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Stephens

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(54) **MEDICATION MANAGEMENT SYSTEMS
AND METHODS FOR HEALTH AND
HEALTH RELATED FACILITIES**

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This patent is subject to a terminal dis-
claimer.

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(Continued)

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G07F 17/00 (2006.01)
G06Q 50/22 (2018.01)
(Continued)

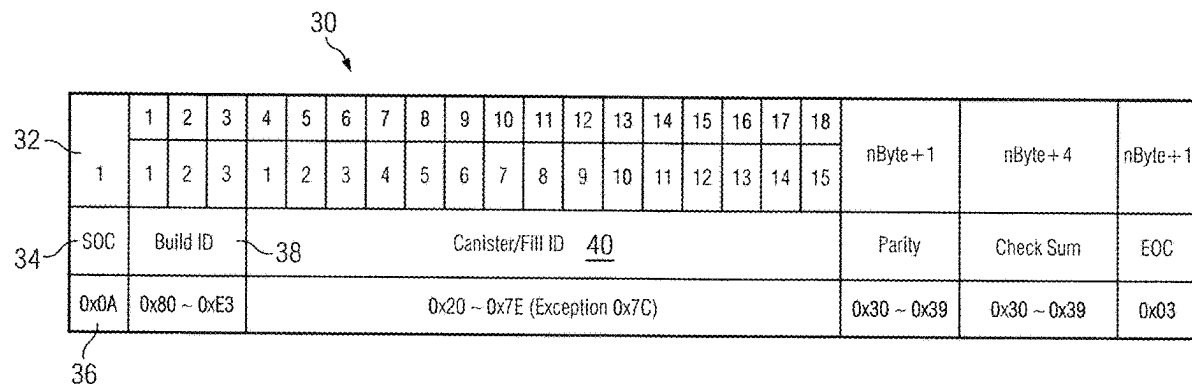
(52) **U.S. Cl.**
CPC **G07F 17/0092** (2013.01); **G06Q 50/22**
(2013.01); **G07F 9/001** (2020.05);
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(57) **ABSTRACT**

A system and method of a medication dispensing. One
system includes a canister including a body for housing and
releasing medication comprising a plurality of individually
provided medication. The canister can further include a
memory comprising canister information for identifying
canisters or content of the canisters, wherein the memory is
configured to provide the canister information to a data
storage configured to identify, based on the canister infor-
mation, a type of medication contained in the body, the data
storage further configured to provide the type of medication
to an automated dispensing machine (ADM) containing the
canister. The canister can be configured to be disposed in the
ADM during operation and to release individually at least a
portion of the plurality of individually provided medication
contained therein, the releasing in response to the ADM
operating to release the type of medication associated with
the canister information.

24 Claims, 16 Drawing Sheets



Related U.S. Application Data

continuation of application No. 16/259,716, filed on Jan. 28, 2019, now Pat. No. 11,217,058, which is a continuation of application No. 15/050,460, filed on Feb. 22, 2016, now Pat. No. 10,192,035, which is a continuation of application No. 13/085,298, filed on Apr. 12, 2011, now Pat. No. 9,268,912.

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G07F 9/00 (2006.01)

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G16H 20/13 (2018.01)

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CPC **G07F 9/002** (2020.05); **G16H 10/60** (2018.01); **G16H 20/13** (2018.01)

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See application file for complete search history.

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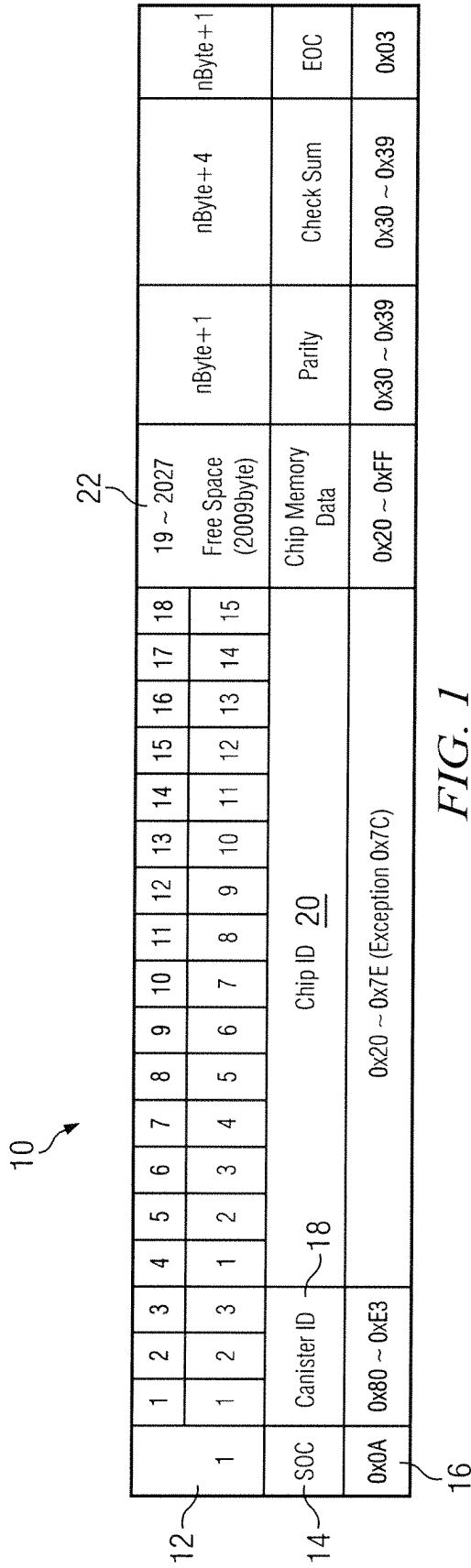


FIG. 1
(PRIOR ART)

