



(22) Date de dépôt/Filing Date: 1996/04/29  
 (41) Mise à la disp. pub./Open to Public Insp.: 1996/11/02  
 (45) Date de délivrance/Issue Date: 2007/08/07  
 (30) Priorité/Priority: 1995/05/01 (US08/431,632)

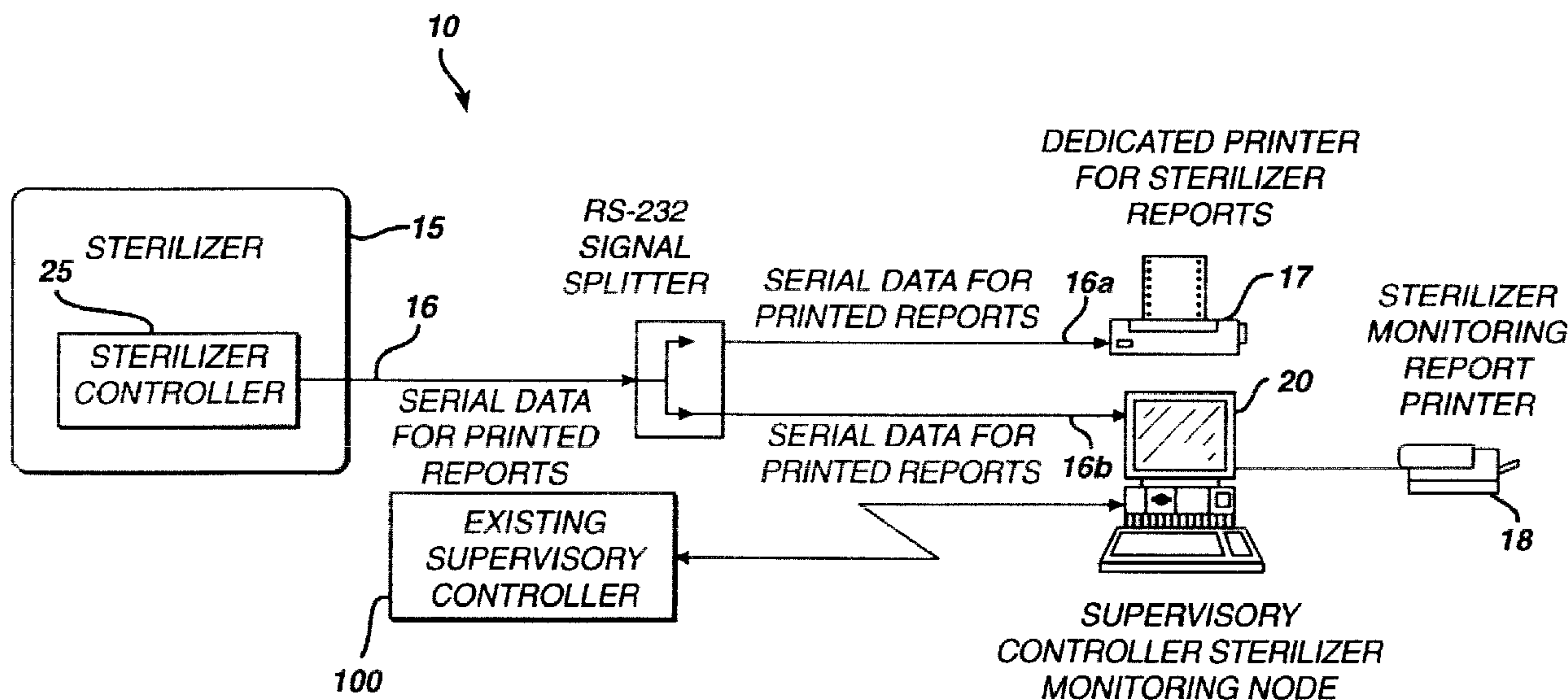
(51) Cl.Int./Int.Cl. *G07C 3/14* (2006.01),  
*A61L 12/00* (2006.01), *A61L 2/24* (2006.01),  
*A61L 2/28* (2006.01)

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(54) Titre : SYSTEME INFORMATIQUE DE CORRELATION POUR LE CONTROLE DE LA QUALITE  
 (54) Title: COMPUTER SYSTEM FOR QUALITY CONTROL CORRELATIONS



(57) Abrégé/Abstract:

A sterilizer data processing system for an automated contact lens manufacturing line that manufactures a plurality of contact lenses defining a lens lot, the manufacturing line including an automated sterilization station for sterilizing the lens lot after their manufacture, the automated sterilization station including a sterilizer process controller for controlling one or more phases of a sterilization process and periodically generating sterilization process data during each sterilization phase includes a device for receiving the sterilization process data from the sterilizer process controller and a device for automatically parsing the sterilization data into text information and sterilizer parameter information and further processing the text information and sterilizer parameter information to automatically generate a sterilization run report associated with a lot number for the sterilized lens lot.

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ABSTRACT OF THE DISCLOSURE

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A sterilizer data processing system for an automated contact lens manufacturing line that manufactures a plurality of contact lenses defining a lens lot, the manufacturing line including an automated sterilization station for sterilizing the lens lot after their manufacture, the automated sterilization station including a sterilizer process controller for controlling one or more phases of a sterilization process and periodically generating sterilization process data during each sterilization phase includes a device for receiving the sterilization process data from the sterilizer process controller and a device for automatically parsing the sterilization data into text information and sterilizer parameter information and further processing the text information and sterilizer parameter information to automatically generate a sterilization run report associated with a lot number for the sterilized lens lot.

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1. Field of the Invention

This invention relates generally to a computer system for a manufacturing facility for the production of ophthalmic contact lenses, and, in particular to a  
5 supervisory system for monitoring the production line processes used in the manufacture of contact lenses in a contact lens fabrication facility, specifically, with the goal of investigating and optimizing the process of contact lens sterilization.

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2. Description of the Prior Art

The direct molding of hydrogel contact lenses is disclosed in U.S. Patent 4,495,313 to Larsen, U.S. Patent 4,680,336 to Larsen et al., U.S. Patent 4,565,348 to Larsen, and U.S. Patent 4,640,489 to Larsen et al.  
15 Essentially, these references disclose an automated contact lens production process wherein each lens is formed by sandwiching a monomer between back curve (upper) and front curve (lower) mold sections. The monomer is polymerized, thus forming

a lens, which is then removed from the mold sections and  
1 further treated and packaged for consumer use.

The manufacturing of contact lenses requires  
tightly controlled conditions and processes, many of which  
are monitored by computers and other control devices.  
5 Much information, in the form of process conditions and  
control data, for e.g., that occur during contact lens  
manufacturing, may be gathered for quality control and  
regulatory approval purposes. However, this entails the  
acquisition of a tremendous amount of data for each  
10 contact lens that is produced, and, additionally, requires  
a means for processing the data acquired in a way that is  
suitable for use by operators, engineers, and supervisors,  
etc., so that they may properly perform their functions.

There is therefore the need to provide a quality  
15 control system that can automatically acquire process  
control data from a plurality of manufacturing process  
controllers that control various aspects of contact lens  
production at process stations in a contact lens  
manufacturing facility, and, that can automatically  
20 process the data for real-time display and archiving  
purposes. More particularly, there is a need for a  
quality control system that can automatically acquire data  
generated from a sterilization controller that controls a  
sterilization process performed to contact lenses that are  
25 individually packaged but not cartoned, and that is  
performed prior to their cartoning.

It would additionally be highly desirable to  
provide a quality control system that can automatically  
gather sterilization process control data for contact  
30 lenses for subsequent generation of sterilizer cycle  
condition records that includes: sterilization run

success/failure indication, lot number, and sterilization  
1 run number from the sterilizer controller. These files  
may be stored in an off-line database storage area and be  
retrieved to analyze the trend of sterilizer performance  
over a long period of time. Furthermore, in accordance  
5 with the inventive processes described herein, these files  
may be processed to automatically generate reports that  
are suitable for compliance with Federal Food and Drug  
Administration ("FDA") record-keeping requirements, and  
are also useful for re-certifying the sterilizer.

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#### SUMMARY OF THE INVENTION

An object of the instant invention is to provide  
a quality control system for a contact lens manufacturing  
15 facility that automatically acquires process control data  
from a plurality of manufacturing process controllers that  
control contact lens production, and, that can  
automatically process the data for real-time display and  
off-line analysis purposes.

20 Another object of the invention is to provide a  
quality control system for a contact lens manufacturing  
facility that implements a sterilization process for  
sterilizing individual contact lens packages after their  
primary packaging in blister packages and prior to their  
25 cartooning.

Still another object of the invention is to  
provide a quality control system for a contact lens  
manufacturing facility that includes a sterilization  
apparatus controlled by a sterilization controller for  
30 sterilizing individual contact lens packages prior to  
their cartooning. Additionally, another object of the

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invention is to provide a quality control system that  
1 gathers sterilization process control data from the  
sterilization controller and subsequently generates  
sterilizer cycle condition records that includes:  
sterilization run success/failure indication, lot number,  
5 and sterilization run number from the sterilizer  
controller.

Yet still another object of the invention is to  
provide a quality control system for a contact lens  
manufacturing facility that includes an apparatus for  
10 secondary packaging of sterilized blister packages  
containing individual contact lenses in cartons.

A further object of the invention is to provide  
a quality control system that incorporates means for  
automatically printing and correlating labelling  
15 information including lot number identification for all  
contact lens packages produced.

The above objects are achieved in a quality  
control system for an automated production line producing  
contact lenses, the production line having a plurality of  
20 contact lens process stations, including an automated  
sterilization station for sterilizing a plurality of  
contact lenses after their manufacture, and a packaging  
station for packaging said lenses after sterilization,  
wherein the system comprises:

25 (a) a first means for receiving contact lens  
data including an associated lot number and lens power for  
a lens lot prior to their manufacture; the lens lot  
defining at least one batch of contact lenses;

(b) a plurality of process controllers for  
30 controlling one or more process stations, each of the

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controllers regulating a plurality of process control  
1 devices at the process stations;

(c) means for tracking movement of the plurality  
of lenses defined by the lens lot from the plurality of  
processing stations to the automatic sterilization station  
5 and the packaging station;

(d) second means for receiving data representing  
the number of lenses that are input to the sterilization  
station and recording sterilization data for each batch of  
the lens lot together with reason codes for contact lenses  
10 lost at the sterilization station;

(e) means for generating a summary report of the  
total number of lenses input to the sterilization chamber  
for a predetermined lens lot and the actual number of  
lenses sterilized and packaged from the lot, the summary  
15 report including lot number, expiration date, power and  
sterilization data for each batch of contact lenses.

Further benefits and advantages of the invention  
will become apparent from a consideration of the following  
detailed description given with reference to the  
20 accompanying drawings, which specify and show preferred  
embodiments of the invention.

#### BRIEF DESCRIPTION OF THE DRAWINGS

25 The foregoing objects and advantages of a  
quality control system for an automated production line  
for producing contact lenses of the present invention, may  
be more readily understood by one skilled in the art with  
reference being had to the following detailed description  
30 of several preferred embodiments thereof, taken in  
conjunction with the accompanying drawings wherein like

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elements are designated by identical reference numerals 1 throughout the several views, and in which:

Figure 1 is an organizational overview of the sterilization monitoring system of the instant invention.

Figure 2 illustrates the hardware configuration 5 of the existing supervisory control system 100 shown interfaced with the sterilizer monitoring node 20 and sterilizer controller 25 of the instant invention.

Figure 3 illustrates a detailed hardware 10 overview of the supervisory controller's sterilizer monitoring node 20 and data flow therein.

Figure 4 is a state data flow diagram showing the internal states of the sterilizer monitoring node while receiving data from the sterilizer controller 25.

Figures 5(a) and 5(b) illustrate, in detail, the 15 steri comm data acquisition process 50.

Figure 6 illustrates the major functional blocks of the steri server process 50.

Figure 7 illustrates the `characterizeLine` 20 algorithm 300 for determining the nature of the data line sent by the sterilization controller.

Figure 8 illustrates the `processDataLine` algorithm 405 for processing the twelve (12) variables of sterilizer phase data from the data line.

Figure 9 illustrates the `makePhaseFileEntry` 25 algorithm 440 to format the process variable information for entry as a line in the corresponding phase file.

Figures 10(a) and 10(b) illustrate the `evaluateTextline` process 450 for processing textual data from the input data line.

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Figure 11 illustrates the **StartOfRunEvent** procedure 466 invoked when a sterilizer run has started from the beginning.

Figure 12 illustrates the **doEndOfRunEvent** procedure 480 invoked when the end of a sterilizer run has been detected.

Figure 13 illustrates the **processAlarmLine** algorithm 500 for processing alarm data from the input data line.

Figure 14 illustrates the **updateAlarmStatus** algorithm 550 for updating alarm status after every line evaluation.

Figure 15 illustrates the **finishSteriRunReport** procedure 485 to generate the sterilizer run report.

Figure 16 illustrates the **doEndOfRunCleanup** procedure 486 to finish the text line processing and print the steri run report.

Figure 17 illustrates the **addDurations** procedure 630 for adding the phase time durations to the sterilizer run report.

Figure 18 illustrates the **addMinMax** procedure 640 for adding the minimum and maximum phase variable data to the sterilizer run report.

Figure 19 illustrates the **openSteriRunReport** procedure for opening the steri run report file.

Figure 20 illustrates the **closeAndPrintRunReport** procedure for closing the steri run report file.

Figure 21 illustrates in detail the **wakeUpCmdFunc** process 280 invoked by the CELLworks system.

Figure 22 illustrates in detail the **endRunReportForTimeout** process 290 that is called to update the steri run report file.

Figure 23 illustrates in detail the  
1 sterilization run report automatically generated for a  
complete sterilization cycle.

Figure 24 illustrates the algorithm for contact  
lens lot number information entry.

5 Figures 25(a) and 25(b) illustrate the  
respective data structures stored in the statistics server  
for the lot information before primary packaging (Figure  
25(a)) and after primary packaging (Figure 25(b)).

Figure 26 illustrates a data flow diagram for  
10 moving a lot by operator request.

Figure 27 illustrates the moveLot algorithm 680  
for tracking lens lot movement throughout the production  
line.

Figure 28 illustrates the start of the lot  
15 reconciliation process which entails the reporting of the  
quantity of lenses input to the sterilizer.

Figure 29 illustrates the procedure for entering  
the number of lenses removed from the secondary packaging  
process.

20 Figure 30 illustrates the lot close out flow  
diagram.

Figure 31 illustrates a table depicting phase  
file data entries in the four sterilizer phase files.

Figure 32 illustrates the  
25 sterilization/secondary packaging lot reconciliation  
sheet.

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**DESCRIPTION OF THE PREFERRED EMBODIMENT**

Illustrated in Figure 1 is a general schematic diagram illustrating the sterilization monitoring system 10 for passively monitoring the sterilization and secondary packaging of contact lens packages. As will be explained in greater detail below, the sterilization monitoring system 10 of the invention is configured specifically to process sterilizer serial data and generate sterilization run reports.

As shown in Figure 1, the sterilization monitoring system 10 comprises a sterilization chamber 15 having a sterilizer control device 25, which, in the preferred embodiment, is a PLC or dedicated process controller that controls the sterilization process and serially broadcasts the sterilization process data 16 and alarm data (when an alarm condition exists) as formatted ASCII characters to a dedicated printer 17, via data line 16a, as well as an intercepting sterilization monitoring node 20, via data line 16b, that is interfaced with an existing contact lens production line supervisor quality control system 100 ("existing supervisor system"). As will be explained in detail below, the sterilization node 20 will process the ASCII sterilizer data and automatically generates a sterilization run report, a portion of which is shown in Figure 22, and explained in detail below, at a second printer 18. The operational details of the existing contact lens production line supervisor quality control system 100 are disclosed in the above-identified U.S. Patent No. 5,461,570 entitled "Computer Program for Quality Correlations", assigned to

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the same assignee as the instant invention. As described in the above-identified U.S. Patent No. 5,461,570 the existing supervisor system automatically acquires process control data from a plurality of programmable and non-  
5 programmable control devices that control and monitor various manufacturing processes, and, automatically processes the data for real-time display and off-line engineering analysis and quality assurance purposes.

Figure 2 illustrates the hardware configuration of  
10 the existing Supervisor system 100 interfaced with the sterilizer monitoring node 20 and sterilizer controller 25 via ARCNET HUB network devices 99a and 99b that support communication between the sterilizer monitoring node, two operator terminals 29a,29b (preferably  
15 manufactured by Dynaterm), and, the existing supervisor system 100. Preferably, the sterilizer monitoring node 20 includes a 33 MHz Intel '486 computer having the following modules for performing the sterilization monitoring process: sterilization communication module  
20 and process 50, sterilization server module and process 60, a time server 70, and a file server 80. Each of these modules will be described in greater detail hereinbelow.

Briefly, the supervisor system 100 interfaces with and obtains control parameter data from the sterilization  
25 monitoring node, and, at least seven programmable logic controllers that control various contact lens manufacturing processes.

These manufacturing processes include: transferring of injection molded front curve lens molds to carrier

pallets as controlled by PLC 31; transferring of injection molded back curve lens molds to carrier pallets as controlled by PLC 32; monomer filling and contact lens mold assembly operations as controlled by PLC 33; the  
5 precure, UV curing, and lens de-mold operations as controlled by PLC 34; transfer of the front curve mold halves containing molded contact lenses to a hydration chamber for contact lens hydration as controlled by PLC  
10 35; post hydration operations including the generation of contact lens inspection data consisting of pass/fail results as determined by an automatic vision system incorporated in an automatic lens inspection station as controlled by PLC 36; and, the primary contact lens packaging and lens package consolidation aspect of the  
15 lens packaging processes including such processes as solution exchange, saline fill, package foil heat seal, etc., which occur about a rotary index (packaging) dial (not shown) as controlled by PLC 37. An eighth PLC, may be provided for controlling various aspects of the  
20 secondary packaging including transfer of packages from the sterilization chamber to a secondary packaging area where the blister packs are labelled and sealed in secondary packaging cartons as described in U.S. Patent No. 5,488,815 entitled "Apparatus and Method for  
25 Sterilization and Secondary Packaging" assigned to the same assignee as the instant invention. Secondary packaging briefly includes the steps of printing/applying bar-coded lot-number on the carton, printing the power and  
and

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expiration date on the carton, inserting the blister  
1 packages into the carton and gluing carton flaps closed,  
verification of the lot number, verification of the power  
and expiration date, weighing the carton, loading and  
closing the case and applying the case label, and, loading  
5 and completing the pallet and applying the pallet label.

In the preferred embodiments, each PLC 31-38 is  
a TI system 545 (Texas Instruments) and may include a TI  
386/ATM coprocessor module for communicating with the  
respective PLC across the backplane or by serial link (not  
10 shown). It is understood that each PLC has its own memory  
and addressing capabilities for storing and updating  
blocks of data.

As shown in Figure 2, other programmable device  
controllers, for example, those manufactured by Yushin  
15 Corp., are provided in a contact lens production line for  
controlling, respectively, the front curve mold machine  
39a which produces the front curve lens molds at a rate of  
eight every six seconds, back curve mold machine 39b which  
produces the back curve lens molds, the primary packaging  
20 machine 39c for producing the contact lens packages in  
which the manufactured contact lens is inspected and  
packaged. Another device controller 39d controls a vision  
system (not shown) that automatically inspects the contact  
lenses prior to their primary packaging.

25 Furthermore, in Figure 2, the existing  
supervisory control system (control system) 100 includes  
five (5) types of processing nodes: a Data Acquisition  
Node 205 for communicating with each of the eight (8)  
programmable logic controllers (PLCs), discussed above, by  
30 means of communication lines 41 and TIWAY adapter card 42,  
and also, for communicating with the device controllers of

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the three mold machines, and the vision inspection machine  
1 by means of an 8-channel serial card 44, shown connecting  
the machines by dedicated asynchronous serial lines  
43a,b,c, and d; a Relational Database Node 210 which runs  
relational database software 212 and includes at least  
5 three 200 megabyte hard disks provide for off-line data  
storage consisting of production records and long-term  
data histories; an Analysis and Routing Node 220 that  
contains most of the software that is used to initiate  
data gathering and processing of raw data from the eight  
10 PLCs, and, that maintains "real-time databases". The  
analysis and routing node 220 comprises modules such as:  
the Statistics Server 225 that stores data within logical  
user defined groups or datasets, is capable of generating  
statistics and (optional) alarms on data sets, and, that  
15 support statistical control charts and other displays; a  
poller 226 which coordinates the acquisition of all data  
from the PLCs, Mold Machines, and the Vision Inspection  
Machine; a C-language Control Server 228 which is a  
companion module to the Statistics server and directs the  
20 Statistics Server to perform statistical functions needed  
to support active displays; and, an alarm control server  
229 which handles and maintains workcell alarms, warnings,  
and exceptions that are activated according to defined  
conditions.

25 The supervisor system 100 further includes four  
or more identically-configured Operator Stations 230 that  
handle the presentation of graphs and displays for the  
operators of the production line including a module 90 to  
support lot information entry and lot changes, and for  
30 performing contact lens lot tracking and reconciliation as  
will be described below; and, an Offline Analysis Node 240



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that provides for analysis of data collected into the Relational Database Node after the data is no longer on-line, i.e., after a given run of the line. As shown in Figure 2, ARCNET interface cards 201,211,221, 231, and, 5 241 are provided for each respective nodes 205,210,220,230,and 240 to support communication between the various nodes via the ARCNET hub 99a. In the preferred embodiment, all of the above-mentioned servers are standard CELLworks software that are commercially 10 available software modules manufactured by FASTech Integration located in Lincoln, Mass.

As mentioned above in view of Figure 2, there are three types of input sources for the existing Supervisor Controller 100: the eight PLCs, the controllers 39a-d for 15 the Injection Molding, Vision Inspection and primary packaging machines, and, data from the sterilization monitoring node 20. The structures of event blocks and data blocks that the supervisory control system 100 reads from each of the eight PLCs and the Vision Inspection and 20 primary packaging machines are described in detail in the above-mentioned U.S. Patent No. 5,461,570. Additionally, as described in detail in the above-identified patent, a relational database is created that is used to store production records and long-term data histories. The 25 existing supervisory controller system 100 provides for on-line and off-line access to this database and includes the mechanism for generating informative graphs including, but not limited to: scattergrams of process parameters vs. contact lens inspection results, 30 histograms of defects by position on pallet, parieto chart of alarm count and duration by machine, time plot of cumulative

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inspection results, measured and calculated parameters  
1 plotted vs. time as a single trend, wherein trend fixed  
time scales are available to show data over minutes,  
hours, days, and weeks.

#### 5 Sterilization Monitoring Node

As shown in the detailed hardware configuration of  
Figure 3, the sterilizer monitoring node 20 interfaces  
with the existing supervisory system 100 by means of an  
10 ARCNET interface card 21 to provide communication with the  
existing supervisor system. An 8-channel serial card 22,  
receives serial data from the sterilizer controller 25  
from a dedicated asynchronous serial line 16b that is  
split from the main sterilizer data line output 16 (Figure  
15 2). A serial port 19 is provided in the node 20 for  
communication of completed lines of sterilization process  
data for complete or incomplete sterilizer runs, messages  
indicating that data is complete or incomplete, or, that  
a sterilizer run was or was not successful, and error log  
20 information and phase file information to the report  
printer 18 for sterilizer run, error log, and phase file  
report generation.

The sterilization monitoring process 10 to be  
described in greater detail hereinbelow comprises two  
25 functional modules: the sterilization communications  
module 50 ("steri comm server") and the sterilization  
server module 60 ("steri server"). Briefly, the  
sterilizer communication server 50 functions to receive  
characters generated by the sterilizer controller via the  
30 8-channel serial card 22. The sterilizer server module 60  
processes all of the inputs and produces reports for a

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variety of users. For instance, as shown in Figure 3, the sterilizer server 60 processes the information to form sterilizer run reports for printing via serial port 19, and, additionally, generates the sterilization run report file, sterilizer phase report file containing sterilizer phase information of the four major sterilizer phases (heat load, exposure, cool load, and cycle complete), and, error log file for error information all for long term disk storage 23. Preferably, the disk storage capacity is at least 324MB but this is easily modifiable to any capacity and may constitute any data storage media. Any alarm or process data point that needs to be immediately acted upon, processed, or stored for later off-line analysis is routed to the existing supervisory controller 100 via ARCNET interface card 21 and ARCNET hub 99a (Figure 1). Details of the sterilization monitoring process 10 and the sterilizer process modules 50, 60 therein will be described in greater detail below.

## 20 **Sterilizer Controller**

Figure 4 illustrates a schematic state diagram showing the internal states of the sterilizer monitor node 20. Barring any alarm errors or communication timeout errors, a normal sterilization run will begin after a batch comprising a quantity ranging from about 1-14,000 individually primary (blister) packaged contact lenses are loaded in the sterilization chamber, in the manner as described in the above-mentioned U.S. Patent No. 5,488,815.

30 As shown in Figure 4, a normal sterilization run comprises five consecutive states or phases: A START phase 161, a

HEAT LOAD phase 162, an EXPOSURE phase 163, a COOL LOAD  
1 phase 164, and, a CYCLE COMPLETE phase 165.

After the node is in a start of run state 160,  
which is initiated at the start of a run when the  
sterilizer process hardware including the controller,  
5 sterilization chamber, and process devices therein are  
initialized, the START PHASE state 161, is entered for  
very short duration while the sterilizer chamber  
temperature is brought up to the process setpoints.  
During the HEAT LOAD phase 162 the sterilization chamber  
10 attains its maximum operating temperature of approximately  
122.5° C., under optimal pressure conditions. In  
preferred operating conditions, the HEAT LOAD phase is for  
a duration of approximately five and one-half minutes (5.5  
min.). During the EXPOSURE phase 163, as shown in Figure  
15 4, the batch of lenses are maximally exposed to  
sterilization conditions for a duration of approximately  
thirty-one (31) minutes. During the COOL LOAD phase 164,  
as shown in Figure 4, a drop of both temperature and  
pressure conditions in the chamber is effected to enable  
20 the batch of lenses to cool for a duration of  
approximately ten (10) minutes. After the COOL LOAD  
phase, the sterilizer enters the CYCLE COMPLETE phase 165,  
where the sterilization process terminates for a time  
duration of under one minute under normal conditions, and  
25 a signal 167 is initiated to put the sterilizer node in a  
standby or wait state 159. After this phase, the trays  
containing the now sterilized lenses are positioned for  
output from the sterilization chamber and the  
sterilization chamber is either put in an idle or rest  
30 state 159 before the next batch of contact lenses is to be  
sterilized.

As shown in Figure 4, and, as explained in  
1 greater detail below, if an abnormal event (Communication  
timeout 168) occurs during any of the START CYCLE, HEAT  
LOAD, EXPOSURE, COOL LOAD, and CYCLE COMPLETE phases, the  
phase may be interrupted as shown by respective lines  
5 160a, 162a, 163a, 164a, and, 165a as shown in Figure 4 to  
indicate that a communication timeout has occurred, i.e.,  
the sterilizer node has not obtained any data for a  
specified time period, which, in the preferred embodiment,  
is approximately one line of data every 60 seconds. If  
10 the event that caused the timeout is rectified, signal 169  
is generated to enable the sterilization monitor node 20  
to again process data from the sterilizer controller 25.  
Since the sterilizer controller 25 was still communicating  
data during the timeout condition, logic built in to the  
15 algorithms explained below will direct the sterilizer node  
20, via signals 161b, 162b, 163b, 164b, and 165b, to resume  
the sterilization monitoring process at the appropriate  
sterilizer phase.

## 20 Sterilization Monitor Processes

As shown in Figures 1 and 3, the sterilizer  
controller 25 provides one way communication with the  
sterilizer monitoring node 20 through an ASCII data stream  
on an RS-232 serial interface, and, the sterilizer  
25 monitoring node 20 is in two-way communication with the  
existing supervisory system 100 through the ARCNET hub  
99a. In the preferred embodiment, during each of the  
above-described sterilization phases, the sterilizer  
controller 25 sends out one line of sterilization process  
30 readings at a frequency of preferably once per minute. If  
an alarm condition exists, as will be explained in further

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1 detail below, the sterilization controller 25 will produce  
and broadcast a line of data at a frequency of once every  
two seconds.

