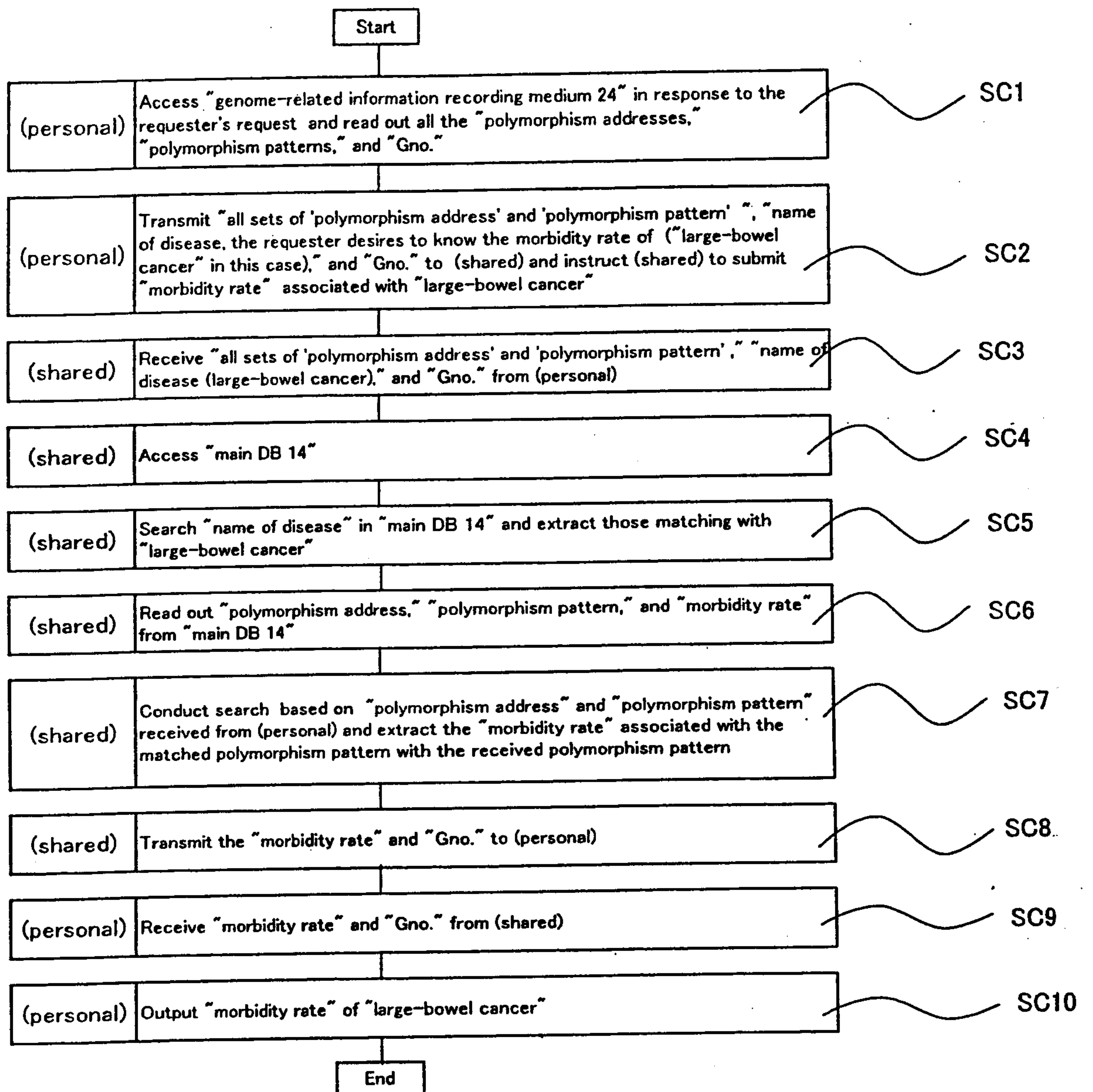
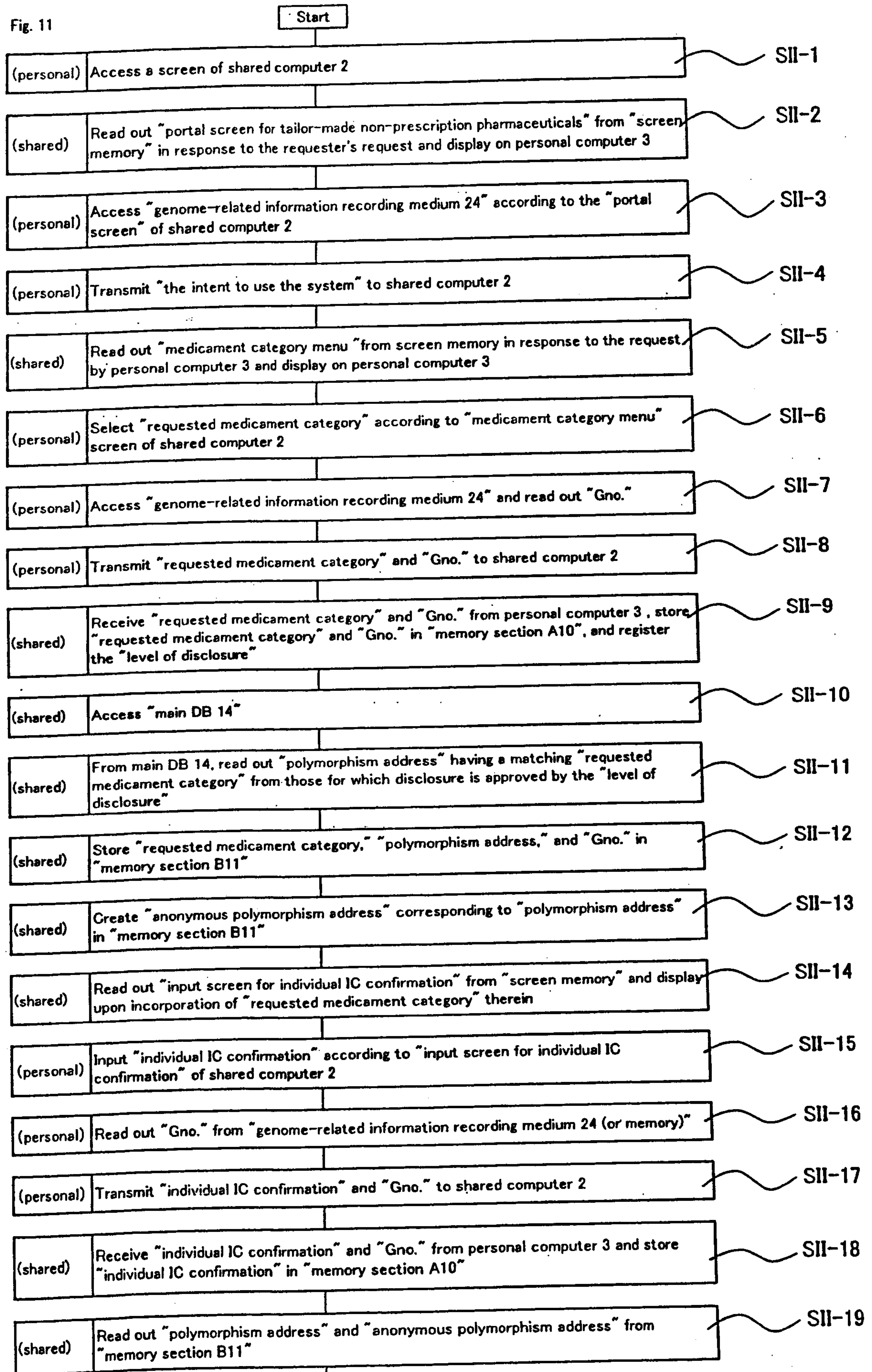