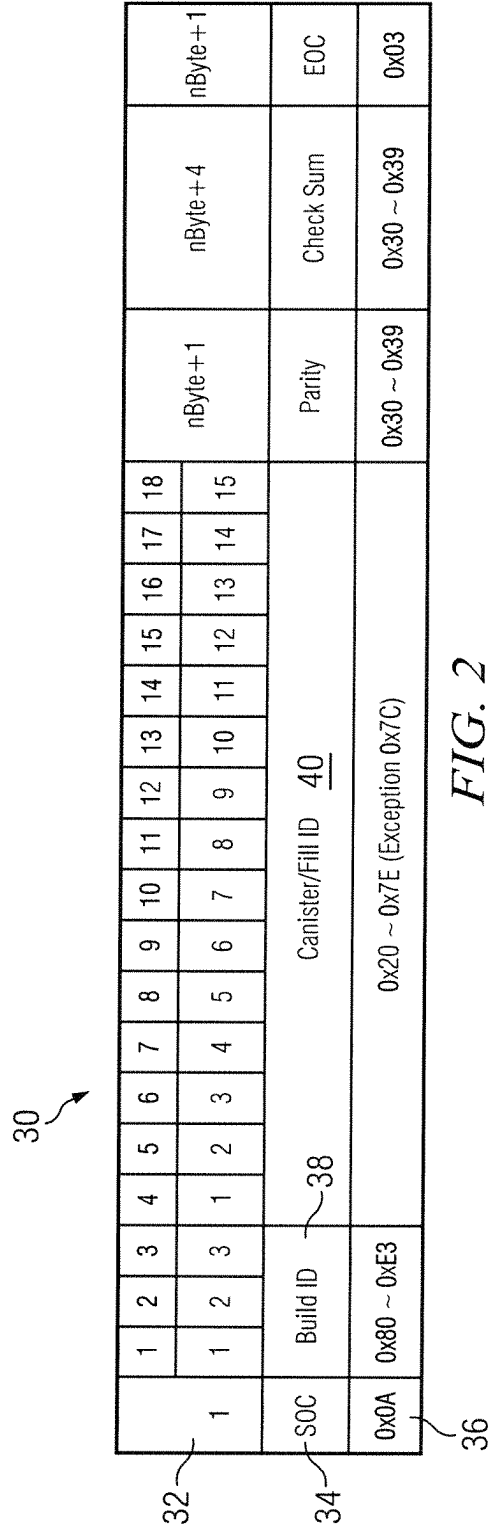


FIG. 2

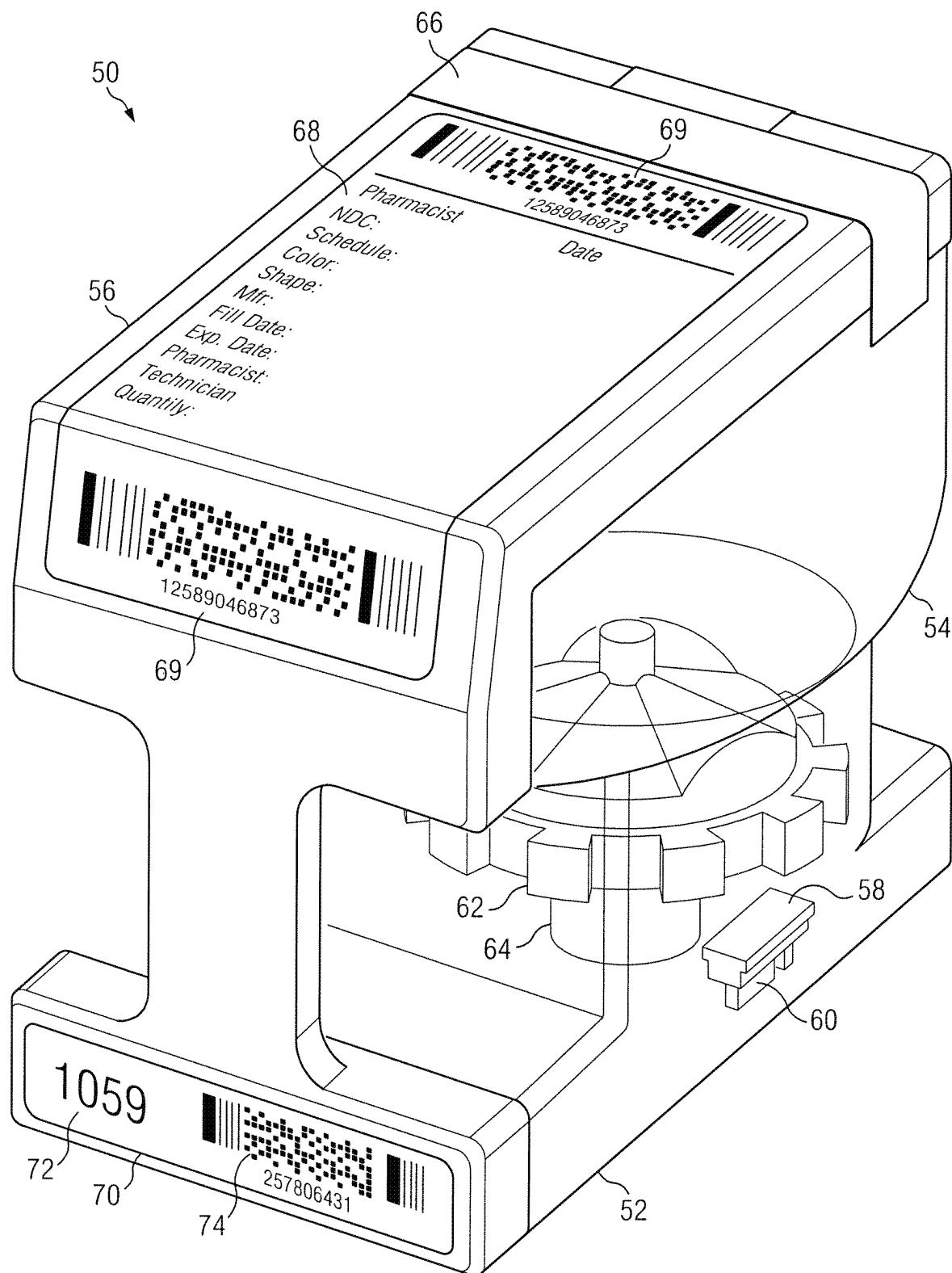


FIG. 3

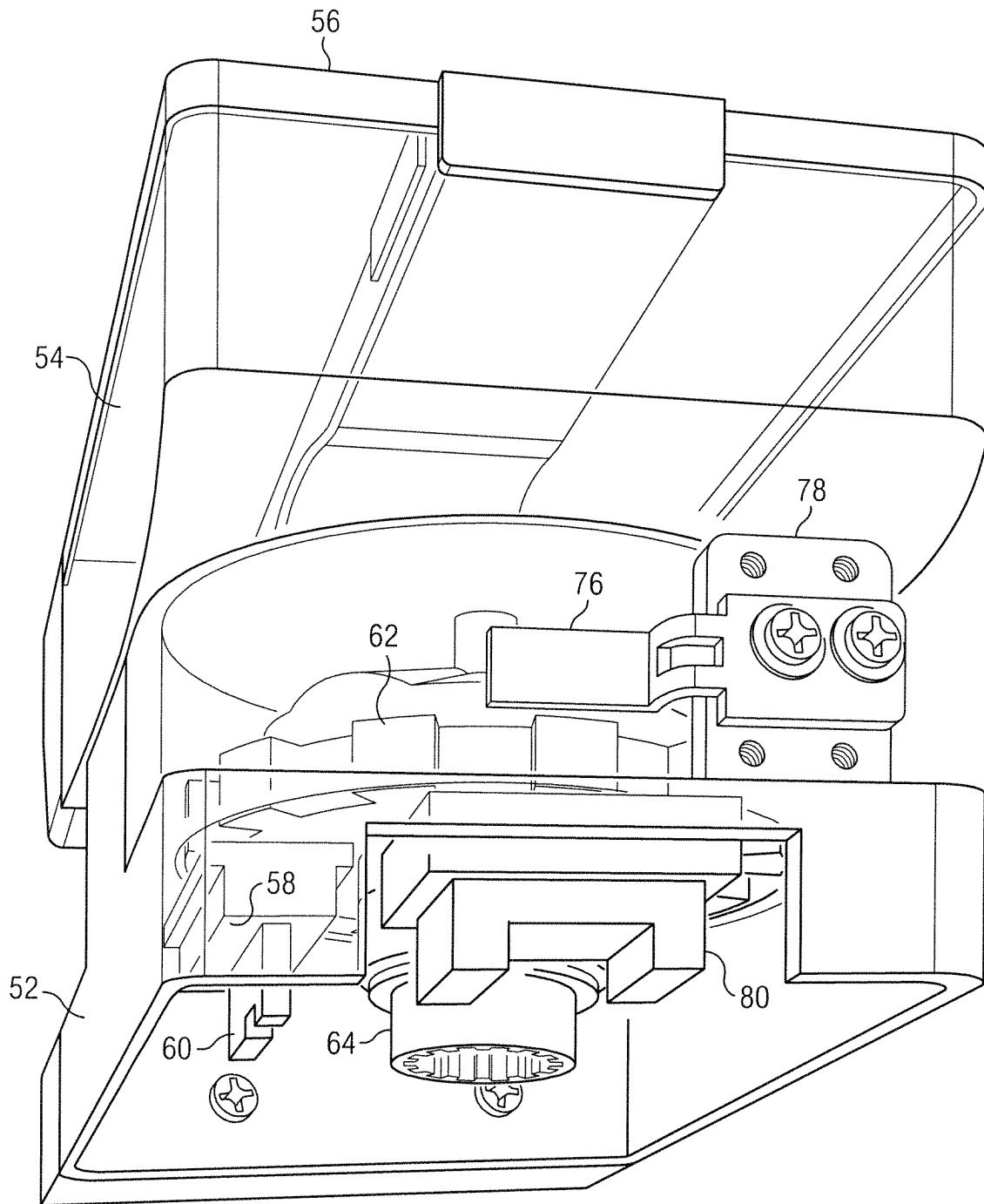


FIG. 4

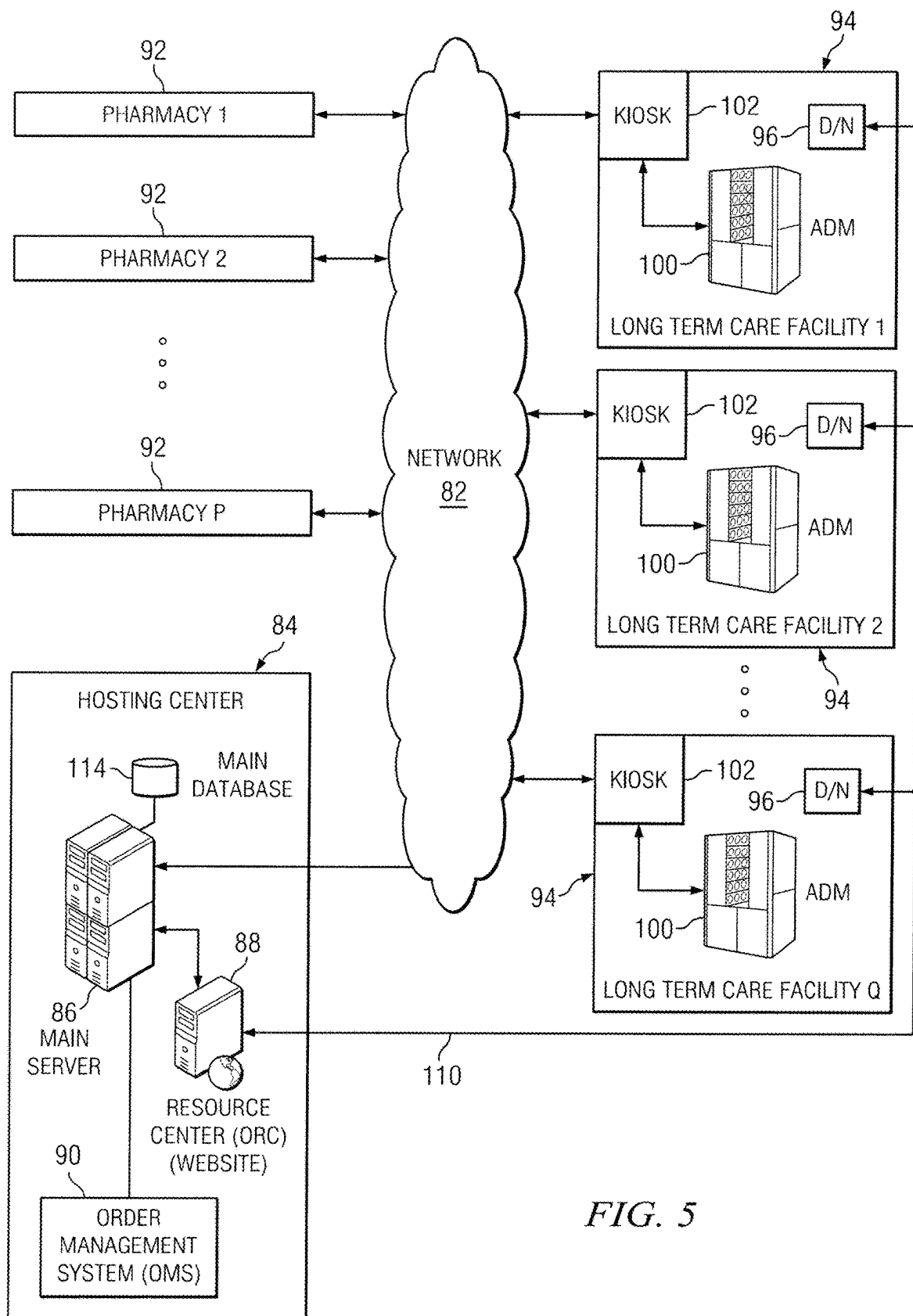
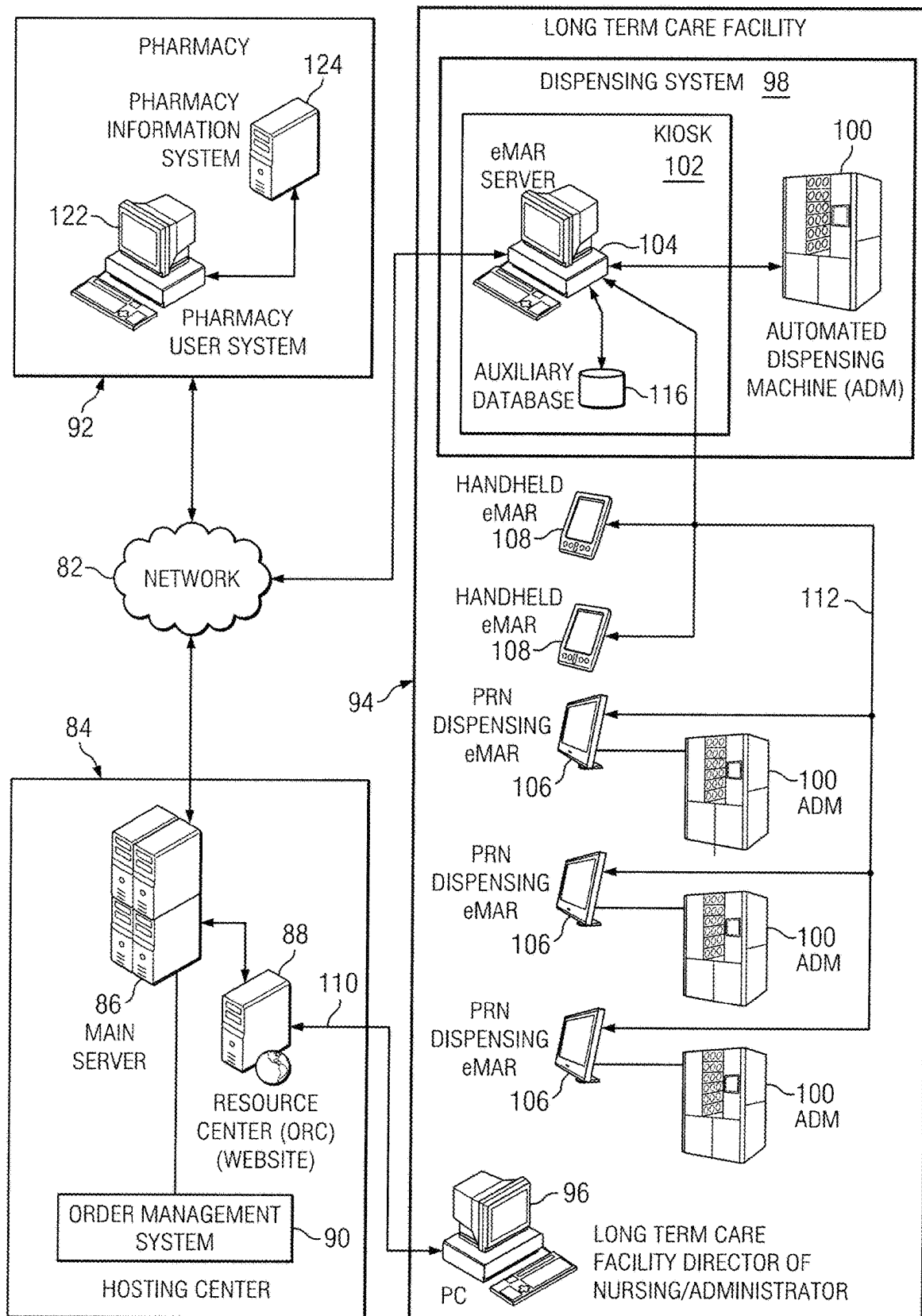