5 With regard to Figures 1 and 3, there generally  
illustrates the flow of data to and from the sterilizer  
monitoring node 20 system for controlling the sterilizer  
monitoring process. As mentioned above, algorithms are  
implemented by each of the sterilization monitoring node  
20 processing modules, i.e., the sterilization  
communications server 50 and the sterilization server 60  
10 to enable passive monitoring of all the data information  
output from the sterilizer and, to communicate the data to  
the data acquisition and analysis nodes of the existing  
supervisor system 100. All information generated from the  
sterilizer controller 25 is serially input to the  
15 sterilization communication server 50 of the sterilizer  
monitor node 20 for data acquisition. Specifically, the  
sterilization controller 25 broadcasts complete lines of  
serial data during each phase of the sterilization process  
to the steri comm server 50. Each line of data will  
20 comprise a number of characters, in the form of twelve  
sterilizer process variable data, alarm information data  
describing an alarm condition, or, textual information.  
The steri server 60 incorporates data processing  
capabilities for processing the input data acquired by the  
25 steri comm server 50 to produce sterilizer run files and  
reports 75, phase files and reports 85, error log files  
95, etc., as will be described in greater detail below.

30 Additionally, as will be explained in greater  
detail below, the following additional information is  
input to the sterilizer server module 60: lot number  
information which is input from the Statistics Server 225

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of the existing supervisor system 100; and, time interval  
1 wakeup data which is generated on a periodic basis by the  
time server 70 of the sterilization node 20 for detecting  
serial communication timeouts. The steri server processor  
60 particularly processes this data as well as the real-  
5 time raw sterilizer process measurement data, to generate  
the following: a sterilization run number for storage and  
subsequent reporting by the statistics server 225 (Figure  
1); alarm messages, for input to the alarm control server  
229 (Figure 1); sterilizer phase file data containing  
10 information for each specific phase of the sterilization  
process for storage in hard disk file 23 (Figure 3);  
Sterilization Run Report file information, which is input  
to the hard disk storage 23 for subsequent generation of  
sterilizer run reports to be described in detail below;  
15 and, sterilizer parameter value information which is input  
to the control server 228 of the existing supervisor  
control system 100 (Figure 1). Each of the above-  
mentioned sterilizer data processing functions will be  
described in greater detail below.

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#### Sterilizer Comm Process

CELLworks system is configured to execute the  
steri comm server 50 when data is to be received. Figures  
5(a) and 5(b) illustrate the steri comm data acquisition  
25 process 50. The first step of calling the Cellworks steri  
comm server 50 is to initialize the serial port hardware  
as indicated as step 110. Next, at step 113 the serial  
port is opened for communication and the input data buffer  
(not shown) is flushed. Next, at step 115, the get\_a\_line  
30 function is called by an infinite loop for acquiring a  
line of serial data from the sterilization controller one

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character at a time. Specifically, as illustrated in  
1 Figure 5(b), at the first step 124, the character count is  
initialized to zero. Next, at step 126 the current  
character count is compared with the size of the input  
buffer (not shown). If the character count is greater  
5 than the input buffer size, then an error message is  
printed at step 127 and no characters are returned for  
processing (step 128). As long as the character count is  
less than the input buffer size, steps 131 and 133 are  
performed for retrieving each successive character (step  
10 131) and comparing the character to determine if it is an  
end of file character (step 133). If the character is not  
the end of file character, then the character is saved in  
the input buffer at step 135, the character count is  
incremented at step 138, and a determination is made as to  
15 whether the character was an end of line character at step  
140. If the character was not the end of line character,  
then the process returns to step 126 to acquire the next  
character. If the character was the end of line  
character, then the process returns the number of  
20 characters in the input line (step 141) and the line of  
data is sent to the steri server's mailbox, i.e., buffer  
location, at step 155 as shown in Figure 5(a). When the  
current character is determined to be the end of file  
character, this indicates that a serious error has  
25 occurred. This may be occur when the serial line is  
disconnected then reconnected. Therefor, if the current  
character is determined to be the end of file character at  
step 133, then, at step 144 of Figure 5(b), the serial  
device is closed, reopened, and initialized. At step 145,  
30 a determination is made as to whether an error occurred  
when the serial port was opened. If no error has

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occurred, then no characters are returned for processing  
1 (step 146). If an error has occurred, then an error  
message is printed at step 149 and the steri comm process  
60 is aborted at step 152.

## 5 Sterilizer Server Process

Figure 6 illustrates the major functional blocks  
of the steri server process 60 which comprises a steri  
server QNX Mailbox System, indicated as element 250. The  
QNX Mailbox System is the primary message routing engine  
10 for the node, and, each server in the sterilizer node 20  
is provided with a mailbox that accepts and sends command  
or data messages. Depending upon the source of the  
message, the steri server process 60 will implement either  
of three functions: startCmdFunc 260; getSteriDataFromMbx  
15 270; and, wakeUpCmdFunc 280, as illustrated in Figure 6  
and explained in further detail below.

The CELLworks system is configured upon startup  
to execute the startCmdFunc 260 when the command to start  
steri server process 60 is received. This function  
20 ensures that the steri server 60 and internal variables  
therein are initialized and that all the QNX communication  
mailboxes (not shown) are setup. Additionally, an entry  
is placed in the error log file 95 (Figure 3) to indicate  
that the system has started up, and, a request is made for  
25 a future wake-up message from the Time Server 70 at a  
prespecified time.

CELLworks is configured to implement the calling  
of the getSteriDataFromMbx functional block 270 whenever  
a message comprising the complete line of data from the  
sterilizer comm process 50 appears in the Steri Server  
30 mailbox. The getSteriDataFromMbx process 270 functions to

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copy the message from the mailbox into local steri server  
1 input buffer (not shown) and initiate the processing of  
the message. Additionally, a check is made to determine  
if the proper message line ending is present. The  
**getSteriDataFromMbx** functional block 270 calls the  
5 **characterizeLine** algorithm 300 for determining the nature  
of the line of sterilizer data information sent from the  
sterilization monitor only if the proper message line  
ending is present. As will be explained below,  
determining the nature of the line of sterilizer data  
10 information and the processing of this data will involve  
one or more of the following functions: **evalTextline**  
indicated as block 440; **processDataLine** indicated as block  
405; **processAlarmLine** indicated as block 500;  
**openSteriRunReport** indicated as block 320; and,  
15 **updateAlarmStatus** indicated as block 550.

As shown in Figure 7, the first step 302 of the  
**characterizeLine** algorithm 300 is to record the current  
time to aid in the detection of a possible communication  
timeout. The next step 304 is to massage the line of data  
20 to remove spurious printer control characters sent by the  
sterilization monitor that could interfere with  
characterization of the line as a line of data. Next, at  
step 306, a determination is made as to if there are any  
printable characters in the line and whether a sterilizer  
run report file 85 has not been opened. If there are  
25 printable characters in the line and a sterilizer run  
report file has not been opened, then the  
**openSteriRunReport** ("steri run report") file algorithm is  
implemented at step 320.

30 The procedure for opening the steri run report  
is illustrated in Figure 19, and as a redundant check the

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1 first step 322 is to determine if the steri run report  
file is already opened. If the steri run report file is  
open the process will return to step 308 of the  
characterizeLine process (Figure 7). If the file is not  
open, then the following steps are performed prior to  
5 opening up the steri run report file at step 330: First,  
at step 324, the minimum and maximum values for each  
variable of each phase of the sterilizer is initialized.  
Then, at step 326, the saved start and stop times for each  
phase are initialized for later calculation of phase  
10 duration explained in detail below. Next, at step 328,  
the initial run report file is set up with a default name  
comprising the current date and time, "date\_time\_RPT", as  
indicated. At step 330, the run report file is opened.  
If it is determined at step 332 that an error has occurred  
15 when opening up the report file, then, at step 334, a  
retry file is set up to open every 15 minutes in the  
preferred embodiment, and, at step 336, the error is  
logged in the log report file. If it is determined at  
step 332 that an error has not occurred when opening up  
20 the report file, then, at step 338, a request is made for  
the previously entered lot number to be sent to this  
process from the statistics server, and the process  
returns to the line characterization algorithm (Figure 7).

25 If a steri run report file has already been  
opened in Figure 7, then communication timeout detection  
is enabled at step 308. The algorithm then proceeds by  
determining at step 310 if the line begins with a time  
signature of the form:

30 hh:mm:ss (hours, minutes, seconds)

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If the line does begin with such a time signature, then  
1 the `processDataLine` algorithm is invoked at step 405  
indicating that the data line contains process variable  
data including twelve (12) variables of sterilizer phase  
data. Else, at step 312, a determination is made as to  
5 whether an alarm condition exists. If an alarm condition  
exists, then the `processAlarmLine` algorithm is invoked at  
step 500 for processing the alarm data and updating the  
alarm control server of the existing supervisor system  
100. If the line of data is determined to be textual in  
10 nature, then the `evalTextLine` algorithm is invoked at step  
450, for processing the text data. Whenever the  
processing of an incoming line has been completed, the  
`updateAlarmStatus` process is called at step 550 to the  
check whether the alarm status has changed.

15 Figure 8 illustrates the `processDataLine`  
algorithm 405 for processing the twelve (12) variables of  
sterilizer phase data from the data line. After removing  
any other remaining printer control characters from the  
line at step 407, the line of data is added to the steri  
20 run report at step 409. Next, the `getDataLinetime`  
algorithm is invoked at step 410 to ensure that, for each  
of the five sterilizer phases, the start time of the  
current operating phase in addition to the phase stop time  
of the previous sterilizer phase has been recorded in the  
25 appropriate phase file. Next at step 412, the 12 process  
variable values of the current data line are isolated and  
saved as text. Then, the `convertAndDoMinMax` algorithm is  
called at step 420 for converting each of the sterilizer  
process readings from text format to floating point  
30 format, and, updating the maximum and minimum values for  
each of the twelve variables for the current sterilizer

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phase. At step 425, it is determined if it is time to update the variables in the phase files. If not, the process returns to the characterizeLine algorithm as indicated at step 427. If it is time to update the variables, then at step 430, a message will be formulated and sent to the control server 228 of the existing supervisor control system 100 to make the twelve floating process variables available for display, trending, and inclusion in the engineering database (not shown) in the manner as explained in detail in the above-mentioned U.S. Patent No. 5,461,570. A determination is then made at step 435 whether the current sterilizer phase is in the heat load, exposure, cycle complete or cool load phase. If the sterilizer is in one of these phases, then the makePhaseFileEntry process is called at step 440. If the sterilizer is not in one of these phases, then a return is made to the calling characterizeLine algorithm 280.

As shown in Figure 9, the makePhaseFileEntry process 440 is called from the Process Data line algorithm 405 to format the process variable information for entry as a line in the corresponding phase file. In the preferred embodiment, there is a separate phase file on disk for each of the four major sterilizer phases. When a new entry is made, it is added to the end of the appropriate phase file as follows: First, at step 445, it is determined if process data has been obtained from the start of the sterilizer run. If not, then the process will return to the processDataLine algorithm at step 446. If the current phase has started from the start cycle, then the following information is gathered in line format: the current lot number at step 448; the sterilizer run

number at step 451; the date at step 452; the time at step  
1 454; an indicator to convey whether an alarm condition  
currently exists or not at step 456; and, the current  
twelve process values for the current sterilizer run at  
step 458. Finally, at step 460, the entire line is  
5 written to the current phase file.

Figure 31 illustrates the format of the above  
described entries of a phase file 876 for storing  
sterilizer phase information. Each row in the table of  
Figure 31 illustrates represents one entry in the phase  
10 file. The lot numbers are entered by the operators on the  
operator stations 230 (Figure 1) of the supervisory  
controller, and, the "mode" entry is a single character  
designation of whether the sterilizer controller indicated  
an alarm at that time or not.

15 Returning to the `characterizeLine` algorithm 300  
as shown in Figure 7, if the data obtained from the  
sterilizer controller is not alarm data or process  
variable data, then the `evalTextline` is called at step 450  
to process the line of text.

20 Figures 10(a) and 10(b) illustrate the  
`evaluateTextline` process 450 for processing textual data  
from the input data line. The first step 461 of the  
evaluate text line process is to remove any remaining  
printer control characters from the line, and, at step  
25 462, adding the line to the steri run report. A  
determination is made at step 463 if the text indicates  
the start of a sterilizer run. If so, then a  
`StartOfRunEvent` procedure is called at step 465 to perform  
the following steps as shown in Figure 11: First, at step  
30 467, an entry into the error log 95 (Figure 3) is made  
indicating that the sterilizer run has started. Then, at

step 468, a flag is set to indicate that the current run  
1 has started from the beginning (start of sterilizer run)  
and, at step 469, that the current state is the start of  
run state. Next, at step 470 of Figure 11, a message is  
sent to the control server of the existing supervisor  
5 system that the current state of the sterilizer is the  
start of run state and the process returns to characterize  
the next line of data (Figure 7).

In Figure 10(a), if the text did not indicate  
the start of a sterilizer run at step 463, then, at step  
10 464, a determination is made if the text indicates the  
beginning of the start phase. If the text indicates the  
beginning of the sterilizer start phase, then the  
doStartPhaseEvent procedure is called at step 465 to set  
a flag that the current state is the sterilizer start  
15 phase, and, to send a message to the control server of the  
existing supervisor system that the current state is the  
sterilizer start phase state before returning to  
characterize the next line of data (step 550, Figure 7).

If the text did not indicate the beginning of  
20 the sterilizer start phase at step 464, then, at step 471,  
a determination is made if the text indicates the  
beginning of the sterilizer heat load phase. If the text  
indicates the beginning of the sterilizer heat load phase,  
then the doHeatLoadPhaseEvent procedure is called at step  
25 472 to set a flag that the current state is the sterilizer  
heat load phase, and, to send a message to the control  
server of the existing supervisor system that the current  
sterilizer state is the heat load phase. Additionally,  
the HeatLoad phase file is opened for writing data thereto  
30 and the procedure returns to characterize the next line of  
data.

1 If the text did not indicate the beginning of  
the heat load phase at step 471, then, at step 473, a  
determination is made if the text indicates the beginning  
of the sterilizer exposure phase. If the text indicates  
the beginning of the sterilizer exposure phase, then the  
5 **doExposurePhaseEvent** procedure is called at step 474 to  
set a flag that the current state is the sterilizer  
exposure phase, and, to send a message to the control  
server of the existing supervisor system that the current  
state is the sterilizer exposure phase. Additionally,  
10 before returning to characterize the next line of data,  
the Heat load phase file is closed and the Exposure phase  
file is opened for writing the data thereto.

If the text did not indicate the beginning of  
the sterilizer exposure phase at step 473, then, at step  
15 475, a determination is made if the text indicates the  
beginning of the sterilizer cool load phase. If the text  
indicates the beginning of the sterilizer cool load phase,  
then the **doCoolLoadPhaseEvent** procedure is called at step  
476 to set a flag that the current state is the sterilizer  
20 cool load phase, and, to send a message to the control  
server of the existing supervisor system that the current  
state is the sterilizer cool load phase. Additionally,  
before returning to characterize the next line of data,  
the Exposure phase file is closed and the Cool load phase  
25 file is opened for writing the data thereto.

If the text did not indicate the beginning of  
the sterilizer cool load phase at step 475, then, at step  
477, a determination is made if the text indicates the  
beginning of the sterilizer cycle complete phase. If the  
30 text indicates the beginning of the sterilizer cycle  
complete phase, then the **doCycComplPhaseEvent** procedure is



called at step 478 to set a flag that the current state is  
1 the sterilizer cycle complete phase, and, to send a  
message to the control server of the existing supervisor  
system that the current state is the sterilizer cycle  
complete phase. Additionally, before returning to  
5 characterize the next line of data, the Cool load phase  
file is closed and the Cycle Complete phase file is opened  
for writing the data thereto. Then, the process returns  
to step 550, Figure 7 to characterize the next line of  
data.

10 If the text did not indicate the beginning of  
the sterilizer cycle complete phase at step 477, then, at  
step 479, a determination is made if the text indicates  
the end of the sterilizer run. If it is the end of a  
sterilizer run, then a `doEndOfRunEvent` procedure is called  
15 at step 480 to perform the following steps as shown in  
Figure 12. First, at step 481 of Figure 12, an entry is  
made in the error log that the sterilizer run has ended.  
Then, at step 482, a flag is set to indicate that the  
current state is the end of run state. Next, at step 483  
20 of Figure 12, the Cycle Complete phase file is closed. At  
step 484, the time of the most recent data line is saved  
as the stop time for the Cycle Complete phase. Then, at  
step 485, the `finishSteriRunReport` procedure is called to  
generate the sterilizer run report as will be explained in  
25 greater detail below. Another procedure indicated as the  
`doEndOfRunCleanup` is performed at step 486 to finish the  
text line processing and print the steri run report as  
explained in greater detail below. Finally, at step 487,  
since it is not known when the next sterilizer run will  
30 begin, the communication timeout detection is disabled.

Referring back to Figure 10(a), if the text did

not indicate the end of the sterilizer run at step 479,  
1 then, at step 488, Figure 10(b), a determination is made  
if the current text line indicates the date. If the  
current line does indicate the date, then the date is  
saved at step 489 and the process returns to the evaluate  
5 text line procedure. If the current line does not  
indicate the date, then, at step 490, a determination is  
made if the current text line contains the cycle count.  
If the current line does contain the cycle count, then the  
count is saved at step 491 and the **makeSteriRunNumber**  
10 procedure is called at step 492 for building the  
sterilization run number and sending the sterilization  
number to the control server of the existing supervisor  
system. In the preferred embodiment, the sterilization  
run number is put together as a combination of the date,  
15 sterilizer number, and the sterilizer cycle count and is  
of the form:

**YP1NNNN**

where Y is the last digit of the year, P1 is the  
20 sterilizer number of the production line, and, the NNNN is  
the sterilizer cycle count. After the sterilization run  
number is obtained, the process returns to the line  
characterization algorithm.

If the current line does not contain the cycle  
count, then nothing is done with the line as indicated as  
25 step 494, and the process returns to the calling  
**characterizeLine** algorithm (step 550, Figure 7).

Figure 13 illustrates the **processAlarmLine**  
algorithm 500 for processing alarm data from the serial  
data line. After removing any other remaining printer  
30 control characters from the line at step 502, a list of

known alarm phrases is searched at step 505 to indicate if  
 1 a known alarm condition exists. Table 1 and Table 2 below  
 indicates various alarm phrases and alarm code data:

	Alarm Text from Sterilizer Controller	Alarm Code
5	CLEAN STEAM LOW	1
	COLD WATER LOW	2
	COMP. AIR LOW	3
	CYCLE ABORT	4
10	DOORS NOT CLSD	5
	DOORS NOT SEALED	6
	FAN FAILURE	7
15	HIGH TEMPERATURE	8
	HIGH WATER	9
	LOW TEMPERATURE	10
	OVER PRESSURE	11
20	PLANT STEAM LOW	12
	POWER FAIL	13
	PRES SENS. ERROR	14
25	PT ISOLATED	15
	SAM FAILURE	16
	STEAM FAILURE	17
30	TEMP SENS. ERROR	18

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1	UNDER PRESSURE	19
	unknown alarm: this code used when there is alarm text that is not recognized	999

5	Condition Detected by Sterilizer Monitor Server	Alarm Code
	Communication timeout with sterilizer controller	01

10 If the alarm condition is unknown, i.e., a match  
is not found between the alarm data phrase and the alarm  
phrase table at step 507, then, at step 508, a  
communication is made to the alarm control server of the  
existing supervisor system that an unknown alarm has been  
received and an active alarm condition is recorded. The  
15 process then proceeds to step 515 to add the alarm line to  
the steri run report. If a known alarm condition exists,  
i.e., a match is found between the alarm data phrase and  
the alarm phrase table at step 507, a determination is  
made at step 509 if the alarm indicates a successful run.  
20 If the indicator is a successful run alarm, it is recorded  
at step 511 and the alarm line is added to the steri run  
report at step 515. If the indicator is not a successful  
run alarm, then, at step 513, a communication is made to  
the alarm control server of the existing supervisor system  
25 which alarm condition has occurred and it is recorded that  
an active alarm condition exists. Finally, at step 515  
the alarm line is added to the steri run report and the  
process returns to the calling characterizeLine algorithm.

30 As mentioned above in view of Figure 7  
illustrating the characterizeLine algorithm, after a line

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is characterized and process variable data, alarm data, or  
1 textual data information is processed, the alarm status is  
updated by invoking the `updateAlarmStatus` process at step  
550. Figure 14 illustrates the `updateAlarmStatus` function  
in detail. The first step, indicated as step 552, is to  
5 determine the status of a parameter (not shown) that is  
supplied indicating if any alarm from the sterilizer  
controller is currently active, and, sending data lines  
every two seconds as described above. If so, then at step  
554, a determination is made whether there is enough time  
10 to clear the alarm or whether the currently active alarm  
should be cleared by force. If not enough time has  
elapsed to clear the alarm, i.e., if the time between the  
last sterilizer data entry and the previous data entry is  
two (2) seconds indicating that the alarm condition should  
15 be cleared by force, then, at step 556, the current  
sterilizer alarm state is set to indicate that no steri  
alarms are active. If there is enough time to clear the  
alarm, i.e., the time between the last sterilizer data  
entry and the previous data entry is greater than two (2)  
20 seconds indicating that the alarm condition data is not to  
be cleared by force, then a determination is made at step  
559 if a communication timeout alarm is active and if  
recent data has been input. As mentioned above, a  
communication timeout alarm occurs when data is received  
25 at an interval greater than one minute, for e.g., when the  
serial data line is temporarily disconnected. If a  
communication timeout alarm is active and recent data has  
been input, then at step 561, the current communication  
alarm state forced to indicate that no communication  
30 alarms are active and is set to provide such indication.  
If a communication timeout alarm is not active or, recent

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1 data has not been input, for e.g., end of a run, a  
determination is made at step 563 if all alarms are  
cleared. If all alarms are currently cleared, then at  
step 565, a message is sent to the alarm control server of  
the existing supervisor system to reset all sterilizer  
5 related alarms. Else, if all alarms are not clear the  
program returns to the calling characterize line process.

Referring back to Figure 12, at step 485, the  
`finishSteriRunReport` procedure 485 is invoked when the end  
of a sterilizer run is detected. As shown in Figure 15,  
10 the first step 610 of the `finishSteriRunReport` procedure  
is to determine if the flag to indicate that the current  
run has started from the beginning (start of sterilizer  
run) had been set as indicated above with respect to step  
468, Figure 11. If the flag indicating that the cycle has  
15 started from the beginning of the sterilization run is not  
set, then, at step 612, a line is added to the  
sterilization run report indicating that the data is  
incomplete, and, at step 614, an entry is made to the  
error log indicating that the data is incomplete. The  
20 process then resumes to step 617. If the flag indicating  
that the cycle has started from the beginning of the  
sterilization run is set, then, at step 611, the  
sterilization run number is added to the steri run report.  
Next, at step 613, a determination is made as to whether  
25 any of the four major sterilization phases has been  
missed. If any of the four sterilization phases has been  
missed in the current run, then a line is added to the  
sterilization run report indicating that the data is  
incomplete (step 612), and, an entry is made to the error  
30 log indicating that the data is incomplete (step 614). If  
none of the four sterilization phases has been missed in

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the current run, then, at step 615, a line is added to the  
1 sterilization run report indicating that the data is  
complete. The next step, indicated as step 617 in Figure  
15, is to determine if the sterilization controller issued  
a successful run alarm indicating a valid cycle as  
5 described above at step 511, Figure 13. If a successful  
run alarm indicating a valid cycle has been generated by  
the sterilization controller, then, at step 619, a line is  
added to the sterilization run report indicating that the  
run was successful. If a successful run alarm indicating  
10 a valid cycle has not been generated by the sterilization  
controller, then, at step 621, a line is added to the  
sterilization run report file indicating that the run was  
NOT successful. Finally, at step 630, a procedure is  
called to add the time durations for each of the major  
15 sterilization phases to the steri run report file, and, at  
step 640, a procedure is called to add the minimum and  
maximum values of the process variables for each of the  
major sterilization phases to the steri run report file.

The `addDurations` procedure 630 of the  
20 `finishSteriRunReport` procedure begins by adding the Phase  
Duration header line of the sterilization run report as  
indicated as step 631 of Figure 17. Next, a pointer is  
set to the first of the four major phases, as indicated at  
step 632. The following steps indicate the printing of  
25 phase durations: At step 633, a label for the phase  
duration is obtained; then, at step 634, the phase  
duration is calculated using the saved phase start and  
stop times obtained at step 326 of the `openSteriRunReport`  
algorithm (Figure 19). Next, a determination is made at  
30 step 635 as to whether the phase duration values for all  
four major sterilizer phases have been processed. If the

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1 phase duration values for all four major sterilizer phases  
have been processed, then, at step 636, the resulting line  
having the calculated phase durations for each of the four  
major sterilizer phases is added to the steri run report  
file and the program returns to step 640 of the  
5 **finishSteriRunReport** procedure (Figure 15). If the phase  
duration values for all four major sterilizer phases have  
not been processed, then steps 633 and 634 will be  
repeated for each successive phase pointed by the pointer  
as indicated at step 638 of Figure 17.

10 The **addMinMax** procedure 640 called by the  
**finishSteriRunReport** procedure begins by adding the  
Min/Max header line to the sterilization run report file  
as indicated as step 641 of Figure 18. Next, as indicated  
at step 642, a line containing the twelve variable names  
15 are added to the run report file. Then, a pointer is set  
to the first of the four major phases, as indicated at  
step 643a. For each phase, a line for the minimum  
readings and a line for the maximum readings are to be  
added to the file. The following steps indicate the  
20 printing of minimum and maximum readings: at step 644, a  
label for the phase's minimum values is added at the  
beginning of a first line of the file; then, at step 645,  
a line containing the twelve minimum values for the  
variables for this phase are added; at step 646, the  
25 resulting line of minimum values is added to the steri run  
report file; at step 647, a label for the phase's maximum  
values is added at the beginning of a second line of the  
file; then, at step 648, a line containing the twelve  
maximum values for the variables for this phase are added;  
30 and, at step 649, the resulting line of maximum values is  
added to the steri run report file. Next, a determination

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is made at step 650 as to whether the Min/max values for  
1 all four major sterilizer phases have been added to the  
steri run report file. If the Min/max values for all four  
major sterilizer phases have been printed, then the  
process returns to step 486 of the doEndofRunEvent  
5 procedure for printing of the sterilizer phase durations.  
Until the Min/max values for all four major sterilizer  
phases have been printed, step 644 through step 649 will  
be repeated for each phase pointed to by the pointer as  
indicated at step 643b of Figure 18.

10 Finally, after performing the addDurations and  
addMinMax values, a return is made to the doEndofRunEvent  
procedure 480 where the doEndOfRunCleanup procedure is  
performed at step 486, as indicated in Figure 12.

Figure 16 illustrates the doEndOfRunCleanup  
15 procedure 486. As shown at step 651, this procedure  
implements the closeAndPrintRunReport procedure for  
closing and printing the sterilization run report, as  
described in further detail below with respect to Figure  
20 steps 660 through 669. Furthermore, the  
doEndOfRunCleanup procedure of Figure 16 will: initialize  
20 all internal variables to no-run-in-progress conditions at  
step 652; invoke the update alarm status procedure at step  
653 and as described above with respect to Figure 14 steps  
552 through 565; close the error log at step 654; and, at  
25 step 655, communicate to the control server that the  
sterilizer is currently in a standby or waiting state.  
Finally, a return is made to step 487 of the  
doEndofRunEvent procedure of Figure 12.