Fig. 10

Poly-morphism address	Poly-morphism type	Poly-morphism pattern	Major classification	Middle classification (medicament category)	Minor classification	Drug type	Level of disclosure 1	Level of disclosure 2
123456	SNP	A	responsiveness to drug	cold remedy	runny nose	a	A (clinical)	B (non-prescription drug)
123456	SNP	G	responsiveness to drug	cold remedy	runny nose	b	A	B
223456	SNP	G	responsiveness to drug	antipruritic	pruritus	(i)	A	B
223456	SNP	A	responsiveness to drug	antipruritic	pruritus	(ii)	A	B
234567	SNP	G	side-effect	cold remedy	runny nose	c	A	B
234567	SNP	A	side-effect	cold remedy	runny nose	d	A	B
334567	SNP	A	side-effect	antipruritic	pruritus	(iii)	A	B
334567	SNP	G	side-effect	antipruritic	pruritus	(iv)	A	B
345678	SNP	C	responsiveness to drug	cold remedy	cough	e	A	B
345678	SNP	T	responsiveness to drug	cold remedy	cough	f	A	B
445678	SNP	T	responsiveness to drug	gastrointestinal drug	stomachache	(I)	A	B
445678	SNP	C	responsiveness to drug	gastrointestinal drug	stomachache	(II)	A	B
456789	SNP	T	side-effect	cold remedy	cough	g	A	B
456789	SNP	C	side-effect	cold remedy	cough	h	A	B
:	microsatellite	14 times	:	:	:	—	disclosure forbidden	—
:	microsatellite	9 times	:	:	:	—	disclosure forbidden	—
:	deletion	G	:	:	:	:	A	—
:	deletion	deletion	:	:	:	:	A	—