FIG. 5

FIG. 6



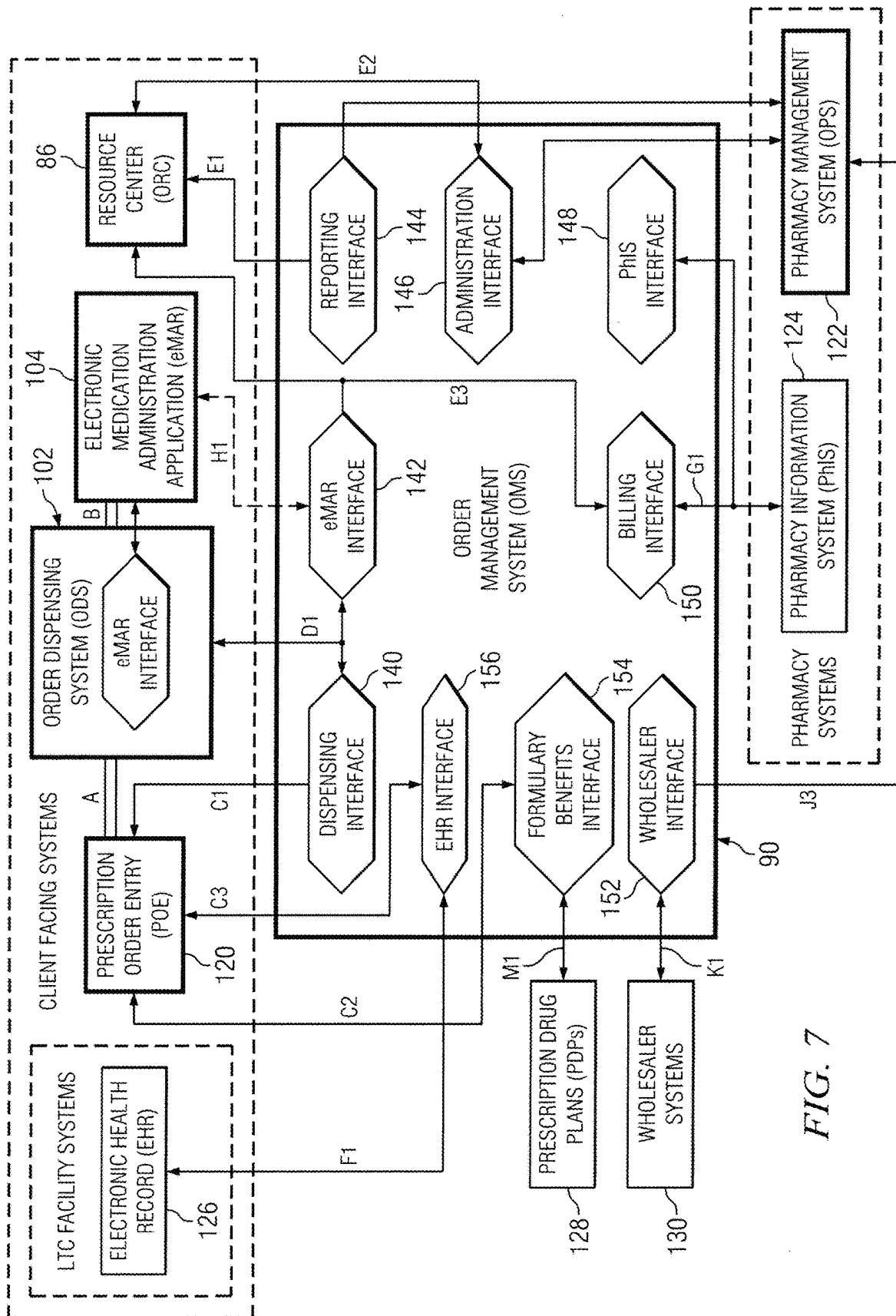


FIG. 7

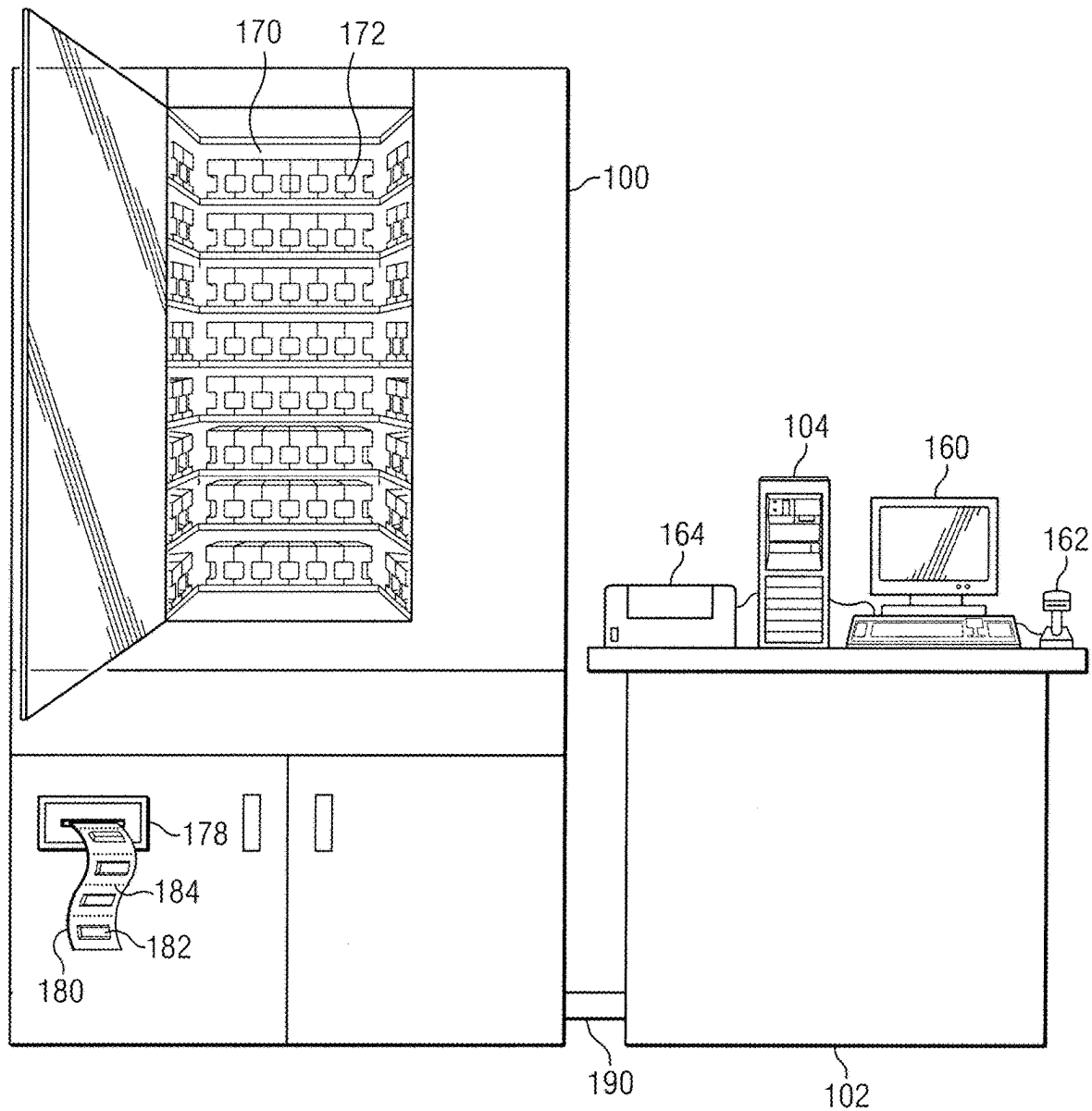


FIG. 8

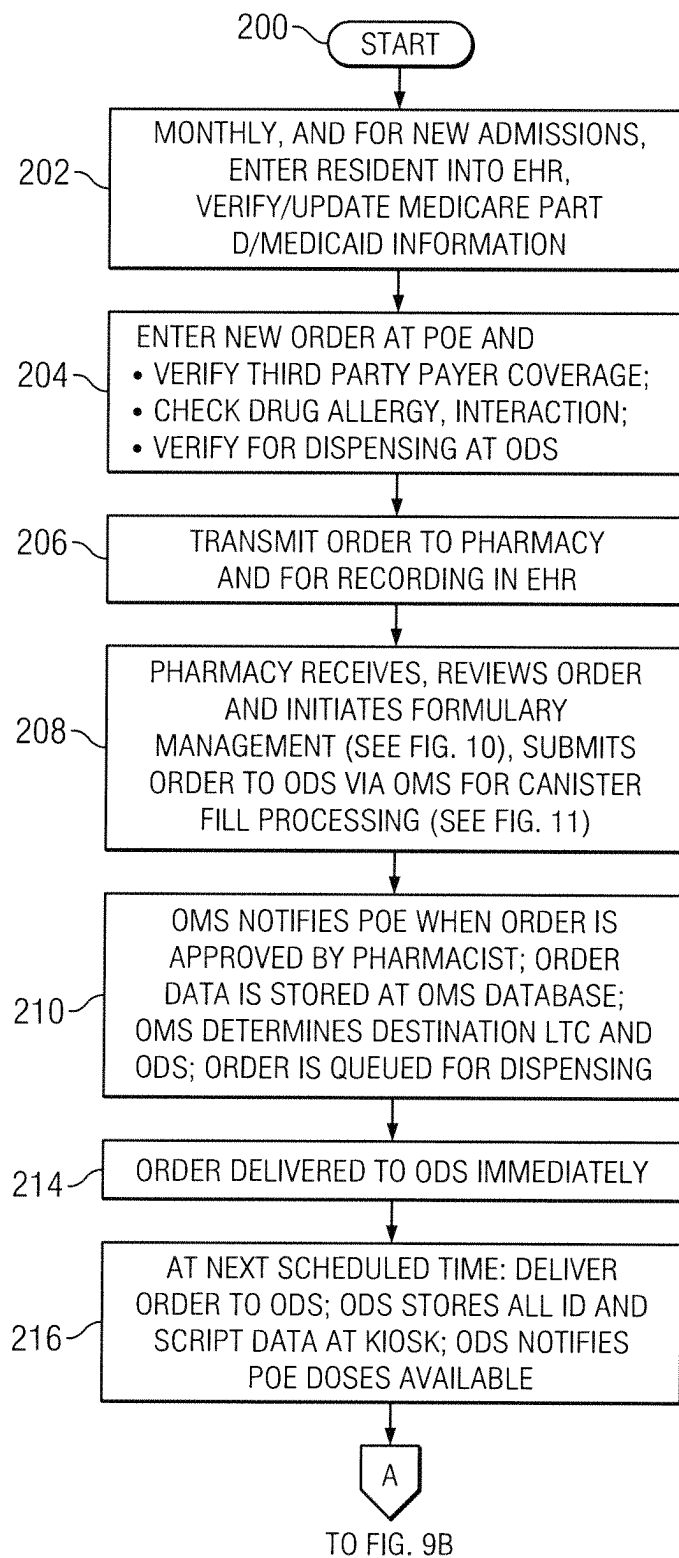
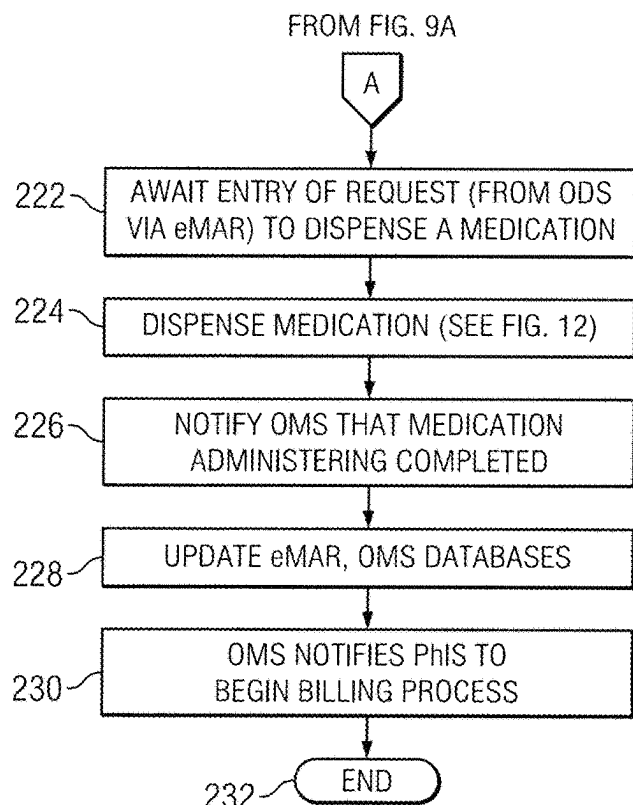
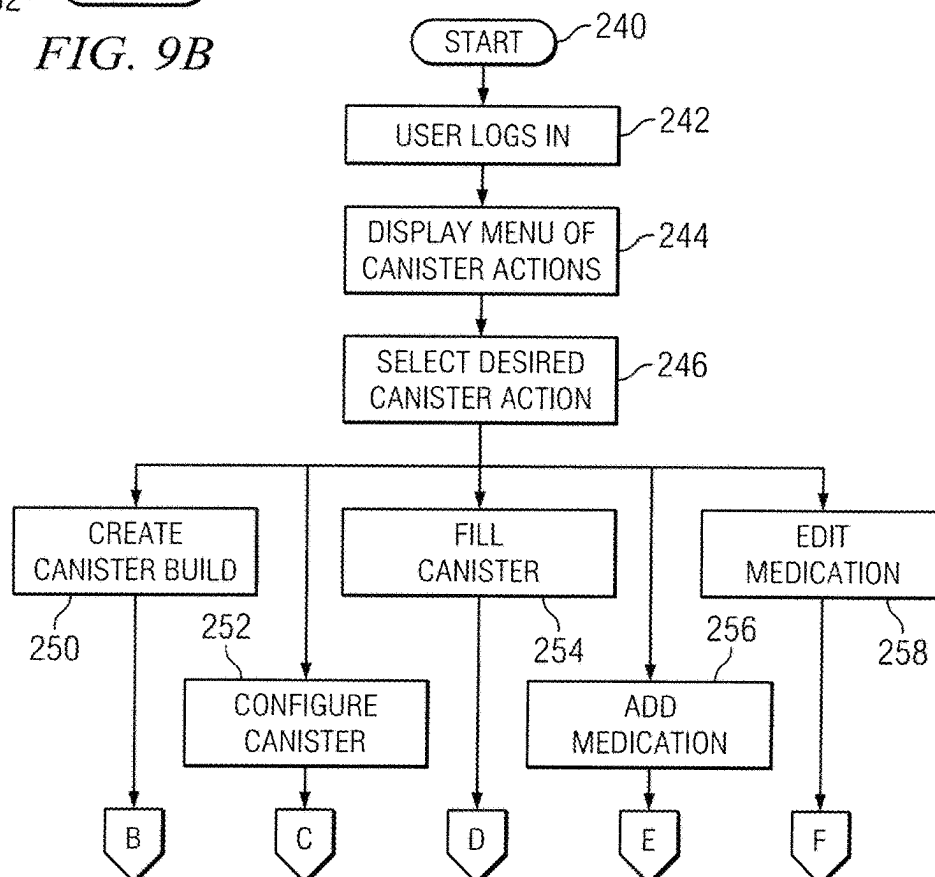