As illustrated in Figure 20, to close and print  
30 the sterilization run report, the first step 660 is to  
close the run report file. As mentioned above, if a

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sterilizer cycle was determined to be incomplete, a  
1 sterilizer run number may not have been obtained for the  
incomplete run. Therefore, at step 662, a determination  
is made if a sterilization run number has been obtained  
for this run. If a sterilization run number has been  
5 obtained for this run, then, at step 665, a new run report  
name is configured that will preferably comprise the  
sterilization run number followed by an underscore and the  
lot number. At step 667, the run report file is renamed to  
the new configuration. Finally, the run report file is  
10 printed at step 669. If a sterilization run number had  
not been obtained for this run, then, at step 663, an  
entry is placed in the error log indicating that there is  
no steri run number and the run report file is printed at  
step 669 with the default file name which is initially  
15 assigned as comprising the date and time as described  
above with respect to step 328, Figure 19. After the run  
report file is printed at step 669, the process returns to  
step 652 of the doEndOfRunCleanup procedure 486 (Figure  
16).

20 The printed sterilizer run report 800 for a  
valid and complete run, as shown in Figure 23, and printed  
by the printer 18 connected to the node 20, consists of:  
the updated steri run report file name comprising the  
sterilization run number followed by an underscore and the  
25 lot number, as indicated as line 810; heading data,  
indicated as lines 815, that includes: the date and time  
a sterilizer run begins, lines 816a, 816b, respectively;  
the name and number uniquely identifying the sterilizer,  
lines 817a,817b, respectively; a cycle counter number to  
30 uniquely identify the sterilizer run, line 818; twelve  
sterilizer process parameters, indicated as lines 819, of

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which the Expose Timer therein indicates the target value  
1 for the duration of the Exposure Phase; the program 822  
which indicates the control program in the sterilizer  
controller; and, the proptime, 824, indicating the column  
headings for the output data lines, indicated as lines  
5 825, that are sent to the existing supervisory controller  
100 and sterilizer monitoring node 20 by the sterilizer  
controller 25. Each of the data lines 825 additionally  
include the time relative to the start of the run,  
followed by a reading for each of the twelve signals  
10 described in the header. Also provided are: the  
sterilization run number indicated as line 812; the lines  
of variable data for twelve process variable data labelled  
V1-V12 and generated once every minute; the durations for  
the four major sterilization phases, indicated as line 835  
15 with label header line 830; the minimum and maximum  
readings observed for each sterilizer variable during each  
of the four major sterilization phases, as indicated as  
lines 845, with a header line 840; the sterilization run  
success failure assessment, indicated as line 850; and, if  
20 all data has been received for the four sterilization  
phases, i.e., for the current run, then the line 855  
indicating that the data is complete. Additionally, a  
line 805 for placing the signature of an operator or  
engineer is provided in the steri run report. Alarm  
25 information (for e.g., fan failure) may also be printed by  
the sterilizer controller and any alarm text is printed on  
a line by itself. The sterilizer controller will surround  
the alarm text with printer control codes that cause the  
text to be printed in red.

30 Although not shown, the dedicated printer 17  
(Figure 1) will print out a sterilization run report

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similar to the report shown in Figure 23 directly from the  
1 sterilizer controller via serial data line 16a. However,  
the report printed by dedicated printer 17 will not have  
the phase duration and minimum/maximum value summaries,  
and success/failure indications as provided in the  
5 sterilizer run report generated by the sterilizer  
monitoring node.

Figure 21 illustrates in detail the  
**wakeUpCmdFunc** process 280, which is a software function  
that is executable whenever the Time Server 70 (Figure 3)  
10 sends a wakeup message to the Steri server process 60. As  
illustrated in Figure 21, the first step 281 of the  
**wakeUpCmdFunc** process is to request the time server to  
send to the steri server 60 another wakeup message at a  
specified time in the future. It is understood that this  
15 is not a continuous process and that the time server 70  
must be requested to send a wakeup message to the steri  
server. The next two steps, indicated as steps 283 and  
284 are to indicate if a communication timeout has  
occurred. Specifically, at step 283, a determination is  
20 made as to whether the steri server is initialized, the  
timeout detection function is enabled, and that there is  
currently no communication timeout alarm that is active.  
If any of these conditions do not exist, then the process  
will return to the CELLworks system as indicated at step  
25 285. If all these conditions exist, then at step 284, a  
determination is made as to if an undue amount of time,  
for e.g., greater than one (1) minute, has elapsed since  
the last receipt of data. A condition such as this would  
occur if the serial data line has been temporarily  
30 disconnected. If the amount of time since the last data  
receipt is not excessive (not greater than 1 minute), the

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process will return to the CELLworks system as indicated  
1 at step 285. If the amount of time since the last data  
receipt is excessive, then, at step 286, a communication  
timeout alarm is indicated as active. Then, at step 287,  
a communication timeout alarm message is sent to the alarm  
5 control server of the existing supervisor system.  
Additionally, at step 288, a communication timeout alarm  
entry is made in the error log. Next, at step 290, an  
**endRunReportForTimeout** process is called as illustrated in  
Figure 22 and described in detail as follows: First, at  
10 step 291, a determination is made as to whether a  
sterilizer run report file is currently open. If a  
sterilizer run report file is not currently open, then the  
process returns to step 295 of the **wakeUpCmdFunc** process.  
If a sterilizer run report file is currently open, then,  
15 at step 292, a line is added to the run report file  
indicating that a communication timeout has occurred, and,  
at step 293, that there is incomplete run data for the  
current sterilizer run. At step 294, an entry is made in  
the error log that the run data is incomplete and the  
20 process returns to step 295 of the **wakeUpCmdFunc** process  
of Figure 21. At step 295, the **doEndOfRunCleanup**  
procedure is called as described above with respect to  
Figure 16. Finally, the communication timeout enable  
function is disabled at step 296 and, at step 297, any  
25 phase file that is currently open, is closed before  
returning to the **wakeUpCmdFunc** process.

#### Lot Tracking and Reconciliation

As shown in Figure 2, lot number and power for  
30 the lenses to be produced will be input by operators at  
any of the four operator stations 230 located along the

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production line. Then, the lot reconciliation and  
1 tracking algorithms 90 that are resident at the stations  
are implemented for calculating and recommending the  
expiration date, lens center thickness, and other  
variables for lot information storage. In the preferred  
5 embodiment, the expiration date is sixteen months from the  
entered system date, but the number of months may change  
and the algorithm is easily modifiable by skilled  
artisans.

As indicated in process flow diagram of Figure  
10 24, an operator will first be queried to enter his/her  
name and password as shown as step 671 which are then  
verified at step 672 by the password files of authorized  
individual names and passwords residing in the data  
analysis node (not shown). Using the operator entry of  
15 lens lot number, power value information and the current  
date information at the operator stations, the lot  
tracking and reconciliation algorithm 90 is implemented.  
Specifically, at step 673, the entered lens lot number is  
retrieved and the format of the lens lot number is  
20 verified. From the verified lot number which contains a  
digit that signifies whether the lot is for revenue or  
trial ("R/T"), a determination is made at step 674 as to  
whether the lot will be for revenue or trial. Next, at  
step 675, the entered lens power value is retrieved and  
25 the validity of the power factor is verified. A  
determination as to whether the power factor is valid is  
made at step 676. If not a valid power factor, the  
operator is prompted to enter a new power factor for  
validation (step 675). If the power factor is valid, the  
30 algorithm will use look-up tables (not shown) to determine  
the product code ("UPC") and lens center thickness

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information for the lot at step 677. It should be  
1 understood that the lot number, power value information,  
lot expiration date, product code and other entries may be  
displayed at the front of the production line, as shown at  
step 678, and, at any of the operating stations until the  
5 product reaches primary packaging. Additionally, the lot  
number, power value information and other entries may be  
changed until the lots product reaches primary packaging.

To aid in tracking lot movement, lot  
reconciliation, and lot changeover procedures, a display  
10 is available at either an operator station or a DynaTerm  
display station 29a,b at the sterilizer node to enable an  
operator to request for display in formatted fields, the  
stored or previously entered lot number data including lot  
number, product code, lens power, lens center thickness,  
15 expiration date, whether the product is for revenue or  
trial, as well as the current location of the lot on the  
line. As will be explained in further detail below,  
besides displaying the previous-entered information, the  
sterilization run number, as provided by the Sterilization  
20 server 60 of the sterilization node 20, may also be  
displayed. It is understood that the each operator  
station shown in Figure 2 has a specialized CELLworks  
program for obtaining operator requests and displaying  
information to the operator in formatted fields.

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#### Lot Tracking

As mentioned above, an operator is capable of  
sequencing the lot to the next part of the line. When the  
lot enters production, six variables representing lot  
30 number information are stored in the Statistics Server of  
the existing Supervisory control system 100. These

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variables are lotnumber, UPC, R/T, Power, Thickness, and  
1 Expiration and is shown having the data structure as shown  
in Figure 25(a). Depending upon the manufacturing zone  
where the lot is located, each data structure for the lot  
is tagged with an "0" or "F" to respectively signify  
5 whether the lot is prior to line, or, is at the front of  
the line (injection molding, lens fab, cure, demold, etc.)  
prior to packaging. When a lot starts into primary  
packaging, the data structure of Figure 25(a) is tagged  
with a "P" to signify that it is in packaging. When the  
10 lot of lenses are in the tray loading area, sterilization,  
or secondary packaging areas, the Statistics server 225  
will store two additional variables onto the data  
structure of Figure 25(a) to form new data structure as  
shown in Figure 25(b). These variables are the input  
15 quantity and loss quantity and are supplied to the system  
during lot reconciliation. When a lot starts into the  
sterilizer tray loader area, the variables are tagged with  
an "L", and when the lot enters the sterilizer they are  
tagged with an "S". When a lot enters secondary  
20 packaging, the six lot information variables are tagged  
with a "C" to signify cartooning. Thus, the feature of  
the present invention is the ability for an operator to  
track movement of a particular lot.

As shown in the data flow diagram of Figure 26,  
25 an operator may request movement of a lot by inputting  
data such as the requested lot 875 being processed on the  
line, and, to which location 880 on the production line to  
which the lot is to be moved. A moveLot algorithm 680 is  
implemented which will process the operator entered data,  
30 as well as lot information data such as: lot number 881,  
product code 882, power 883, lens center thickness 884,



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1 expiration date 885, whether the product is for revenue or  
trial 886, and, total lens input to the particular  
location of the line 887 and the total lenses lost 888 (as  
will be explained below).

5 Figure 27 illustrates the moveLot algorithm 680  
for tracking lens lot movement throughout the production  
line. As shown as step 680 in Figure 27, the source  
location of the lens lot on the production line is  
retrieved. Next, at step 683, a determination is made as  
10 to whether the lot number is to be changed, for instance,  
when a lot is to be split prior to entry in the sterilizer  
15 as a preventative measure to eliminate defective  
batches rather than a whole lens lot when there exists a  
problem in the line. If the lot number is not to be  
changed, then, at step 685, the lens lot and lot  
15 information data such as: lot number, product code, power,  
lens center thickness, expiration date, and whether the  
product is for revenue or trial, is moved to the next line  
location and tagged with a "P", "L", "S" or "C" indicator  
depending at which point on the line the lens lot has  
20 moved to. If the lot number is to be changed, then, at  
step 687, the lens lot number is retrieved from the  
operator and the format of the lens lot number is verified  
at steps 689 and 690. If the entered lot number is not  
valid, the operator is prompted to enter a new power  
25 factor for validation (step 687). If the lot number  
entered is valid, the algorithm will proceed to update the  
above-identified lot information at step 692.

#### Lot Reconciliation

30 Due to the fact that product may be lost during  
manufacture, or, may be removed for quality assurance

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purposes, it is necessary to account for this product and  
1 the reasons for their loss or removal prior to secondary  
packaging. Lot reconciliation is the process whereby each  
lens of a particular lot that leaves primary packaging,  
hereinafter indicated as Zone 1, and enters sterilization  
5 (sterilization tray loading), hereinafter indicated as  
Zone 2, is accounted for at the time that the lot is  
available for secondary packaging, (i.e., sterilization  
tray unloading, cartooning, check weighing, and  
labelling), hereinafter indicated as Zone 3.  
10 Particularly, a lot reconciliation sheet 890, illustrated  
in Figure 32, is generated that will indicate the number  
of lenses input to the sterilizer (line 891) and number of  
lenses available for secondary packaging (lines 899) as  
well as verify the difference that is equal to the number  
15 of lenses that have been lost or removed at each  
particular zone (lines 892,893,894 and 895). The lot  
tracking and reconciliation algorithm 90 in the existing  
supervisor control system 100 supports data entry and  
calculations for Lot Reconciliation.

20 Figure 28 illustrates the start of the lot  
reconciliation process which entails the reporting of the  
quantity of lenses input to the sterilizer 15 after  
primary packaging. Before this quantity could be entered  
at the operator terminals 230 or the DynaTerm Operating  
25 consoles 29a,b (Figure 2), a determination is made at step  
702 to determine whether the lot has cleared from primary  
packaging, i.e., have been packaged in blister packs and  
readied for sterilization. If not, then an error message  
will be displayed at step 704 that an input quantity can  
30 not be entered until the lot has cleared primary packaging  
(Zone 1). If the lot has cleared primary packaging, then

the operator is prompted to enter his/her initials and the  
1 actual quantity of lenses ready for sterilization at step  
705. Next, at step 706, the lot information is updated to  
reflect the new variable, quantity input, with the lot  
number.

5 Just as a number representing the quantity of  
lenses entering sterilization is entered, the number of  
lenses removed from the sterilization (Zone 2) and  
secondary packaging (Zone 3) areas must be recorded and  
entered. As shown in Figure 29, at steps 710 and 720, an  
10 operator may enter the number of product lost, the  
particular zone where the product was lost, the reasons  
why the product was lost or removed, and, the  
sterilization run number, all at the operator terminals or  
the DynaTerm Operating consoles. This information is all  
15 referenced with a particular lot number that is input from  
the sterilization server. This process may be repeated  
several times while a lot is in one zone, for e.g., when  
there are multiple incidents of lost lenses while in the  
zone.

20 Specifically, at step 710, the zone number for  
the current lot location is retrieved and a verification  
is made at step 713 as to whether the zone number is  
valid. If not, then the operator will be prompted to  
enter the a correct zone number. If the lot number is  
25 valid, then a determination is made at step 716 as to  
which zone number was entered. If the current lot is in  
Zone 2 or 3, then the operator is prompted to enter the  
Sterilization run number at step 718 before entering the  
number of lenses removed at step 720. If the current lot  
30 is in Zone 1, then the operator is prompted to enter the  
number of lenses removed at step 720. In response to

entering the number of quantity removed, at step 723, the  
 1 operator station or Dynaterm displays a reason selection  
 list for the operator to enter the particular reason(s)  
 why the lens packages were removed from production at step  
 725. As there may be multiple sterilization runs per lot,  
 5 product may be lost for a number of reasons per  
 sterilization run. Look-up tables (not shown) having  
 reason codes and their definitions are available on each  
 of the operator stations or DynaTerm consoles. Table 2  
 below details some of the reasons for lens removal:

REASON CODE	DEFINITION
01	Removed_by_QA
02	Lenses_to_Distribution
03	Foil_Tear_Mechanical
04	Foil_Tear_Foil_Size
05	Incorrect_Power_on_Foil
06	Incorrect_Expiration_Date_on_Foil
07	Illegible-Print_on_Foil
08	Misaligned_Print
09	Misaligned_Foil
10	Misaligned_Perforation
11	No_Solution
12	Low_Solution
13	Incorrect_Lot_Number_on_Foil

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1	14	Incomplete_Seal
	15	Perforation_Tear
	16	Blown_Seal/Package
5	17	Barcode_Not-Printed_In_Proper-Place
	18	Lot_Number_Not_In_Brackets
	19	Incorrect_#_Digits_in_Lot#_Barcode
	20	Incorrect_#_Digits_in_UPC_Barcode
10	21	Invalid_Barcode#_(Lot Number)
	22	Invalid_Barcode#_(UPC)
	23	Incorrect_Check_Digit
15	24	Spots_On-Barcode
	25	Void(s)_On_Barcode
	26	Low_Grade_Barcode_Scan_Verification
	27	Excess_Glue
20	28	Tabs_Unsealed
	29	Tabs_Unparallel
	30	Ink_Smears_Carton
25	31	Damage_To_Carton
	32	Foreign_Matter
	33	Cartoner_Rejected_at_Checkweigher
	34	Misaligned_Label
30	35	Print_Out_of_Shaded_Area_Carton

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1	36	Incorrect_Case_Label_Info
	37	Jam_Array_Destroyed
	38	Cartoner_Jam_Carton_Destroyed
5	39	Conveyor_Transfer_Jam_Carton_Destroyed
	40	Jam_into_Belt_Array_destroyed
	41	Jam_out_of_Belt_Array_Destroyed
	42	No_Print_Foil
10	43	No_Perf_Foil
	44	Extra_Lens
	60	Supply_Your_Own_Reason

15

At step 727 in Figure 29, a determination is made as to whether the selection is made from the list displayed on the operator station, or, whether the operator entered a new reason for lens package removal. If the selection is made from the displayed list, then the operator is prompted to enter his/her initials at step 731. If the operator entered a new reason for lens package removal, the description entered by the operator, which may be up to 34 characters, is retrieved at step 729 prior to operator initials entry at step 731. The next step 735 updates the lot reconciliation data which has the data structure depicted as shown in Figure 25(b) explained above.

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After secondary packaging is complete, the last action to be performed during lot reconciliation is to close out the lot as shown in the process flow diagram of

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Figure 30. As shown in Figure 30, an operator will first  
1 be queried to enter his/her name and password as shown as  
step 740 which are then verified at step 742 by the  
password files. The updated lot information data is  
retrieved at step 745 and a determination as to if the lot  
5 has cleared the prior zone is made at step 747. If the  
lot has not cleared the prior zone, then an error is  
reported at step 749 and the lot will not be closed out.  
If the lot has cleared the prior zone, then the master lot  
(or split lot) information is obtained by operator entry  
10 at step 751. Then, at step 754, lot reconciliation  
calculations are performed to total number of lenses lost  
or removed. Next, at step 756, a determination is made  
if the total number of lens packages lost or removed is  
greater than the total number of lens packages input. If  
15 the total number of lens packages lost or removed is  
greater than the total number of lens packages input then  
an error message is reported at step 758. If the total  
number of lens packages lost or removed is greater than  
the total number of lens packages input then a  
20 determination is made at step 759 whether the total lost  
or removed is more than one percent (1%) of the total  
number input. If the total number of lens packages lost or  
removed is greater than one percent, then an error message  
is reported at step 761. If the total number of lens  
25 packages lost or removed is less than one percent, then a  
determination is made at step 764 whether the total lost  
or removed is equal to or less than one percent of the  
number of lenses input. If the lens package quantity  
input equals the quantity output, then the lot is closed  
30 out at step 768. If the total number of lens packages  
lost or removed is less than one percent of the quantity

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input, then the total loss of less than one percent is  
1 reported to the operator at step 769, and the operator is  
prompted for closure action at step 771. If the operator  
elects to close the master lot (or split lot) at step 771,  
then the master lot (or split lot) is closed at step 768.  
5 Then, a lot reconciliation report is created for storage  
and printing as indicated at step 775.

While the invention has been particularly shown  
and described with respect to the preferred embodiments  
thereof, it will be understood by those skilled in the art  
10 that the foregoing and other changes in form and details  
may be made therein without departing from the spirit and  
scope of the invention, which should be limited only by  
the scope of the appended claims.

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The embodiments of the invention in which an exclusive property or privilege is claimed are defined as follows:

1. A quality control system for an automated production line producing contact lenses, said production line having a plurality of contact lens process stations, including an automated sterilization station for sterilizing a plurality of contact lenses after their manufacture, and a packaging station for packaging said lenses after sterilization, wherein the system comprises:

(a) a first means for receiving contact lens data including an associated lot number and lens power for a lens lot prior to their manufacture; said lens lot defining at least one batch of contact lenses;

(b) a plurality of process controllers for controlling one or more process stations, each of said controllers regulating a plurality of process control devices at said process stations for manufacturing said contact lenses;

(c) means for tracking movement of said plurality of lenses defined by said lens lot from a said plurality of processing stations to said automated sterilization station and said packaging station;

(d) means for receiving data representing the number of lenses that are input to said packaging station together with reason codes for contact lenses lost at said automated sterilization station;

(e) means for generating a summary report of the total number of lenses input to said automated sterilization station for a predetermined lens lot and the actual number of lenses sterilized and packaged from said lot, said summary report including lot number and lens power data for each batch of contact lenses.

2. The quality control system as claimed in Claim 1, wherein said lens lot comprises a plurality of batches of lenses, said sterilization station sterilizing said one batch of contact lenses at a time and generating a sterilization cycle run number for each batch sterilized.

3. The quality control system as claimed in Claim 2, wherein said summary report reconciles the number of lenses input to said automated sterilization station with the number of lenses packaged from said batch for each sterilization cycle run number.

4. The quality control system as claimed in Claim 2, wherein each lens lost or removed has associated therewith a reason code, said summary report generating means including the lens package removal code with a sterilization cycle run number.

5. The quality control system as claimed in Claim 2, further including means for automatically calculating a thickness specification and product code for a given lens power.

6. The quality control system as claimed in Claim 5, wherein the means for calculating a thickness specification and product code includes a look-up table.

7. The quality control system as claimed in Claim 2, further including means for automatically calculating an expiration date for a given lens lot.

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8. The quality control system as claimed in  
1 Claim 2, wherein said tracking means includes means for  
displaying the location of a particular lot on said  
production line.

5 9. A sterilizer data processing system for an  
automated contact lens manufacturing line for  
manufacturing a plurality of contact lenses defined by a  
lens lot, said contact lens manufacturing line including  
an automated sterilization station for sterilizing said  
10 lens lot after their manufacture, said automated  
sterilization station including a sterilizer process  
controller for controlling one or more phases of a  
sterilization process, said process controller  
periodically generating sterilization process data during  
15 each said sterilization phase, said system including:

(a) means for receiving said sterilization data  
from said sterilizer process controller; and,

(b) means for automatically parsing said  
sterilization data into text information and sterilizer  
20 parameter information, said means further processing said  
text information and sterilizer parameter information to  
automatically generate a sterilization run report  
associated with a lot number for said sterilized lens lot.

25 10. The sterilizer data processing system for an  
automated contact lens manufacturing line as claimed in  
Claim 9, wherein said processing means further conveys  
said sterilizer parameter information to data acquisition  
devices in said automated contact lens manufacturing line  
30 for display and storage thereof.

11. The sterilizer data processing system for an  
1 automated contact lens manufacturing line as claimed in  
Claim 9, wherein said process controller generates alarm  
condition information, said processing means further  
conveying said alarm condition information to data  
5 acquisition devices in said automated contact lens  
manufacturing line for display thereof.

12. The sterilizer data processing system for an  
automated contact lens manufacturing line as claimed in  
10 Claim 9, wherein said processing means further includes  
means for evaluating the success or failure of a  
sterilization run based on said text information and  
sterilizer control parameter information for indication on  
said sterilization run report.

15  
13. The sterilizer data processing system for an  
automated contact lens manufacturing line as claimed in  
Claim 9, wherein said processing means further includes  
means for evaluating whether a complete set of data has  
20 been obtained for the current sterilizer run.

14. The sterilizer data processing system for an  
automated contact lens manufacturing line as claimed in  
Claim 9, wherein said text information includes the  
25 current operating phase of said sterilization process.

15. The sterilizer data processing system for  
an automated contact lens manufacturing line as claimed in  
Claim 9, wherein said sterilizer parameter information  
30 includes a plurality of isolated process values generated

by said sterilization process controller during each operating phase of said sterilization process.

16. The sterilizer data processing system for an automated contact lens manufacturing line as claimed in Claim 9, wherein said processing means further includes means for determining a phase duration time for each said operating phase of said sterilization run, wherein said phase duration time is included in said sterilization run report.

17. The sterilizer data processing system for an automated contact lens manufacturing line as claimed in Claim 9, wherein said processing means further includes means for determining a minimum value and maximum value of isolated process values for each said sterilization phase, wherein said minimum and maximum of said isolated process values are included in said sterilization run report.

18. The sterilizer data processing system for an automated contact lens manufacturing line as claimed in Claim 15, wherein said processing means further includes means for generating phase files for storing phase file information for each of said sterilizer operating phases.

19. The sterilizer data processing system for an automated contact lens manufacturing line as claimed in Claim 9 wherein said sterilizer process controller generates sterilizer data at a first pre-specified time interval, said sterilizer data processing system further including means for determining whether said receiving

means receives said sterilization data from said  
1 sterilizer process controller within said time interval.

20. The sterilizer data processing system for  
an automated contact lens manufacturing line as claimed in  
5 Claim 19 wherein said means for determining whether said  
receiving means receives said sterilization data from said  
sterilizer process controller within said time interval  
includes a time server for generating a wakeup message  
during each said first time interval.

10  
21. The sterilizer data processing system for  
an automated contact lens manufacturing line as claimed in  
Claim 20, wherein said processing means generates a  
communication timeout alarm message in the event that said  
15 sterilization data is not received during each said first  
time interval.

22. The sterilizer data processing system for  
an automated contact lens manufacturing line as claimed in  
20 Claim 9 wherein said receiving means is a communication  
server.

23. The sterilizer data processing system for  
an automated contact lens manufacturing line as claimed in  
25 Claim 9 wherein said processing means is a sterilization  
server.

24. The sterilizer data processing system for  
an automated contact lens manufacturing line as claimed in  
30 Claim 11 wherein said sterilizer process controller

generates alarm condition information at a second  
1 prespecified time interval.

25. A method for processing data generated in  
an automated contact lens manufacturing line for  
5 manufacturing a plurality of contact lenses defined by a  
lens lot, said contact lens manufacturing line including  
an automated sterilization station for sterilizing said  
lens lot after their manufacture, said automated  
sterilization station including a sterilizer process  
10 controller for controlling one or more phases of a  
sterilization process, said process controller  
periodically generating sterilization process data during  
each said sterilization phase, said method including the  
steps of:

15 (a) receiving said sterilization data from said  
sterilizer process controller; and,

(b) automatically parsing said sterilization  
data into text information and sterilizer parameter  
information; and,

20 (c) further processing said text information and  
sterilizer parameter information to automatically generate  
a sterilization run report associated with a lot number  
for said sterilized lens lot.