Fig. 11



*

Fig. 12

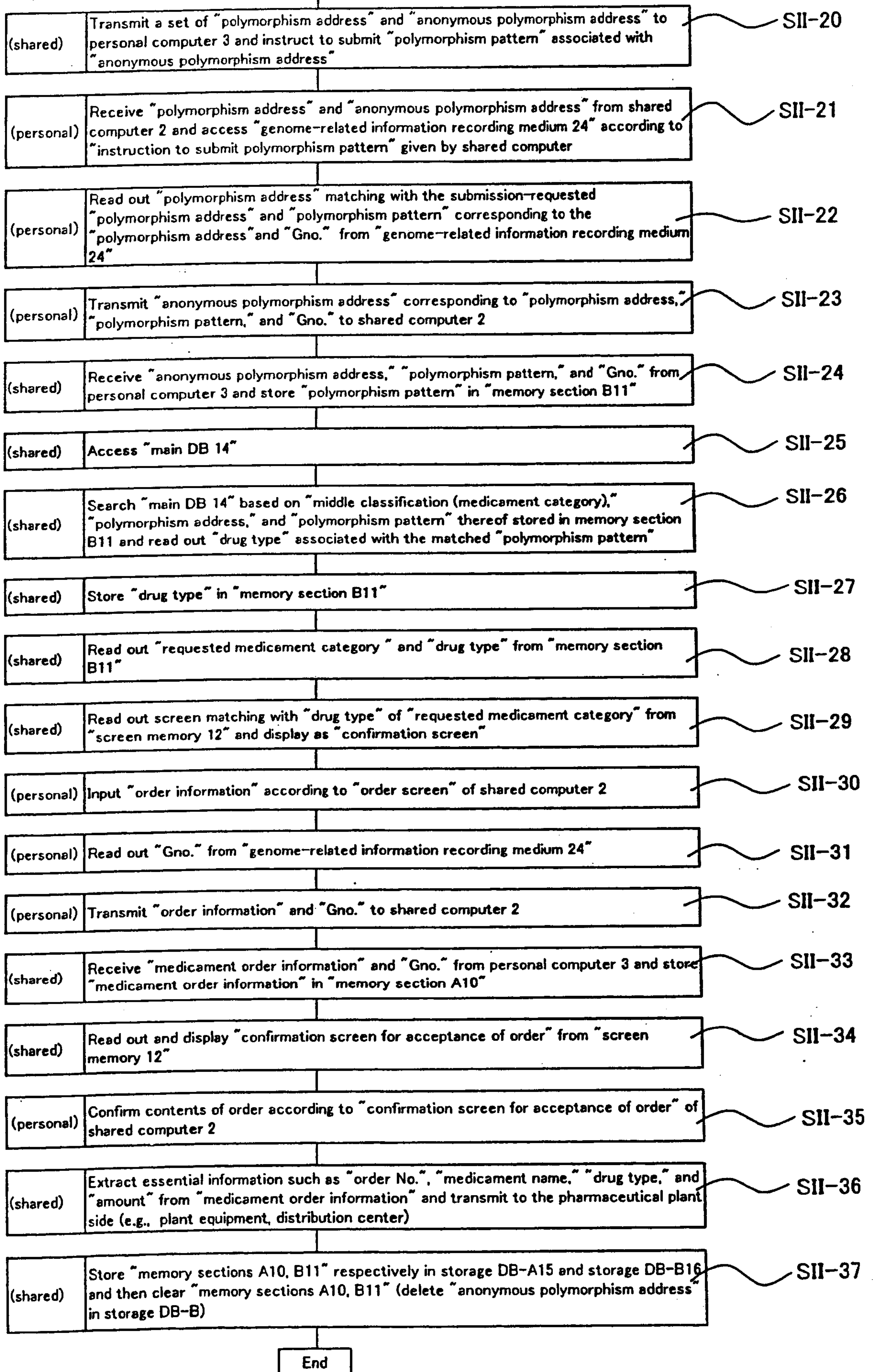


Fig. 13

Welcome to ○○ Pharmaceuticals' "My first-aid kit"

- What is "My first-aid kit" ?
 - It is a system for providing medicament which relatively suits your diathesis based on your genomic information.
 - Please be aware of the need for you to provide a part of your genomic information in order to carry out the above ※.
- ※ You are requested to provide genomic information only on "efficacy" and "side-effect" of the desired medicament from your genomic information.
- Please follow the following procedures when obtaining medicine through "My first-aid kit"
 - ① Insert your "G card" into the "G card reader"
 - ② Click "OK."

Fig. 14

○○ Pharmaceuticals' "My first-aid kit" Selection of medicine

- Please select a medicine you need. (more than one may be selected)

☒ cold remedy	☐ cough medicine
☐ antipyretic agent	☐ antipruritic agent
☐ antidiarrheic agent	☐ gastrointestinal drug
☐ medicine for pollinosis	☐ pain killer

Fig. 15

○○ Pharmaceuticals' "My first-aid kit" About "informed consent"

- Please let us confirm your consent regarding the following:
 - You are requested to provide your genomic information to ○○ Pharmaceuticals to prepare "cold remedy" that relatively suits your diathesis regarding only:
 - ① genomic information on the "efficacy" of "cold remedy"; and
 - ② genomic information on the "side-effect" of "cold remedy"
 - The genomic information you provide to ○○ Pharmaceuticals is made anonymous and then used only for the preparation of "cold remedy" that suits your diathesis.
 - Please click "OK" below if you consent to the above.

Fig. 16

○○ Pharmaceuticals' "My first-aid kit" About medicine suitable for your diathesis

● "Cold remedy" relatively suitable for your diathesis is as follows:

① Name of medicine:「△△△」 「a,c,e,g...」 type
 <it works at once and is less likely to cause drowsiness or constipation>

② Benefits: Relief of headache, cough, runny nose, sore throat, chillness, fervescence, aching joint, and aching muscle

③ Ingredients: *****,*****,****,*****

④ Dose: Three tablets at a time for those over 15 years old, two tablets at a time for those from 11 to 14 years old, one tablet at a time for those from 5 to 10 years old

⑤ Direction: Preferably take within 30 minutes after meal, three times a day

⑥ Precautions: Refer to "Precautions for use" on the next page

Please click "order screen" below to order the above medicine.

Next page
(Precautions for use)

Fig. 17

G no.	Requested medicament	I C	Order information
0001	cold remedy	○	
0002	:	:	:
:	:	:	:

Fig. 18

G no.	Requested medicament	Polymorphism address	Anonymous polymorphism address	Polymorphism pattern	Drug type
0001	cold remedy	123456	1	A	a
0001	cold remedy	234567	2	G	c
0001	cold remedy	345678	3	C	e
0001	cold remedy	456789	4	T	g
:	:	:	:	:	:

Fig. 19

Order No.	Medicament name	Type	Number of tablets	Number of boxes	
111111	△△△	a,c,e,g...	100 tablets	1 box	
:	:	:	:	:	:

Fig. 20

Poly-morphism address	Poly-morphism type	Poly-morphism pattern	Major classification	Middle classification (category of disease)	Minor classification (name of disease)	Annotative information on polymorphism pattern (morbidity rate) (responsiveness, side-effect)	Level of disclosure (disclosability) 1	Level of disclosure (disclosability) 2
123456	SNP	A	morbidity rate	hypertension	hypertension	4	A (clinical)	B (medical examination)
123456	SNP	G	morbidity rate	hypertension	hypertension	2	A	B
223456	SNP	G	morbidity rate	cancer	large-bowel cancer	4	A	B
223456	SNP	A	morbidity rate	cancer	large-bowel cancer	2	A	B
234567	SNP	G	morbidity rate	cancer	stomach cancer	Z	A	B
234567	SNP	A	morbidity rate	cancer	stomach cancer	Z	A	B
334567	SNP	A	responsiveness to drug	respiratory organs	asthma	a type	A	—
334567	SNP	G	responsiveness to drug	respiratory organs	asthma	b type	A	—
345678	SNP	C	responsiveness to drug	diabetes	I type	c type	A	—
345678	SNP	T	responsiveness to drug	diabetes	I type	d type	A	—
445678	SNP	T	morbidity rate	digestive organs	gastric ulcer	Z	A	B
445678	SNP	C	morbidity rate	digestive organs	gastric ulcer	Z	A	B
456789	SNP	T	morbidity rate	liver	hepatopathy	1	A	B
456789	SNP	C	morbidity rate	liver	hepatopathy	4	A	B
500001	SNP	A	morbidity rate	circulatory organs	arrhythmia	3	A	B
500001	SNP	G	morbidity rate	circulatory organs	arrhythmia	4	A	B
500500	SNP	C	morbidity rate	cancer	stomach cancer	Y	A	B
500500	SNP	T	morbidity rate	cancer	stomach cancer	Y	A	B
600001	SNP	A	morbidity rate	respiratory organs	asthma	1	A	—
600001	SNP	G	morbidity rate	respiratory organs	asthma	4	A	—
600600	SNP	T	morbidity rate	cancer	stomach cancer	Y	A	B
600600	SNP	C	morbidity rate	cancer	stomach cancer	Y	A	B
700001	SNP	C	drug side-effect	renal diseases	O type	g type	A	—
700001	SNP	T	drug side-effect	renal diseases	O type	h type	A	—
700700	SNP	T	morbidity rate	digestive organs	gastric ulcer	Y	A	B
700700	SNP	C	morbidity rate	digestive organs	gastric ulcer	Y	A	B
800001	SNP	T	morbidity rate	cancer	lung cancer	2	A	B
800001	SNP	C	morbidity rate	cancer	lung cancer	4	A	B
800800	SNP	T	morbidity rate	lipid	hyperlipemia	4	A	B
800800	SNP	C	morbidity rate	lipid	hyperlipemia	1	A	B
900001	microsatellite	14 times	morbidity rate	inmedicable disease	OO disease	5	A	—
900001	microsatellite	9 times	morbidity rate	inmedicable disease	OO disease	1	A	—
900900	deletion	G	morbidity rate	leukemia	△ type	1	A	—
900900	deletion	deletion	morbidity rate	leukemia	△ type	5	A	—
:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:

Fig. 21

Polymorphism address Z	Pattern Z	Polymorphism address Y	Pattern Y	Morbidity rate
234567	G	500500, 600600	C, T	2
234567	G	500500, 600600	C, C	2
234567	G	500500, 600600	T, T	4
234567	G	500500, 600600	T, C	2
234567	A	500500, 600600	C, T	2
234567	A	500500, 600600	C, C	2
234567	A	500500, 600600	T, T	1
234567	A	500500, 600600	T, C	2
445678	T	700700	T	4
445678	T	700700	C	2
445678	C	700700	T	2
445678	C	700700	C	4
:	:	:	:	:

Fig. 22

I

Gno.	Date of birth
0001	**,**,****

II

Polymorphism address	Polymorphism pattern	Comment
000001	G	
000002	T	
:	:	:
123456	A	
:	:	:
223456	G	
:	:	:
234567	G	
:	:	:
334567	G	
:	:	:
345678	C	
:	:	:
445678	T	
:	:	:
456789	T	
:	:	:
50001	A	
:	:	:
500500	T	
:	:	:
600001	A	
:	:	:
600600	T	
:	:	:
700001	C	
:	:	:
700700	T	
:	:	:
800001	T	
:	:	:
800800	C	
:	:	:
900001	14 times	
900900	G	
:	:	:

III

Anamnesis
infantile asthma
gout
pollinosis
gastric ulcer
atopy
hypertension
diabetes

IV

Characteristics	Record
blood type	type A
body height	
weight	
vision	
running ability	
psychological test	
:	:
:	:

V.....

.....
(clinical record, etc.)
.....
.....
.....
.....
.....
.....
.....
.....