FIG. 9A

*FIG. 9B**FIG. 11A* TO FIG. 11B TO FIG. 11C TO FIG. 11D TO FIG. 11E TO FIG. 11F

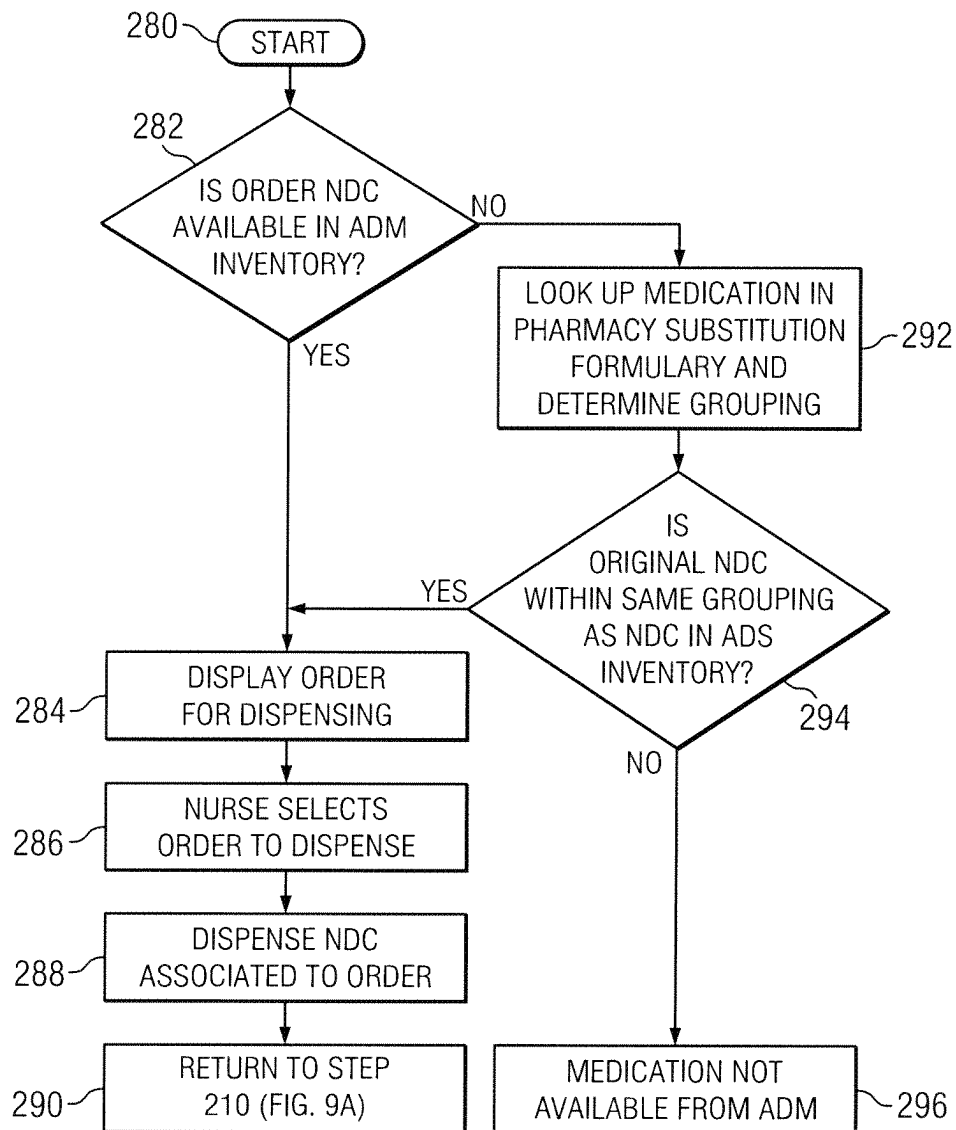
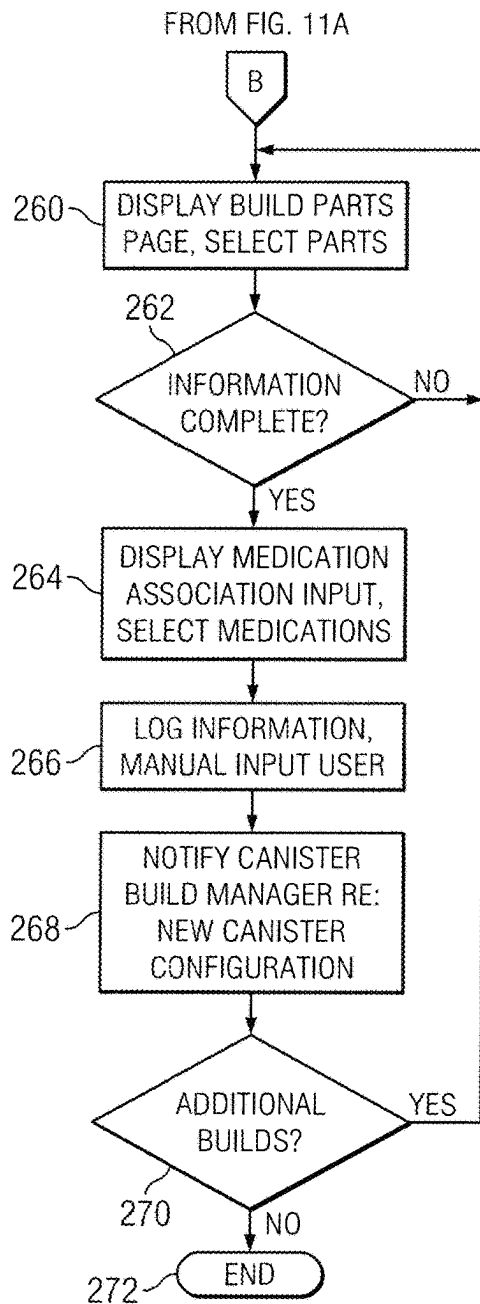
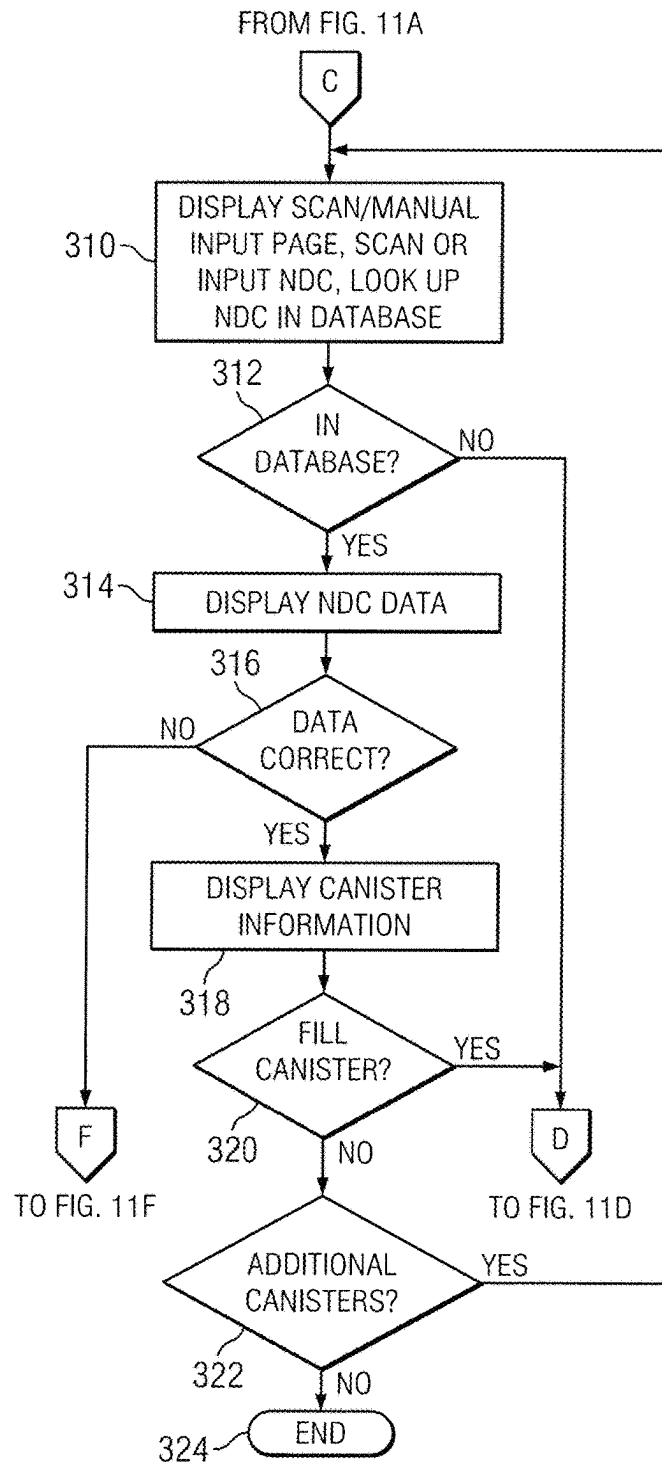
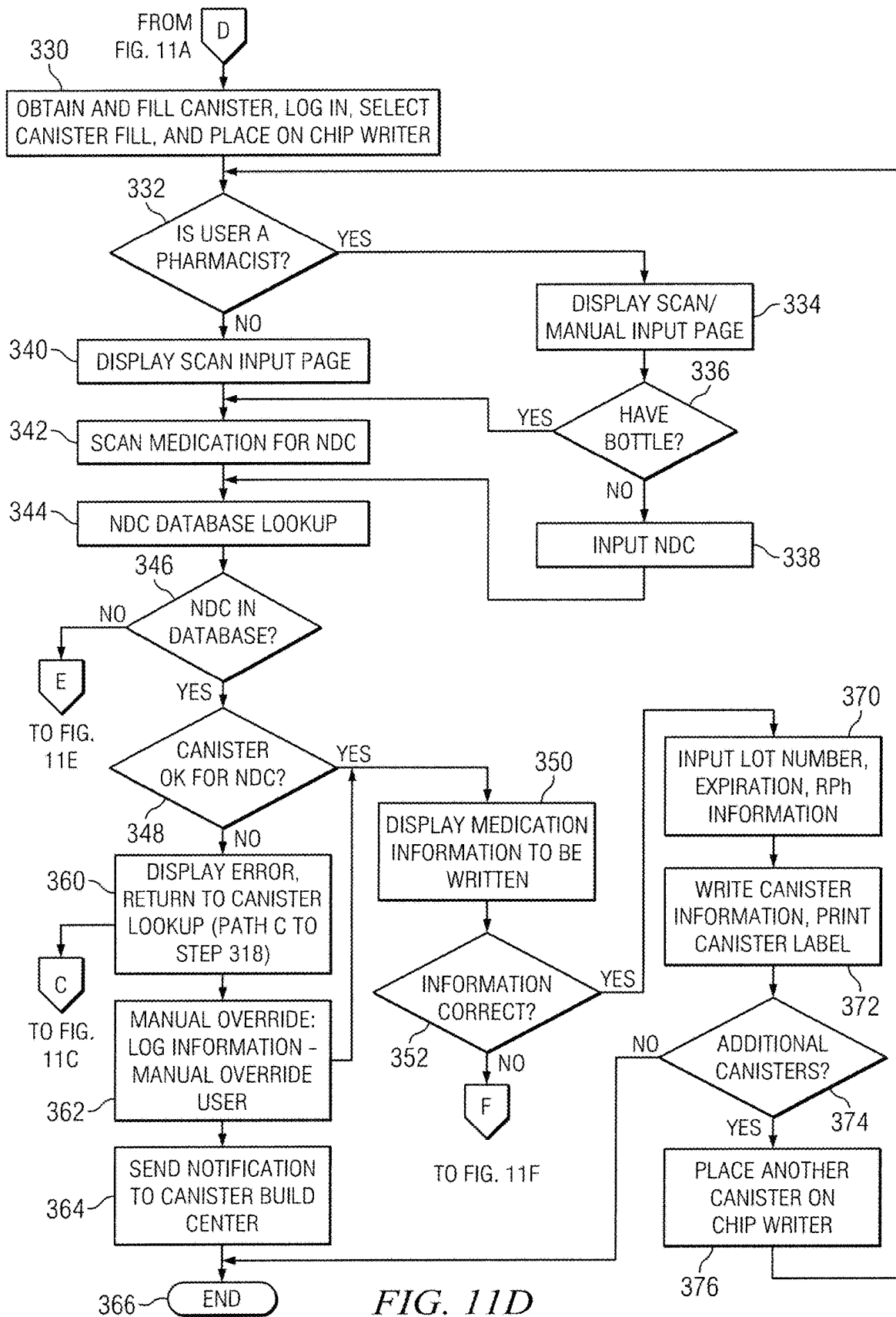
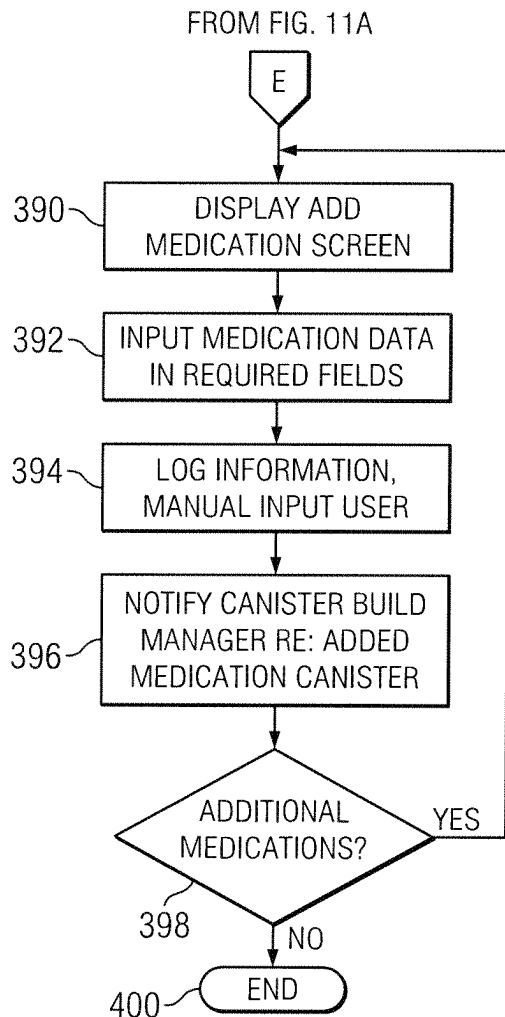
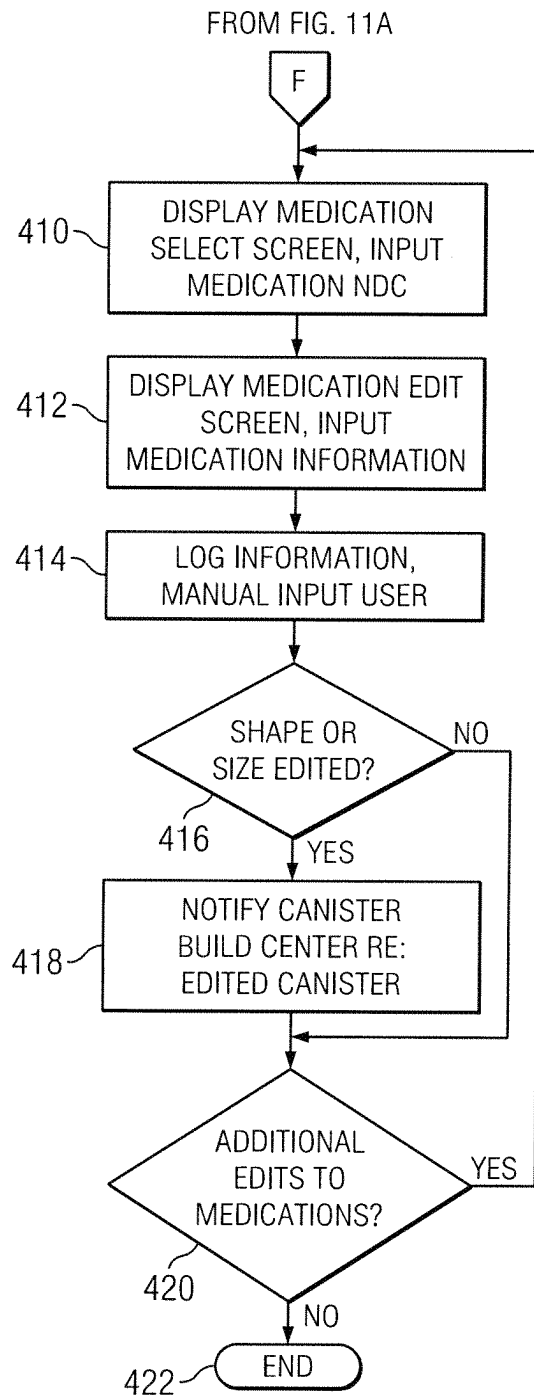


FIG. 10

*FIG. 11B**FIG. 11C*



*FIG. 11E**FIG. 11F*

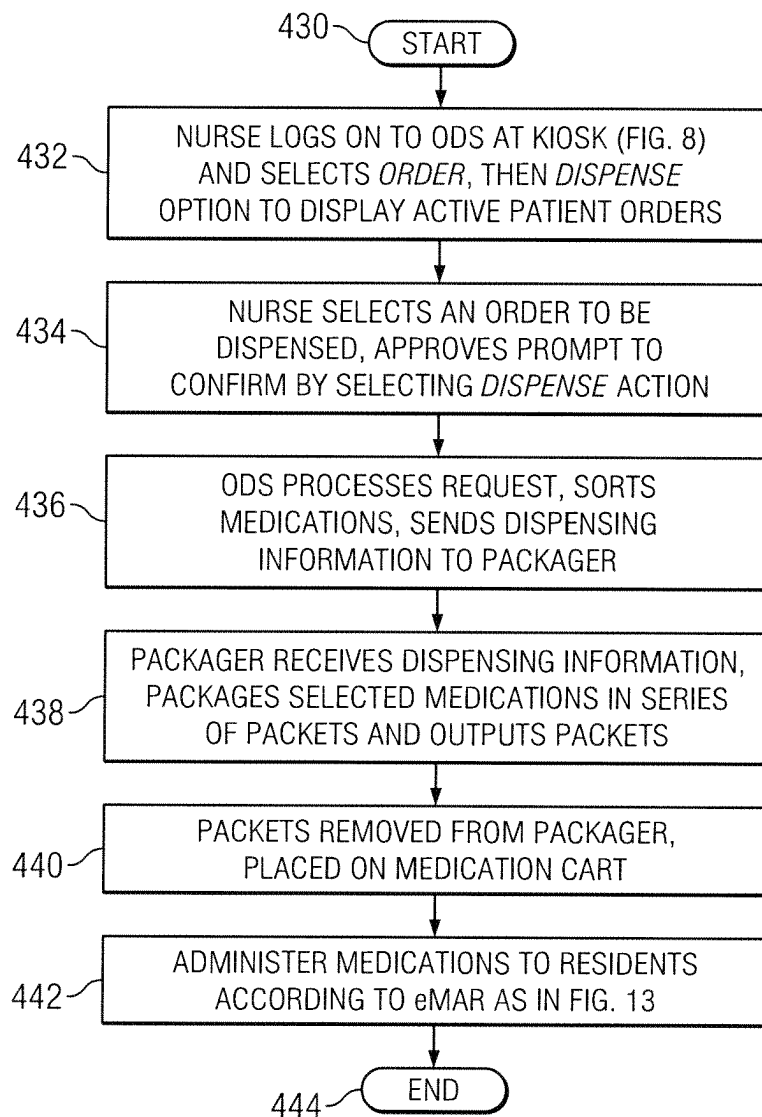
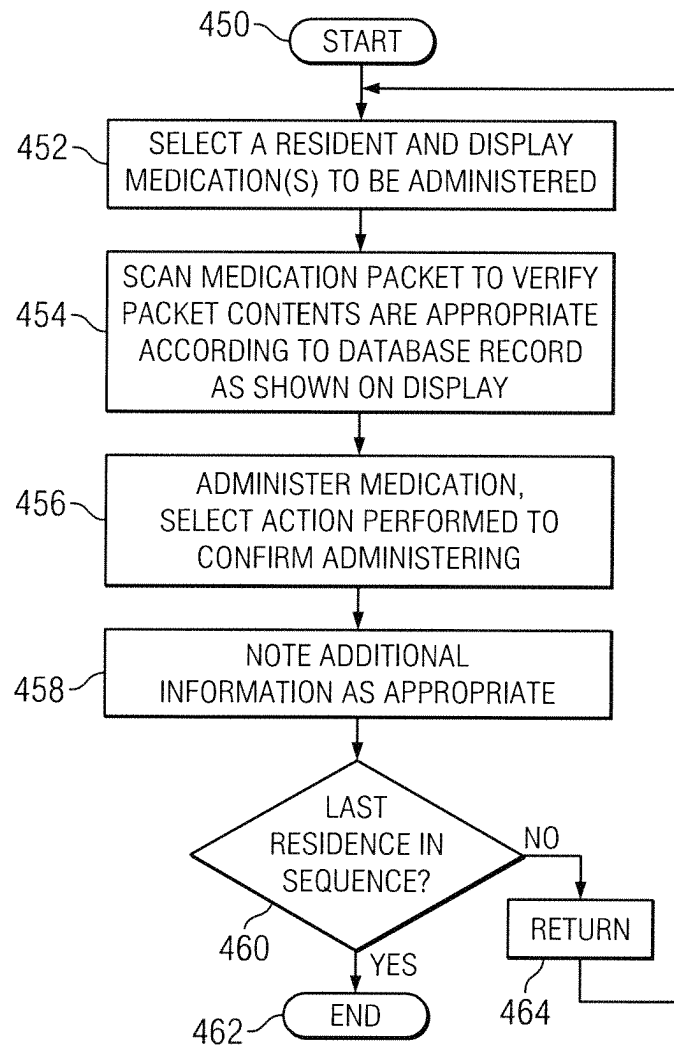