25

30

35

**FIG. 1**

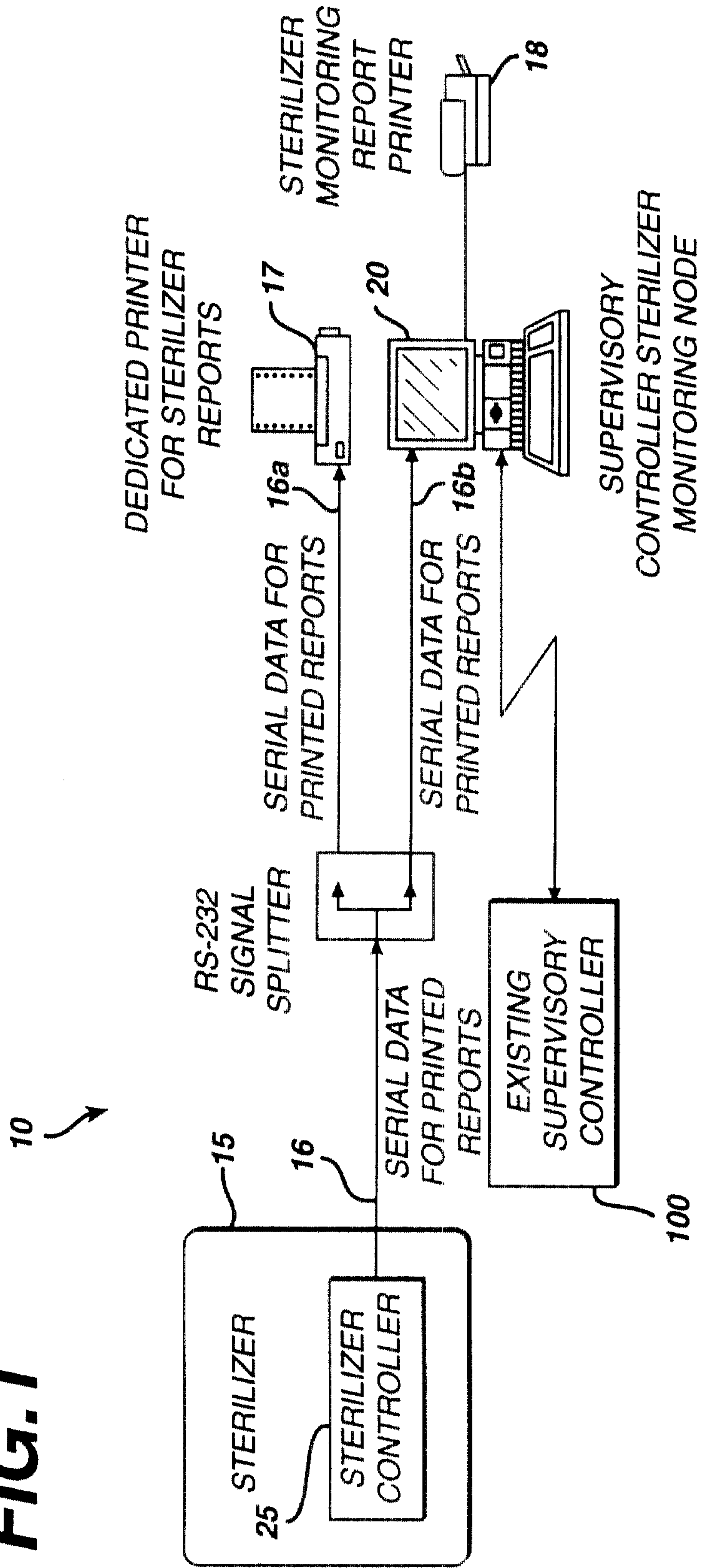
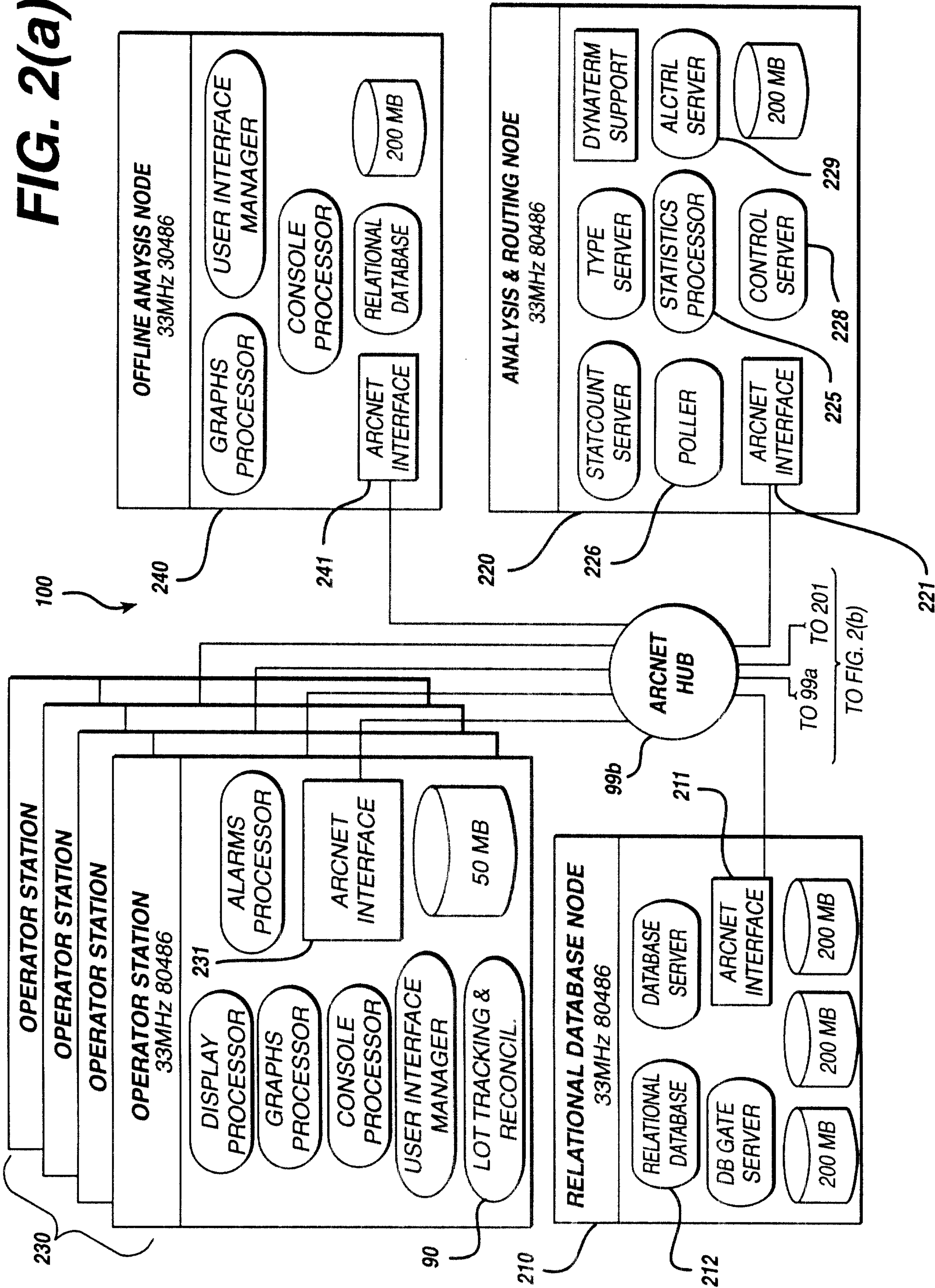




FIG. 2(a)



**FIG. 2(b)**

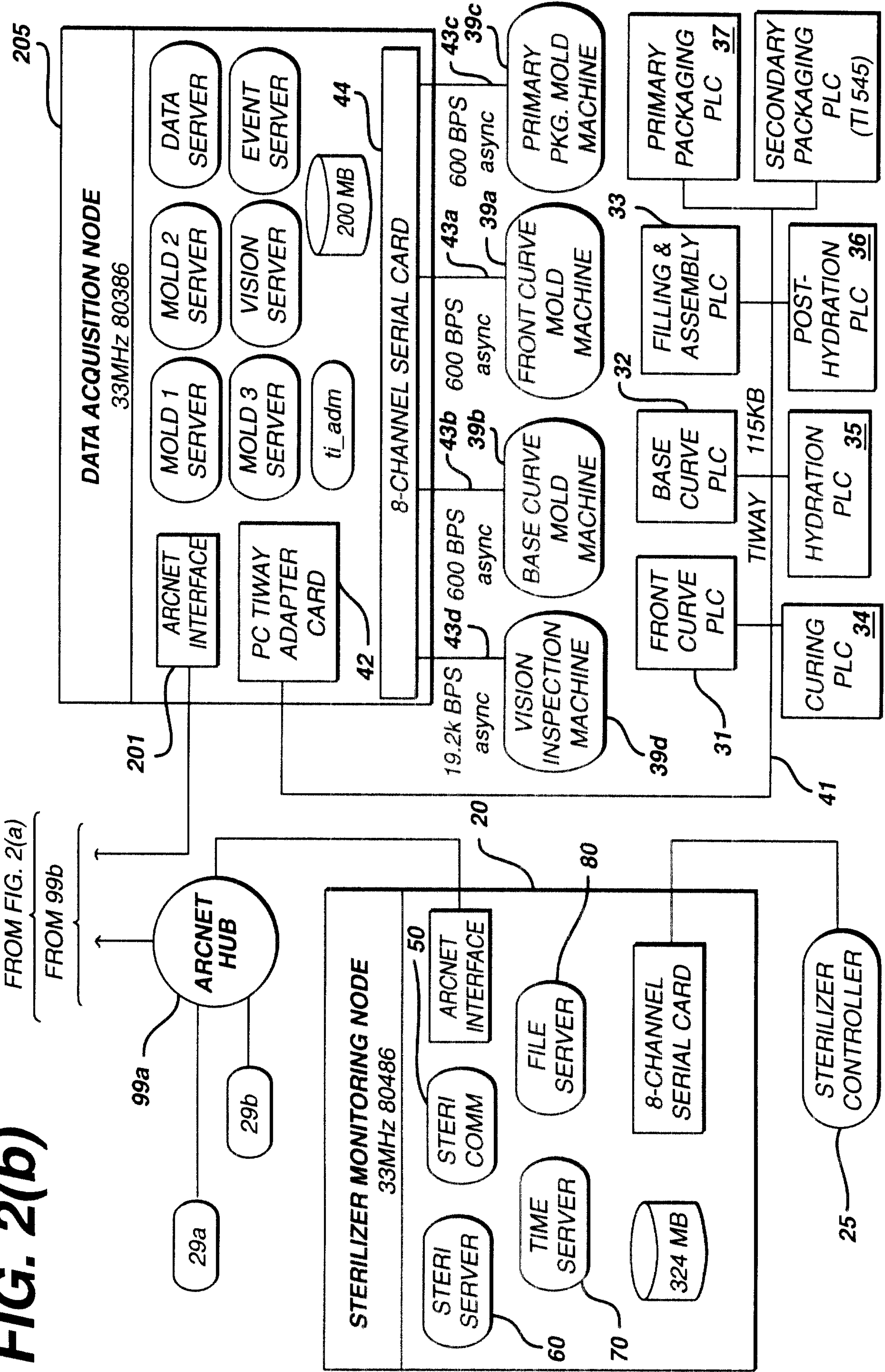
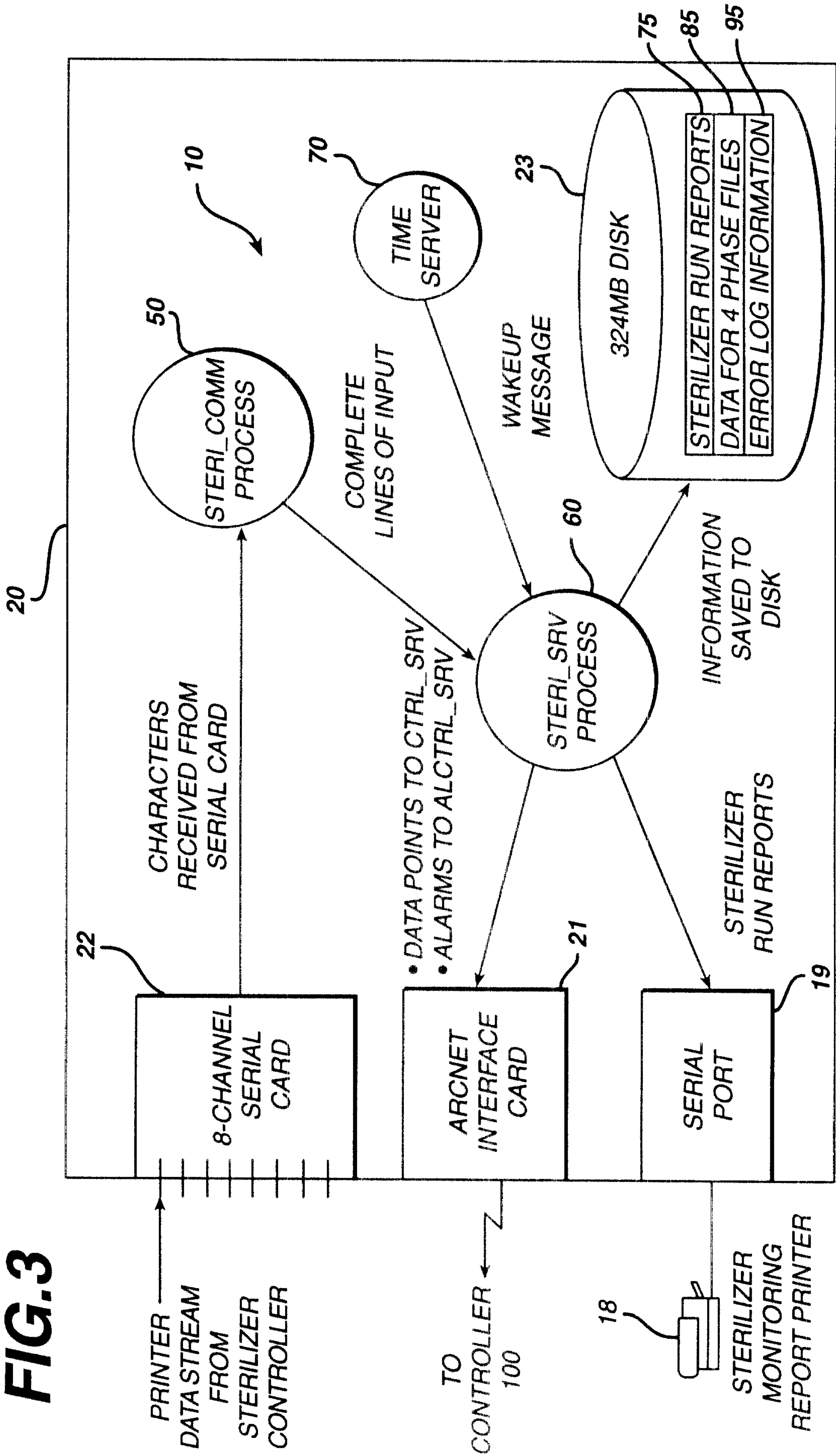
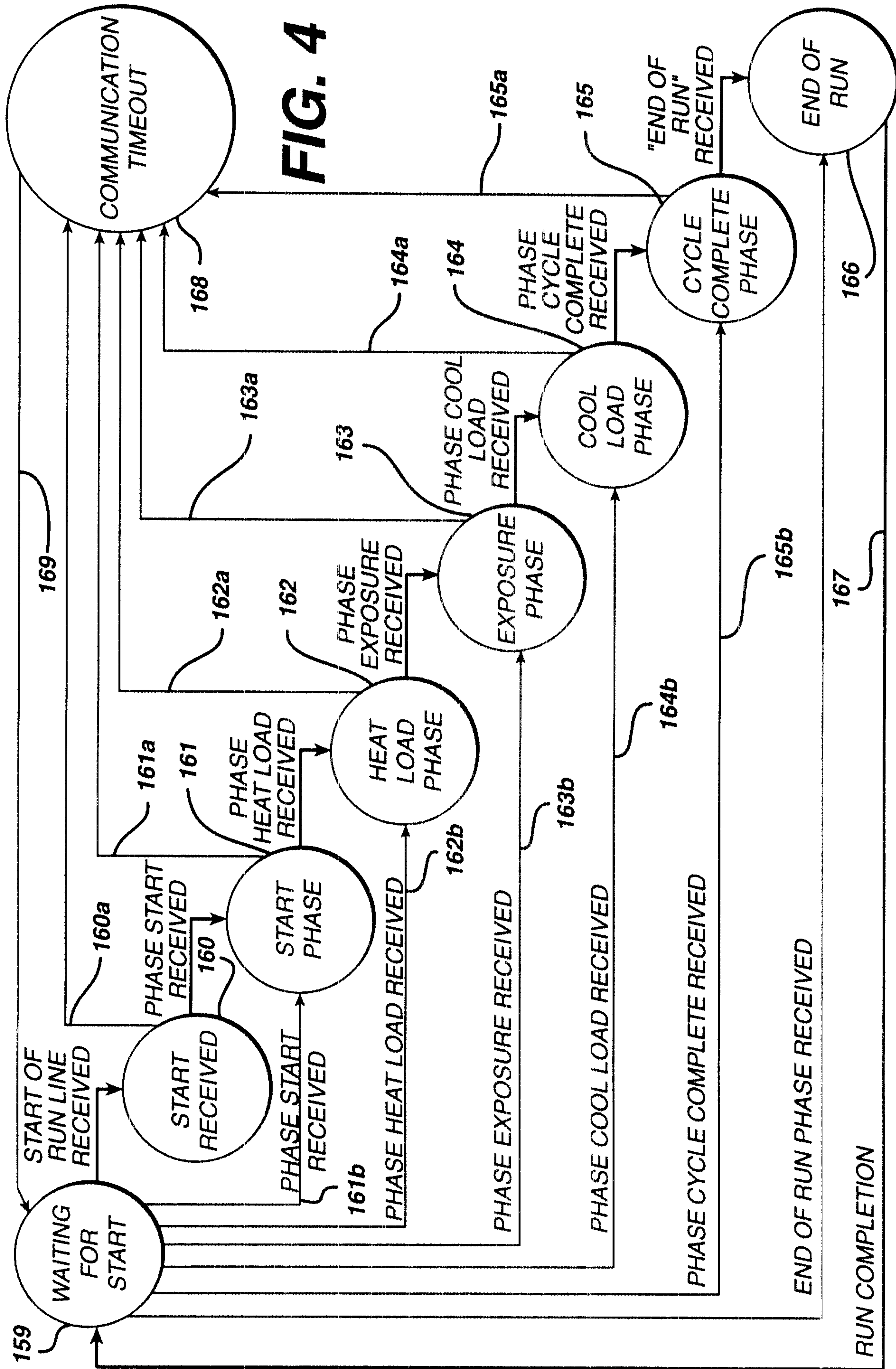
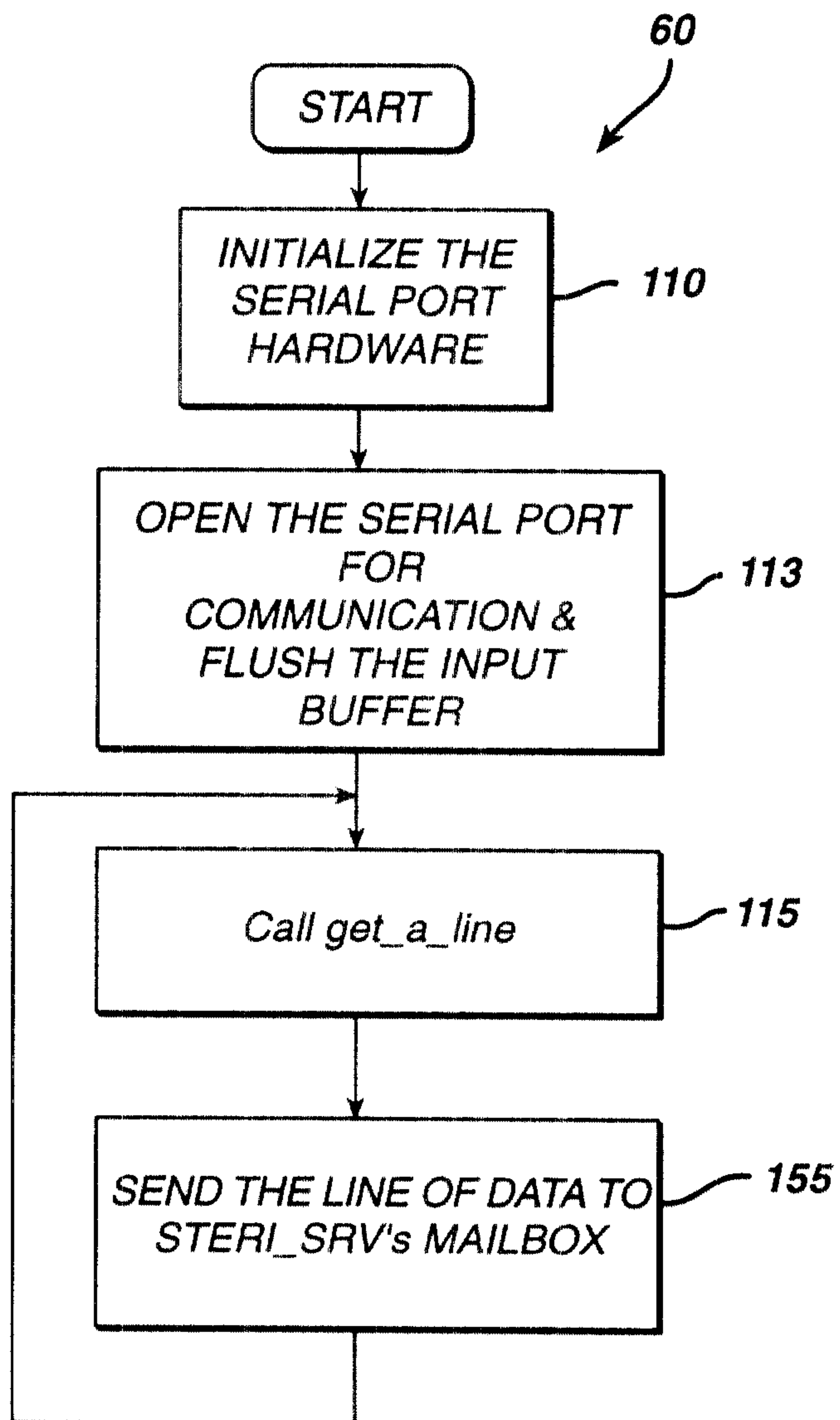