Fig. 23

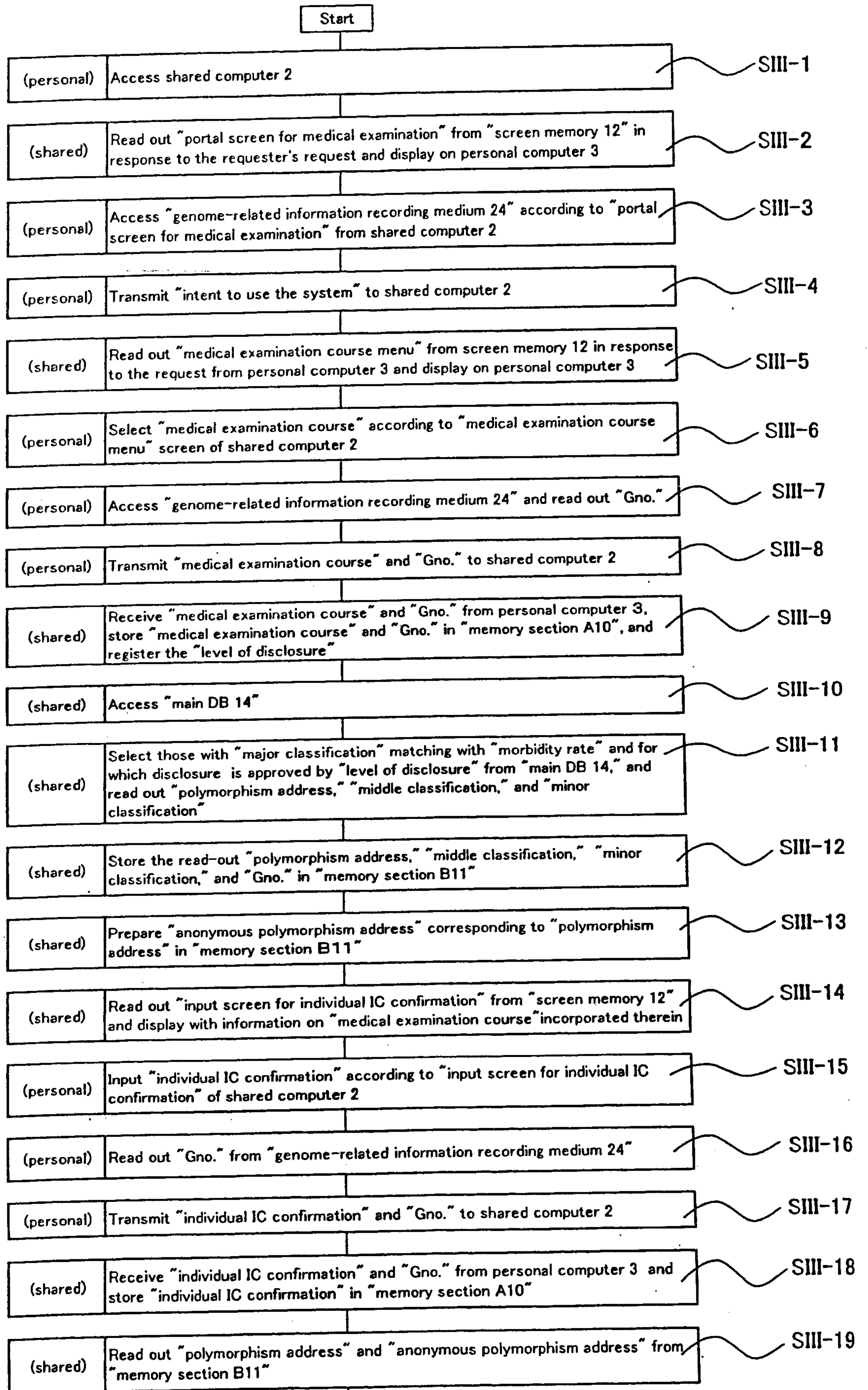


Fig. 24

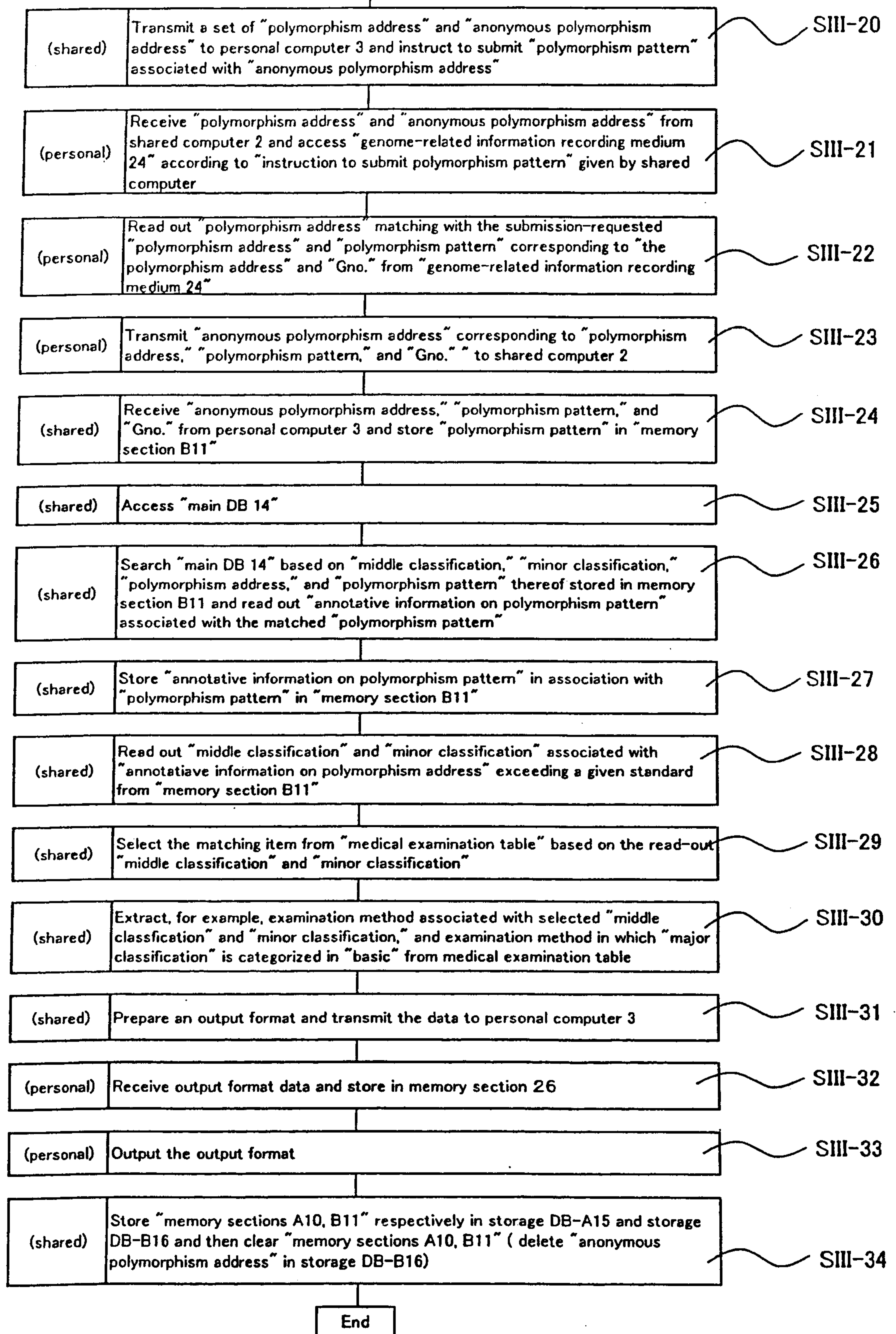


Fig. 25

<OO Medical Care Center>
 Welcome to "My check-up !" portal site
 ~Why not get a medical examination suitable for your diathesis ? ~

● What is "My check-up !"?
 • "My check-up !" is a preventive medical care system which automatically selects examination items for diseases you need to be aware of with respect to your living habits due to your diathesis based on your genomic information and performs periodic medical examination and guidance for better living, thereby leading to the prevention, early diagnosis, and treatment of diseases.

