

ASSESSMENT OF COMPUTER USE ADVANCES IN HOSPITAL PATIENT CARE ANALYSIS

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ABSTRACT

The global financial crisis of the past few years has brought to light a number of disheartening circumstances and has heightened public awareness of the necessity of utilizing the world's finite resources more judiciously. The provision of healthcare has been significantly impacted by the numerous reforms and cuts to services implemented by countries currently experiencing economic difficulties, despite the fact that many consider access to healthcare to be a fundamental right. Consequently, the healthcare industry is currently investigating innovative materials and procedures to minimize expenses without sacrificing quality. The entire medical procedure, from diagnosis to therapy, is dependent on technology. It is possible to contend that they are indispensable to the healthcare system. The content they convey is scant and fragmented, which is why any attempt to optimize them is likely to fail. The complex process of manufacturing medical equipment is delineated in this article. This doctoral thesis endeavors to aid the medical device industry in the development of germane solutions and the optimization of its operations by gaining a deeper understanding of the relationship between medical devices and other components of the healthcare system, including regulatory demands and reimbursement decisions. Consequently, the industry would be more prepared to provide patient support. In order to facilitate assimilation of the "medical device system" and its constraints, the recommended approach was graphically represented and disseminated.

Keywords: Optimization, Medical devices, computers, Hospitals.

INTRODUCTION

This has far-reaching consequences for healthcare administration, funding, and management at all levels. By way of illustration, the advent of rapid diagnostic testing has allowed health personnel in low-resource locations to diagnose patients with little to no training. As a result of these tests, it is now possible to screen

populations at risk for infection quickly, which reduces the time and effort required to achieve an accurate diagnosis and the likelihood that patients' conditions would deteriorate during that period. In addition, there was an improvement in the precision of the diagnosis, which led to less antibiotic overprescribing and more treatment adherence for patients. For a brief while, barbers also performed surgical procedures, and local the blacksmiths have been crafting medicinal implements since ancient times. Despite advancements in healthcare and allied sectors, the creation of new medical devices continued to be a pioneering and, at times, chaotic process. The rapid development of medical devices is a considerable problem, notwithstanding their tiny proportion of total health expenditure. This industry has some very high standards and regulations to meet. Before very recently, businesses could charge what they were worth for their efforts as long as their technology satisfied the requirements established by authorities and customers alike. While new Chinese and Indian manufacturers are entering the market with their ideas for inexpensive, stylish designs, the economic crisis is also affecting the industry. Medical device companies are always on the lookout for fresh ways to improve their wares, just like their rivals in the car, consumer electronics, and telecom sectors. A new product development process (NPDP) aims to serve as a guide for researchers and innovators as they traverse the "valley of death" that separates the academic and business spheres. A lot of people think that developing a new product is a tough undertaking since there are so many unknown dependencies, technological issues, and departments that need to be considered and addressed. Even while it does something no one has ever done before, it's also quite similar to other things that have been mass-produced using the same techniques, thus it's not really original. According to Browning et al., there is a need for rigorous post-launch testing if the outcomes of product development initiatives are to be believed as they are sometimes difficult to quantify. In addition to originality and innovation, activities include nonlinearity, iteration, and the simultaneous occurrence of several events. Despite much evidence to the contrary, many continue to hold the belief that creativity and innovation are difficult to systematize. In addition to being iterative and nonlinear, the processes often occur simultaneously(Alexander ,2019).

BACKGROUND OF THE STUDY

The danger a product poses to buyers influences its distribution route. It is essential that all US-made medical devices be both effective and "reasonably" safe, meaning that the benefits of following the product's instructions far exceed any potential risks. The FDA's responsibility in this matter is clear. The only thing manufacturers need to establish to have their devices certified in Europe is that the device is safe and functions as intended. If a European manufacturer wanted to market a laser to treat arrhythmia—an irregular heartbeat—they would have to prove it was safe to use on cardiac tissue. The manufacturer of the laser, on the other hand, would be required to prove that it cuts heart tissue and treats arrhythmia before selling it in the US market. The United States often demands randomized and controlled studies,