FIG.3

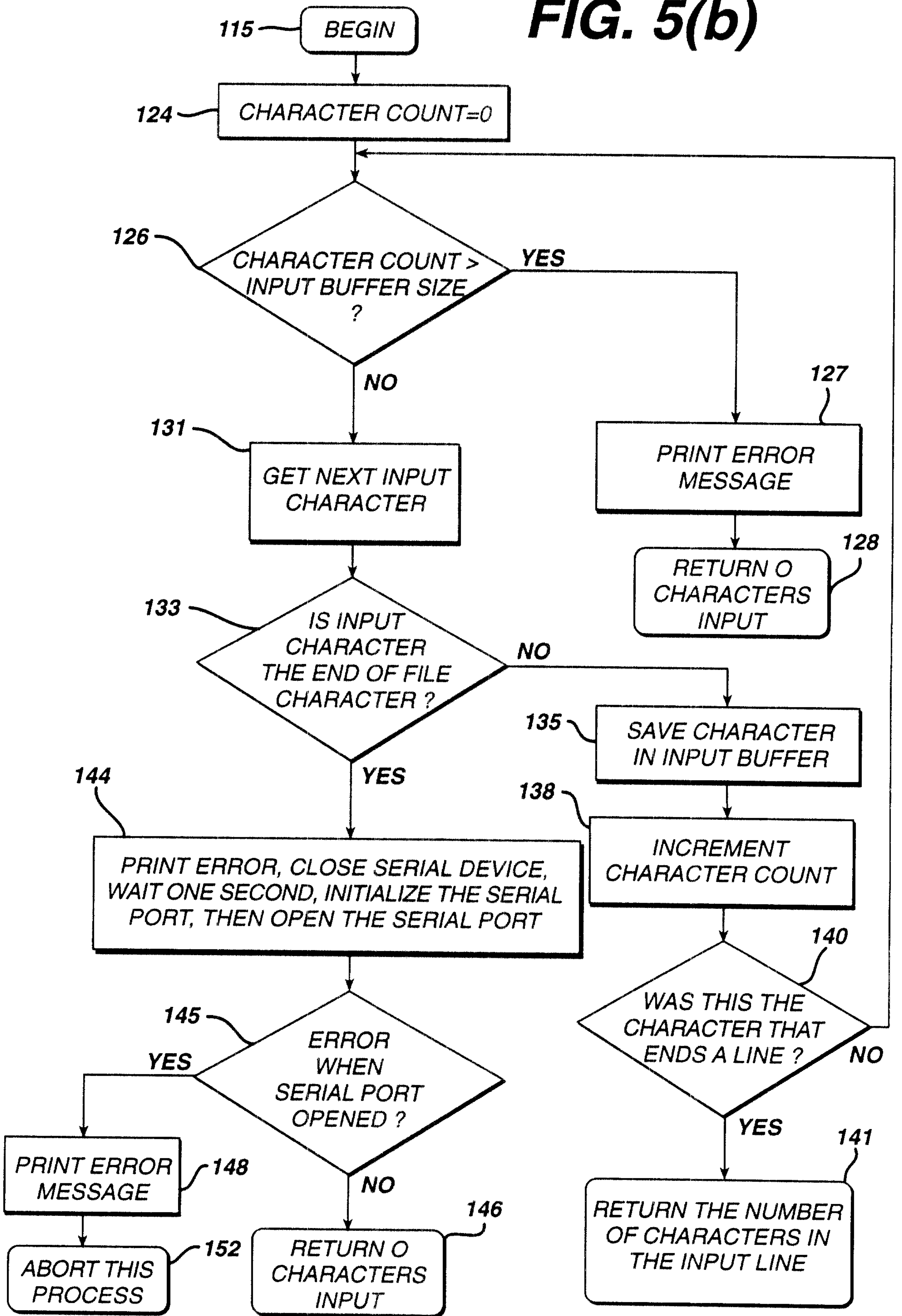




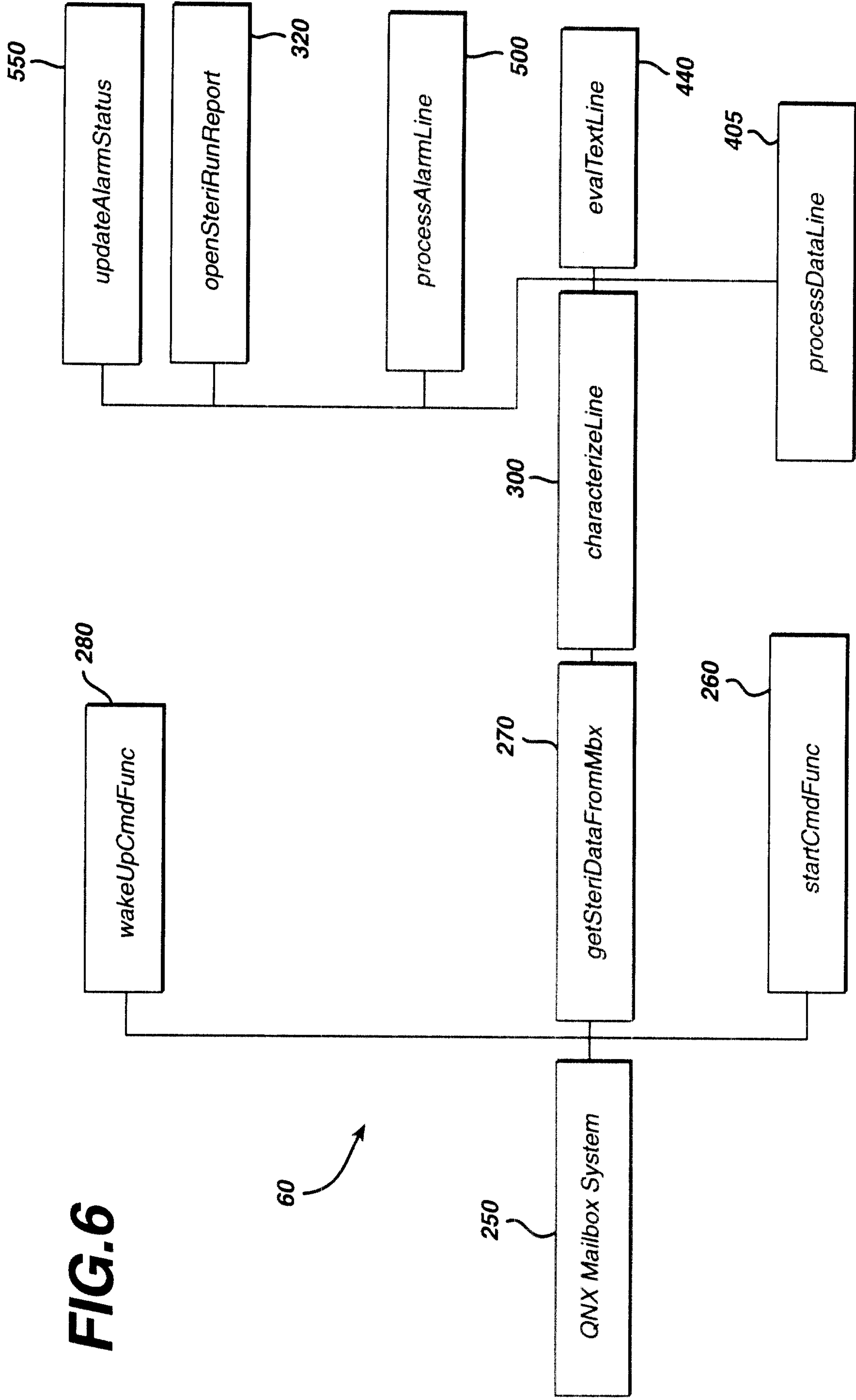
**FIG. 5(a)**

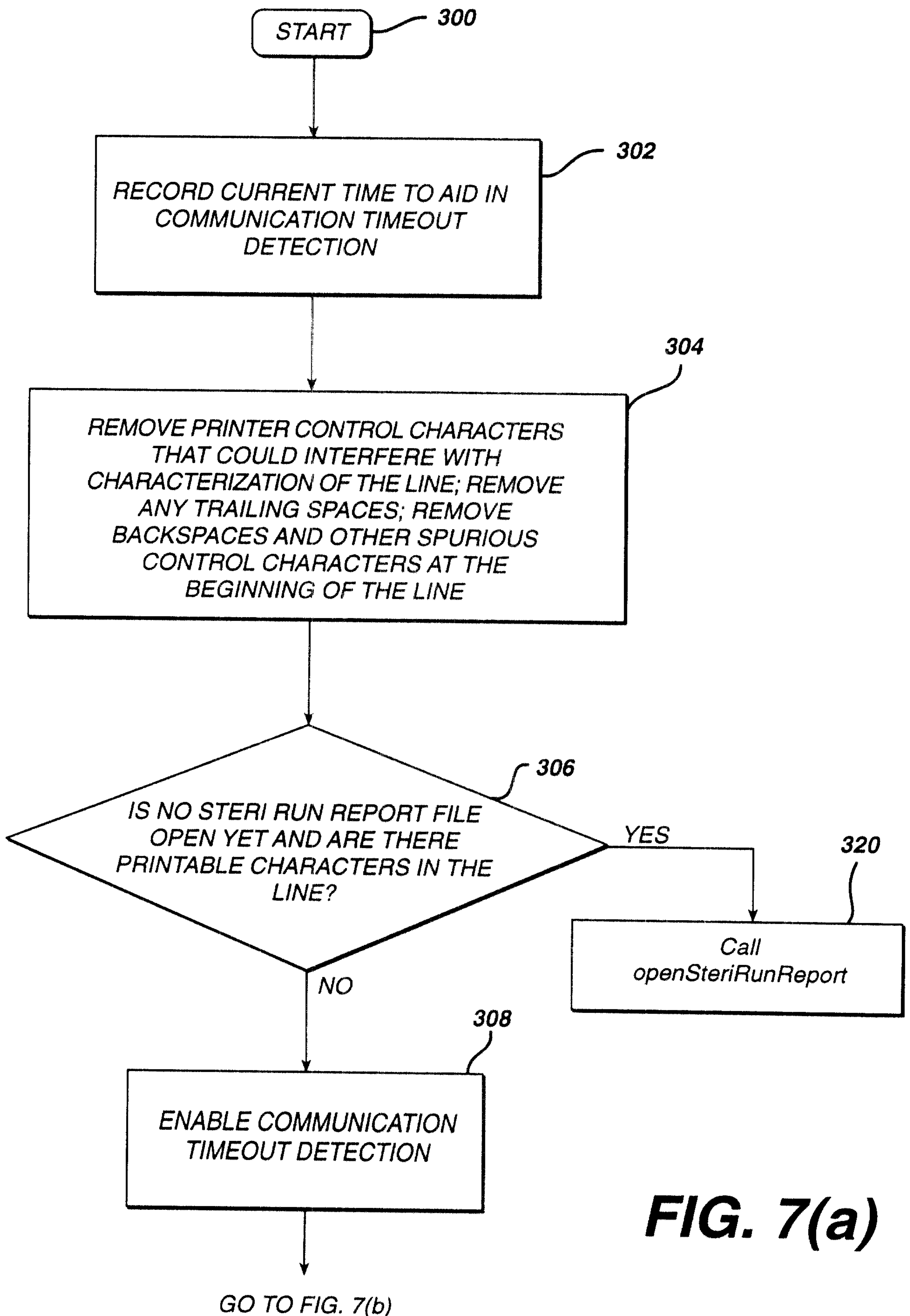


**FIG. 5(b)**



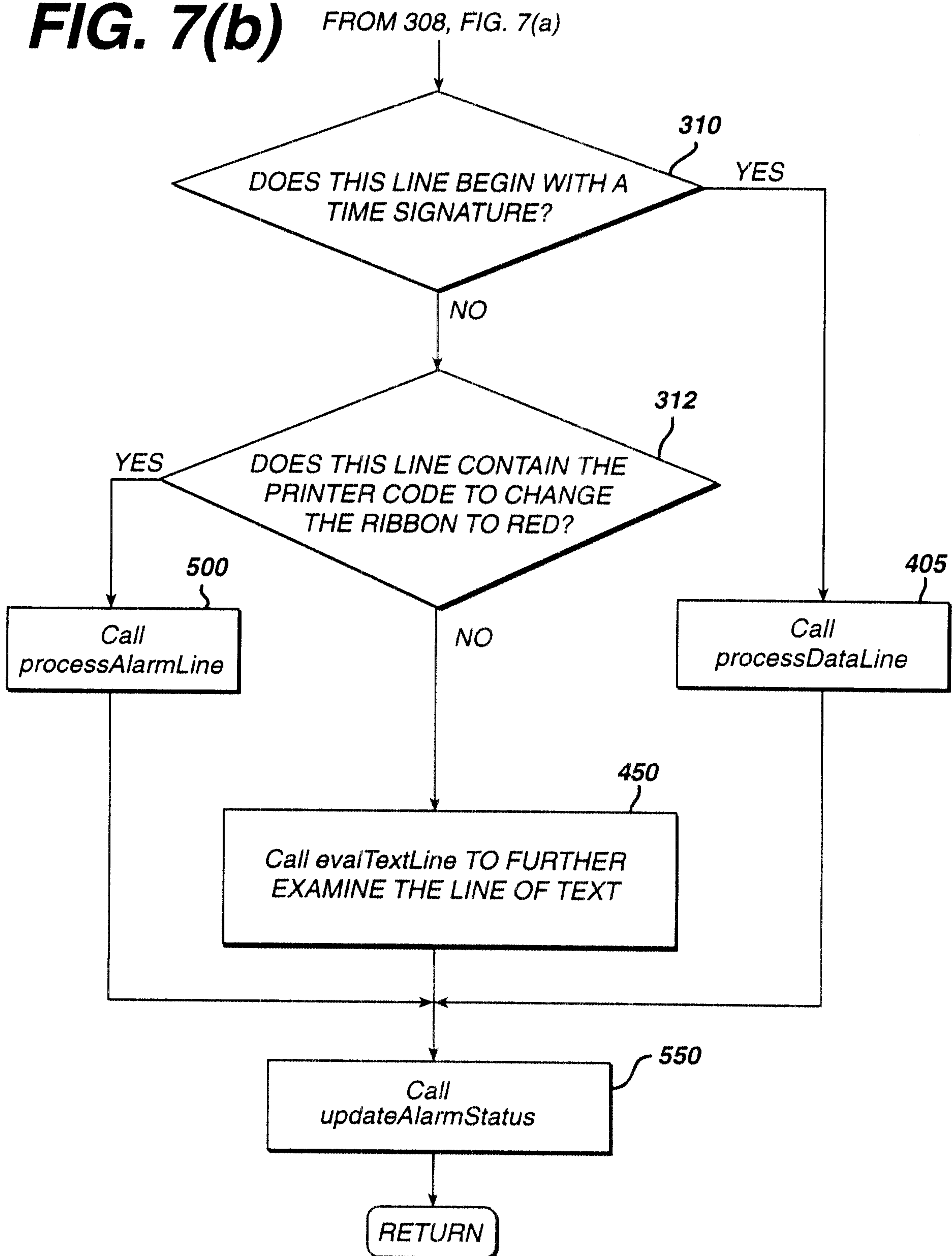
**FIG. 6**

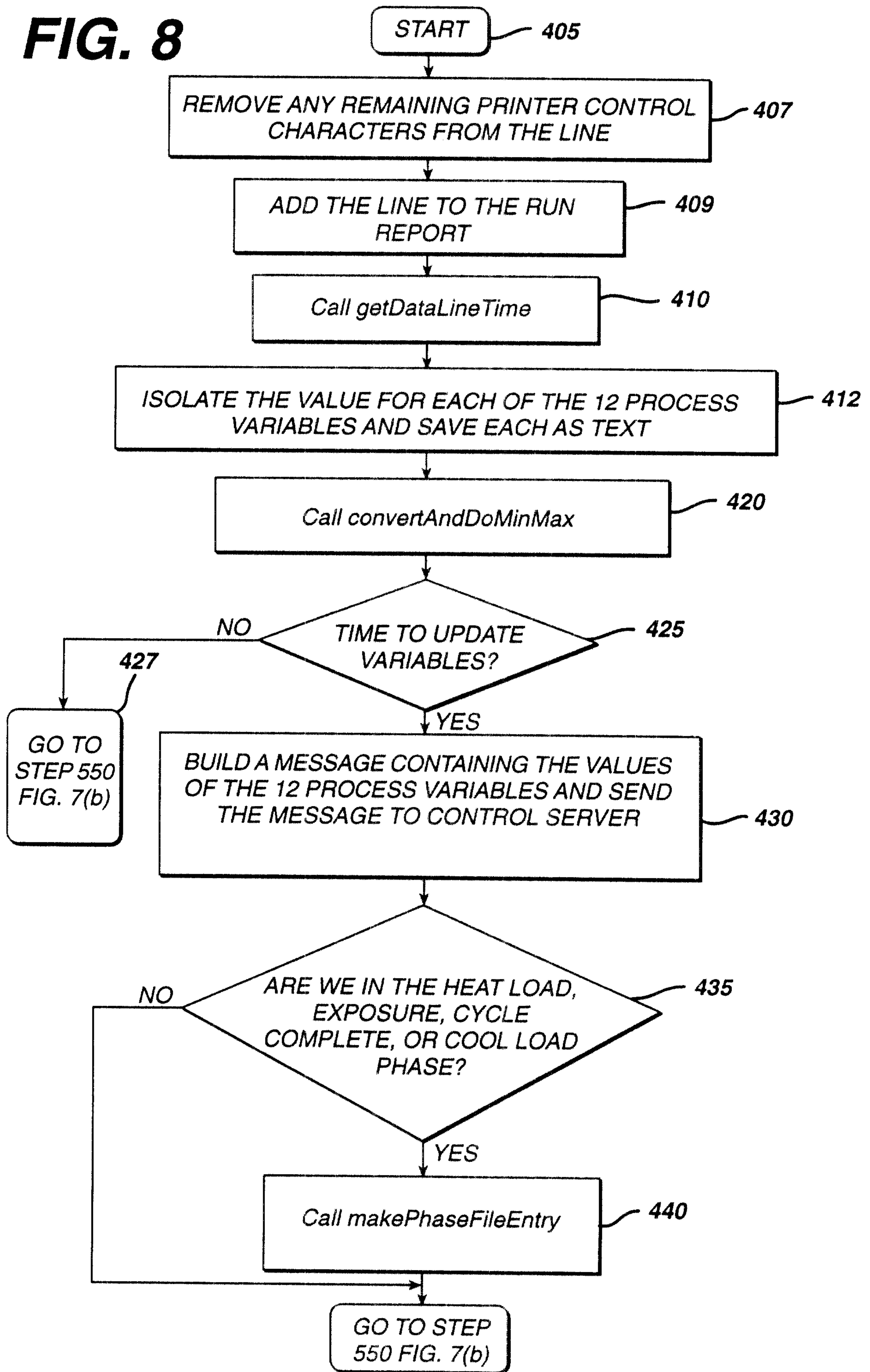


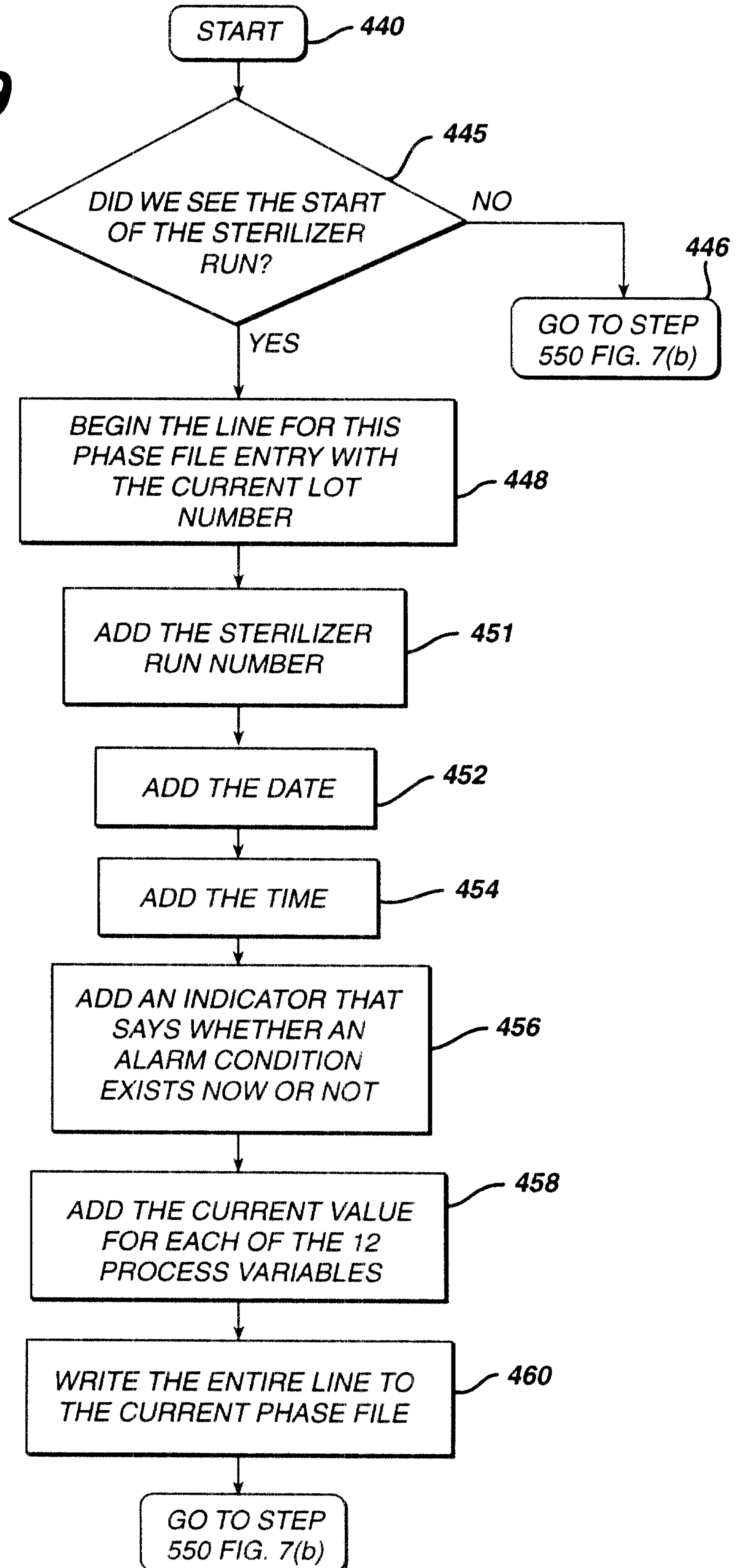


**FIG. 7(a)**

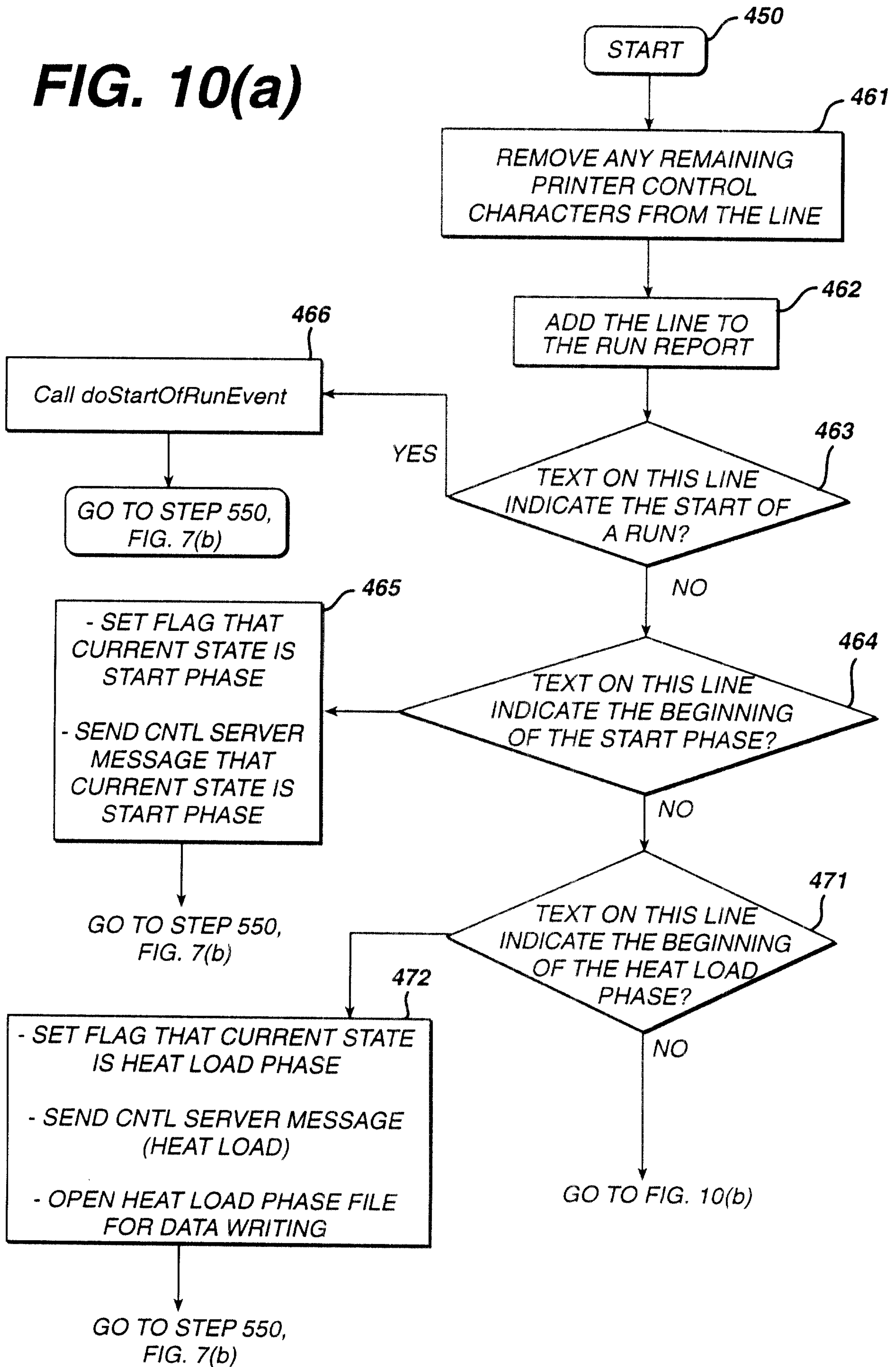


**FIG. 7(b)**

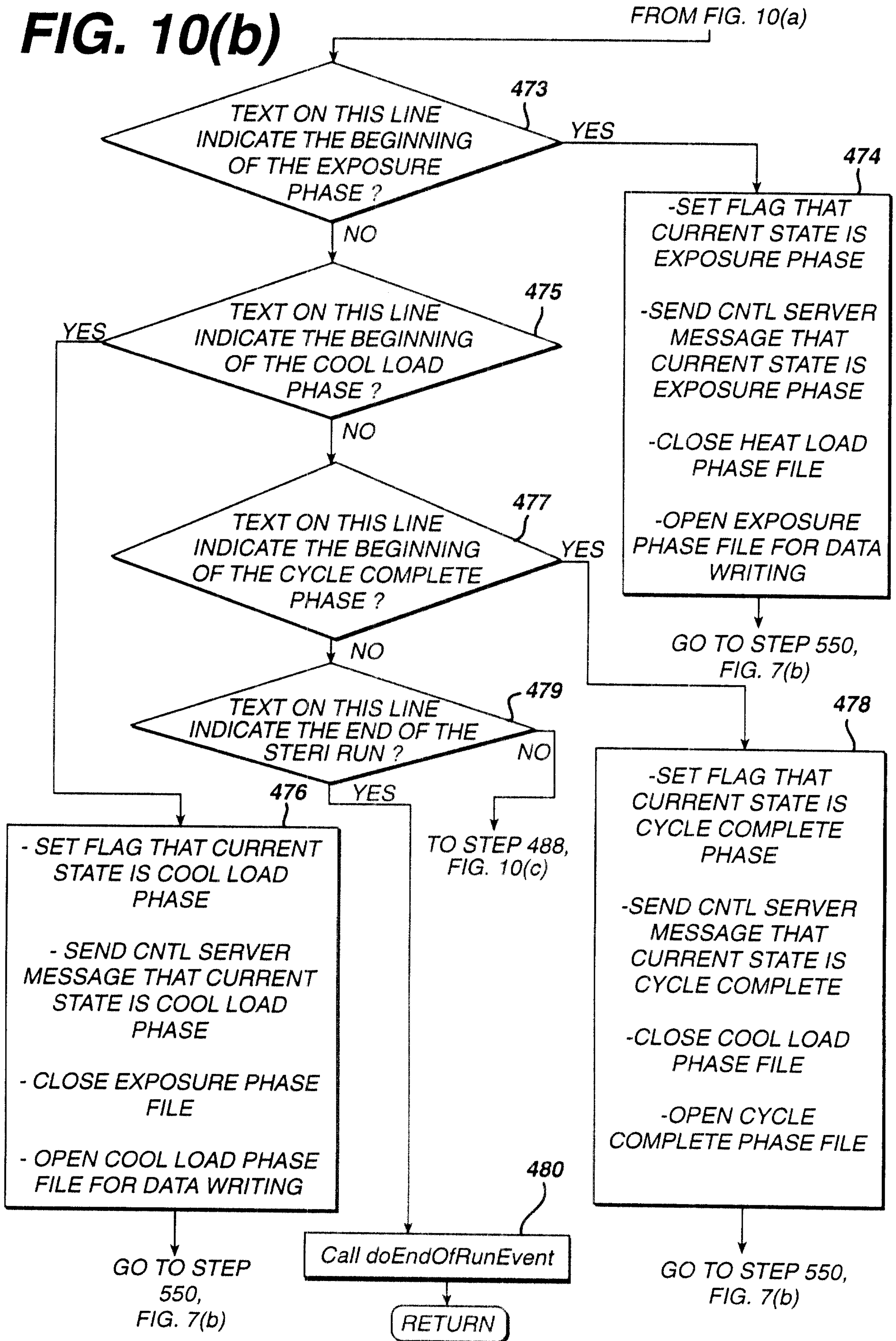
**FIG. 8**

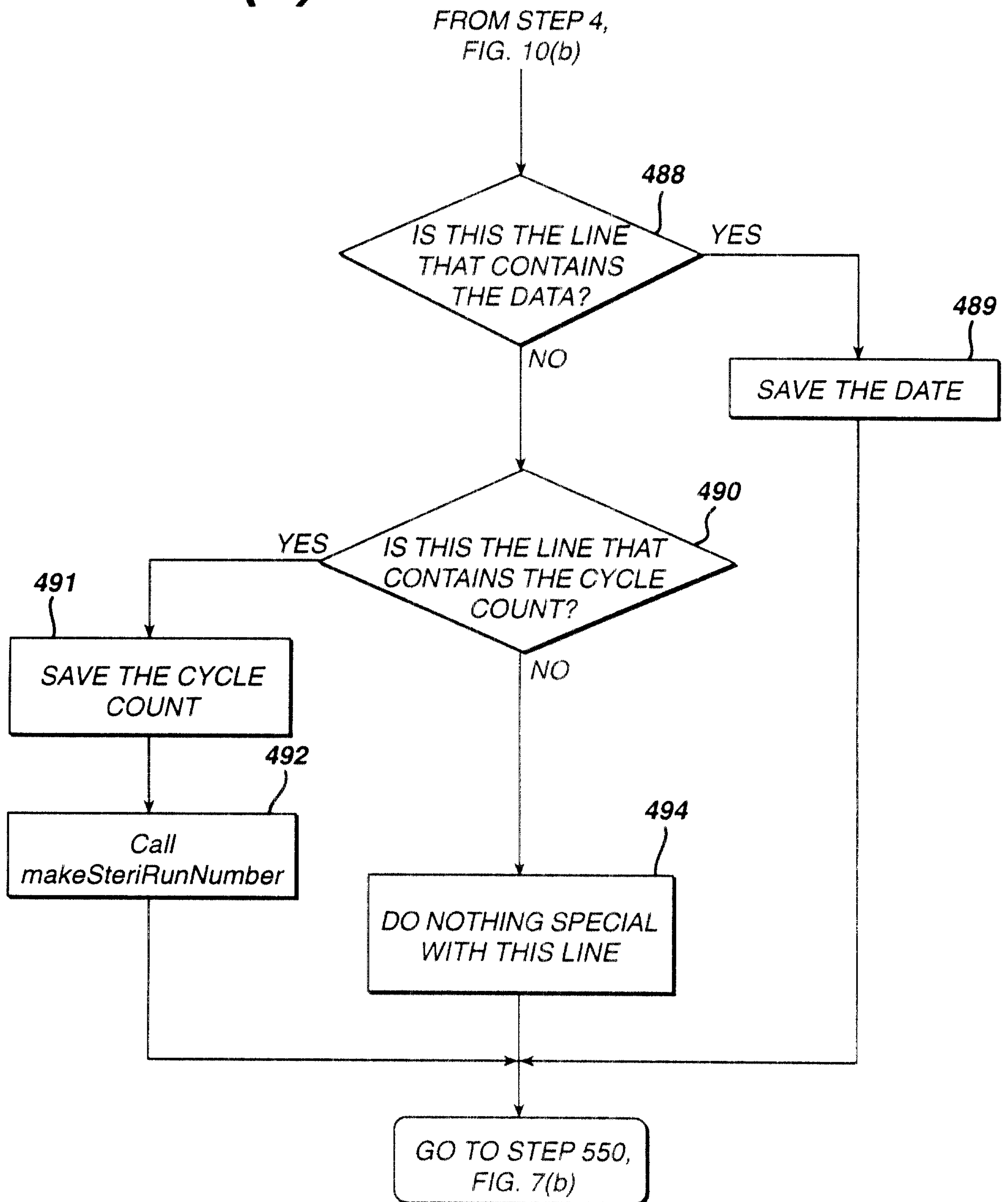
**FIG. 9**

**FIG. 10(a)**

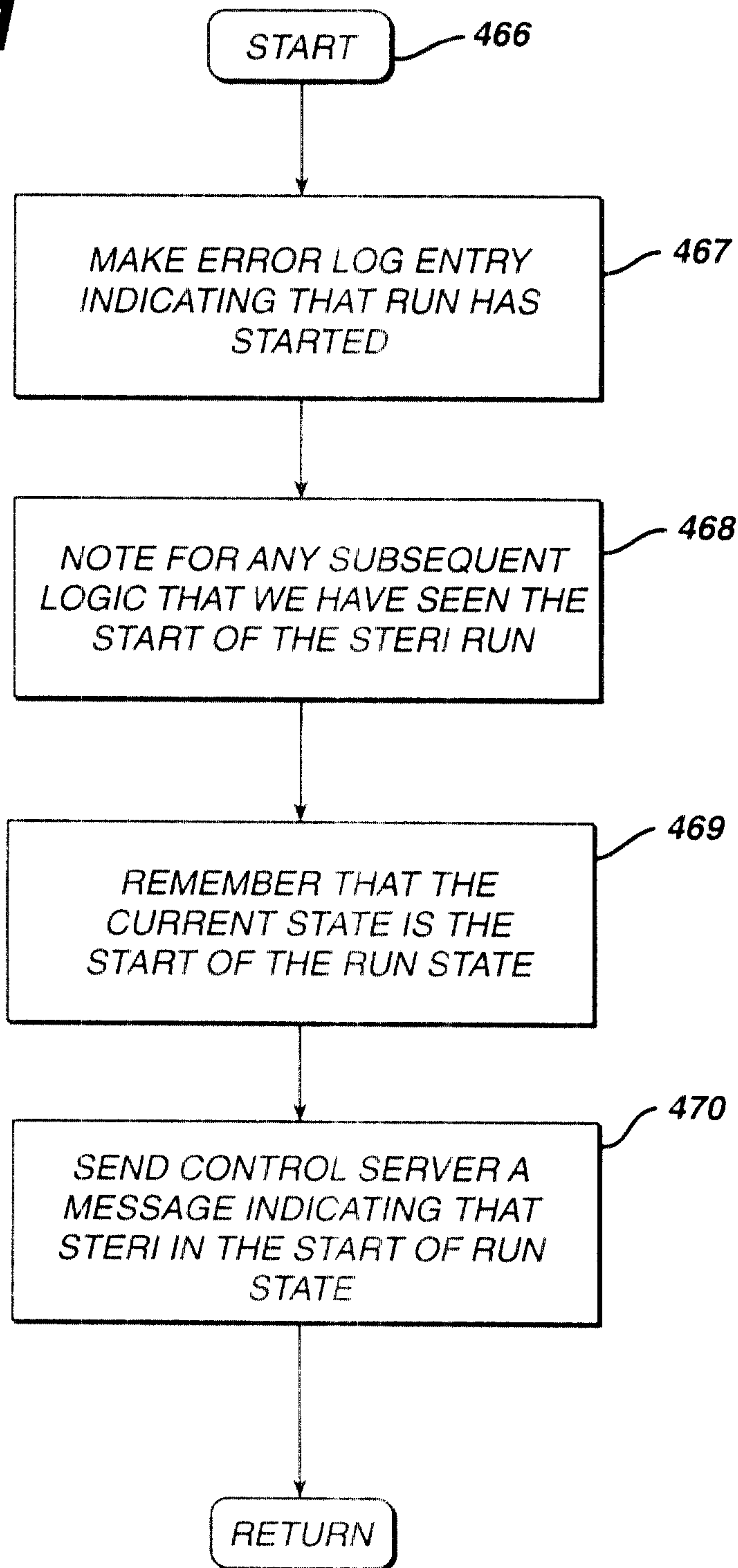


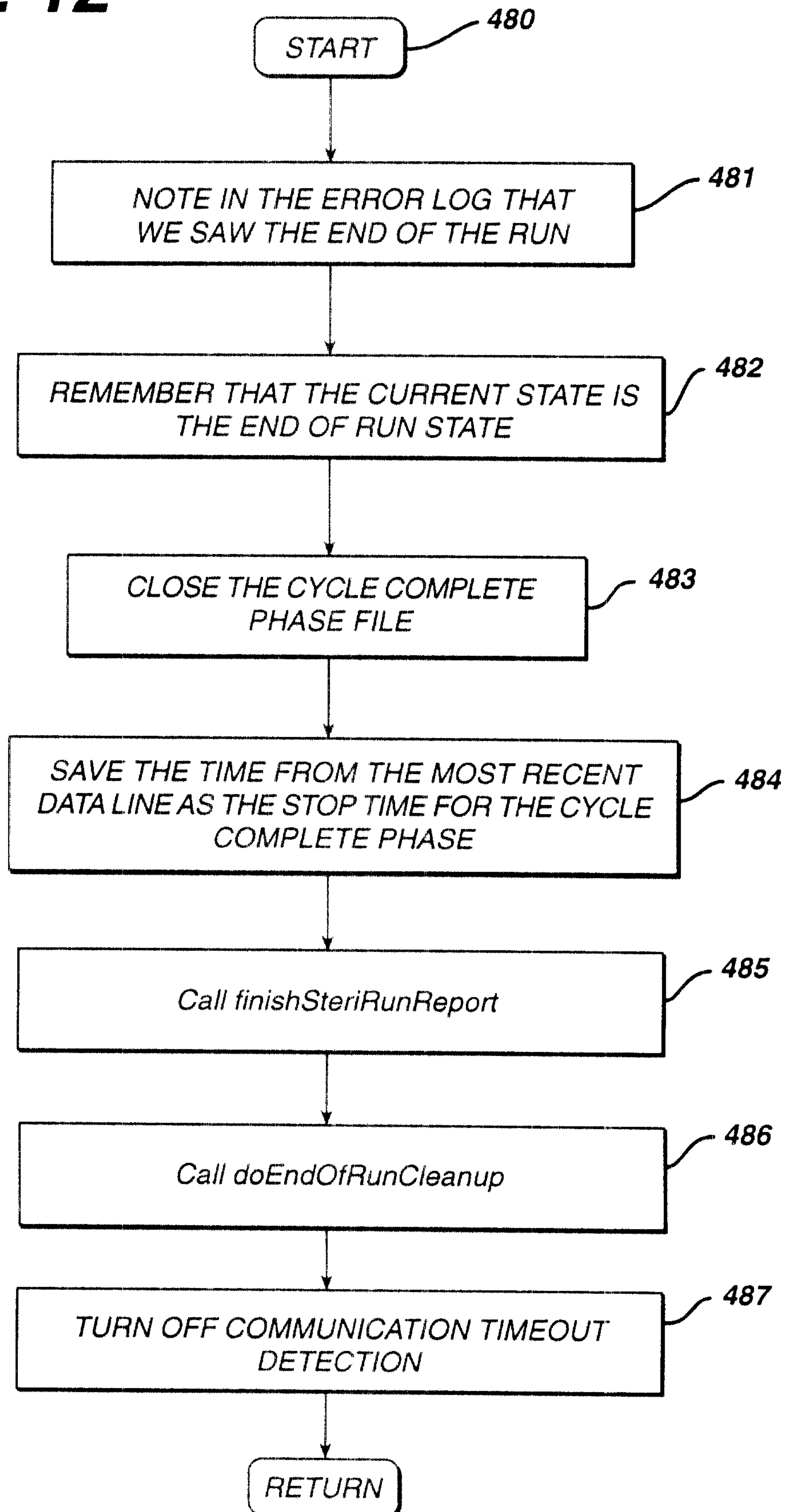
**FIG. 10(b)**



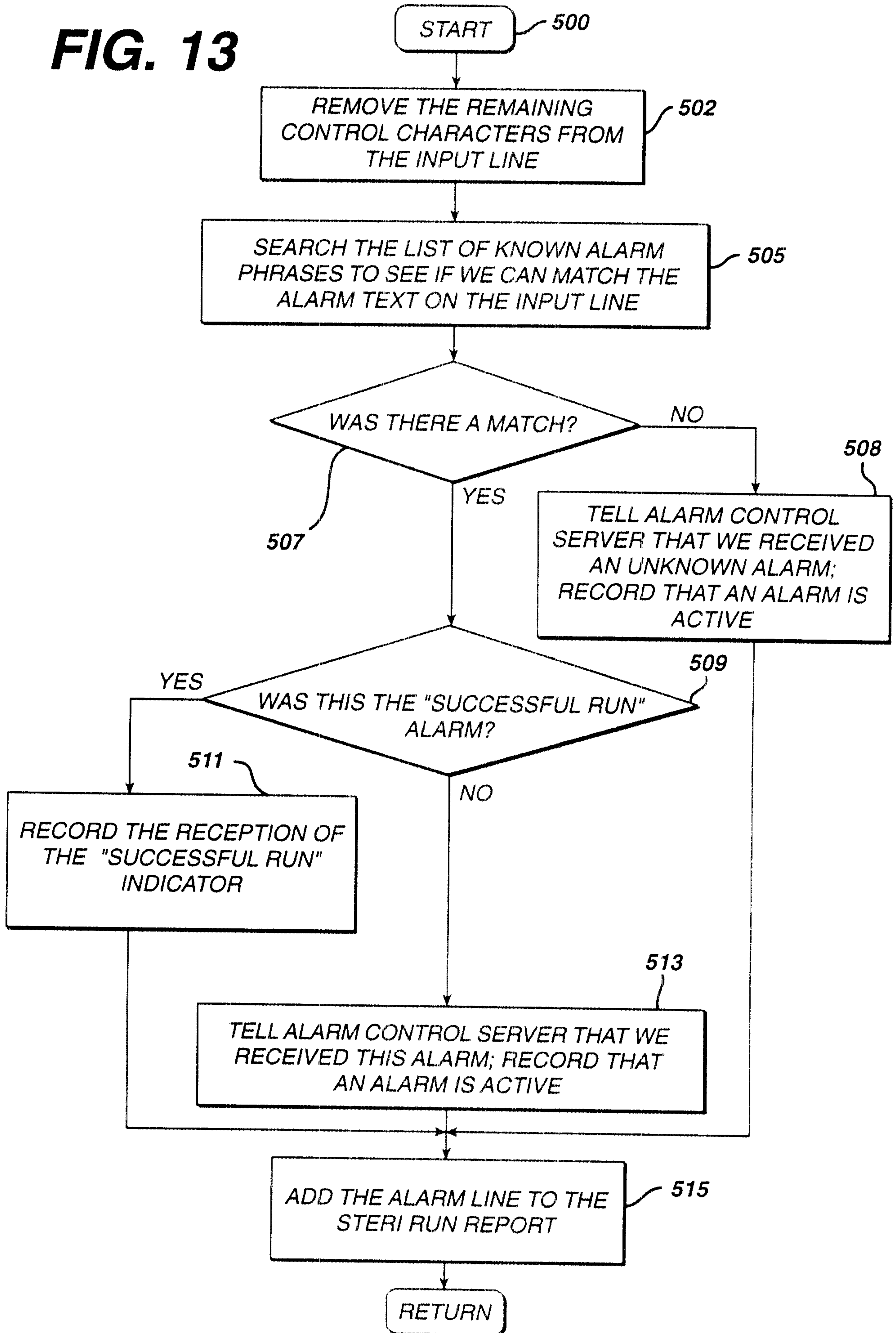
**FIG. 10(c)**

**FIG. 11**

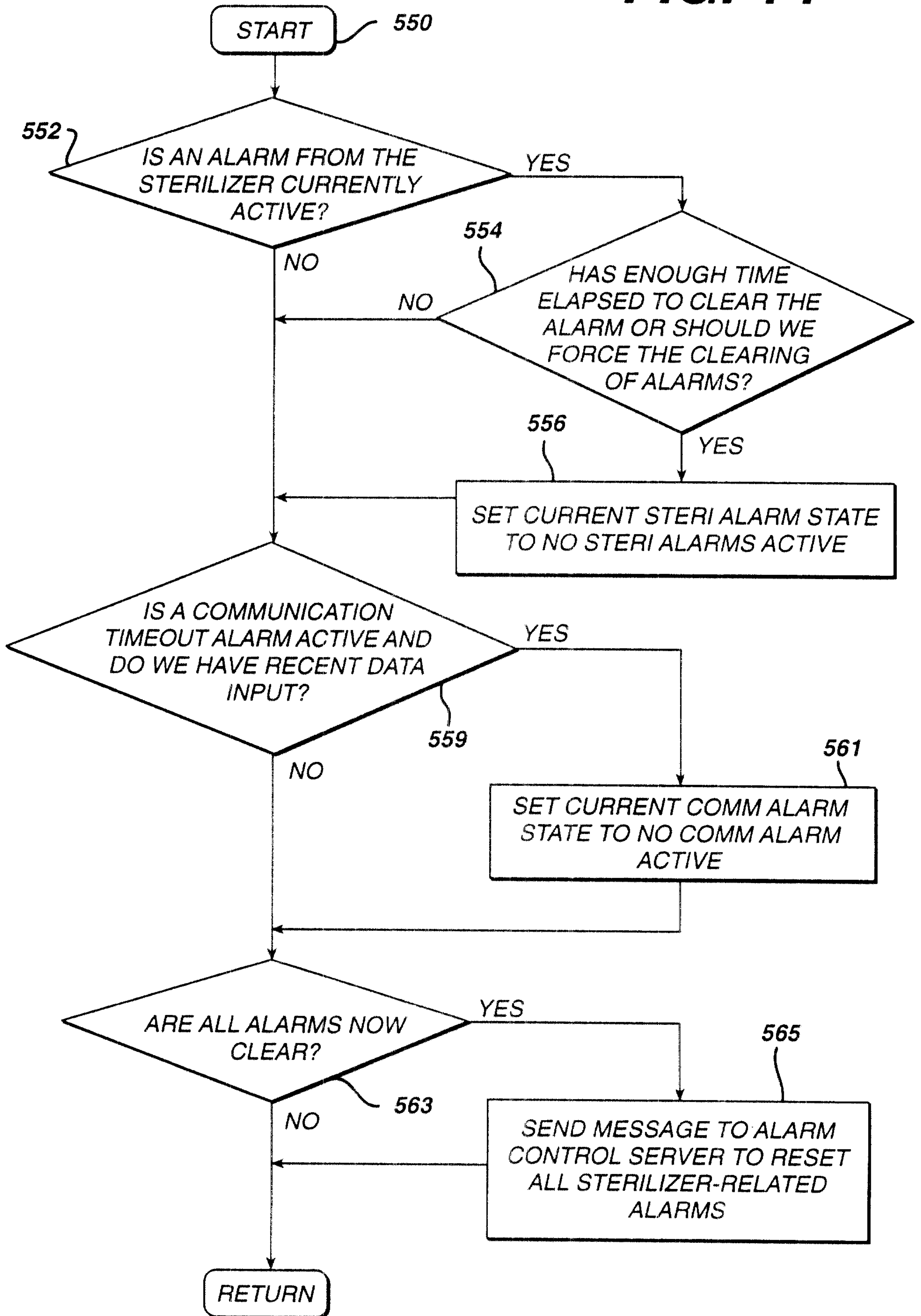


**FIG. 12**

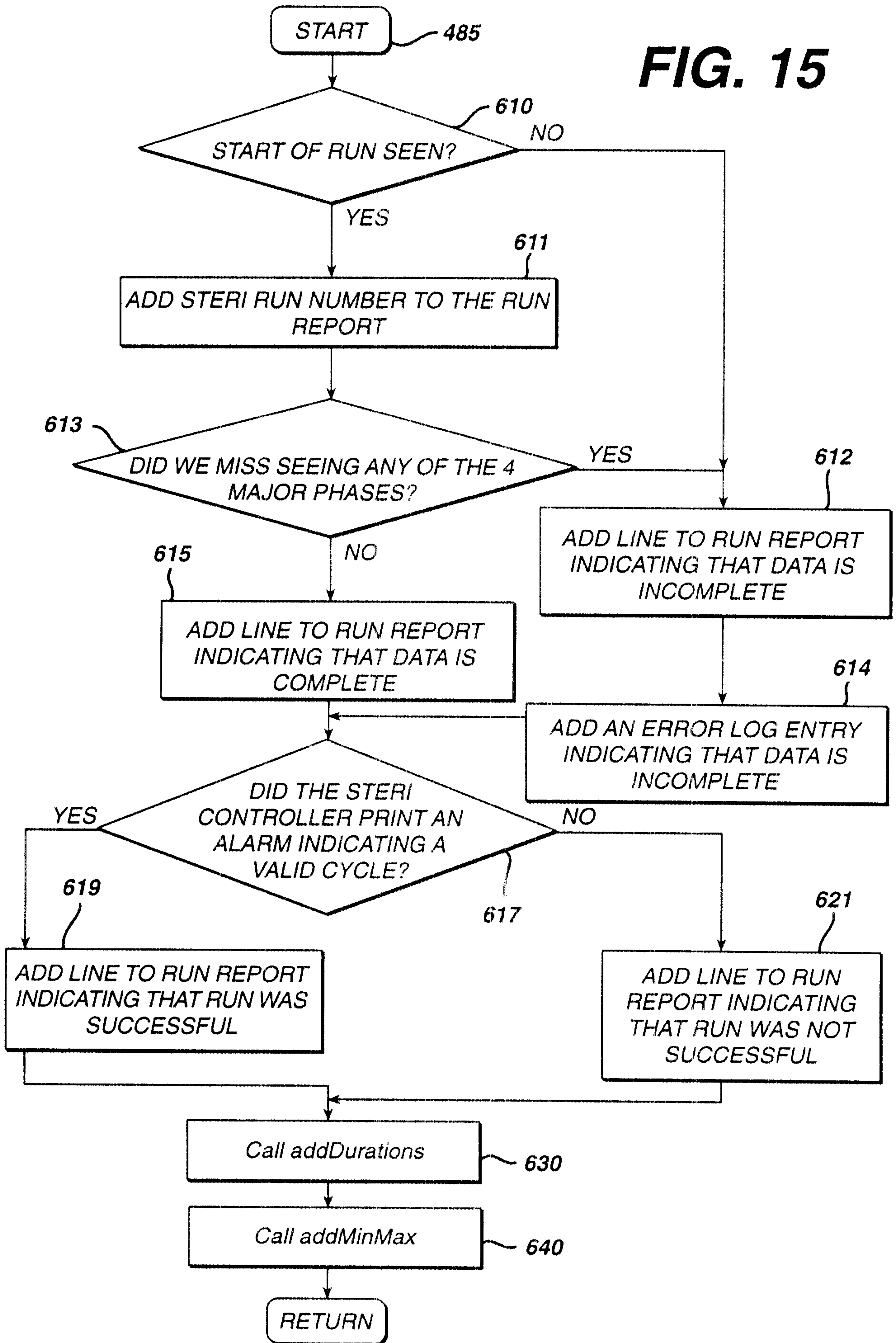


**FIG. 13**

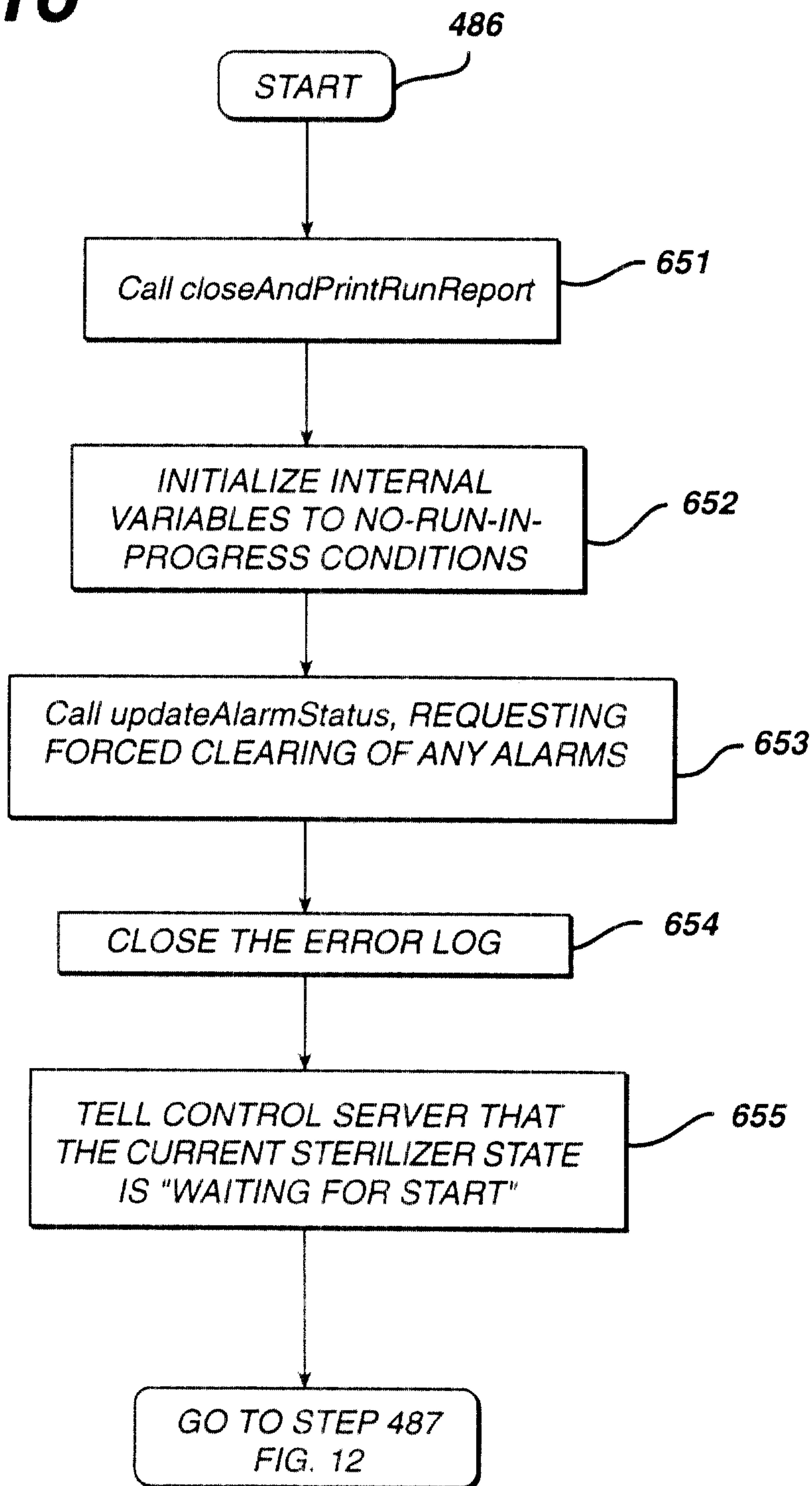
# FIG. 14

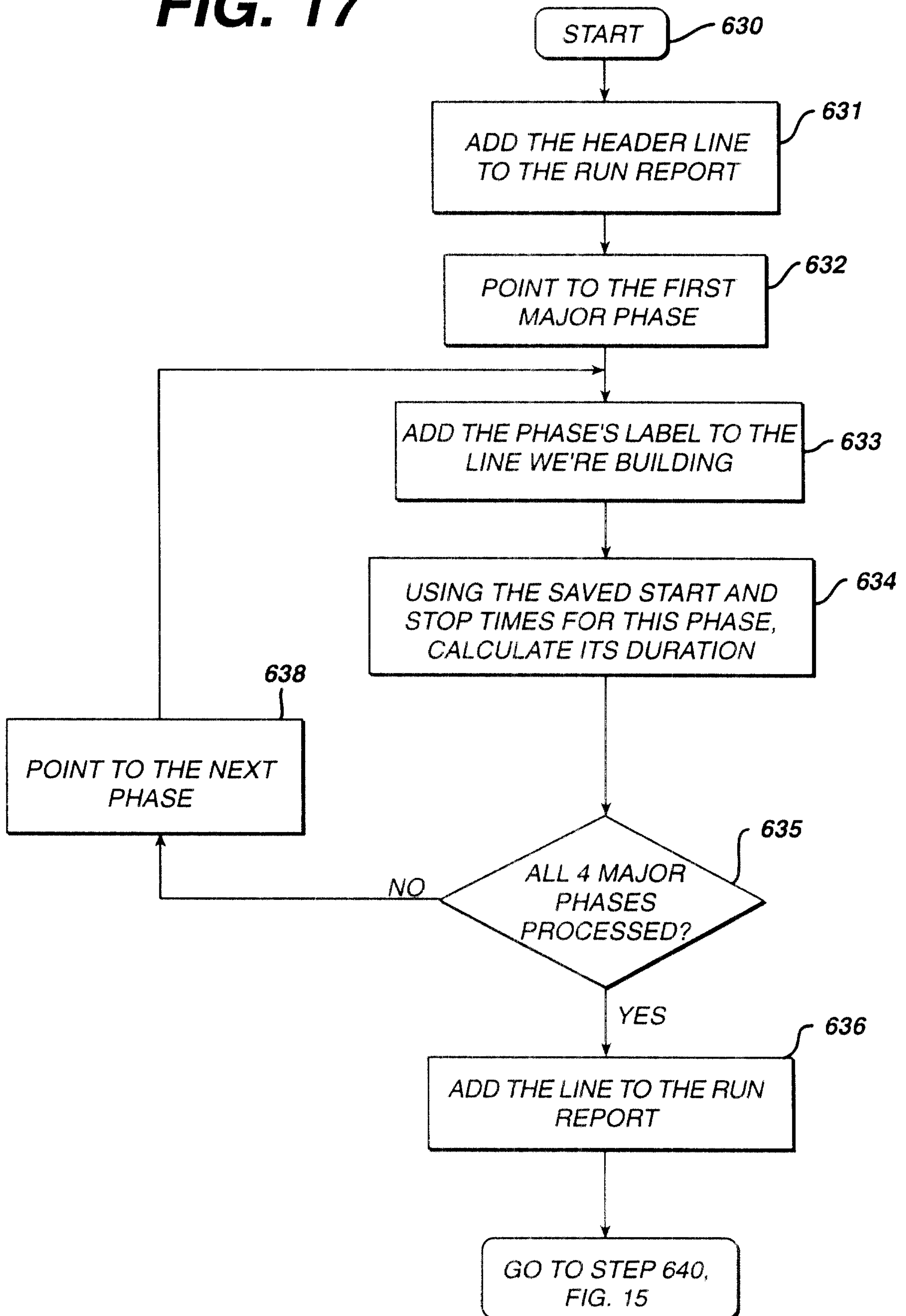


**FIG. 15**

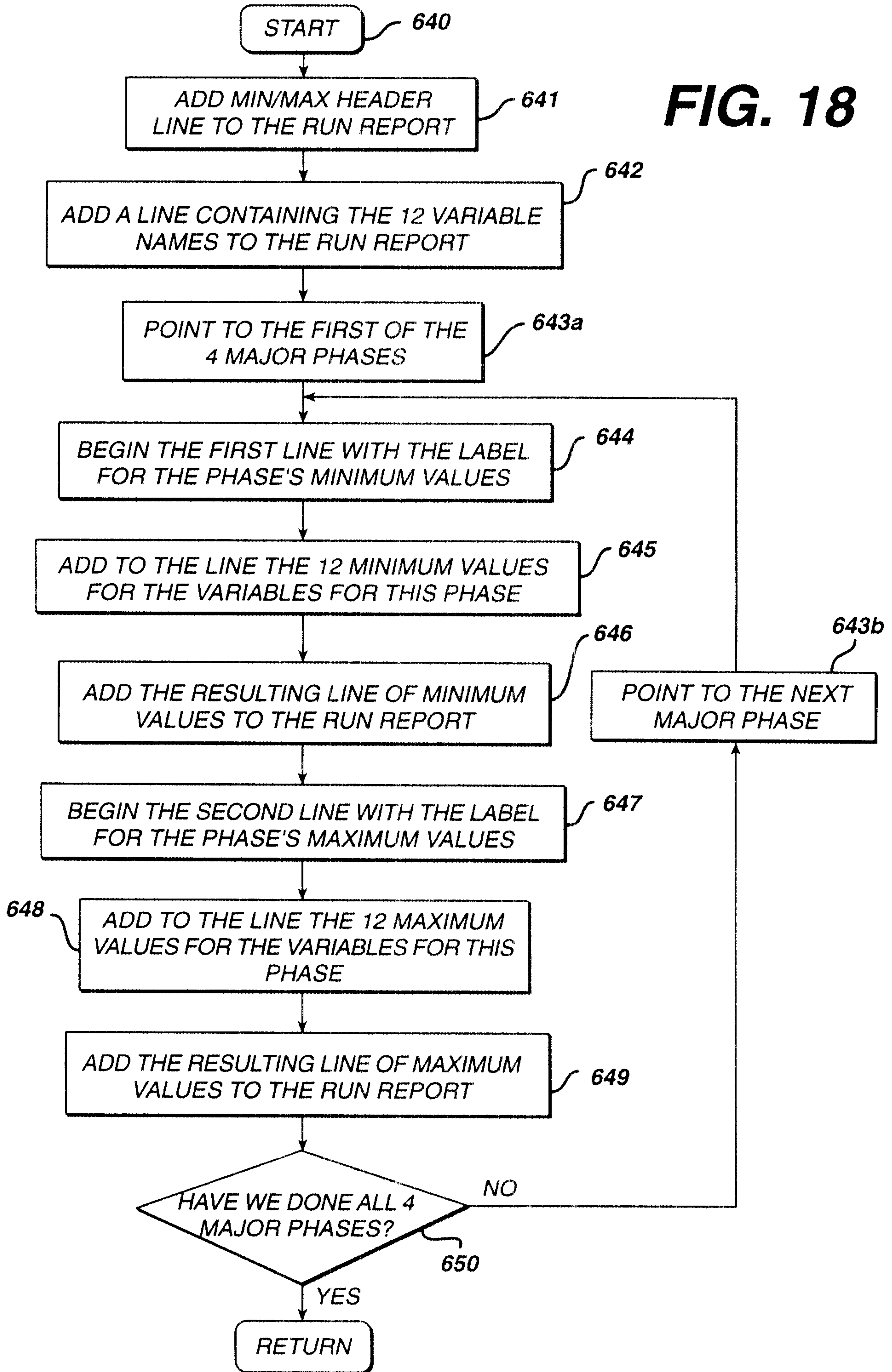


**FIG. 16**

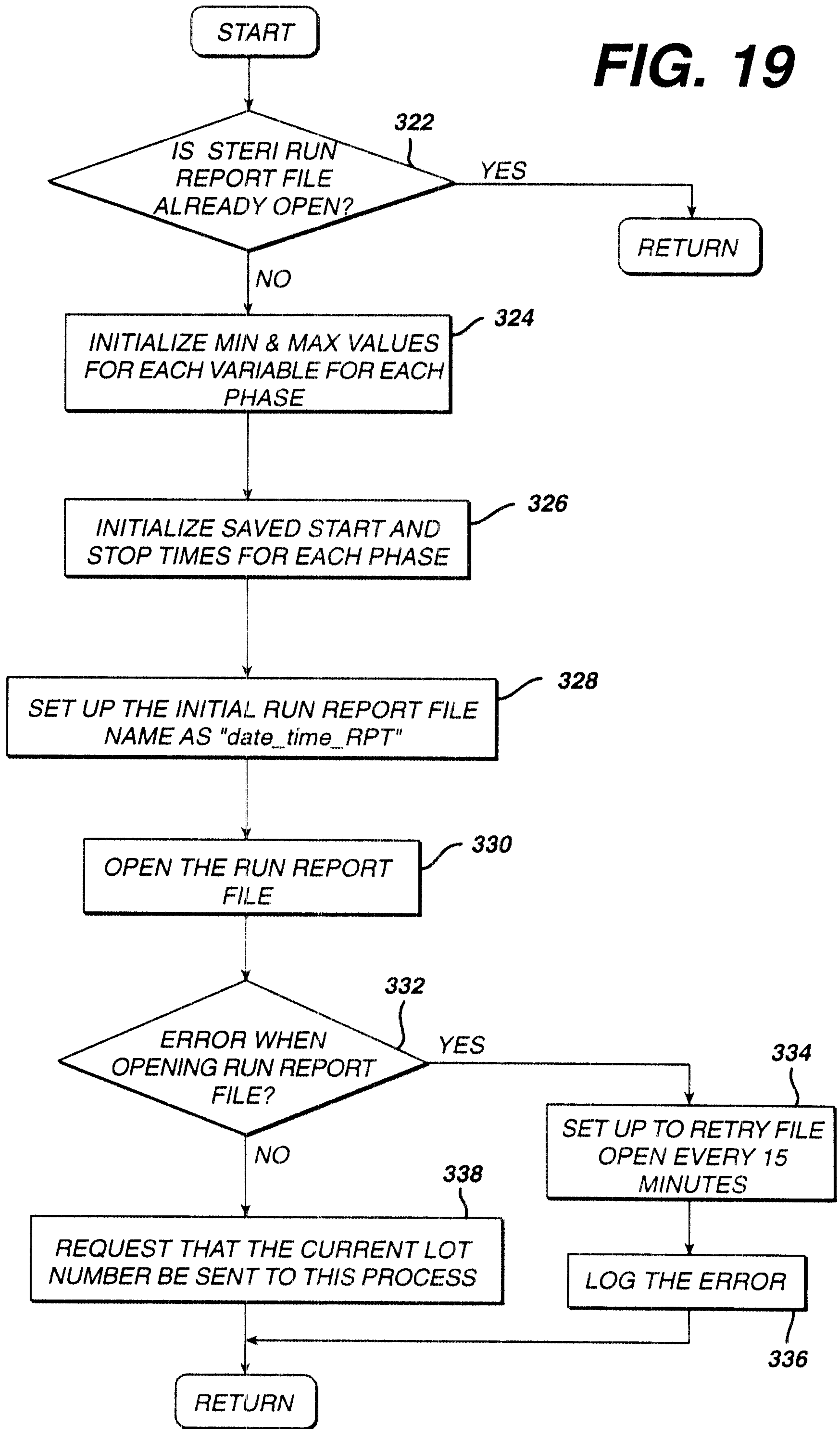


**FIG. 17**

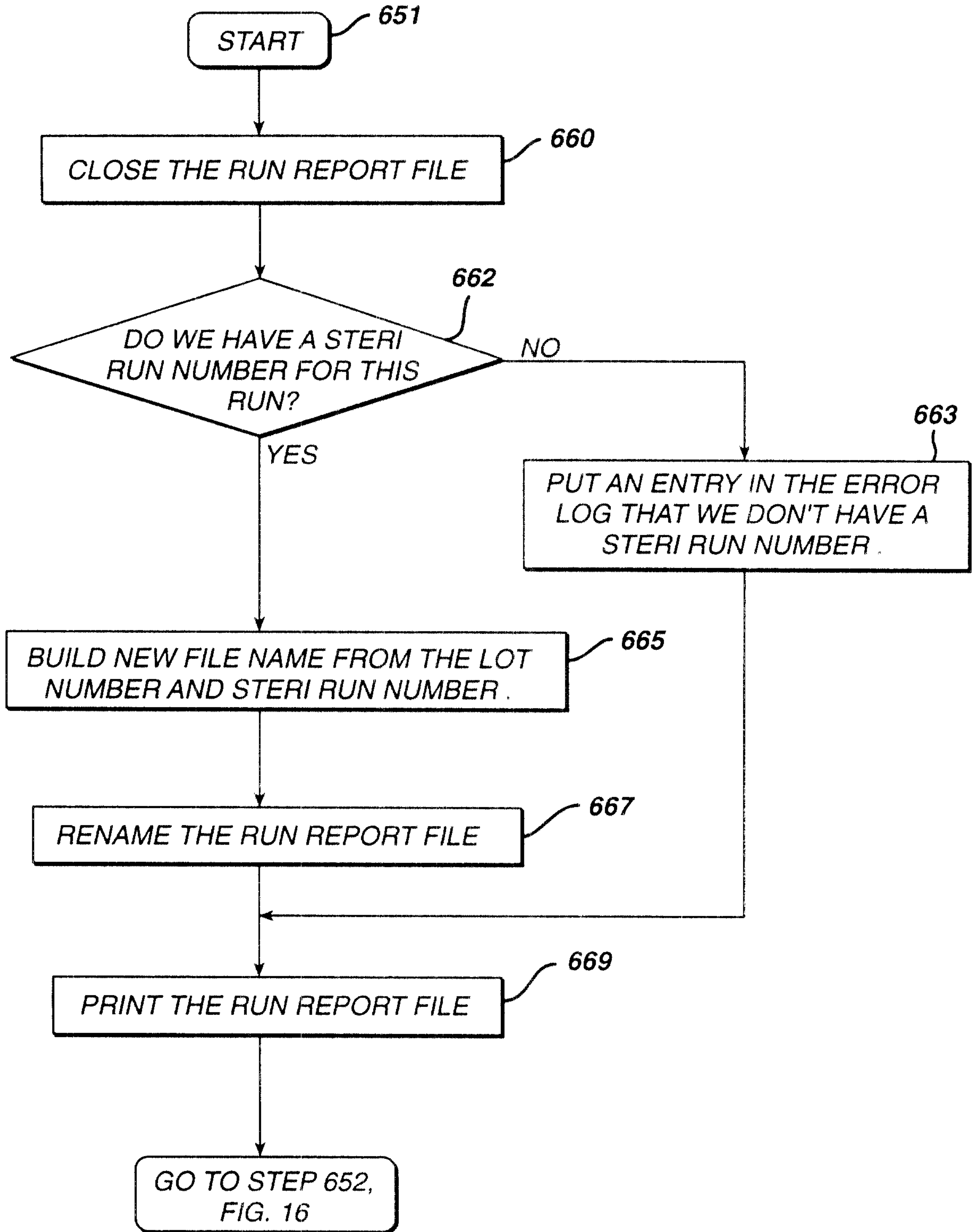
**FIG. 18**



**FIG. 19**

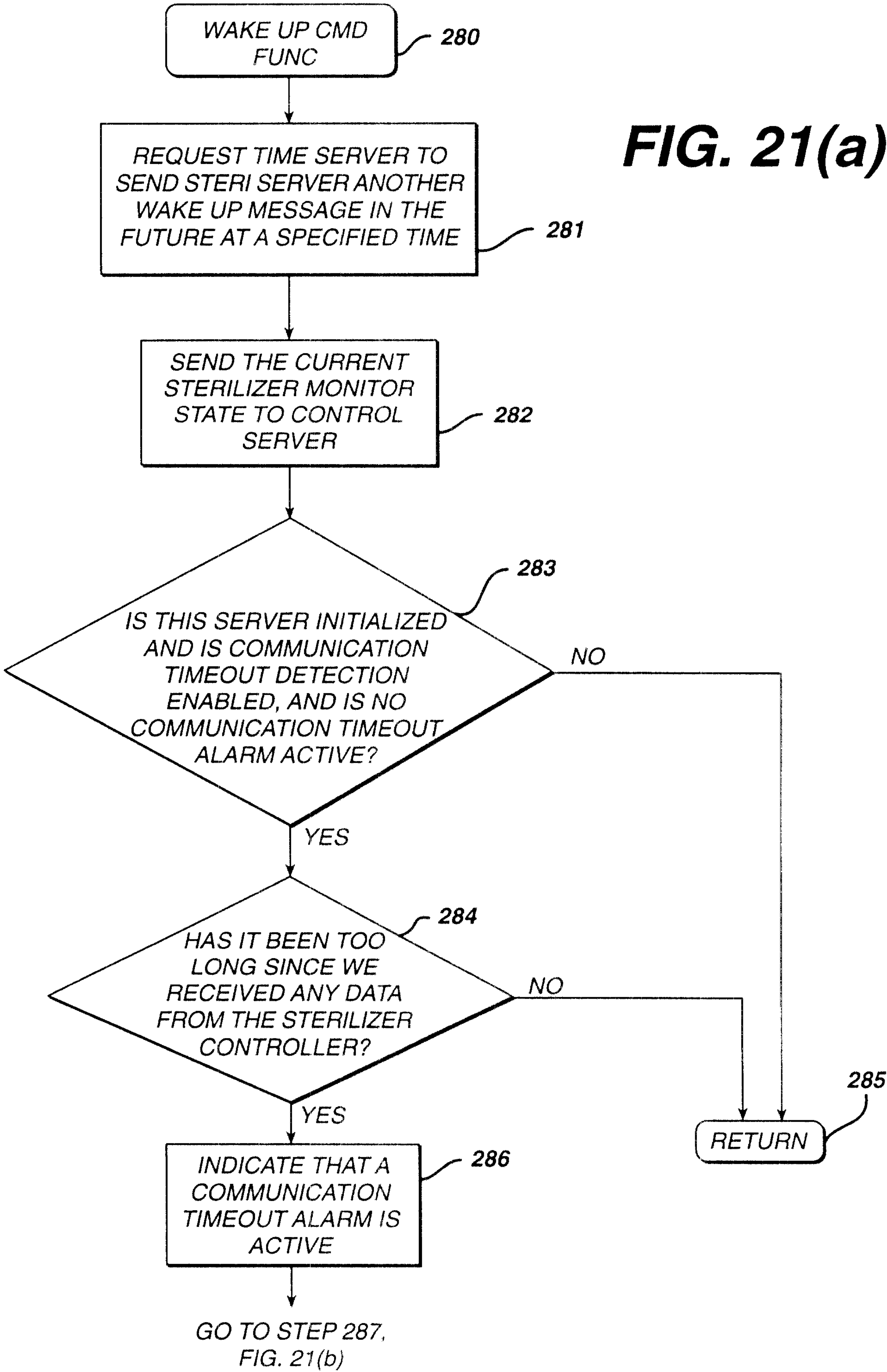


**FIG. 20**

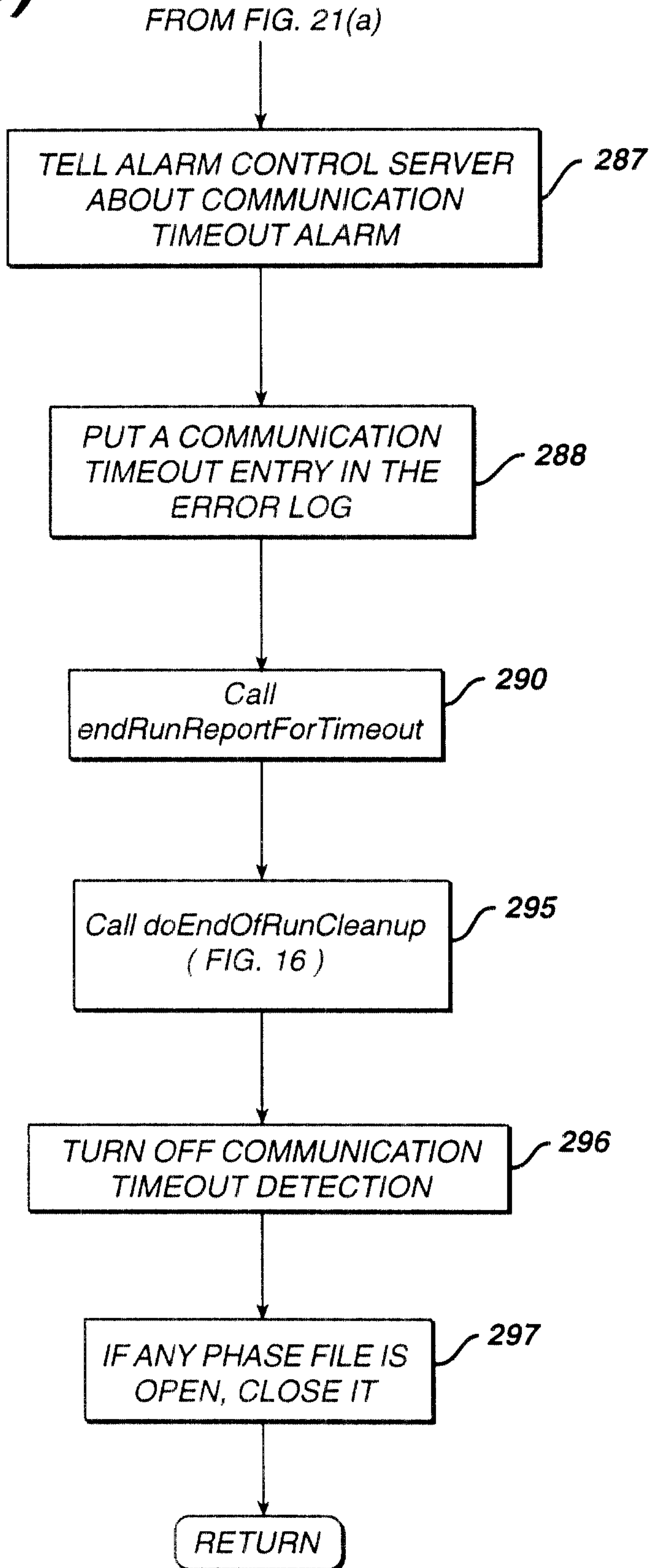