• Please be aware of the need for you to provide the part of your genomic information relating to medical examination items * to undergo a "My check-up !"

* Detailed "examination items" are displayed on following screens.

• Please click "OK" below upon confirmation and consent to the above.

Fig. 26

<OO Medical Care Center>
"My check-up !" Check-up course menu
 ~ Please click "OK" upon selection of a desired course ~

● "Entrusted check-up course"
 • In addition to "basic examination items *", this course automatically selects examination items associated with diseases you are susceptible to due to your diathesis from among various cancers, liver disfunction, heart disease, brain disease, hypertension, diabetes ...* and performs examination and guidance for better living.

○ "Entrusted + total cancer check-up course"
 • In addition to the above "Entrusted check-up course," this course performs a series of examinations associated with various cancers* including "cancers" you are less likely to have due to your diathesis

○ "Total check-up course"
 • In addition to "basic examination items *", this course performs a series of examinations associated with various cancers, liver disfunction, heart disease, brain disease, hypertension, diabetes ...*

Fig. 27

Major classification	Middle classification (category of disease)	Minor classification (name of disease)	Examination method	Examination items	Examinee's preparation	Inquiry items	...
basic	infectious disease	infectious disease	blood sampling (hematology)	white blood cell count	—	—	...
basic	anemia	anemia	blood sampling (hematology)	red blood cell count	—	*****	...
basic	anemia	anemia	blood sampling (hematology)	hemoglobin content	—	*****	...
:	:	:	:	:	:	:	:
basic	liver function	hepatopathy	blood sampling (biochemistry)	total protein level	—	*****	...
basic	liver function	hepatopathy	blood sampling (biochemistry)	total bilirubin level	—	*****	...
:	:	:	:	:	:	:	:
basic	sugar	diabetes	urinalysis (paper test)	urinary sugar	*****	*****	...
:	:	:	:	:	:	:	:
basic	hypertension	hypertension	hemo-dynamometer	blood pressure level	*****	*****	...
morbidity rate	hypertension	hypertension	hemo-dynamometer	blood pressure level	—	*****	...
morbidity rate	cancer	large-bowel cancer	stool examination (protein chip)	OO protein	collection of stool	*****	...
morbidity rate	cancer	stomach cancer	blood sampling (protein chip)	× × protein	—	*****	...
morbidity rate	cancer	lung cancer	blood sampling (protein chip)	△△ protein	—	*****	...
:	:	:	:	:	:	:	:
morbidity rate	liver	hepatopathy	blood sampling (biochemistry)	ZTT	—	*****	...
morbidity rate	liver	hepatopathy	blood sampling (biochemistry)	LOH	—	*****	...
:	:	:	:	:	:	:	:
morbidity rate	circulatory organs	arrhythmia	electrocardiogram (rest)	myocardial action potential	—	*****	...
:	:	:	:	:	:	:	:
morbidity rate	lipid	hyperlipemia	blood sampling (biochemistry)	neutral fat	—	*****	...
:	:	:	:	:	:	:	:

Fig. 28

<〇〇 Medical Care Center>

About "informed consent" in "My check-up ! Entrusted check-up course"

~ Please let us confirm your consent to the following. ~

① In order to perform "My check-up! Entrusted check-up course", you will first provide us, 〇〇 Medical Care Center, with only a part of your genomic information with which your "susceptibility" to various cancers, liver disfunction, heart disease, brain disease, hypertension, diabetes ... can be distinguished from your genomic information.

② The genomic information you provide is made anonymous and then checked with the "causal relation between polymorphism information and morbidity rate" prepared by "****" to determine your susceptibility to the diseases in ① above.

Examination items associated with diseases with a "susceptibility" exceeding a given standard※ are selected as examination items peculiar to you and are examined in addition to general examination items, and reflected in to guidance for better living.

※ Basic calculation of "susceptibility" and details on setting of standard are displayed by following screens

③ Regarding diseases associated with examination items not selected in "Entrusted check-up course", susceptibility is merely relatively low, and it does not indicate that there is no morbidity rate.

Also, you do not necessarily suffer from the disease selected, and instead, you may be able to prevent it by awareness and life style improvements.

● Please click "OK" below upon confirmation and consent to the above.

OK Cancel

Fig. 29

Gno.	Middle classification (category of disease)	Minor classification (name of disease)	Polymorphism address	Anonymous polymorphism address	Polymorphism pattern	Morbidity rate
0001	hypertension	hypertension	123456	1	A	4
0001	cancer	large-bowel cancer	223456	2	G	4
0001	cancer	stomach cancer	234567	3	G	4
0001	cancer	stomach cancer	500500	4	T	-
0001	cancer	stomach cancer	600600	5	T	-
0001	cancer	lung cancer	800001	6	T	2
0001	digestive organs	gastric ulcer	445678	7	T	4
0001	digestive organs	gastric ulcer	700700	8	T	-
0001	liver	hepatopathy	456789	9	T	1
0001	circulatory organs	arrhythmia	500001	10	A	3
0001	lipid	hyperlipemia	800800	11	C	1
:	:	:	:	:	:	:

Fig. 30

Gno.	Name of check-up	IC confirmation
0001	entrusted	○
0002	:	:
:	:	:

Fig. 31

Gno.	Chart No.	Name	Sex	Date of birth	Date of examination	Medical institution admitting examinees			
Category of examination	Examination method	Examination purpose	Mor- bidity	No.	Examination items	Category	Inquiry items	Preparation	Inquiry result/ remarks
1. Blood sampling	hematology	infectious disease		1	white blood cell count	basic	*****	*****	
		anemia		2	red blood cell count	basic	*****	*****	
				3	hemoglobin content	basic	*****	*****	
				4	hematocrit value	basic	*****	*****	
		coagulation system		5	platelet count	basic	*****	*****	
	biochemistry	hepatopathy		6	total protein level	basic	*****	*****	
				7	total bilirubin level	basic	*****	*****	
				8	GOT	basic	*****	*****	
				9	GPT	basic	*****	*****	
				10	ALP	basic	*****	*****	
				11	γ -GTP	basic	*****	*****	
				12	cholinesterase	basic	*****	*****	
		hyperlipemia		13	neutral fat	basic	*****	*****	
				14	total cholesterol	basic	*****	*****	
				15	HDL cholesterol	basic	*****	*****	
				16	LDL cholesterol	basic	*****	*****	
		nephropathy		17	urea nitrogen	basic	*****	*****	
				18	creatinine count	basic	*****	*****	
		diabetes		19	blood glucose level	basic	*****	*****	
		electrolyte		20	serum sodium (Na)	basic	*****	*****	
				21	serum potassium (K)	basic	*****	*****	
				22	serum chlor (Cl)	basic	*****	*****	
	protein chip	stomach cancer	4	23	** protein	specific	*****	*****	
2. Urinalysis	paper test	diabetes		24	urinary sugar	basic	*****	*****	
		nephropathy		25	urine protein	basic	*****	*****	
				26	urine occult blood	basic	*****	*****	
		hepatopathy		27	urobilinogen	basic	*****	*****	
3. Stool examination	orthotolidine	digestive organs	4	28	stool occult blood	basic	*****	*****	
	protein chip	large-bowel cancer	4	29	OO protein	specific	*****	*****	
4. Electro- cardiogram	rest	arrhythmia		30	myocardinal action potential	basic	*****	*****	
5. Blood pressure	hemo- dynamometer	hypertension	4	31	blood pressure level	basic	*****	*****	
6. Chest X-ray	X-ray	lung disease		32	lung X-ray image	basic	*****	*****	
7. Eyeground	retinal camera	diabetes, arteriosclerosis		33	retinal camera image	basic	*****	*****	
8. Visual acuity/audibility	direct examination	visual acuity/audibility		34	direct visual acuity/audibility test	basic	*****	*****	
9. Measurement	direct measurement	obesity index		35	body height, weight	basic	*****	*****	

Fig. 32

Classification	Polymorphism address		Polymorphism pattern		Individual property type	Compatible type	Level of disclosure (disclosability)
	1	2	1	2			
Given property	1000	2000	A	C	a	b	A
Given property	1000	2000	G	T	a	b	A
Given property	1000	2000	G	C	b	a	A
Given property	1000	2000	A	T	b	a	A
Other property	3000	4000	G	C	(i)	(i)	B
Other property	3000	4000	deletion	T	(ii)	(ii)	B
:	:	:	:	:	:	:	:

Fig. 33

I

Gno.	Date of birth
0001	****.****

II

Polymorphism address	Polymorphism pattern	Comment
0001	G	
0002	T	
:	:	:
1000	A	
:	:	:
2000	C	
:	:	:
3000	deletion	
:	:	:
4000	T	
:	:	:

III

Anamnesis
infantile asthma
gout
pollinosis
gastric ulcer
atopy
hypertension
diabetes

IV

Characteristics	Record
blood type	type A
body height	
weight	
vision	
running ability	
psychological test	
:	:
:	:
:	:
:	:
:	:

V.....

.....
(clinical record, etc.)
.....
.....
.....
.....
.....
.....
.....
.....
.....