in contrast to Europe that is satisfied with literature reviews, small-scale clinical trials, or laboratory testing. Additionally, the proof that is required varies among areas. These factors influence the seemingly disparate times it takes to bring new items to market and the amount of testing that medical devices are required to complete. European medical technology association research shows that new medical technologies are usually introduced in Europe five years before they reach Japan and two years before they hit the US market. It premieres in the US two years before it does in Japan. In the United States, the data used to determine whether a product is permitted for sale is made public by the Food and Drug Administration (FDA). The Food and Drug Administration also has a responsibility to protect the public's health. European approvals and the facts supporting them are kept secret, and manufacturers are held responsible for selecting and working with a recognized authority. Manufacturing must commence in Europe before the approval process can commence. Reason being, it calls for an in-depth analysis of the device and its manufacturing procedure. Once the design is finalized, certification may begin in the US as the device itself is the only item that needs to be examined. This is a major issue since it makes changing the device's manufacturing process impossible. All class III and certain class II devices in the US must undergo post-marketing surveillance studies prior to FDA clearance. This category encompasses all tools used in surgery and medicine. As to Salhie, once items reach the European market, firms are compelled to implement a "vigilance system" to assess their quality control (Bajwa, 2019).

The purpose of the research

One of the many goals of maintaining a patient's medical history is to facilitate communication between the many members of the medical care team, such as doctors, nurses, and allied health workers. This document aims to simplify the process of treatment continuation for patients by acting as a readily accessible reference. because proof that medical treatment was obtained at the medical institution must be shown.

LITERATURE REVIEW

The current population of Britons aged 65 and over is 10.3 million, however the ONS projects that figure will have risen to 12.7 million by 2018, based on research by Rutherford and colleagues . People aged 65 and over will eventually account for 12.7 million, up from 10.3 million in 2010. Based on projections made public by the Office of National Statistics , the population of people aged 80 and older is anticipated to more than double by mid-2037, going from around 2.5 million to 6 million. It is a tremendous increase when contrasted with the current elderly population. It is believed that the current old population will play a significant part in the enormous expansion. Rising healthcare costs are often attributed, in part, to the ever-

increasing proportion of the population aged 65 and higher. Although these are distinct matters, healthcare expenditure is connected to both the supply and demand for services and the actual use of healthcare by patients. When we think that a patient's health and ability to reach their objectives will be improved by receiving treatment, we say that they are "needing" medical care. To determine the target degree of improvement, one may utilise any one of many viewpoints and measures. Variations in infant mortality rates among regions and the country, as well as patient and physician perspectives, and health status markers like blood pressure are all part of this. When attempting to predict consumer actions, one must take the existence of a demand into account . Demand is affected by a number of factors, including the desire and ability of patients and the nation to pay for certain healthcare treatments. Certain medical treatments are seeing unprecedented demand. How individuals make use of healthcare services is regulated by an ever-changing process. The procedure takes a number of things into account, one of which is the availability and demand for healthcare services. A measure of the demand for healthcare may be the frequency with which hospitals are filled to capacity. Spending on healthcare is directly correlated to consumption, which satisfies healthcare requirements. The Directorate-General for Economic and Financial Affairs of the European Commission predicts that, as the population ages, these services will receive much more funding in the future. Predictions of a longer median age of population are the most probable factor propelling this expenditure surge. The elderly are more prone to suffer from acute and chronic ailments, making medical treatment and the associated social care services more costly on average compared to younger persons. Many nations and health organisations are beginning to fret about the potential impact of future demographic shifts on medical and social care expenditures, as reported by the European Commission's Directorate-General for Economic and Financial Affairs(Robinson,2021).

Research Question

- i) How do computers rate in terms of health technology?
- ii) What are computers used for in the health care field?
- iii) What kinds of things do medical gadgets make?
- iv) How many different kinds of computers are there in the health technologies sector?