FIG. 12

*FIG. 13*

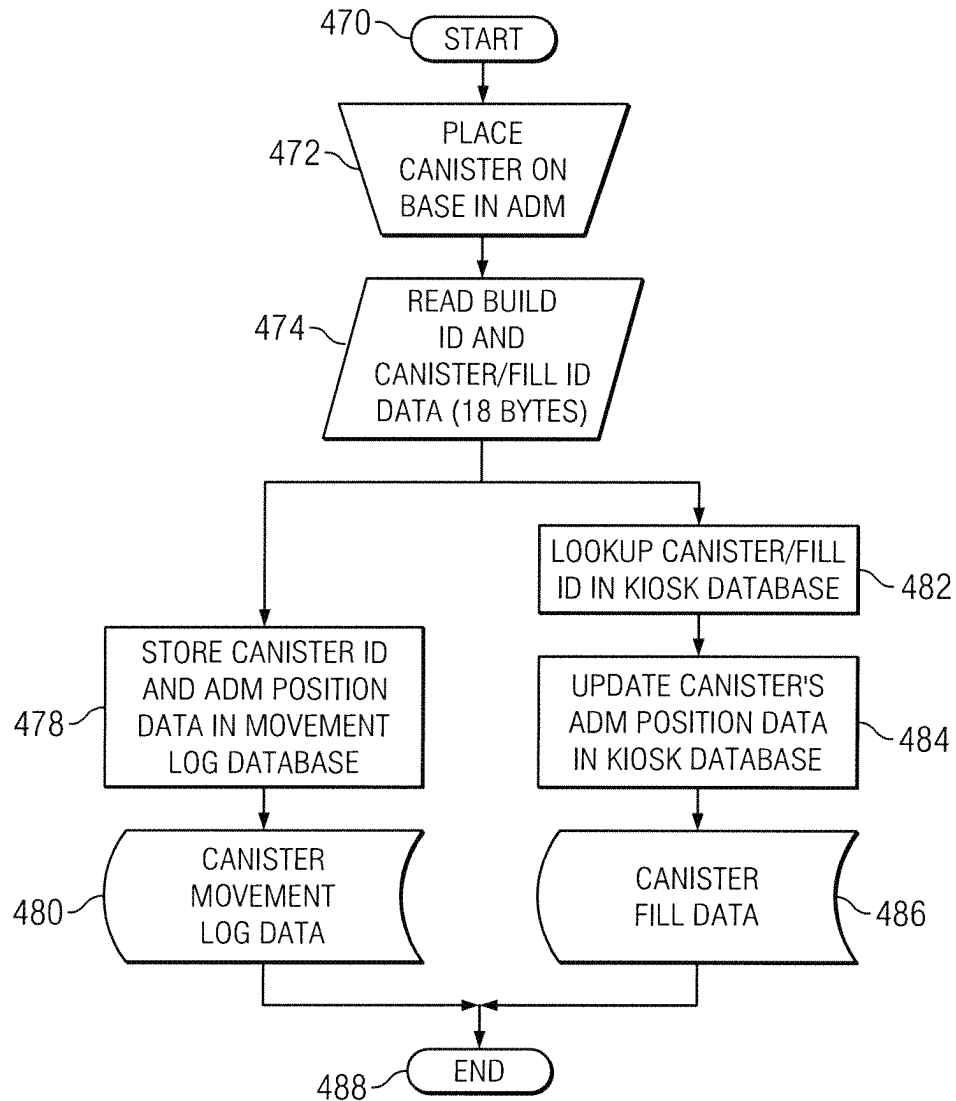


FIG. 14

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MEDICATION MANAGEMENT SYSTEMS AND METHODS FOR HEALTH AND HEALTH RELATED FACILITIES

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation of U.S. patent application Ser. No. 17/012,829, filed Sep. 4, 2020, and entitled "MEDICATION MANAGEMENT SYSTEMS AND METHODS FOR HEALTH AND HEALTH RELATED FACILITIES," which is a continuation of U.S. patent application Ser. No. 16/259,716, filed Jan. 28, 2019, and now issued as U.S. Pat. No. 11,217,058 and entitled "MEDICATION MANAGEMENT SYSTEMS AND METHODS FOR HEALTH AND HEALTH RELATED FACILITIES," which is a continuation of U.S. patent application Ser. No. 15/050,460, filed Feb. 22, 2016, and now issued as U.S. Pat. No. 10,192,035 and entitled "MEDICATION MANAGEMENT SYSTEMS AND METHODS FOR HEALTH AND HEALTH RELATED FACILITIES," which is a continuation of U.S. patent application Ser. No. 13/085,298, filed Apr. 12, 2011, and now issued as U.S. Pat. No. 9,268,912 and entitled "On Site Prescription Management System and Methods for Health Care Facilities," which claims priority to U.S. Provisional Patent Application No. 61/323,125, filed on Apr. 12, 2010, and entitled "On Site Prescription Management System and Methods for Health Care Facilities," all of which are hereby incorporated by reference for all purposes.

TECHNICAL FIELD

The disclosure generally relates to medication delivery in health and health related facilities using automated pharmacy dispensing systems and more particularly to automated medication prescription and dispensing, inventory management and control, and information processing in facilities including but not limited to resident health care facilities such as nursing homes, assisted living centers, retirement homes and senior communities with health care facilities, long term care facilities, prisons, and the like.

BACKGROUND

Management of automated prescription medication delivery in a resident health care facility, which may be referred to generally herein as a long term care facility or LTC, particularly those with in-house pharmacies and/or dispensaries involves multiple participants and a variety of complex operations. These include communicating prescriptions, dispensing medications as directed and on time, keeping accurate and timely records of the medications dispensed, receiving new and replenishment prescriptions into a facility or pharmacy-controlled management system, processing payment for the medications from a variety of providers, managing and tracking the inventory of drugs to be dispensed, providing accurate and timely feedback about the dispensing and actual administration of the medication, stock levels and rates of dispensing of the items in inventory, scheduling and requesting reorders, managing and monitoring of controlled substances, performing drug interaction and allergy analyses, all with little or no centralized control system to facilitate management and ensure compliance with regulatory and standard operating procedures required for these diverse operations.

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These tasks must be performed accurately and efficiently to ensure safety and security of the patients and their medications, and minimize loss and waste of controlled substances and unused medications.

Traditional pharmaceutical delivery systems that employ written and spoken communication of information and hand-carried delivery are frequently characterized by very substantial inefficiencies and opportunities for errors, occasionally with grave consequences. At nearly every step of the way from writing a prescription, getting it filled and records generated, delivering it to a long term health care facility and providing for administration of the medication at the appointed time to the correct patient, opportunities for mistakes abound because the process involves numerous people handling the records and the medication for each instance of delivery and administration to fulfill the prescription. Further, the flow of information associated with management of the process takes place slowly, in serial fashion, and involves frequent handling of paper records.

In recent years a number of systems have been developed for use in various kinds of in-patient or resident health care facilities to automate the process. Automated dispensing machines have been developed to package and dispense medications to personnel tasked with administering the medications to patients on predetermined schedules according to the "script," the instructions accompanying the prescription. Portions of the process may be brought under computer control to manage the flow of information. Although computers are used to a substantial extent to handle the volume of data associated with dispensing prescription medications, the control and packaging of the medications, the patient records for the medications prescribed by physicians, processing payment by third parties, etc., inefficiencies and the opportunities for errors and substantial waste remain.

One ongoing problem in many systems is the amount of time delay that occurs from the instant a prescription is issued by a physician until the initial dose is dispensed and administered to the patient. The consequences of delay and errors in administering the drugs prescribed can be serious and even catastrophic. Accordingly there is a need for system and methods that minimize this delay and include controls that minimize error in the processes involved in the delivery, management, dispensing and administering the medications.

Another problem with conventional systems is the lack of timely feedback regarding the status of processes at each step from incoming prescription to administering the prescribed dose at the appointed time, and the inability of conventional systems to operate in a forward-looking mode, that is, to know what medications are needed in the future according to present prescriptions so that inventory levels may be regulated and medications reordered and distributed to patient locations before an out-of-stock condition occurs. Present systems are "reactive," that is, they are designed to react to out-of stock or par level conditions. A consequence of the reactive operation is that residents may have to go without a dose at the prescribed time because the medication was not dispensed on time due to an out-of stock condition caused by sudden increases in usage or improper inventory control. A missed dose can have serious consequences to some residents or patients. Thus, what is needed is a "proactive" system that knows when a supply of a medication will be emptied and provides for distribution of the replenishment on a schedule that maintains the dispensing of the medications in an uninterrupted flow, even in instances of sudden increases in usage.

A related problem with conventional systems is the opposite condition when the processes of dispensing medications are not sufficiently controlled—i.e., inventory levels are not well-matched to prescription requirements or the tracking of medications dispensed and administered is insufficient. The result is substantial quantities of medications that are not administered and must be discarded because, for safety and security reasons, there is no provision for re-stocking the unused medications. This circumstance causes waste that requires controlled procedures for destruction of the unused substances. Elimination of such waste would result in reduced costs to the long term care facilities, resident patients and their families, insurance providers and taxpayers, and enhance both safety and efficiency of the prescription process.

Yet another deficiency of existing systems is an inability of various stations or functional entities of the system to interact with each other or to be organized so that their functional operations are coordinated to minimize lost opportunities to respond to current and anticipated conditions because prescription data is not available or the status of a dispensing process is not known, etc. Accordingly there is a need for synchronizing the operations and processes in the system so that process flows make use of all available data and are coordinated with each other to ensure efficient and accurate dispensing and administering of medications. Especially important would be the ability to provide direct communication between prescription and order entry and the dispensing processes and apparatus, and the dispensing apparatus and the records associated with dispensing and administering of the medications.

While the use of automated packaging and dispensing machines located in the long term care facility or pharmacy may provide some efficiencies, and alleviate some of the bottlenecks, unless they are used to maximum advantage, few of the aforementioned problems are actually solved. The problems are more in the nature of information flow than automating some portion of the packaging or handling of prescription medications. In one example the data that defined the identity and contents of each canister in a dispensing machine occupied more than two kilobytes of data. One typical example of this prior art technique is illustrated in FIG. 1, which depicts the layout of the canister memory space **10** having a first row **12** specifying the byte population of the memory, a second row **14** identifying the data stored in the memory **10**, and a third row **16** defining the hexadecimal address of the various items of data. Of interest are the data sections of the second row: canister ID **18** (3 bytes), the Chip ID **20** (15 bytes), and, most significantly, the Chip Memory Data **22** that describes the prescription medication contained in the canister (more than two kilobytes). When two kilobytes of data is multiplied by several hundred times—the number of canisters in a typical dispensing machine—the processing time takes on significant dimensions, becoming a substantial cause of processing delay, and dispensing errors due to the delay in processing the data in the prescription processing and dispensing operation.

In another example of the problems encountered when using automated packaging and dispensing machines, consider that in such systems the containers from which the medications are dispensed, often called canisters, must be available in a variety of sizes and configurations, each adapted to a particular physical size and shape of a medication unit, pill, or tablet. The result is each medication can only be contained in one configuration of canister. Some medications having the same formula are available in multiple sizes and shapes; others in only one size and shape; still

other generic forms of the same drug, available from different manufacturers, may require a variety of different configurations. Further, the size and shape of medications are subject to change. This variety of configurations complicates the process of determining the most efficient container or canister configuration. The result can be inefficiency and compromise, which is wasteful of resources, more susceptible to errors, and often wasteful of medications because accommodating this variety is cumbersome, time-consuming, and expensive.

SUMMARY

Accordingly there is provided a system for controlling dispensing of medications in a health care facility coupled to a network, comprising: an order management system (OMS) and an associated database operating on a hosted (central) remote server coupled to said network, said OMS serving said health care facility via said network; a prescription order entry (POE) system coupled via said network to said order management system; a medication dispensing system installed in said health care facility and coupled via said network to said order management system; and an electronic medication administering record (eMAR) coupled through said medication dispensing system via said network to said order management system.

In another embodiment a method is provided for automated dispensing of prescription medications to one or more health care facilities via associated remote pharmacies, comprising the steps of: providing a central hosted remote server having a system database and coupled to a network and operating an order management system (OMS) containing interface links to a plurality of remote operative units; providing at least one medication dispensing system in each health care facility, wherein each said medication dispensing system includes an automated prescription drug packaging and dispensing machine (ADM) and a work station coupled thereto, and each said automated dispensing machine (ADM) is coupled to said central hosted server; coupling a prescription order entry system (POE), accessible from said health care facility, to said central remote server via said network; including in said automated dispensing machine (ADM) a plurality of medication canisters for storing and dispensing prescription drugs therefrom; and associating with each said canister a memory device having a storage capacity not exceeding 48 bytes.

Further, a system for control of prescription drug dispensing is provided comprising: a database and a prescription order entry (POE) system coupled respectively via a network to an order management system (OMS), said OMS serving an in-patient health care facility via said network; an automated drug packaging and dispensing machine (automated dispensing machine or ADM) installed in said health care facility and coupled to said OMS; and an electronic medication administering record (eMAR) coupled to said OMS through said automated dispensing machine.

In other aspects, the automated dispensing machine comprises a cabinet containing a plurality of individually removable canisters for storing an inventory of prescription medications selected according to resident's needs; a processor responsive to communication with said OMS and said eMAR and configured for control of said canisters; and a canister memory associated with each canister and accessible to said OMS and said eMAR, said canister memory location containing a build code ID and a canister fill ID, wherein said build code ID comprises a code representing a particular medication and a species thereof, and said canister

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fill ID comprises a code representing said canister and a suffix code representing each instance of filling said canister.

In another aspect, the system further comprises a mechanism for packaging a medication for a patient and a mechanism for dispensing said medication to a patient care worker.

In another aspect, the system further comprises a computing device having a display and data entry provision for requesting dispensing of and recording administration of a prescription medication to a resident patient.

In another aspect, the computing device further comprises a scanning device for reading an encoded label.

In an alternate embodiment, a method is provided for dispensing prescription drugs in a health care facility, comprising the steps of stocking an automated dispensing machine (ADM) installed at said health care facility and linked to an order management system (OMS); entering patient prescription information into a prescription order entry (POE) system in communication with said order management system; compiling a comprehensive patient medication record in an electronic medication administration record (eMAR) contained in a handheld eMAR device coupled to said ADM; and linking said eMAR through said ADM to said OMS.

In another aspect, the method comprises the steps of updating said eMAR with each dispensing operation from said ADM; and updating a database coupled to said OMS according to said eMAR.

In another aspect, the method comprises the step of updating a re-order queue according to said data stored in said database coupled to said OMS, including scanning label information from said eMAR device for entry into said eMAR.

In yet another alternate embodiment, a method is provided for reducing delay in processing item data in an inventory control system, comprising the steps of formatting a data transfer message to exclude said item data to be transferred; limiting the contents of said data transfer message to a build code identification statement and a canister fill identification code statement; and storing said item data to be transferred in a database location identified by said build code statement, wherein said build code identification (ID) comprises a code representing a particular stock item and a species thereof, and said canister fill identification (ID) comprises a code representing said canister and a suffix code representing each instance of filling said canister.

In an alternate embodiment a method for coordinating the processing of data items among participating entities in the system wherein the data being processed is synchronized by a combination of user tokens (who), data identifiers (what), and unique IDs for the entities in the system (where).

DESCRIPTION OF THE FIGURES

The accompanying drawings facilitate an understanding of the various embodiments:

FIG. 1 illustrates one example of a prior art memory format for prescription medications;

FIG. 2 illustrates an example of a more efficient memory format for a prescription medication according to at least a first embodiment as disclosed herein;

FIG. 3 illustrates a front perspective pictorial view of a medication canister for a dispensing machine for use in at least the first embodiment as disclosed herein;

FIG. 4 illustrates a rear perspective pictorial view of the medication canister of FIG. 3;

FIG. 5 illustrates a general system block diagram according to one embodiment disclosed herein;

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FIG. 6 illustrates a system block diagram according to a variation of the embodiment of FIG. 5;

FIG. 7 illustrates a system block diagram of the basic architecture of one embodiment disclosed herein;

FIG. 8 illustrates an automated prescription dispensing machine and an associated work station disclosed herein, for use with the embodiments of FIGS. 2 through 7;

FIG. 9A illustrates a first portion of a flow chart for the basic operation of the system of FIGS. 6 and 7;

FIG. 9B illustrates a second portion of the flow chart for the basic operation of the system of FIGS. 6 and 7;

FIG. 10 illustrates a flow chart of one embodiment of a process 280 for formulary substitution as disclosed herein;

FIG. 11A illustrates a flow chart of initial steps to begin a medication canister configuration and fill process according to at least one embodiment as disclosed herein;

FIG. 11B illustrates a flow chart of a process to create a canister build as disclosed herein;

FIG. 11C illustrates a flow chart of a process to configure a canister as disclosed herein;

FIG. 11D illustrates a flow chart of a process to fill a canister as disclosed herein;

FIG. 11E illustrates a flow chart of a process to add a medication canister as disclosed herein;

FIG. 11F illustrates a flow chart of a process to edit a medication canister as disclosed herein;

FIG. 12 illustrates a flow chart of a process to prepare medications for administering them to residents of a long term care facility as disclosed herein;

FIG. 13 illustrates a flow chart of a process for administering medications to residents of a long term care facility as disclosed herein; and

FIG. 14 illustrates a simplified flow diagram for reading medication data from a canister memory as disclosed herein.