Fig. 34

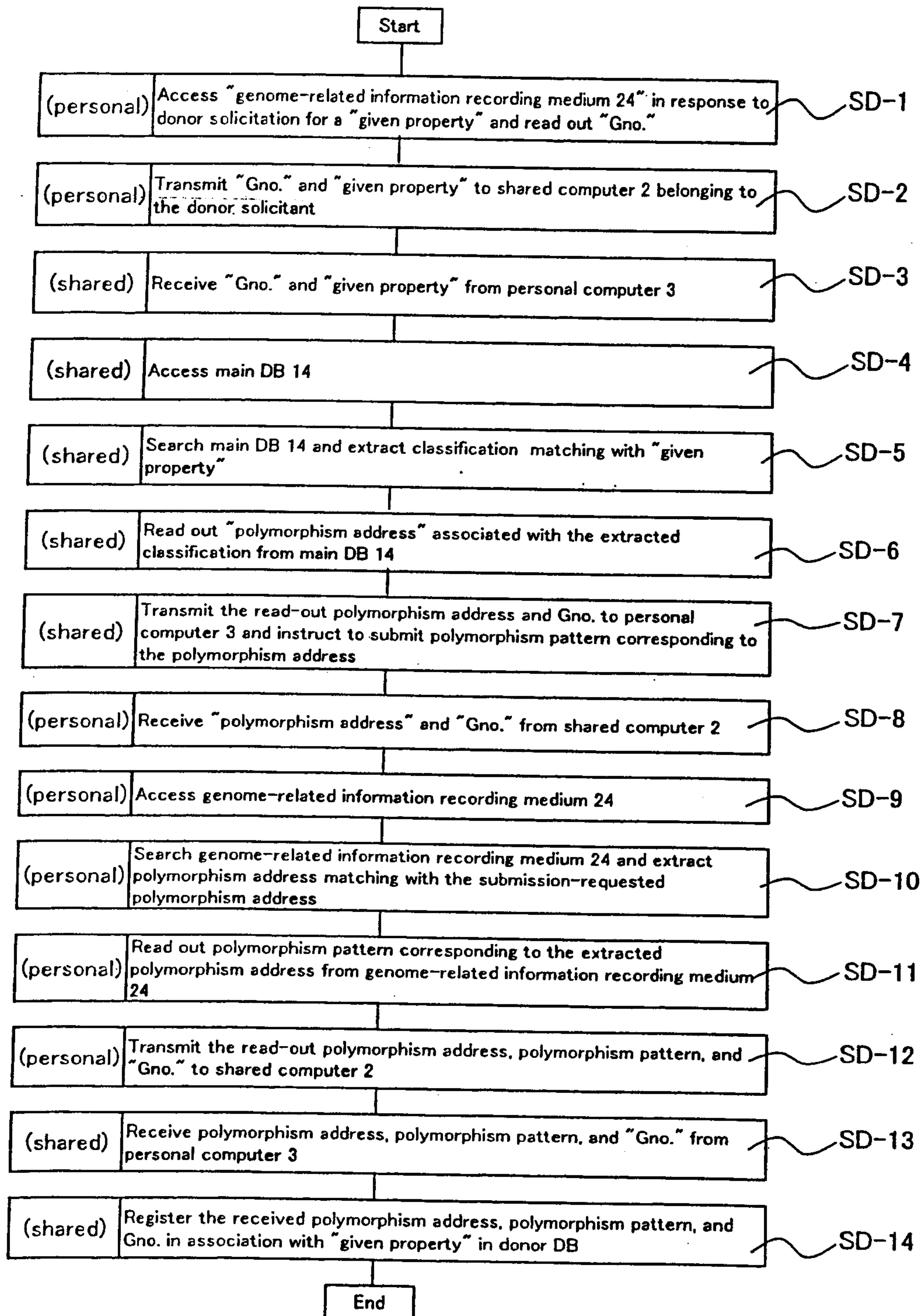


Fig. 35

Gno.	Classification	Polymorphism address		Polymorphism pattern		...
		1	2	1	2	
0001	Given property	1000	2000	A	C	...
0002	Given property	1000	2000	G	T	...
0003	Given property	1000	2000	G	C	...
0004	Given property	1000	2000	A	T	...
:	:	:	:	:	:	:

Fig. 36

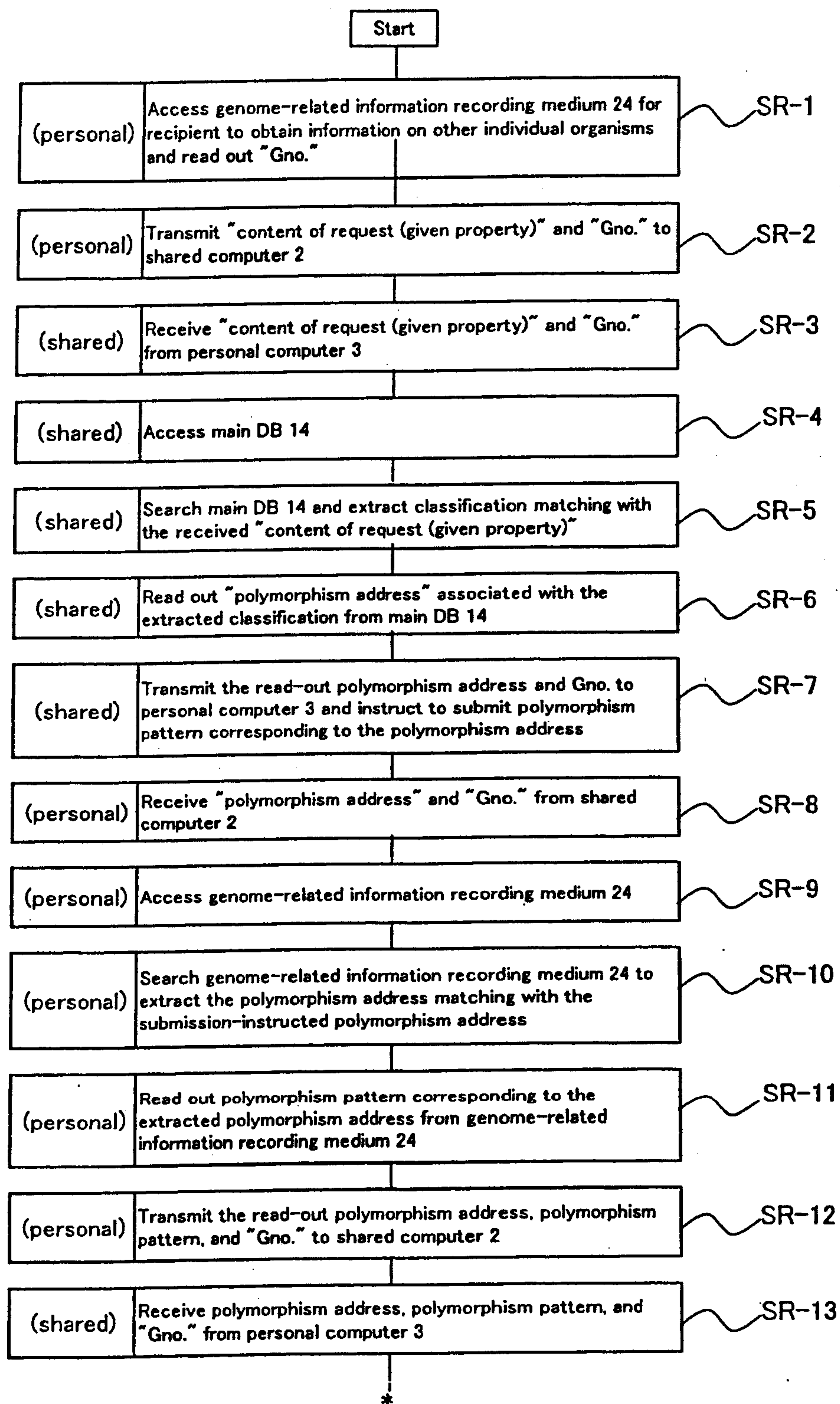


Fig. 37

