

Scientific Article

Implementation of Magnetic Resonance Imaging-Guided Radiation Therapy in Routine Care: Opportunities and Challenges in the United States



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Abstract

Purpose: Magnetic resonance image (MRI)-guided radiation therapy with the 1.5 Tesla magnetic resonance linear accelerator (MR-Linac) is a rapidly evolving and emerging treatment. The MR-Linac literature mainly focused on clinical and technological factors in technology implementation, but it is relatively silent on health care system-related factors. Consequently, there is a lack of understanding of opportunities and barriers in implementing the MR-Linac from a health care system perspective. This study addresses this gap with a case study of the US health care system.

Methods and Materials: An exploratory, qualitative research design was used. Data collection consisted of 23 semistructured interviews ranging from clinical experts at the radiation therapy and radiology department to insurance commissioners in 7 US hospitals. Analysis of opportunities and barriers was guided by the Nonadoption, Abandonment, Scale-up, Spread and Sustainability framework for new medical technologies in health care organizations.

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Results: Opportunities included high-precision MR-guidance during radiation therapy with potential continued technical advances and better patient outcomes. MR-Linac also offers opportunities for research, professional, and economic development. Barriers included the lack of empirical evidence of clinical effectiveness, technological complexity, and large staffing and structural investments. Furthermore, the presence of patients with disadvantaged socioeconomic background, and the lack of appropriate reimbursement as well as regulatory conditions can hinder technology implementation.

Conclusions: Our study confirms the current literature on implementing the MR-Linac, but also reveals additional challenges for the US health care system. Alongside the well-known clinical and technical factors, also professional, socioeconomic, market, and governing influences affect technology implementation. These findings highlight new connections to facilitate technology uptake and provide a richer start to understanding its long-term effect.

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Introduction

Defining the outcomes of new medical technology upfront implementation is important to identifying its potential benefits and obstacles.¹⁻³ Insight into comparative effectiveness of new medical technology to current standard of care is critical to ensure patient access to high-quality health care at the lowest possible cost levels.⁴⁻⁸ Many new medical technologies, however, lack a comprehensive effectiveness evaluation before being implemented in the clinical setting, hence, without proven effectiveness.^{5,6,9} This is problematic because the implementation of unproven medical technologies may threaten the quality of health care and increase costs.¹⁰ Consequently, payers and policymakers are reluctant to approve or reimburse costly medical innovations.

Despite these concerns, especially with regard to medical technology still in development, it can be difficult to estimate the relative effectiveness, nor its long-term effect on the patient, provider and payer.¹¹ This is particularly relevant for convergent technologies, which tent to cross disciplinary boundaries and are becoming dominant in health-related domains.¹² For instance, the convergence between biology and informatics knowledge has led to bioinformatics and personalized medicine innovations.¹³ The involvement of different disciplines means that convergent medical innovation and its potential value, cannot be understood and implemented without intertwined institutional practices.^{14,15} Early assessment of crucial factors of technology implementation is therefore needed for an initial understanding of its long-term impact.¹⁶

A modern example of a new developing and convergent technology in the field of radiation oncology is the 1.5 Tesla MRI-guided linear accelerator (MR-Linac) system, which combines high-precision, real-time MRI external beam radiation therapy (MRgRT).¹⁷⁻¹⁹ As a result, the irradiation plan can be adjusted at any time based on these changes. This allows more targeted delivery of radiation to the tumor and avoids the healthy tissue surrounding the tumor compared with most conventional radiation therapy techniques.¹¹⁻¹³ Therefore, the MR-Linac may reduce radiation therapy induced toxicity and

improve tumor control outcomes.^{17,19} The MR-Linac also allows therapy in fewer sessions with a higher dose of radiation therapy, also called hypofractionation, hence, permitting a shorter treatment.^{20,21} This can be beneficial in a case like localized prostate cancer²²⁻²⁴ because traditional external beam radiation therapy often varies from 5, 20, to 39 treatment sessions.²⁵

Alongside technical and clinical opportunities, the MR-Linac may also offer positive professional and economic prospects as identified by a Dutch implementation study.¹⁹ For instance, technology adopters may gain a more efficient treatment, a higher hospital quality profile with potential financial benefits, and improve their competence and technical expertise as well as multidisciplinary collaboration. However, technology implementation may also deal with technical complexities, substantial staffing and structural investments, and the presence of patient referral patterns and professional silos. In addition, we still know very little about the actual patient benefit of MRgRT due to the current lack of empirical evidence of clinical effectiveness.

The lack of effectiveness evidence has different implications across health care systems because the dynamics of stakeholders, organizational procedures, dominant existing routines, professional identities, and legal and regulatory standards in technology implementation vary between countries.^{1,26-28} For instance, developing and acting upon evidence-based practices is perceived to be more challenging in health care systems with large private payers such as those in the United States, than in publicly funded health care systems as in Europe or Canada.⁵ A fragmented and disjointed public and private health care hybrid in the United States can result in substantial variations in health care delivery.⁵ This is particularly relevant for MR-Linac systems as several have been installed worldwide, with a considerable number currently operational in the United States (US). Hence, it is interesting to study how to facilitate the implementation of the MR-Linac technology in specific health care systems such as the US. In the present study, we aim to identify the opportunities and barriers to the implementation of the MR-Linac technology in US hospitals.