DETAILED DESCRIPTION

In the following detailed description of the present invention, various structural components illustrated in the drawings will be identified by a reference number. When the same component appears in more than one figure, the same reference number will generally be used. However, in some cases, such as when the embodiment is a different one, a different reference number may be used in a particular drawing. The description is to be read in conjunction with the drawings. The term health care facility as used herein may mean any facility that provides health care. The facility may include dispensing of medications to patients or residents of the facility. Examples include long term care facilities, nursing homes, prisons, certain hospitals, retail facilities, and the like. The term "dispensing machine" is used synonymously with "automated prescription drug dispensing machine" and "automated dispensing machine (ADM)."

The invention to be described herein concerns automated pharmacy dispensing systems and methods. Disclosed herein are a number of distinct embodiments including: (1) a system for control of prescription drug dispensing comprising a novel combination of an automated prescription drug packaging and dispensing machine in a health care facility plus off-site management and control coordinated by a main server coupled via a global communications network to the operating entities in the system; (2) a method for dispensing prescription drugs in a health care facility, that is, the use of the novel system identified above; (3) a method for reducing delay in processing data in a prescription drug order and dispensing system of a health care facility that

relies in part on re-defined contents of data that is incorporated into each canister installed in the packaging and dispensing machine of the system; and (4) a method for coordinating or synchronizing the operating entities in the system with respect to the data being processed during the operation of the above prescription drug dispensing system. The synchronization system ensures that all entities in the system, including the stations or kiosks in the health care facilities connected to the network, are operating on the correct data. In addition, (5) the system is fully scalable and can accommodate multiple health care facilities, multiple pharmacies, and multiple automated dispensing machines in each health care facility.

In a first illustrative embodiment, the overall system is illustrated in schematic block diagram form in FIG. 5, which shows the basic structure of the system as depicted by its primary operating components. In general, the system may accommodate multiple health care facilities such as long term care facilities (LTCs) and pharmacies, all connected to a main server operated by a hosting center as a typical example. FIG. 6 presents a similar schematic block diagram showing the structures of the long term care facilities and the pharmacies, and the functional relationships among the operating entities and components of the system. FIG. 7 illustrates in block diagram form the architectural relationships of the functional components of the system. As will be explained, the portions of FIG. 7 having a bold line outline depict new structures created specifically for the present invention. FIG. 8 depicts a prescription drug dispensing system that includes one form of an automated prescription drug packaging and dispensing machine, which will often be referred to as an automated dispensing machine (ADM) herein, and a kiosk or work station equipped with a PC or server, a scanner, and optionally, a printer.

In a second embodiment, FIGS. 9A and 9B depict a basic operation of the system. FIGS. 10 through 14 illustrate other basic aspects of the methods of using the system of FIGS. 5, 6, 7 and 8 to dispense prescription medications in a health care residence or facility. Some of these operating relationships may be understood by referring to FIGS. 11A through 11F. Several specific communicating links and signal flows in FIG. 7 that represent new flows of information in pharmaceutical management processing systems are depicted by highlighted lines.

In a third embodiment shown in FIG. 2, reduction of the data content of the canister memory chip to the combination of a build code ID and a canister fill ID, each requiring only relatively small amounts of data, enables a very substantial reduction in processing time resulting in much less inherent delay in processing the system inventory, the tracking of dispensing problems, and the dispensing of prescriptions and prescription information. The canisters themselves are described in FIGS. 3 and 4, and the process of using the modified canister system is illustrated in FIG. 8. Detailed data regarding the prescription drug information associated with the build code ID and the canister fill ID is stored in the main database accessed by the management system and referred to only as needed.

The build code ID may comprise a four digit code representing the structural configuration (stored in a database) of the canister that is compatible with a particular group of medications. Medications are supplied in a large variety of forms, sizes, shapes, etc. ("attributes"). The structural components of each canister are adapted to accommodate a particular group of such attributes. The canister fill ID may comprise a 12 digit code that uniquely identifies the canister. In one example, the code may represent when (the

date and time) the build code ID is first assigned and a three digit suffix code representing each instance of filling said canister with said medication. Other data associated with the canister fill ID may be stored in a database location corresponding to the canister fill ID. Other code types or length for the build ID and canister fill ID codes may be used in variations of the exemplary embodiment described herein.

In a fourth embodiment, a method for coordinating data being processed by the various entities in the system is provided through assignment of ID information to enable synchronization of the data of the various stations on the network as needed. It is this coordination process that enables the system communication efficiencies leading to substantially improved processing speed and the ability to proactively manage the forward-looking inventory control aspects of the system.

Beginning with the detailed description, FIG. 2 illustrates an example of a more efficient canister memory format for a prescription medication according to at least one embodiment disclosed herein. The figure depicts the layout of the canister memory space 30 as disclosed herein having a first row 32 specifying the byte population of the memory in each of the identified groups, a second row 34 describing the data content stored in the memory 30, and a third row 36 defining the hexadecimal value ranges of the various items of data. Of interest are the data content sections of the second row: a Build ID 38 (3 bytes), and the Canister/Fill ID 40 (15 bytes). It is important to notice that the "Chip Memory Data 22" of the prior art, which used in excess of 2,000 bytes of data is no longer stored in the canister memory 30. This is because the inventor realized that such detailed information about the prescription drug contained in the canister, which occupied more than two kilobytes of data, no longer needed to be stored in the canister itself. This data may be stored in a database location and accessed as needed outside the most important real time sequences for operation of the system as will be subsequently described. The result of this insight is a very substantial reduction in processing time leading to a major improvement in efficiency and responsiveness of the system to the needs of the health care facility.

FIG. 3 illustrates a front perspective pictorial view of a medication canister for an automated prescription drug dispensing machine (ADM) for use in the illustrative embodiment of the present invention. The canister 50 may be assembled from interchangeable parts including a transparent main body 52 that includes a transparent hopper section 54, a lid 56, a memory receptacle 58, a memory chip 60, a division block 62 with drive hub 64, and a security tape 66. The division block 62 is a rotating distributor that indexes one position for each medication pill, tablet or capsule to be dispensed. The rotation is provided by a drive mechanism within the dispensing machine to be described that is coupled to the drive hub 64. The drive hub 64 has an internal spline to engage the drive shaft of the dispensing machine in a non-slip manner. The engagement of the drive shaft and the splined drive hub 64 occurs when the canister 50 is selected for dispensing and moved into position in engagement with the drive mechanism. Others of the interchangeable parts of the canister 50 will be described for FIG. 4.

The canister 50 may include a detailed medication fill label 68 attached to the lid 56 of the canister with printed specifications of the prescription drug medication and a machine-scannable portion of information 69 such as a bar code or the like. The canister fill label 68 may further include a duplicate machine-scannable portion of information 69 wrapped over the upper front edge of the canister as shown

to enable scanning the label when the canister is installed in a dispensing machine. The canister may further include a build label **70** that has a four digit build code **72** and bar coded build information **74**. The build label **70** is preferably printed when the canister **50** is constructed and may be permanently affixed to the canister **50** throughout its life. The build information is scanned whenever the canister is handled so that the system always knows the identity and location of the canister and can thereby reference its contents with database information whenever the build label **70** is scanned. The medication fill label **68** is preferably printed and affixed to the canister **50** each time the canister is prepared for installation in an automated dispensing machine. Locating the scannable build code label **70** on the front, handle portion of the canister facilitates rapid identification and logging of the canister regardless of where it is in the system or how it is being handled. The canisters are prepared in a pharmacy, which may usually be located at the health care facility or off-site, following any maintenance on the canister before use or reuse. Maintenance may include removal of old labels, replacement of worn or broken parts, inspection, cleaning, etc. before restocking for reuse. In some cases the canisters may be modified or configured to accommodate different medication attributes—that is, different form, shapes, and sizes of the tablet, pill or capsule that will be stored therein for dispensing. A new build ID is assigned to the canister when it is reconfigured.

FIG. **4** illustrates a rear perspective pictorial view of the medication canister **50** of FIG. **3**. In this view from slightly below the canister **50** the main body **52**, transparent portion **54** and lid **56** remain visible, along with the division block **62**, the drive hub **64**, the memory receptacle **58** and the memory chip **60**. The memory chip **60** may preferably be a printed circuit board component having connection traces on one of both of its faces for mating with corresponding conductors when inserted into a receptacle as the canister is installed in the dispensing machine. Thus, the memory chip **60** is connected into the circuitry of the dispensing machine so that its contents may be read during operation of the machine to access the canister (i.e., bring it into a dispensing position), dispense a unit (or the required number of units) into a package, and arrange the packages in a predetermined sequence for dispensing to the person who will administer the medication to the resident intended to receive the unit(s) of medication. Although this process is generally operated under programmed control the facility may dispense medications whenever it chooses, for example in PRN or demand circumstances.

Also visible in FIG. **4** are a partition **76**, a partition fix block **78**, and a guide block **80**. The partition **76** is a flexible arm, fixed to the fix block with screws as shown. The fix block **78** includes several threaded holes to permit adjustment of the partition **76** in a vertical direction relative to the lower portion of the division block **62**. The flexible arm includes a small tab (not shown) on the side of the partition **76** facing the division block **62** and acts to assist in dispensing a unit of medication during operation of the dispensing machine. The guide block **80** is fixed to the underside of the main body **52** of the canister and acts as a key to locate in the transport mechanism (not shown) of the dispensing machine. The transport mechanism can accommodate several different sizes (capacities) of canisters, each with its own keyed receptacle to match the guide block **80** of the appropriate canister **50**. The interchangeable parts of a canister **50** may include at least the main body **52**, the division block **62**, the partition **76**, and the guide block **80**.

The canisters illustrated in FIGS. **3** and **4** may generally contain a 10 to 30 day supply of prescription medications of an individual resident or patient of a health care facility. The quantity of medications depends strongly on their costs, so the costs of maintaining an inventory is an important consideration. The system may accommodate supplies of up to 90 days if needed. In one alternative, for example, a canister may contain a larger quantity of medications of a type to be dispensed to more than one resident in an area served by the dispensing machine. The dispensing machine provides for secure storage, convenient packaging, and accurate dispensing of prescription medications on schedule, “STAT” (immediately), or on demand (also referred to as “PRN” or as the resident or patient requests), automatically or under computer control, under the supervision or administration of a director of nursing in the typical facility.

FIG. **5** illustrates a general system block diagram according to one embodiment of the present invention. A hosting center **84** is shown that includes a main server **86** and a resource center (ORC) **88**. The main server **86** operates an order management system (OMS) **90**. The main server **86** further includes a main database **114** coupled thereto. The order management system **90** is a comprehensive software system that runs on the main server **86** and includes a suite of interfaces to a number of other functional units of the system, in addition to the communications and other house-keeping operations provided for in the order management system **90**. The main server **86**, order management system **90**, and database **114** together form the operating hub of the prescription drug ordering and dispensing system of the present invention. The resource center **88** is coupled to the main server **86** and provides a website presence for the functional units of the system to deposit and obtain information concerning operations of the system. These units typically are located in the hosting center **84**, which in general may be provided by firms outside of and not otherwise connected or related to the health care facilities or pharmacies participating in the dispensing system described herein.

Communication between the main server **86** and the functional operating units of the system occurs in the illustrative embodiment through connections to a global communications network **82** such as the Internet. The other principal functional operating units of the system coupled to the network **82** include one or more pharmacies **92**, designated as pharmacy **1**, pharmacy **2**, etc., and pharmacy **P**; and one or more long term care (LTC) facilities **94**, designated similarly as LTC **1**, LTC **2**, etc., and LTC **Q**. Long term care facilities **94** have been selected in this exemplary system to illustrate the principle concepts disclosed herein. Persons familiar with the prescription drug processing and delivery systems in current use will appreciate the generality of the concepts described herein and recognize their applicability to other types of resident health care facilities such as nursing homes, assisted living centers, rehabilitation facilities, and even prison systems where health care is provided. All of these functional operating units coupled to the network **82** are equipped with communications interfaces to enable the required exchanges of data and commands for the system to perform its operations.

Each LTC facility **94** preferably includes a director of nursing or administration **96**, which may be represented by a workstation PC (not shown), an automated prescription drug dispensing machine (ADM) **100**, and a kiosk **102** coupled directly to the ADM **100**. As will be described, the

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kiosk **102** may be a workstation having a PC with display and a scanning device, and perhaps a server, database, printer, etc. in each LTC **94**.

FIG. **6** illustrates a system block diagram according to a variation of the embodiment of FIG. **5**, wherein further details of the pharmacy **92** and a long term care facility **94** in a typical system according to the present invention are illustrated. The network **82** and the hosting center **84** containing the operating units of the main server **86**, database **114**, resource center **88**, and the order management system **90** are as illustrated in FIG. **5**. The pharmacy **92** in this embodiment may include a pharmacy management system **122** and a pharmacy information system **124**, each coupled to the other and operated by software applications residing on their respective PCs or server. Each of these units may be independently coupled via the network to the main server **86** as will be described. The operative portion of the long term care facility **94** is its prescription drug dispensing system **98**. The dispensing system **98** includes the automated dispensing machine **100**, the kiosk **102** coupled to it that may include an eMAR server **104** operating on the kiosk, along with the other components mentioned for the embodiment of FIG. **5**, including an auxiliary database **116**, a work station with display, a scanning device, etc. FIG. **6** also depicts a LTC Director of Nursing work station **96**, which may be coupled directly to the resource center website **88** via a link **110** through the network **82**. A long term care facility **94** may include a plurality of automated dispensing machines **100**, each under the control of a respective kiosk **102** or a PRN dispensing eMAR **106**—a laptop PC or handheld computer (alternately, slave kiosk)—as shown, each of which is connected to the eMAR server **104** at a kiosk **102** via a local communications bus **112**, which may be internal to the dispensing system **98**. Connected to the same communications bus **112** may be additional handheld or tablet PCs **108**. These multiple ADMs **100** and PCs **106**, **108** may be needed in larger resident patient facilities. Thus, the system can be scaled to include operation with a plurality of dispensing machines **100** from, for example, a single kiosk having a workstation or eMAR server **104**, auxiliary database **116**, scanning device, etc., as illustrated and will be more fully described regarding FIG. **8** herein below.

FIG. **7** illustrates in schematic form a system block diagram of the basic architecture of one illustrated embodiment disclosed herein. The units represented by block outlines are essentially software providing various functions operating within the system, including communication of data via their respective connections to the network **82** as described. Functional units depicted as a block having a bold outline contain software written especially for the present invention. Further, it will be appreciated that the order management system **90** is an essential and central piece of the system because it provides the overall management and control of the system; that is, it integrates the functions of all the operating units connected to the order management system **90** into the seamless operation necessary to provide an efficient and responsive prescription drug dispensing system that overcomes the inefficiencies and inadequacies of the prior art dispensing methods. These advantages will become apparent in the description that follows. The software of the various operating units runs on various PCs or servers as described for FIGS. **5** and **6**. In one section of FIG. **7** several functional units are surrounded by a dashed line and designated “Client Facing Systems.” This means these units are located in or accessible from the long term care facilities **94** via the network **82**. Other functional units are located in a pharmacy system **92**, which may be located

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either outside the long term care facility **94** or within the LTC facility. Still other functional units may typically be external to the system because they represent other entities with which the present system interacts, such as the prescription drug plans **128** and wholesaler systems **130**.