METHODOLOGY

Scientists carried out a detailed cross-sectional investigation. Because it was a cross-sectional research, all that was required was a single, affordable point in time to

collect data. Due to time constraints and limited resources, the researcher used a quantitative approach. After determining a sample size of 1234 using Rao-soft software, 1400 questionnaires were sent, 1356 were returned, and 31 were excluded owing to incomplete surveys. There were 1,325 participants in the study. All prospective respondents were contacted for the survey using a random sampling process. While waiting for their medical devices to be ready, individuals who chose to participate in the study were told about it and the researcher addressed any questions they had. When a respondent was unable to read or write, or was wheelchair-bound, the researcher read aloud the survey questions and answered categories before recording their responses on the form. People were requested to complete and return questionnaires all at once in certain areas.

Study Area

The study was conducted by health technology on medical devices.

Data Collection

For this study, the researcher used a quantitative approach by administering a survey. Down below, you'll find the specifics of the interview and survey methodology. Questions about participants' extracurricular involvement in healthcare methodologies: an in-depth research based on medical equipment served as a control. Questionnaires often make use of a rating system called a "Likert scale" to gauge respondents' thoughts, feelings, and perspectives. In this kind of survey, people are asked to rate how strongly they agree or disagree with a statement or issue. The potential answers vary from "strongly agree" to "agree," but "did not answer," "disagree," and "strongly disagree" are also common. It is common practice to use numerical codes to denote answer categories; however, these codes must be established for each individual research; for example, 5 = highly agree, 4 = agree, etc. Researchers viewed respondents' gender and age range as part of the demographic data used in the study. All of it is included in the demographic information.

Sample

Interviews and data collecting methods were used to compile the study's findings. A total of 1400 questionnaires were issued; 1356 were returned; 31 were discarded due to incompleteness; the sample size was 1234, as computed using Rao-soft software. The total number of questionnaires used for the research was 1,325; 772 were male and 553 were female.

Quantitative Method Research

A wide range of phenomena may be described and explained by quantitative research, which makes use of mathematical representations and data manipulation. The use of numerical representations allows for this to be achieved. In recent decades, this phenomena has found extensive use across a variety of fields, with unique applications in fields as diverse as geology, physics, biology, and sociology. In addition, according to Cohen (2018), "quantitative research" refers to social studies that use an empirical approach and derive conclusions based on facts. When discussing social science studies, the phrase "quantitative research" is often used. From the author's point of view, an empirical assertion is a claim about the actual situation as opposed to an opinion on how things should be. As a counterpoint, they bring forth the idea of "ought to be." Quantitative research often uses numerical representations to state empirical claims, which means that empirical assessments are necessary. To review, an empirical evaluation is a methodical effort to determine how well the programme or policy matches up with a predetermined standard. One way to do this is to compare the results of the evaluation to a standard that has already been set (Chouliaras, 2021).

Study Design

The study lasted five months, from January to December of that year, and it was a thorough cross-sectional study. It took place while this probe was underway. There was no break in the action for the whole of the session. Because the data collection for the cross-sectional design was done at a single point in time, the technique was simplified and the related expenses were considerably reduced. This allowed for the faster implementation of the cross-sectional design. The researcher decided that a quantitative method would be the best way to conduct her analysis due to her time and budget constraints. This include members of the academic community, students, public servants, and business and corporate leaders. Stratified random sampling was the method used by the researchers to interview employees at each of the following organisations. The staff members that have been asked to fill out the survey. Information about the research is made available to the participants, and those who are interested may fill out and submit a questionnaire whenever it is convenient for them. At the same time, the researcher is present in the waiting area to address any inquiries that may have arisen due to the provided information. For the sake of speed, they disseminated surveys in a variety of contexts.

Study Area

The study was conducted by health technology on medical devices.