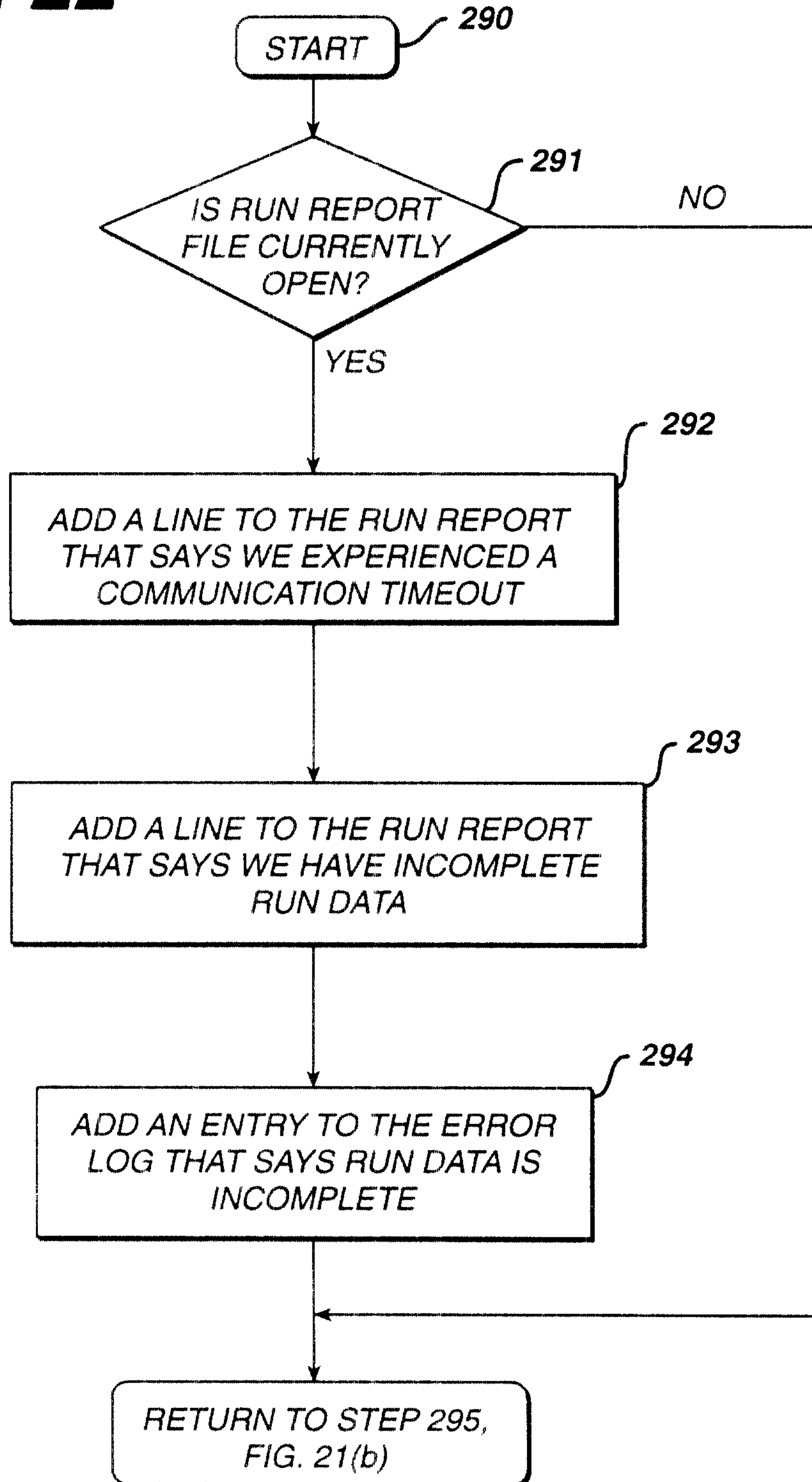
**FIG. 21(a)**



**FIG. 21(b)**



**FIG. 22**



# FIG. 23(a)

STERILIZATION REPORT NUMBER

800

816a	DATE	:	_____
816b	PROCESS START	:	_____
817a	AUTOCLAVE NAME	:	_____
817b	AUTOCLAVE NUMBER	:	_____
818	CYCLE COUNTER	:	_____
	PARAMETERS	:	_____
	FAN SPEED HIGH	:	_____
	OVERPRESSURE	:	_____
	ALARM VAR HIGH	:	_____
	ALARM VAR LOW	:	_____
	TEMP VAR DELAY	:	_____
	FAN SPEED ALM	:	_____
	GRAV DIS TEMP	:	_____
	EVACUATE SLOW	:	_____
	STEAM REG-DRAIN	:	_____
	EXPOSE TIMER	:	_____
	RADIATOR COOLING	:	_____
	UNDER PRESS ALM	:	_____
	OVER PRESS ALM	:	_____

815

819

SIGNALS

1	:	_____
2	:	_____
3	:	_____
4	:	_____
5	:	_____
6	:	_____
7	:	_____
8	:	_____
9	:	_____
10	:	_____
11	:	_____
12	:	_____

TO FIG. 23(b)

# FIG. 23(b)

FROM FIG. 23(a)

800

PROGRAM: 800

PROGTIME	V1	V2	V3	V4	V5	V6	V7	V8	V9	V10	V11	V12
START												
00:00:00	X	X	X	X	X	X	X	X	X	X	X	X
HEAT LOAD												
00:00:00	X	X	X	X	X	•	•	•	•			
00:00:49	X	X	X	X	X	•						
00:01:22	X	X	X	X	X	•						
EXPOSURE												
00:05:	X	X	X	X	X	•	•	•	•			
00:05:49	X	X	X	X	X	•						
00:06:49	X	X	X	X	X	•						
COOL LOAD												
00:36:27	X	X	X	X	X	•	•	•	•			