Continuing with FIG. **7**, the order management system **90** includes the following interface programs for communicating with the various external functional units in the long term care facility **94** (or facilities **94**) or elsewhere. As shown in the drawing, each of these interface programs connects to communication links with other interface programs internal to the order management system **90**, or with functional operating units outside the order management system **90** via communication through the network **82**. These links are designated with capital letters and numerical symbols as shown in FIG. **7**. A dispensing interface **140** is linked via the link **C1** to the prescription order entry (POE) **120** and to the eMAR interface **142** respectively in the order management system **90** and the order dispensing system (ODS) **102** in the LTC **94** via a link **D1**. The eMAR interface **142** is further coupled with the eMAR functional unit **104** in the LTC via link **H1** and also to the resource center (ORC) **86** and the billing interface **152** internal to the order management system **90** via the link **E3**.

Further in the order management system **90** is located a reporting interface **144** that is connected via the link **E1** to the resource center **86** and via the link **J1** to a pharmacy management system (OPS) **122**. An administration interface **146** in the OMS **90** is connected via the link **E2** to the resource center **86** via the link **E2** and to the pharmacy Management system **122** via the link **J2**. The billing interface **152** is also connected via the link **G1** to a pharmacy information system (PhIS) **124**. The wholesaler interface **154** is shown coupled to the pharmacy management system **122** via the link **J3** and to one or more wholesaler systems **130** via the link **K1**.

Continuing with the OMS **90** interfaces, a formulary benefits interface **156** is coupled via the link **M1** with the prescription drug plans (PDPs) **128** and via the link **C2** with the prescription order entry functional unit **120**. An electronic health record (EHR) **158** is connected via a link **F1** and the network to an EHR **126** in the long term care facility **94**, and to the prescription order entry (POE) unit **120** via the link **C3**.

The foregoing lines of communication identified by the letters **C**, **D**, **E**, **F**, **G**, and **H** (including **C1**, **2**, **3**; **D1**; **E1**, **2**, **3**; **F1**; **G1**; and **H1**) between the order management system **90** and the functional units across the network **82** form a combination not known to exist in the prior art. That is, management and control of these on site functional units in a health care facility, via communication over a network such as the Internet from an off-site main server **84** running a suite of software such as the order management system (OMS) **90** of the present invention is not known to exist prior to the invention thereof as disclosed herein. There are many benefits to this architecture as will become clear in the description which follows.

Further, the on-site communication links identified as the order entry bus **A** and the eMAR bus **B**, respectively connecting the order dispensing system (ODS) **102** with the prescription order entry unit **120** and the electronic medication administration record (eMAR) **104** form a structure unique to prescription dispensing system in use in resident health care facilities. These linked-together functional entities in the health care facility provide substantial enhancements to the efficiency and accuracy of the administration of prescription medications in health care facilities that provide

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health care to their residents in addition to their other basic needs. Providing the benefits of this novel combination of communication capabilities required substantial software innovation to overcome obstacles that have stood in the way of achieving these improvements until the insights leading to the present invention, even despite the existence of automated dispensing machines, which hereto fore had been limited to basic, stand alone operation requiring significant human intervention to ensure accurate and timely administration of medications, without the benefits of linking their operation to the other functional entities involved in the dispensing of prescription drugs under machine control.

FIG. 8 illustrates an automated prescription dispensing machine and an associated work station according to the present invention for use with the embodiments of FIGS. 2 through 7. Depicted in the drawing are an automated prescription drug dispensing machine 100 and a kiosk 102 coupled together by a communications bus 190 for conveying both control and data between these units. The kiosk 102, in one form as a cabinet but which could as easily be a desk or table, supports a workstation or PC 160 having a display and other well-known attributes of a PC. Connected to the workstation may be a scanning device 162 such as a bar code scanner as shown in the present illustrative embodiment. The scanning device 162 may be used for scanning encoded indicia printed on a canister label or a medication package label or any other identification data printed in documents, personnel ID cards or badges, for example. Other devices having the ability to scan and read an image (visual or audible) or data, including similar functional devices not yet invented, may be used. The kiosk 102 may further support a server 104 or a printer 164, or other apparatus including software applications that may be added or used to upgrade the dispensing capability of the system. One item that may be included in the arrangement of equipment supported in the kiosk 102 is an auxiliary database 116 as shown in FIG. 6 herein above. An auxiliary database 116 may be used to store data particular to one automated dispensing machine or one group of machines among a plurality of dispensing machine groups in a health care facility, in order to realize operating efficiencies or other operational improvements to the system.

Continuing with FIG. 8, an automated prescription drug dispensing machine (ADM) 100 is illustrated in simplified form with its cabinet door opened to reveal banks 170 of dispensing canisters 172 therein. The canisters may be as described in FIGS. 3 and 4. After canisters 172 are loaded with the medication and are ready for dispensing, they may be plugged into one of the canister banks 170 in any open position. Mechanisms (not shown) within the ADM 100 cabinet provide for dispensing a unit of medication into a package prior to ejection from the machine. An ejection port 178 is provided in a convenient location on the dispensing machine 100. The ejection port 178 includes an opening from which a strip of medication packages 180 is fed under machine control. The strips of transparent packages may include a printed label 182 and may be separated by cross-wise perforations 184 to enable separating the packages prior to administration to a resident.

FIG. 9A illustrates a first portion of a flow chart 200 for the basic operation to provide for ordering and dispensing a prescription drug in a long term care facility by the system whose architecture is illustrated in FIG. 7. The steps will include references to the unit of the system, as shown in FIGS. 6 and 7, that is active or where the step is carried out. The process begins at step 202 in which, both monthly and for any new resident being admitted, the resident is entered

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into the electronic health record (EHR) 126 and information regarding Medicare part D or Medicaid is verified and updated as necessary. This step may be carried out in the long term care (LTC) facility 94. In the following step 204, any new order for a prescription medication is entered at the prescription order entry (POE) 120 unit, which may be located in the pharmacy 92, the LTC 94, or the office (not shown) of a health care provider. This step includes verifying third party payer (insurance or other health care benefit) coverage, checking for drug allergies of the resident and possible drug interactions with other prescriptions the resident requires, and verifying that the new prescription is approved for dispensing at the order dispensing system (ODS) 98 on site at the LTC resident's location. The flow then advances to step 206 to transmit the prescription order to the pharmacy 92 and to enter the information into the EHR 126 for the resident at the LTC 94. The EHR 126 may be located typically in the main database 114 at the hosting center 84 or the auxiliary database 116 associated with the order dispensing system 98 at the LTC 94.

Continuing with FIG. 9A at step 208 the pharmacy 92 receives and reviews the order, then initiates formulary management processing to determine whether the prescription requires a brand name (coded "B") or branded generic (coded "BG") medication or a generic product (coded "G") may be substituted for the prescribed drug. Subsequently the order is submitted to the order dispensing system (ODS) 98 through the order management system (OMS) 90 to process the filling of packets preparatory to dispensing as will be described in FIG. 11 to follow. After step 208, the process advances to step 210 in which the OMS 90 notifies the prescription order entry (POE) 120 when the order has been approved by a pharmacist. Then, the order data is stored in the main database 114, and the OMS 90 determines the destination LTC 94 and the order dispensing system 102 to receive the order, and the order is queued for dispensing at the designated ODS 98 in the LTC 94. At the ODS 98 in step 212 the order proceeds to step 216 for delivery to the ODS 98 and installation into the automated dispensing machine 100 for dispensing and administering at the next scheduled time. The ODS 98 stores all of the identification and script data at the kiosk 102, then notifies the POE 120 that the order is queued for dispensing. Step 216 just described proceeds to step 222 along the path "A" in FIG. 9A, to FIG. 9B to be described next.

FIG. 9B illustrates a second portion of the flow chart for the basic operation of the system of FIGS. 6 and 7. Dispensing of medications occurs according to a predetermined schedule within the daily routine of the long term care or other health care facility. In the illustrated example the operation of the dispensing process may be initiated and controlled from the workstation 104 at the kiosk 102 associated with the dispensing machine 100 that will be utilized. The dispensing and administering of the medications may generally be carried out by a medical aide or assistant under the supervision of the director of nursing 96 or other registered nurse at the LTC 94. In step 222 the system awaits entry of a request from the ODS 98, sent by the electronic Medical Administration Record or eMAR 104 that the schedule calls for dispensing and administering of medications at a scheduled time. Accordingly, in step 224 the medical aide or assistant will operate the automated dispensing machine (ADM) 100 to package and dispense one or more packages of medications to be administered at a scheduled time as described further in FIG. 12. Typically, the ADM 100 will package a plurality of doses into a strip of separate packages separated by perforations. The doses in

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each successive package may then be administered to one resident after another who reside in close proximity, for example. Each administering of a dose is then recorded in the eMAR **104** by the medical aide or assistant before proceeding with the next dose. This action includes, step **226**, notifying the OMS **90** that the administering of the prescribed dose has been completed and the record in the database **116** may be updated as depicted in step **228**. At the conclusion of this process the OMS **90** notifies the pharmacy information system (PhIS) **124** in step **230** that the billing process may be carried out, after which the process ends in step **232**.

FIG. **10** illustrates a flow chart of one embodiment of a process **280** for formulary substitution according to the present invention. It is described elsewhere that configuring the canisters **50** at the pharmacy **92** and utilizing a compact set of ID data in the canister memory chip **60** provide very substantial improvements in operating and processing speed, efficiency, and error reduction in the prescribing and dispensing of prescription medications in resident health care facilities. One of the exemplary enhancements is the ability to manipulate formularies and reconfigure the canisters accordingly, a capability that does not exist when the canisters are configured by the canister manufacturer. All that the manufacturer can do is manufacture canisters for every type of medication, including the wide variety of formularies that are available. In the present invention, in which the canisters can be configured by the pharmacy, the canisters can be configured on the spot for a particular formulary that is suitable to be dispensed to a resident, even when the pharmacy must respond in real time circumstances. FIG. **10** illustrates a process by which this may be accomplished in the system of the present invention.

The process for FIG. **10** begins at step **282** after advancing from step **208** (see FIG. **9A**) to perform the sub-process of formulary substitution, a subset of formulary management illustrated herein in the exemplary system. Recall that prescription medications are classified as branded, branded generic, or generic, depending on what source of manufacture may be accepted for administering to the resident. Thus, the process opens in step **282** by determining the formulary requirement of the "script" and responding to the query "Is order NDC available in the ADM inventory?" In other words, is this NDC stocked in the ADM **100**? The script is a statement specifying the requisite formulary, the amount of the dose, how often and when the dose is to be administered. If the response in step **284** is YES, the script permits a substitution of "B," "BG," or "G," then a list of the medications available having the same status or classification will be displayed on the display of the workstation **160** at the kiosk **102** to enable a selection to be made by the nurse in step **286** and entered into the record, followed by step **288** to dispense the order. The ODS receiving the order advances to step **290** to return to step **210** in FIG. **9A**. Returning to step **282**, if the NDC ordered is not available in the ADM **100** inventory, the flow goes to step **292** to look up the medication in a pharmacy substitution formulary and determine the appropriate grouping, B, BG, or G. Then, in the query step **294**, if the original NDC is within the same grouping as the NDC in the ADM **100** inventory (a YES response), the flow advances to step **284** and proceeds as described previously. If, in step **294**, the NDCs do not match because the formulary medication is not available in the ADM **100**, then the process goes to step **296** to exit, and the process ends.

The automated dispensing machine (ADM) **100** as described previously is an automated machine for packaging

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and dispensing prescription medications under computer control. Medications for a supply of doses to be stored in the machine are deposited in individual canisters **50** as described in FIGS. **3** and **4** herein. A typical ADM **100** may contain approximately 200 canisters, for example. The actual number depends on the volume of the main body of the canisters, which may vary depending on the size and form of the individual doses to be contained in the canister **50**. The supply quantity of medications for an individual canister **50** may be determined by the needs for some interval of time, e.g., one month, or by the needs for one or more residents in the LTC **94**. The processes for configuring, filling, and changing the contents of the canisters is illustrated in FIGS. **11A** through **11F**.

FIG. **11A** illustrates a flow chart **240** of initial steps to begin a medication canister configuration and fill process according to one embodiment of the present invention. The flow begins at user log in at step **242** and the display of a menu of actions that involve the configuration, filling, or modifying the fill or information affecting the canisters in step **244**. In step **246** the user may select the desired action from the menu, which lists five different actions, respectively: create canister build; configure canister; fill canister; add medication to the canister; and edit the medication information of the canister. These actions are respectively designated as steps **250**, **252**, **254**, **256**, and **258**. Selecting any one of these options directs the user along respective paths B through F of FIG. **11A** to corresponding processes illustrated in FIGS. **11B** through **11F** to be described herein below.

FIG. **11B** illustrates a flow chart of a process to create a canister build according to the present invention. A canister build is the process for configuring a canister for a specified medication. As described in FIGS. **3** and **4**, the configuration of a canister may be adapted to the quantity, size, and shape of the individual doses of the medication to be stored or contained therein. The flow enters step **260** from path B, displays build parts on a displayed page, and enables selection of build parts appropriate to the prescription order being fulfilled. Following the selection of parts, a query step **262** as to whether the needed information is complete provides a YES alternative to step **264** to display a medication association input and select the medication that is to be deposited into the canister. Step **262** also provides a NO alternative to return to step **260** if the information is not complete. Following the selection of medication(s) for deposit into the canister in step **264**, the process advances to step **266** to log the information about the canister and provide for the user's manual input of information about the canister build. In the following step **268**, the canister build manager is notified of the new canister configuration in step **268**. Next, if there are any additional canister builds to process, as queried in step **270**, the flow returns to step **260**; if not, the flow ends at step **272**.

FIG. **11C** illustrates a flow chart of a process to configure a canister according to the present invention. The flow enters step **310** from path C, displays a scan/manual input page to enable the user to enter or scan the NDC for the prescription medication, and look up the NDC information in the main database **114** or, in some cases, the auxiliary database **116** as shown in FIG. **7**. If the information sought is in the database (YES, in step **312**), it is displayed in step **314**, followed by a check query "Data Correct?" in step **316**, and if YES to that query, advances to step **318** to display the canister information. If the canister is to be filled with the designated quantity, the YES response to the query in step **320** is entered and the process exits to path D to be described in

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FIG. 11D. Returning to step 312, if the response to the query “Is the NDC data in the database?” is NO, the process also exits to path D. Further, in step 316, if the response to the query “Is the data correct?” is NO, then the flow exits to path F to be described in FIG. 11F. Returning now to step 320, if the response to the query “Fill canister?” is NO, then the flow proceeds to step 322, a query as to whether any additional canisters are to be filled. If YES, the flow returns to step 310; if NO, the process ends at step 324.

FIG. 11D illustrates a flow chart of a process to fill a canister according to the present invention. Canisters are generally filled at a pharmacy 92 that stocks the medications and is under the supervision of a registered pharmacist as required by law. The present invention includes the ability to configure the canisters 50 at the pharmacy 92, at the time the canister is to be filled. In conventional systems the configuration of the canisters is accomplished to order by the canister manufacturer before shipment to the pharmacy operator. The pharmacy must stock all of the different configurations of the canisters that it needs to fulfill orders for the prescription medications from the long term or other health care facilities that it services. Because of the very wide variety of formularies, forms, shapes and sizes of the medications, the pharmacy must maintain an extensive and expensive inventory of configured canisters. The present invention overcomes this problem by stocking the interchangeable component parts of the canisters—a much smaller number of SKUs (stock keeping units). The efficiencies gained from this technique are substantial because each canister can be configured as needed, thus avoiding the inventory of configured canisters not being used. Further, canisters can be readily modified from one configuration to another with minimal effort, thus minimizing the likelihood that a dispenser is out of service because a particular canister is not available. Moreover, quality control, inspection, repair and maintenance operations are significantly more efficient and less likely to cause delays because all of these operations can take place at the pharmacy where the canisters are configured or filled.