HYPOTHESIS

Electronic Health

A patient's or a population's medical history kept in a structured digital database is known as an electronic health record (EHR). Multiple healthcare facilities may access and use these records. Information networks and exchanges allow for the sharing of records across enterprise-wide information systems that are network-connected. Electronic health records (EHRs) may include a wide variety of information, such as patient demographics, medical history, allergies and medications, vaccination status, lab findings, imaging studies, vital signs, personal data (such as weight and age), and financial details. Electronic health records (EHRs) have been hailed for decades as a game-changer in healthcare quality. These days, doctors aren't only utilizing EHRs for patient charts; they're also incorporating patient data into care management programmes to boost quality outcomes. Electronic health records (EHR) compile patient data into a single database, which is then used to aid in the development of "new treatments or innovation in healthcare delivery" that ultimately achieve healthcare's stated objectives. By integrating various forms of clinical data from the system's health records, doctors have been able to better detect and categorize individuals with chronic illnesses. By using data and analytics, EHRs may enhance the quality of treatment by reducing hospitalizations among patients at high risk. The goal of electronic health record systems is to keep track of a patient's condition throughout time and to save that data properly. Finding a patient's old paper medical records is a thing of the past thanks to this technology, which also helps keep data current, accurate, and readable. The patient and physician are able to communicate freely while enjoying the benefits of "privacy and security." Having just one editable file lowers the possibility of data duplication, increases the likelihood that the file is up-to-date, makes paperwork less likely to go misplaced, and saves money. Electronic medical records (EMRs) are superior than paper records when it comes to collecting medical data for the purpose of studying potential patterns and long-term changes in a patient. This is because digital information is searchable and stored in a single file. EHRs and EMRs, if widely used, may also make it easier to conduct population-based studies of medical records(Rittenhouse,2020).

Patient Care In Hospitals

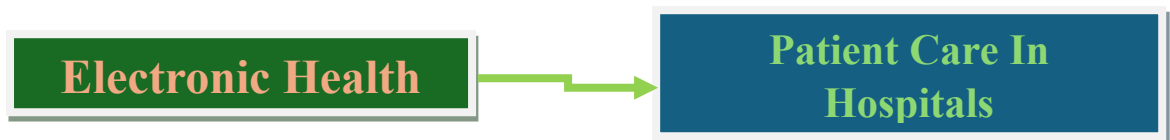
Patients are better equipped to manage their healthcare needs when they have a healthcare management plan and follow the plain directions of their trusted physicians and caretakers. This allows patients to better control their situation. Individuals are more likely to be empowered to take an active part in reaching their own healthcare goals when they are provided with optimum treatment alternatives and when health care systems that include patient-focused services into their design are likely to give such possibilities. It is of the utmost importance to be vigilant in the monitoring and treatment of long-term health conditions such as cardiovascular disease, diabetes, hypertension, and others. The patient care management system, which provides medical professionals with a detailed plan and community resources, makes it feasible for them to effectively handle situations like this. A healthcare facility will continue to work towards bettering the health and well-being of its patients even when those individuals are not physically present at the institution

providing treatment. The use of practice-based strategies is the most important aspect of care management when it comes to improving the health of societies who are dealing with complicated or long-term medical difficulties. Software designed specifically for care management is becoming more popular among healthcare companies as a means of enhancing their care management operations. The requirements of large healthcare systems could be satisfied by a number of different solutions, while the requirements of primary care clinics may be satisfied by other solutions. Contrary to popular belief, the integration of electronic health records is absolutely necessary in every circumstance(Melnick,2021).

H0: There is no significant relationship between Electronic Health and Patient Care in Hospitals

H1: There is a significant relationship between Electronic Health and Patient Care in Hospitals

CONCEPTUAL FRAMEWORK



RESULTS

During the interviews, it was found that most hospitals (reported 19 times) give the job of doing an early review to both employees and outside experts. Only three of the people who took part said that the early review tasks were all run by an outside expert. On the other hand, fifteen of the people who answered said that an early review was only done by people hired by the company. Following the interviews, it was found that the main reason companies seek outside help for the early evaluation of medical equipment is because their own employees lack the necessary skills (N=18). In 18 of the 20 hospitals that were looked at, this was found to be true. It was said by six of the participants that that would be the best way to save money and two said that it would also be the best way to save time. As a result, some of the members said it would be better to have evaluators from outside the group. More than half of the work that goes into internal assessments is done by people who are specifically assigned to do them (N = 22) or by people who work in specialised departments within the company (N = 14), like the Sales and Marketing and Research and Development departments. Six of the people who answered said that their companies do an early review by giving different people different tasks to make the process go more smoothly. Two of the participants finally said that they didn't fully understand how an early review worked at their business. The people

who took part were honest enough to say this.