00:36:49	X	X	X	X	X	•						
00:37:49	X	X	X	X	X	•						
CYCLE COMPLETE												
00:46:04	X	X	X	X	X	•	•	•	•			
VALID CYCLE												

825

TO FIG. 23(c)

**FIG. 23(c)**

FROM FIG. 23(b)

800

805

SIGNATURE : ..... 812  
 STERILIZER RUN NUMBER : ..... 855  
 A COMPLETE SET OF DATA WAS OBTAINED FROM THE STERILIZER CONTROLLER.  
 THE STERILIZER CONTROLLER REPORTED A VALID CYCLE. 850

PHASE DURATIONS (M:MM:SS) 830

835

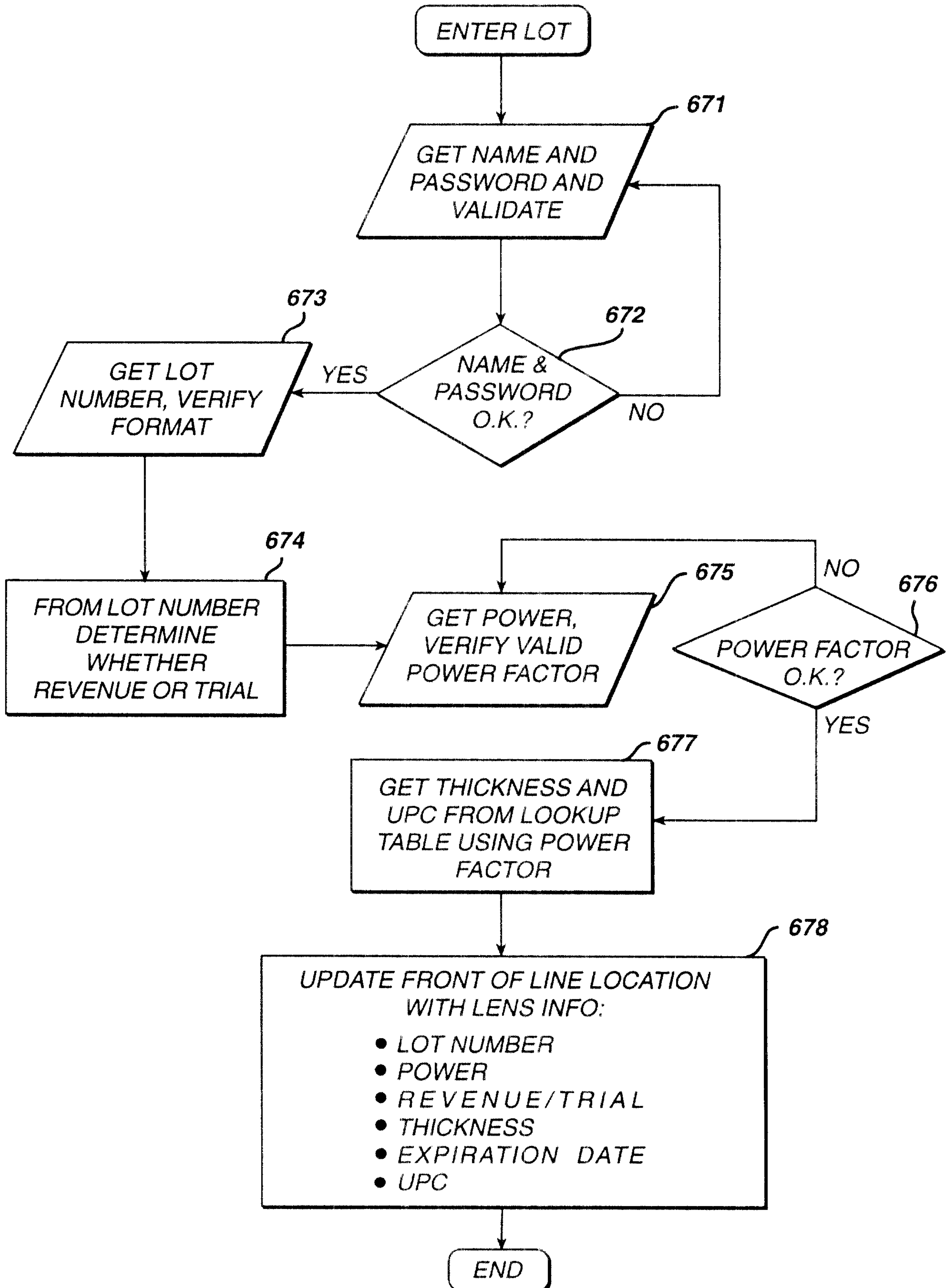
HEAT LOAD qq.rr EXPOSURE ss.tt COOL LOAD uu.vv CYCLE COMPLETE ww.xx

MINIMUM AND MAXIMUM VALUES 840

	1	2	3	4	5	6	7	8	9	10	11	12
HEAT MIN	X	-	-	-	-	-	-	-	-	-	-	-
HEAT MAX	X	-	-	-	-	-	-	-	-	-	-	-
EXPS MIN	X	-	-	-	-	-	-	-	-	-	-	-
EXPS MAX	X	-	-	-	-	-	-	-	-	-	-	-
COOL MIN	X	-	-	-	-	-	-	-	-	-	-	-
COOL MAX	X	-	-	-	-	-	-	-	-	-	-	-
COMP MIN	X	-	-	-	-	-	-	-	-	-	-	-
COMP MAX	X	-	-	-	-	-	-	-	-	-	-	-

845

**FIG. 24**



**FIG. 25(a)**

LOT NUMBER	UPC	R/T	POWER	THICKNESS	EXPIRATION
------------	-----	-----	-------	-----------	------------

**FIG. 25(b)**

LOT NUMBER	UPC	R/T	POWER	THICKNESS	EXPIRATION	INPUT QTY	LOSS QUANTITY
------------	-----	-----	-------	-----------	------------	-----------	---------------