Continuing with FIG. 11D, the flow begins on path D to step 330 to obtain and fill a canister, the user logs in to the system, selects “canister fill” on the workstation screen, and places the canister 50 on a chip writer (not shown). As shown in FIGS. 3 and 4, the canister memory chip extends slightly below the underside of the canister 50, enabling it to be inserted into a receptacle on the chip writer for writing the data thereto. In the next step, the system queries the user to ascertain whether the user is a registered pharmacist. If YES, the process advances to step 334 to display the scan/manual input page, wherein the user may select which of these modes—scan or input manually—will be used. If scan is selected a scanning device such as (but not limited to) a bar code scanner may be used. In the next step 336 the system determines whether the user has a bottle of the medication to be utilized, and if YES, the flow goes on to step 342. If, however, the user does not have a bottle, the flow advances to step 338 to scan or input the NDC, the Nation Drug Code, to the system, followed by the step to look up the NDC in the database 114. Returning to step 332, if the user is not a pharmacist, the system displays the scan input page on the screen of the workstation in step 340 before advancing to step 342 to scan the medication for its NDC.

Following step 344, the system determines whether the NDC is found in the database 116? If NO, the process exits along path E to FIG. 11E. If YES, the process steps to step 348 to determine whether the canister is OK for NDC, that is, whether its configuration matches or is compatible with

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the drug attribute information for the particular pill, capsule, tablet, etc. If YES, then the flow advances to step 350 to display the medication information to be written into the database and the build ID and Canister/Fill Ids written onto the canister memory chip. If this information is NOT correct, then the process at step 352 exits along path F to FIG. 11F to edit the information. Returning to step 352 if the information is correct, the process proceeds to step 370 in which the Lot Number, expiration information, and the registered pharmacist information is input to the system, followed in step 372 by writing the canister information and printing the canister label. Next, the flow advances to determine whether additional canisters need to be filled in step 374 and if the response is NO, the process ends at step 366. If the response is YES, the process prepares to place another canister on the canister chip writer in step 376 and then flows back to step 332 to re-enter the fill processing procedure.

Returning to step 348, if the canister is determined to NOT be suitable for the NDC that the user wishes to deposit in the canister, then the flow proceeds to step 360 wherein the system displays the error. The user has an option to return to the “Canister Lookup” step 318 in FIG. 11C or select a manual override as in step 362. If the “manual override” is selected at 362, the system will send a notification of the error and the manual override to the canister build center in step 364 before ending the routine in step 366.

FIG. 11E illustrates a flow chart of a process to add a medication canister according to the present invention. Entry to the process proceeded along path E following selection in step 246 of the action to “Add Medication” at step 256. At step 390 the display screen for adding a medication appears on the workstation display, enabling the user to input the medication data in step 392 in the respective fields for the medication to be added. This information may include for example a mnemonic, the NDC, the name of the drug, its strength, unit, manufacturer, color, shape, marking, and schedule for administering. The added information is logged and the user input manually in step 394, followed by step 396 to notify the canister build manager regarding the added medication. Next, in step 398 a query occurs whether there are any other additional medications to input. If YES, the flow returns to step 390 and the process repeats; if NO, the process ends.

FIG. 11F illustrates a flow chart of a process to edit a medication canister according to the present invention. It is similar to the process for adding a medication and proceeds as follows. Entry to the process proceeded along path F following selection in step 246 of the action to “Edit Medication” at step 258. At step 410 the display screen for editing a medication appears on the workstation display, enabling the user to input the medication data in step 412 in the respective fields for the medication to be added. This information may include for example a mnemonic, the NDC, the name of the drug, its strength, unit, manufacturer, color, shape, marking, and schedule for administering. The edited information is logged and the user input manually in step 394, followed by a query step 416 whether the size or shape of the medication unit is to be edited. If YES, the flow proceeds to step 418 to notify the canister build center regarding the edited canister; if NO, the flow advances ahead of step 418 to another query step 420 whether there are any additional medications. If YES, the process returns to the step 410 to repeat the edit routine. If NO, the process ends.

FIG. 12 illustrates a flow chart of a process 430 to prepare medications for administering them to residents of a long term care facility according to the present invention. The

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first step 432 occurs as the nurse logs on to the order dispensing system (ODS) 98 at the kiosk 104 (See FIG. 8) and selects Order, then the Dispense option displayed on the workstation screen of the kiosk 104 to display active patient or resident orders. Then in step 434 the nurse selects an order to be dispensed, approves a prompt to confirm by selecting the Dispense option. Next, in step 436 the ODS 98 processes the request, sorts the medications, sends dispensing information to the packager in the automated dispensing machine (ADM) 100. Upon receiving the dispensing information the packager, in step 438 packages selected medications in a series of individual packets delineated by cross-ways perforations to facilitate ease of tear-off separation, and outputs the packets from the port 178 on the ADM 100. The nurse or assistant may then remove the strip of packets from the ADM 100 in step 440 and place the strip 180 on a medication cart (not shown) for delivery to residents scheduled to receive the medications in the strip 180 of packets. In the last step 442 the medications are administered to residents according to the eMAR as will be depicted in FIG. 13 to follow.

FIG. 13 illustrates a flow chart of a process 450 for administering medications to residents of a long term care facility according to the present invention. The process begins at step 452 in which an assistant selects a resident on a display of a workstation to display the medication(s) to be administered. Next, in step 454 the assistant scans the medication packet to verify the packet contents are appropriate according to the database record as shown on the display. Then the flow proceeds to step 456 to administer the medication, select the action performed to confirm the administering of the medication. The process continues at step 458 with an opportunity to record additional information as appropriate, and concludes with a query step 460 whether the medication last administered is the last resident in the scheduled sequence? If the result is YES, the process ends; if the result is NO, the process returns via step 464 to the beginning of the process 450 at step 452.

FIG. 14 illustrates a simplified flow diagram 470 for reading medication data from a canister memory 60 according to the present invention to highlight the reduction in processing and dispensing time, and errors in dispensing afforded by the modified memory contents. Beginning at step 472, the canister 50 is placed on a base in the ADM 98 that contains a keyed receptacle (not shown) that matches the guide block 80 (See FIG. 4) on the underside of the canister 50. In step 474 the Build ID and Canister/Fill ID data (18 bytes total) are read. At this point the process follows two parallel paths simultaneously. Before proceeding, some terms need to be defined. A "canister movement log" contains data about the canister 50 that tracks the historical position of the canister 50 in the Automated Dispensing Machine (ADM) 100. "Canister movement data" represents the contents of the database where the movement data is stored. "Canister/fill data" represents the contents of the database where the canister/fill data is stored. "Fill data" is the data associated with the Fill ID of the canister.

Continuing with FIG. 14, advancing from step 474 proceeds along two parallel paths. In a first path, the flow advances to step 478 to store the canister ID and ADM 100 position data (the location in the ADM of the canister) in the movement log database. Step 480 represents the stored movement log data in the auxiliary database, which may reside in the server 104 or the workstation 160 at the kiosk 102, and the process ends at step 488. In the second path following step 474, the canister/fill ID is looked up at step

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482 in the kiosk database, which is synonymous with the auxiliary database 116 in FIG. 6. Then the canister position data representing the canister's position in the ADM 100 is updated in the kiosk or auxiliary database 116. Step 486 represents the updated canister/fill data in the auxiliary database 116, and the process ends at step 488.

While the invention has been shown in only several of its forms, it is not thus limited but is susceptible to various changes and modifications without departing from the spirit thereof.

What is claimed is:

1. A configurable canister of a medication dispensing system for use in an automated dispensing machine (ADM), the configurable canister comprising:

a body for housing and releasing medication comprising a plurality of individually provided doses of medication; and

a memory comprising a unique identifier corresponding to canister information that correlates to (i) a structural configuration of the configurable canister and (ii) a content of the configurable canister that includes a type of the medication in the configurable canister, wherein the memory is configured to provide the unique identifier to a computer that includes a data storage configured to identify, based on the unique identifier the type of medication contained in the body of the configurable canister and a quantity of the medication remaining in the configurable canister,

wherein the configurable canister is configured to be disposed in the ADM during operation and to release individually at least a portion of the plurality of individually provided doses of the medication contained therein, the releasing in response to the computer providing one or more instructions to the ADM operating to release the type of medication associated with the unique identifier.

2. The configurable canister of claim 1, wherein the data storage is at least one of a remote database or a local memory, and wherein the remote database is remote from the medication dispensing system, and wherein the local memory is disposed within the ADM and remote from the configurable canister.

3. The configurable canister of claim 1, wherein the memory is configured to be operable with circuitry of the ADM to operate with the ADM and provide to the ADM the unique identifier for identifying the configurable canister.

4. The configurable canister of claim 1, further comprising a first label thereon, the first label containing specifications of the medication contained therein, and wherein additional canister information is provided on at least one of the first label and a second label.

5. The configurable canister of claim 1, wherein the body comprises at least one interchangeable part configured to be exchanged for a different interchangeable part to accommodate a different type of medication being stored in the body.

6. The configurable canister of claim 5, wherein the data storage comprises a record associating (i) the canister information for identifying the canister with (ii) the type of medication contained in the body, and wherein the type of medication stored in the data storage is configured to be modified in the data storage to identify the different type of medication being stored in the body.

7. The configurable canister of claim 1, wherein the body comprises at least one adjustable part configured to be adjusted to accommodate a different type of medication being stored in the body.

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8. The configurable canister of claim 1, wherein the plurality of individually provided medication is in a form of a pill, tablet, caplet, or capsule, and wherein the type of medication is based in part on dimensional attributes of the medication.

9. A method for providing a configurable canister with a memory thereon, the method comprising:

storing a first dataset in a data storage (i) separate from the memory and (ii) not in the memory, the first dataset comprising medication information identifying at least a type of medication to be stored in the configurable canister for dispensing by an automated dispensing machine (ADM) in communication with the data storage;

storing a second dataset in the data storage (i) separate from the memory and (ii) not in the memory, the second dataset comprising a code corresponding to a structural configuration of the configurable canister, the structural configuration of the configurable canister corresponding to attributes of the type of medication to be stored in the configurable canister for dispensing by the ADM; configuring the configurable canister in accordance with the structural configuration, wherein at least the type of medication in the first dataset is associated with the structural configuration in the second dataset and the association is stored in the data storage (i) separate from the memory and (ii) not in the memory;

identifying, from the memory of the configurable canister, data representing a canister identifier stored thereon; storing, in the data storage, the canister identifier that was read from the memory of the configurable canister; associating, in the data storage, (a) the canister identifier, (b) the structural configuration, and (c) the type of medication;

storing the association of (a) the canister identifier, (b) the structural configuration, and (c) the type of medication in the data storage; and

providing the canister identifier from the memory of the configurable canister, wherein the canister identifier is configured to be read from the memory of the configurable canister by the ADM, and wherein the ADM is configured to retrieve the type of medication in response to the identifying the canister identifier.

10. The method of claim 9, wherein the data storage is at least one of a remote database or a local memory, and wherein the remote database is remote from the ADM, and wherein the local memory is disposed within the ADM and remote from the configurable canister.

11. The method of claim 9, further comprising:

incrementing a counter stored on the memory, a value of the counter corresponding to a number of times the configurable canister has been filled, wherein the code is a four digit code, and wherein configuring the configurable canister in accordance with the structural configuration comprises adjusting the configurable canister.

12. The method of claim 9, wherein the attributes of the type of medication are selected from one or more of a form of a medication, a size of the medication, and a shape of the medication, and wherein the configurable canister is configured to be filled with the medication having the attributes associated with the structural configuration of the configurable canister.

13. The method of claim 10, wherein the configurable canister is reconfigurable to a different structural configuration associated with a different type of medication having different attributes than the type of medication, and wherein

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the method further comprises replacing the structural configuration in the second dataset with the different structural configuration.

14. A configurable canister management system comprising:

a configurable canister comprising a memory, wherein the memory is configured to communicate with a data storage and stores a canister identifier comprising (i) first data identifying the configurable canister, and (ii) second data corresponding to a number of times that the configurable canister has been attached to a filling station and filled with one or more medications, wherein the filling station is communicably coupled with the data storage, and wherein the data storage stores third data identifying a structural configuration of the configurable canister and associates the structural configuration with (i) the one or more medications and (ii) the canister identifier.

15. The configurable canister management system of claim 14, wherein the memory comprises a memory control circuit configured to provide the canister identifier to an automated dispensing machine (ADM) to which the configurable canister is coupled following disconnection from the filling station.

16. The configurable canister management system of claim 15, wherein the data storage is at least one of a remote database or a local memory, and wherein the remote database is remote from the ADM, and wherein the local memory is disposed within the ADM and remote from the configurable canister.

17. The configurable canister management system of claim 14, wherein the configurable canister further comprises a first label identifying the one or more medications and the canister identifier, and wherein the configurable canister further comprises a second label identifying the structural configuration associated with the configurable canister.

18. A method of filling and deploying a configurable canister with a medication, the method comprising:

connecting the configurable canister to a filling station, wherein the configurable canister comprises:

a main body for housing and releasing a plurality of medication comprising a plurality of individually provided medication;

a memory coupled to the filling station and containing (i) canister information for identifying the configurable canister, and (ii) a fill for tracking a number of times the configurable canister is filled with the medication;

reading, by the filling station, the canister information;

writing, to a data storage, an update associated with the canister information, the update comprising data corresponding to a medication identifying code of the medication for filling into the configurable canister;

disconnecting the configurable canister from the filling station and connecting the configurable canister to an automated dispensing machine (ADM);

reading the canister information from the configurable canister; and

retrieving, from the data storage, additional data associated with the canister information, the additional data comprising the medication identifying code of the medication in the configurable canister.

19. A method of configuring configurable canisters to dispense a medication, the method comprising:

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providing, by a filling station to a data storage, a medication identifying code of the medication for filling into a configurable canister;

retrieving, from the data storage, a canister configuration corresponding to the configurable canister indicated to be compatible with attributes of the medication associated with the medication identifying code;

retrieving a set of canister identifiers, each canister identifier of the set of canister identifiers is associated with an unavailable configurable canister that (i) is presently unavailable and (ii) has another canister configuration that is available for use with the filling station;

in response to retrieving no available configurable canister associated with the set of canister identifiers, retrieving a part list associated with at least one medication identifying code corresponding to at least the canister configuration compatible with the attributes of the medication;

in response to retrieving the part list, providing data corresponding to human-readable instructions for changing at least one part of at least one unavailable configurable canister associated with the set of canister identifiers, wherein providing the data causes the at least one unavailable configurable canister to have a new canister configuration corresponding to compatibility with the attributes of the medication;

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reading, by the filling station, a canister identifier of the at least one unavailable configurable canister having the new canister configuration; and

storing the canister identifier to additional data associated with the new canister configuration to document that the at least one unavailable configurable canister having the new canister configuration is now compatible with the attributes of the medication.

20. The method of claim **19**, wherein the unavailable configurable canister is at least one of an empty configurable canister or an out-of-service configurable canister.

21. The configurable canister of claim **1**, wherein the ADM does not write to the memory of the configurable canister.

22. The configurable canister of claim **1**, wherein the unique identifier also includes information that correlates to a number of fills of the canister.

23. The configurable canister of claim **1**, wherein the medication is all the same medication and the individually provided doses of the medication are all the same.

24. The configurable canister of claim **1**, wherein the configurable canister is configured by changing the structural configuration to dispense a plurality of different medications.

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