- How well a person does a number of jobs that are linked to a general evaluation

When people were asked to pick the first step in the evaluation process based on six generally understood stages of medical device development, it seemed that clinical context and market assessment were the most important ones. This was because they were asked the question. And this is because, for the vast majority of firms, both of these evaluation processes begin with the ideation step. In terms of doing financial and health economic assessments as well as stakeholder analysis, only 18% of businesses start these steps while the idea is still being developed. This means that most businesses choose to wait until later in the process. This picture (Figure 2a), which came from the Dutch medical products industry, shows the different steps of an early review that began in four different areas.

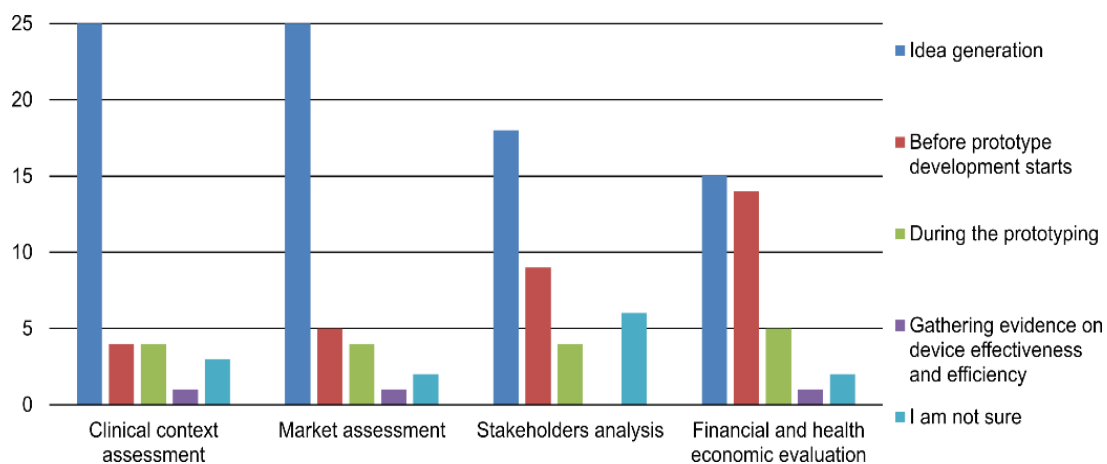
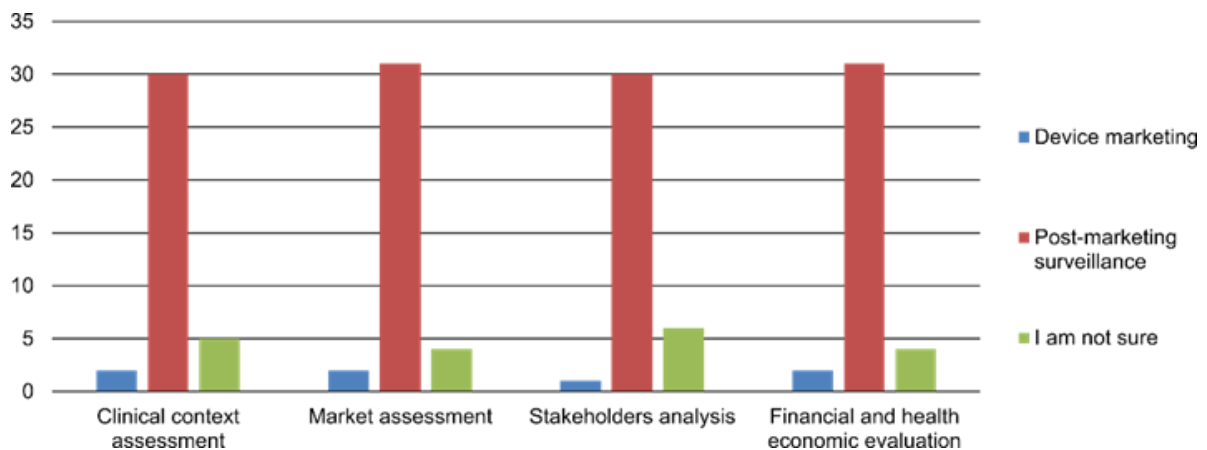


Figure 2. This figure provides an overview of the beginning time of an assessment activity within each of the four categories of an early assessment, as suggested by the participants in the interview.