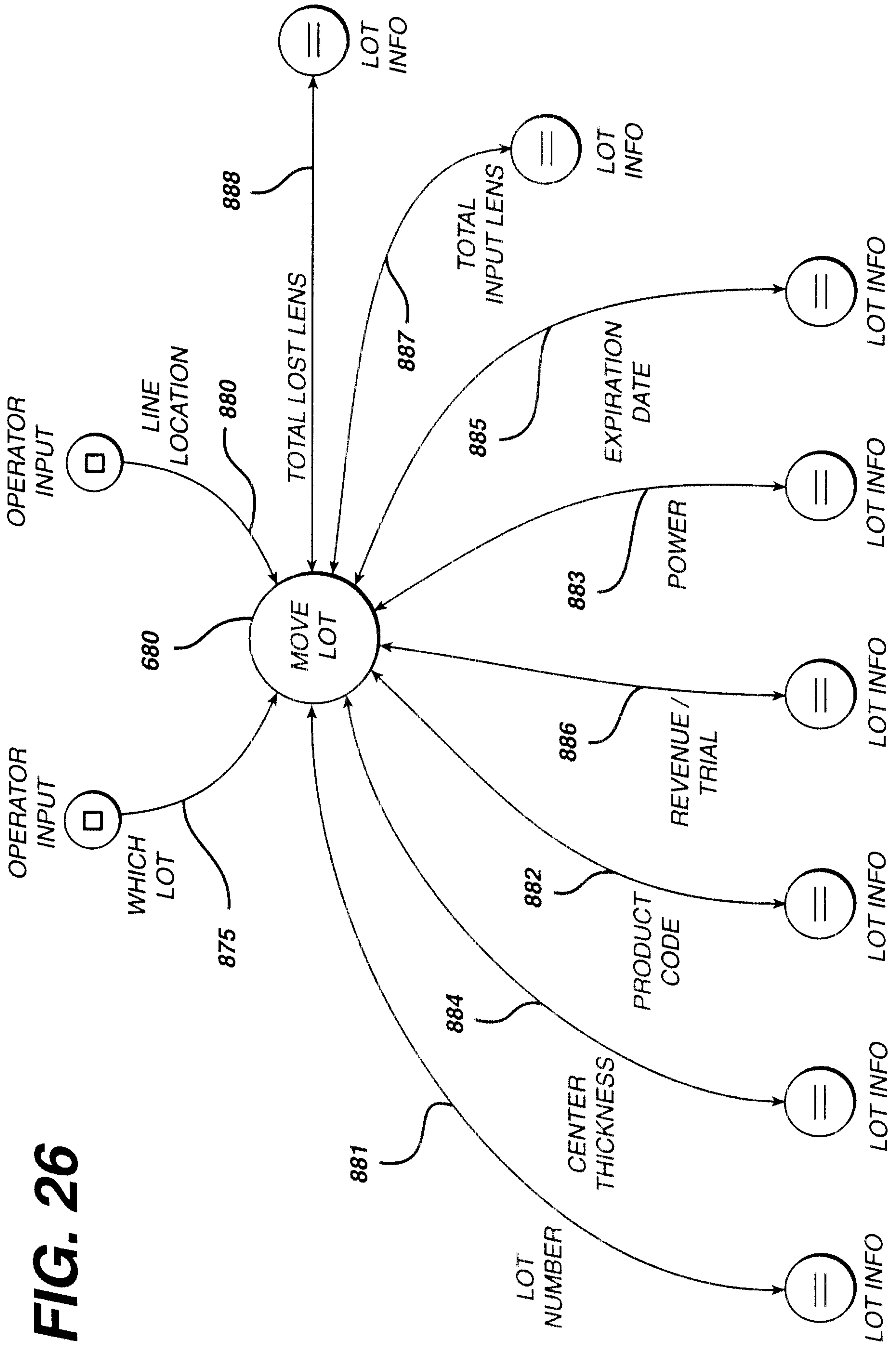
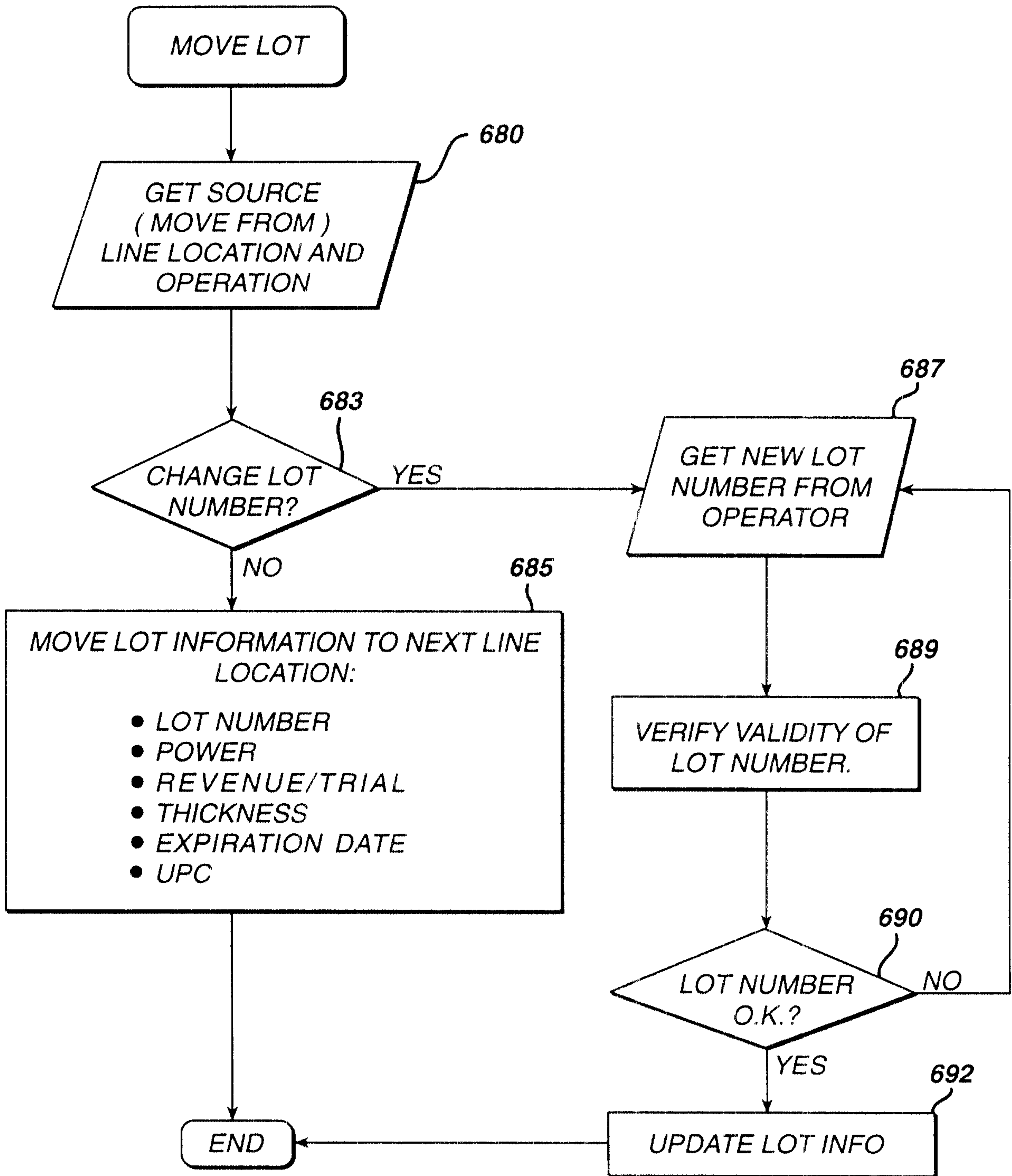
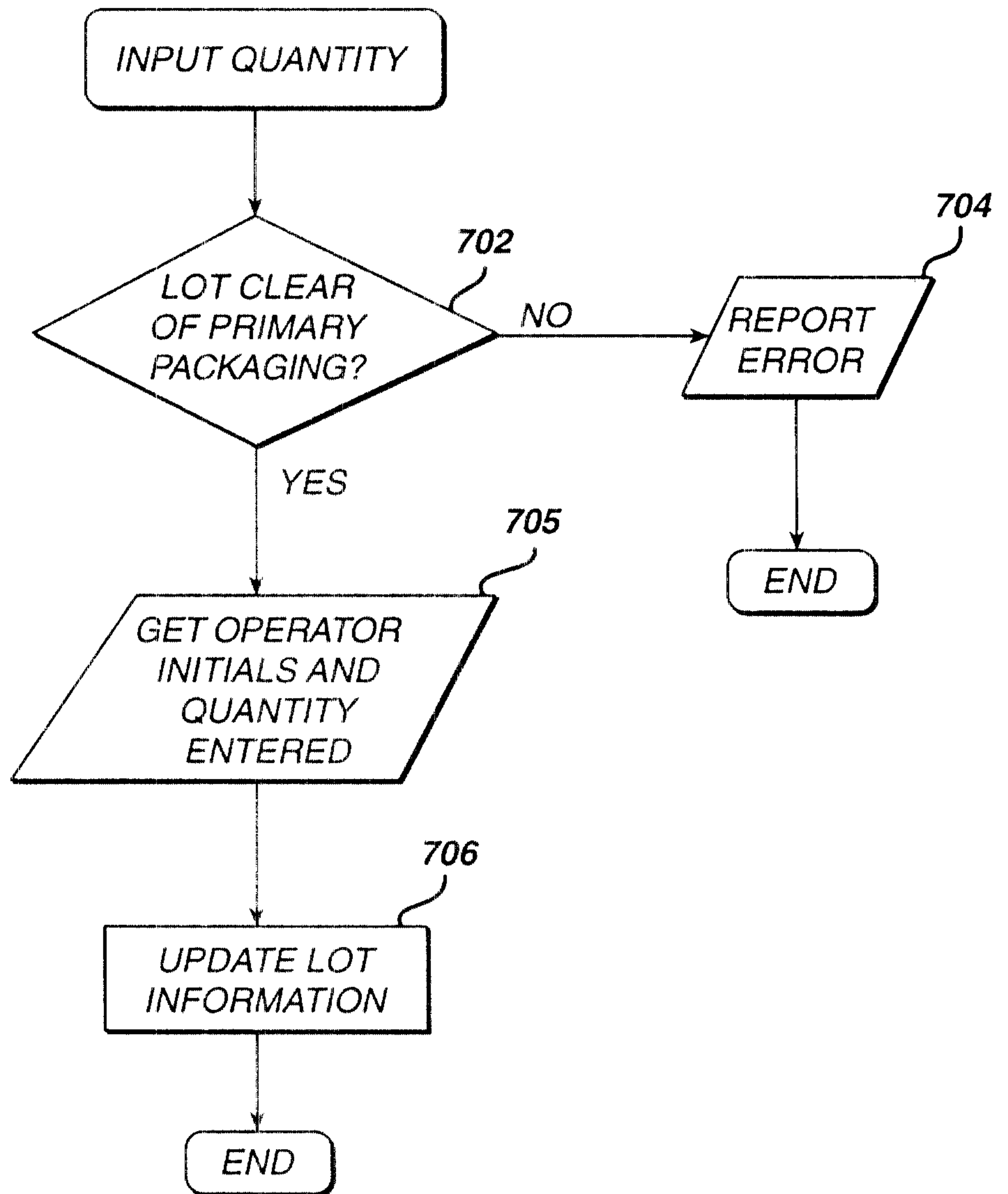


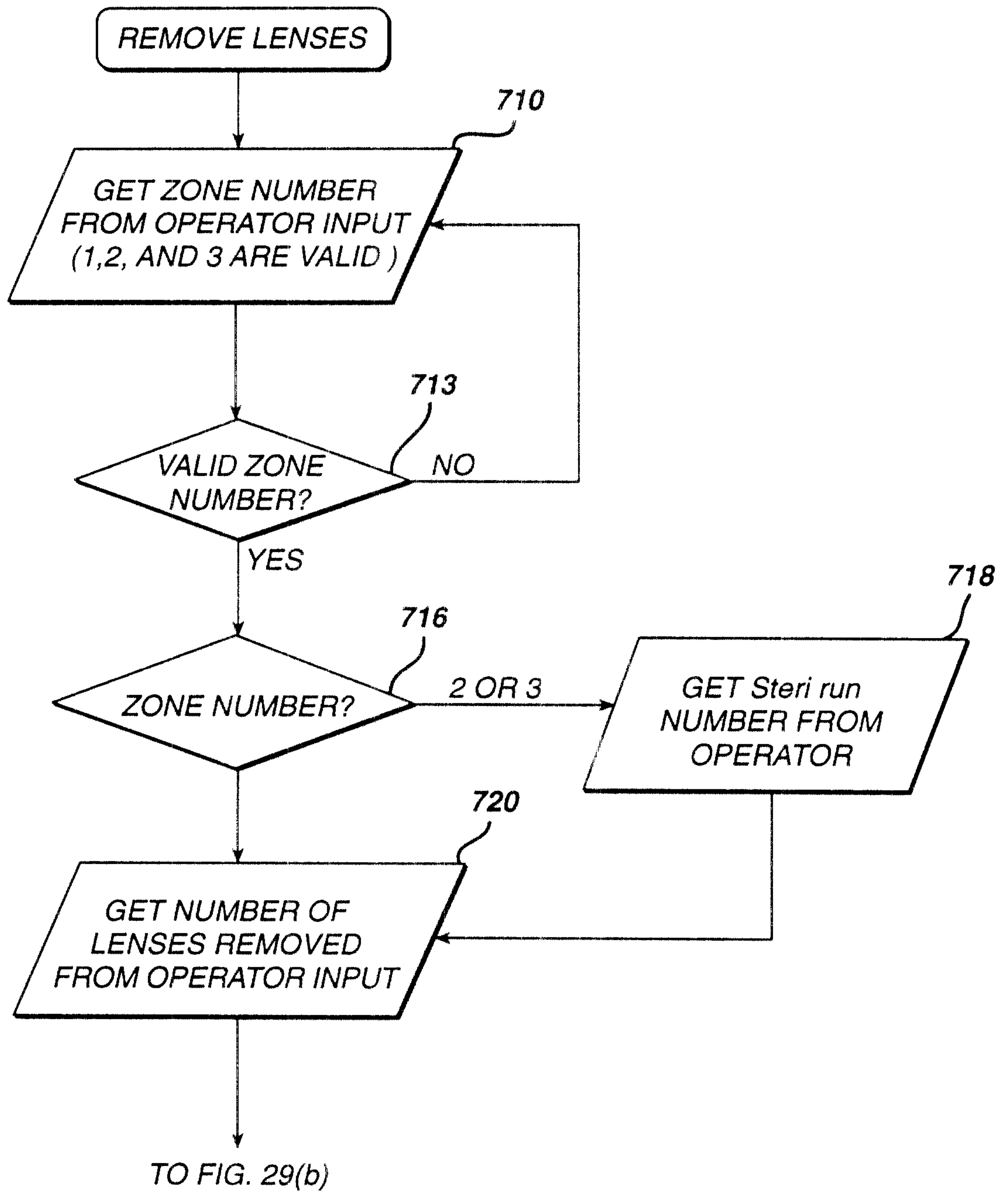
FIG. 26

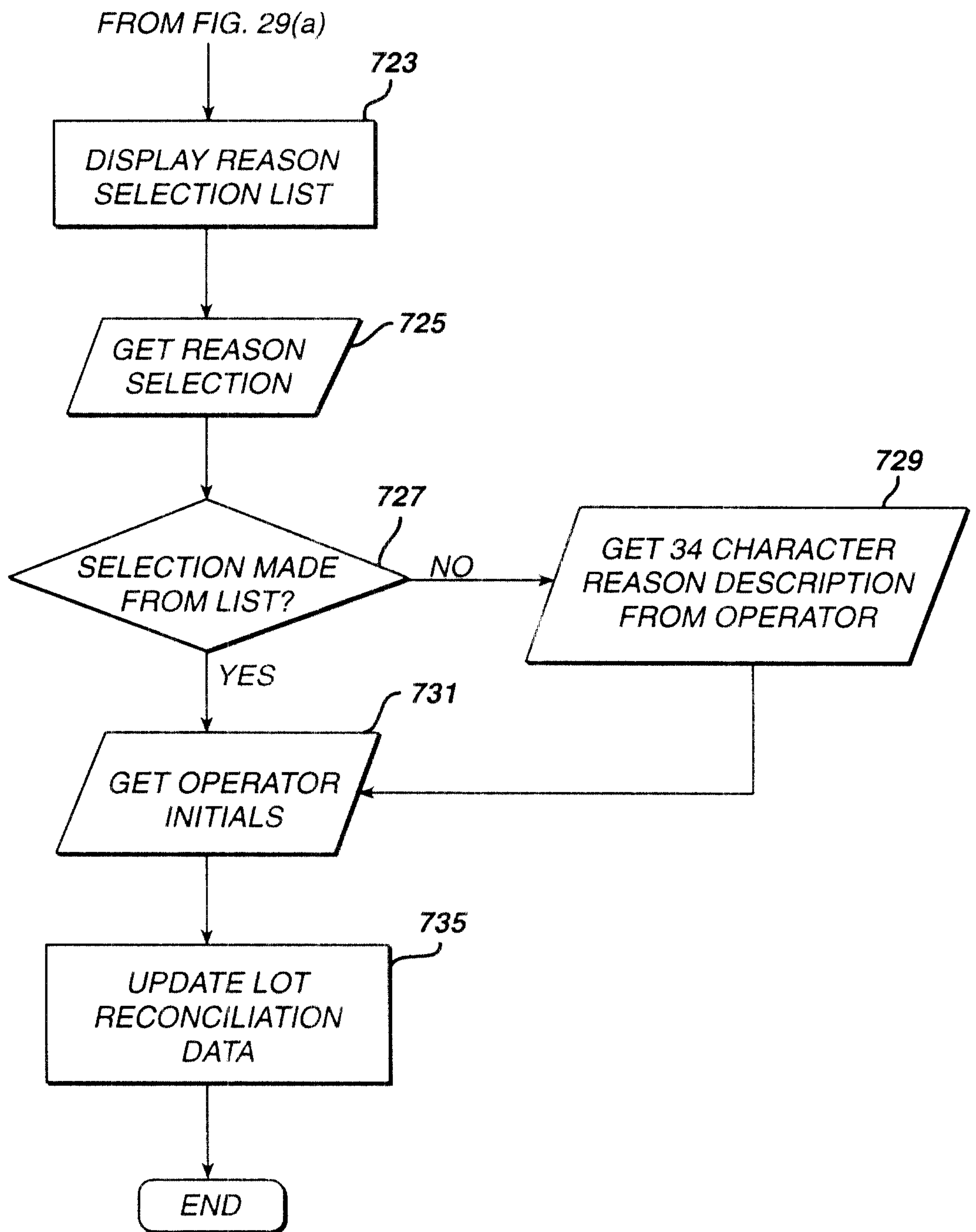
# FIG. 27



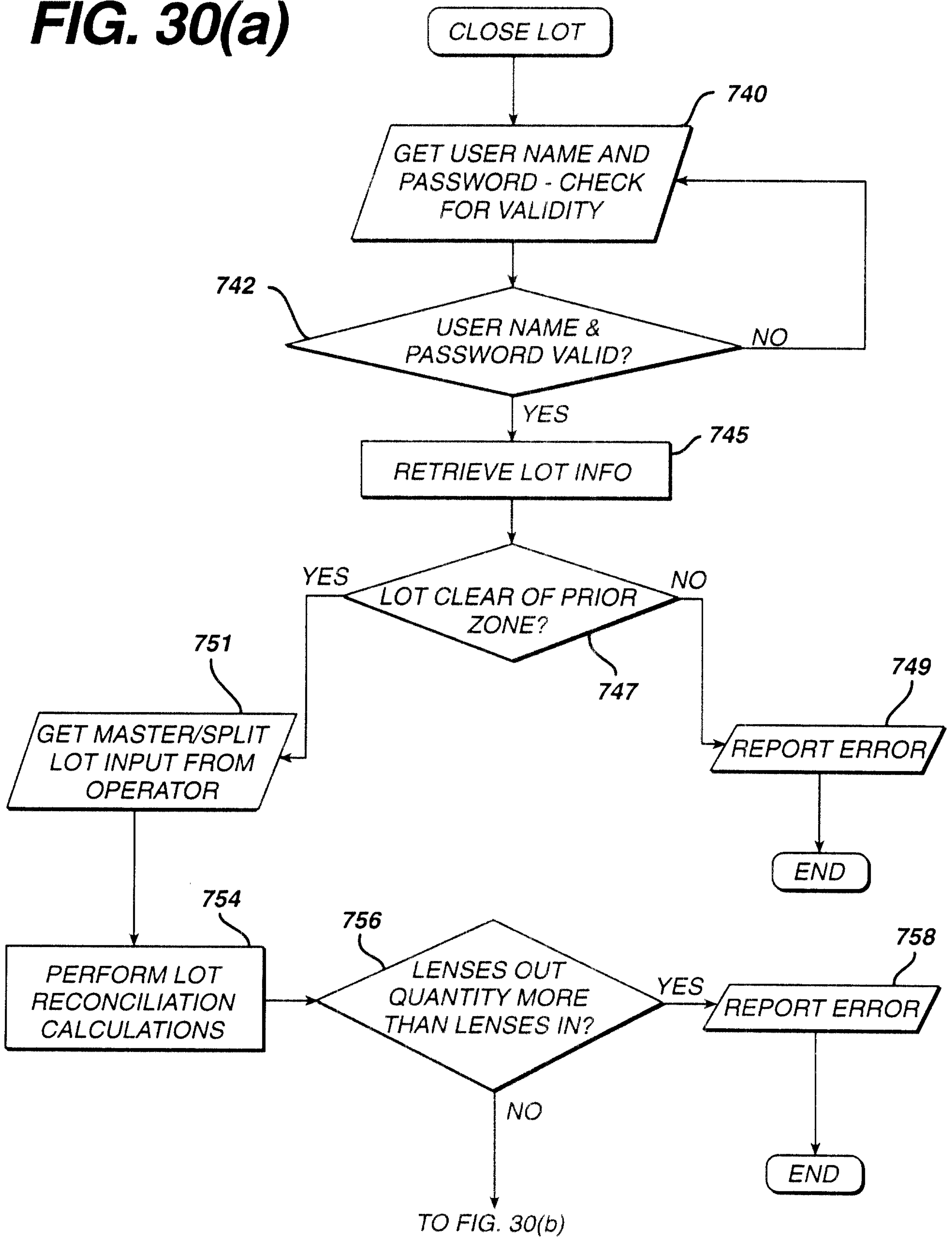
**FIG. 28**



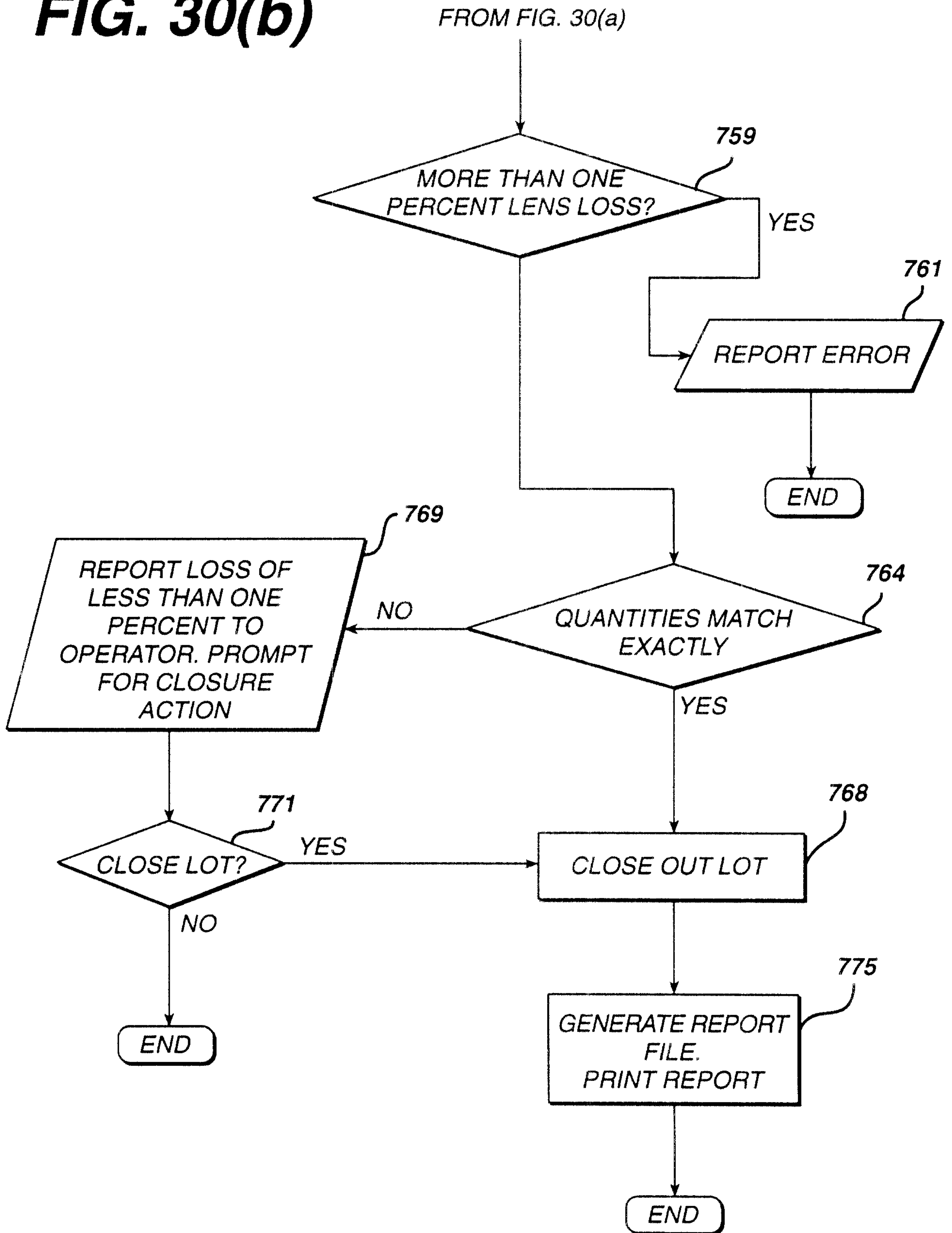
**FIG. 29(a)**

**FIG. 29(b)**

**FIG. 30(a)**



# FIG. 30(b)







# FIG. 32

## STERILIZATION / SECONDARY PACKAGING LOT RECONCILIATION SHEET

890



STERILIZATION RUN # (s)

MASTER LOT # _____	_____	_____
LENS POWER + - _____	_____	_____
MASTER SPLIT (CIRCLE)	_____	_____
REVENUE TRIAL (CIRCLE)	_____	_____
	_____	_____
	_____	_____

891  
 # OF LENSES ENTERING ZONE #1 (STERI TRAY LOAD) -A- \_\_\_\_\_

892  
 SIGNATURE \_\_\_\_\_ DATE \_\_\_\_\_  
 # OF LENSES REMOVED AT ZONE #1 (STERI TRAY LOAD) -B- \_\_\_\_\_

893  
 SIGNATURE \_\_\_\_\_ DATE \_\_\_\_\_  
 # OF LENSES REMOVED AT ZONE # & # (STERI TRAY UNLOAD / CARTONING & CARTON CHECKWEIGH/ LABELING) -C- \_\_\_\_\_

894  
 SIGNATURE \_\_\_\_\_ DATE \_\_\_\_\_  
 # OF LENSES REMOVED BY QUALITY ASSURANCE -D- \_\_\_\_\_

895  
 SIGNATURE \_\_\_\_\_ DATE \_\_\_\_\_  
 # OF LENSES EXITING ZONE # 4 (CARTON CHECK WEIGH/LABELING) -E- \_\_\_\_\_

SIGNATURE \_\_\_\_\_ DATE \_\_\_\_\_

899  
 TOTAL OF ALL LENSES LEAVING THE CELL -F- \_\_\_\_\_  
 B + C + D + E

LOT RECONCILIATION -G- \_\_\_\_\_  
 A - F

VERIFIED BY \_\_\_\_\_ DATE \_\_\_\_\_  
 QA TECHNICIAN

LOT TRANSFER DATE \_\_\_\_\_