When asked to suggest when the assessment activities would end, the majority of the participants (reported 30-31 times) answered that all aspects of an assessment activity will continue until the post-marketing monitoring is completed. Figure 2b provides a summary of the times when assessment activities in four different areas of an early assessment were scheduled to conclude, based on the information provided by the participants in the interviews. Figure 2 provides a summary of the closing time of an assessment activity within each of the four domains of an early assessment, as reported by the people who took part in the interview.

DISCUSSION

The information technology sector has been very successful in creating financial value for end users during the last 50 years. While client electronics and the Internet drove business process automation in the 1980s and 1990s, mainframes and personal computers drove industrial automation in the 1960s and 1970s. With the development of digital data platforms, technological breakthroughs in massive data sets, cloud computing, mobile, and social networks since 2000 have expedited the digital transformation. Intelligent enterprises will be made possible by the advancement of smart technologies like big data, artificial intelligence (AI), deep learning, and the Internet of Everything (IoE). To convert unorganized data into semantically standardized, organized data sources, intelligent, adaptable, adaptive medical information extraction techniques must be combined with highly accurate rule-based algorithms. Clinical data warehouses should be expanded into complete knowledge discovery systems that leverage ontologies for semantic reasoning techniques, rather than only depending on them for coding and code translation. These methods result in the creation of fresh data and hypotheses. Predictive analytics is essential rather than just offering analytical data. Both better care delivery and training for healthcare personnel in clinical data mining and machine learning systems are necessary. Furthermore, ontologies are essential to knowledge systems because they provide codification services and make it possible to categorize illnesses and medical concepts based on their similarities and differences. Many businesses are now looking for methods to reduce expenses. It may be argued that the safest and most dependable approach to managing data is to create and maintain copies of software and data on-site. These days, with all the advantages of cloud computing, things may be different. Cloud computing in healthcare may provide better patient data usage and preservation, lower storage costs, quicker innovation cycles, simpler collaboration, and more opportunities for telemedicine. The adoption of cloud-based technology by the healthcare sector will vary depending on the four stages of digitalization. As a result, software versions that are both cloud-based and on-premise will continue to be used in combination(Huang,2018).

CONCLUSION

In this section, we provide a succinct synopsis of our research, including its introduction, framework construction, method selection, results, and analysis. The study's limitations, implications, and recommendations for more research are then thoroughly discussed. As healthcare organisations depend more and more on technological solutions, productivity, effectiveness, and knowledge gains should be anticipated. Scholars and medical professionals are intrigued by the Internet of Things' (IoT) many applications in the field of healthcare. Hospitals have unique challenges in implementing and using new technologies, even with the potential advantages of the Internet of Things (IoT) for healthcare delivery (Glover et al., 2020). The study's goals were to: (1) assess the rate of IoT adoption in one Indian hospital; (2) pinpoint the obstacles to IoT adoption; and (3) find out how the hospital intends to get beyond these obstacles. By analyzing the challenges hospitals in poor nations have when trying to use IoT and offering suggested solutions to these issues, this study aims to close a knowledge gap. First, a comprehensive literature review was conducted to learn more about the topic of healthcare adopting technology advancements. Our analysis of the most recent research revealed several well-known procedures and obstacles to the adoption of new technologies. In the early talks of technology adoption theories, the user adoption model, UTAUT, and the TOE framework were presented. After that, we shall describe how the Internet of Things (IoT) functions in an organisational setting that is more extensive(Copley,2019).

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