Methods and Materials

We conducted a qualitative study with semistructured interviews, a method most appropriated to make sensitive issues, attitudes, opinions, and experiences of individuals explicit.²⁹

Data collection

We used the Nonadoption, Abandonment, Scale-up, Spread and Sustainability (NASSS) framework of new technologies to explore the factors of success and failure of technology adoption in health care organizations.¹ The NASSS framework considers 7 domains: (1) the targeted clinical indication; (2) the technology to be implemented; (3) the value proposition; (4) the adopter system (patient, technology user and other staff); (5) the providing organization; (6) the wider institutional social context; and (7) organizational resilience and technology development over time.¹

We conducted 23 semistructured interviews in 2020 to identify the opportunities and barriers to MR-Linac implementation in 7 US hospitals (Table 1). A questionnaire was designed based on the NASSS framework (Appendix E1). Interviewees included radiation oncologists, urologists, radiologists, medical physicists, radiation therapy technologists, and dosimetrists, as well as nonclinical hospital administrators, including strategic and financial department managers as well as insurance councils. We included interviewees from academic and community-based hospitals, to avoid professional biases. The study was approved by the local medical university research ethics board and all study participants provided verbal consent.

The research objective was explained in the written invitation and at the start of each interview. All interviews were audio-recorded and transcribed using Sonix transcription software. Interviews lasted between 45 and 60 minutes and were carried out face-to-face and via telephone and Skype. Audio recordings and transcript of interviews are anonymized as well as confidential and therefore not publicly available.

Data analysis

Interview transcripts were analyzed using NVivo software. We applied open coding based on the research objective and ended with axial coding, identifying areas of theoretical interest and common themes. Developed codes were validated by a second reviewer. To include variation in findings and increase construct validity, we triangulated the findings across different respondents and with literature.

Results

We identified 4 opportunity and 5 barrier categories for implementation of the MR-Linac technology in US hospitals.

Opportunities

We first describe the clinical opportunities before moving on to the technological, adopter, and economic opportunities. Figure 1 shows the percentages of the interview cohort who discussed the opportunities, by main theme and subtheme.

Clinical opportunities

Interviewees expected a large added-value from MR-Linac implementation in clinical practice as a result of technical advancements. Main clinical opportunities included: (1) potential reduced toxicity and improved tumor control due to more precise and targeted radiation therapy; (2) possibly improved understanding of individual treatment response; and (3) potentially improved patient comfort and quality of life. Furthermore, interviewees also discussed that superior tumor delineation with real-time imaging and hypofractionation could improve treatment compliance and hospital visits given the potential of less therapy fractions.

Technological opportunities

The MR-Linac was perceived by interviewees to be a sophisticated technology with potential technical advances which could allow for both diagnostic and therapeutic opportunities. For instance, one interviewee referred to the potential of adaptive treatment planning for adaptive contour propagation and rapid dosimetric reconstruction which would allow for smaller treatment uncertainty margins and avoidance of dose to healthy tissues. This explains the possibility of hypofractionation. There was general agreement among interviewees that these opportunities may evolve as software development continues (eg, application of deep learning, enabling potential technology developments).

The opportunity of real-time adaptive MRI-imaging during the course of radiation therapy could allow for increased data collection including anatomic and functional imaging data of both the tumor and surrounding organs. This data is perceived to allow greater understanding to safely deliver the most effective dose to individual tumor biology rather than its stage and location. As a result, interviewees expected new treatment options in radiation oncology and in related medical disciplines.

Table 1 Overview of roles and affiliations of respondents

Respondent	Position	Seniority	Affiliation	Additional roles	Method	Duration (min)
R1	Head of Imaging and Oncology department	Full professor	AMC1	Research on functional imaging	In person	44
R2	Radiation therapy technologist	Senior	AMC1	Research on MR-Linac	In person	43
R3	Insurance commissioner	Senior	AMC1		In person	46
R4	Head of IT for Oncology	Senior	AMC1	Research on functional imaging	In person	44
R5	Medical physicist	Senior	AMC1		In person	41
R6	Nuclear medicine physician	Senior	AMC1		In person	39
R7	Member Board of Directors	Senior	AMC1		In person	44
R8	Member Board of Directors	Senior	AMC2		In person	44
R9	Radiation oncologist	Senior	AMC3		In person	39
R10	Radiation oncologist	Senior	AMC4		In person	39
R11	Radiation oncologist	Senior	AMC2	Head of Radiation Oncology department, Research on MR-Linac	In person	45
R12	Radiation oncologist	Senior	AMC2	Research on MR-Linac	In person	53
R13	Radiation oncologist	Senior	AMC2	Research on MR-Linac	In person	39
R14	Radiation oncologist	Senior	AMC5	Research on MR-Linac	Virtual	41
R15	Urologist	Senior	AMC4	Head of Urology department	In person	40
R16	Urologist	Senior	MC1		In person	29
R17	Radiation therapy technologist	Senior	AMC1		In person	41
R18	Radiologist	Professor	AMC1		In person	35
R19	Radiation oncologist	Senior	AMC4	Head of Radiation Oncology department	Virtual	39
R20	Radiation oncologist	Senior	AMC2	Research on MR-Linac	In person	45
R21	Radiation oncologist	Full professor	AMC2		In person	48
R22	Radiation oncologist	Senior	MC2		In person	41
R23	Market access associate	Senior	Manufacturing company		Virtual	46

Abbreviations: AMC = academic medical center; MC = (nonacademic) medical center.

Professional opportunities

Implementation of the MR-Linac allows different possibilities for professional development. First, using MR-Linac encourages the development of new competencies and welcomed novel responsibilities for staff members because of the technological knowledge required to run both an MRI machine and linear accelerator. For instance, radiation oncologists need to gain understanding of the

potential information obtained from different MRI sequences and response assessment. Furthermore, interdisciplinary relationships could be improved because of the increased cooperation between radiology and radiation oncology experts. These opportunities are expected to allow retention of staff and additional recruitment. As an example, one interviewee stated: “I think rolling out new technology is always important for your recruitment and retention of the highest-level physicians in the hospital.”

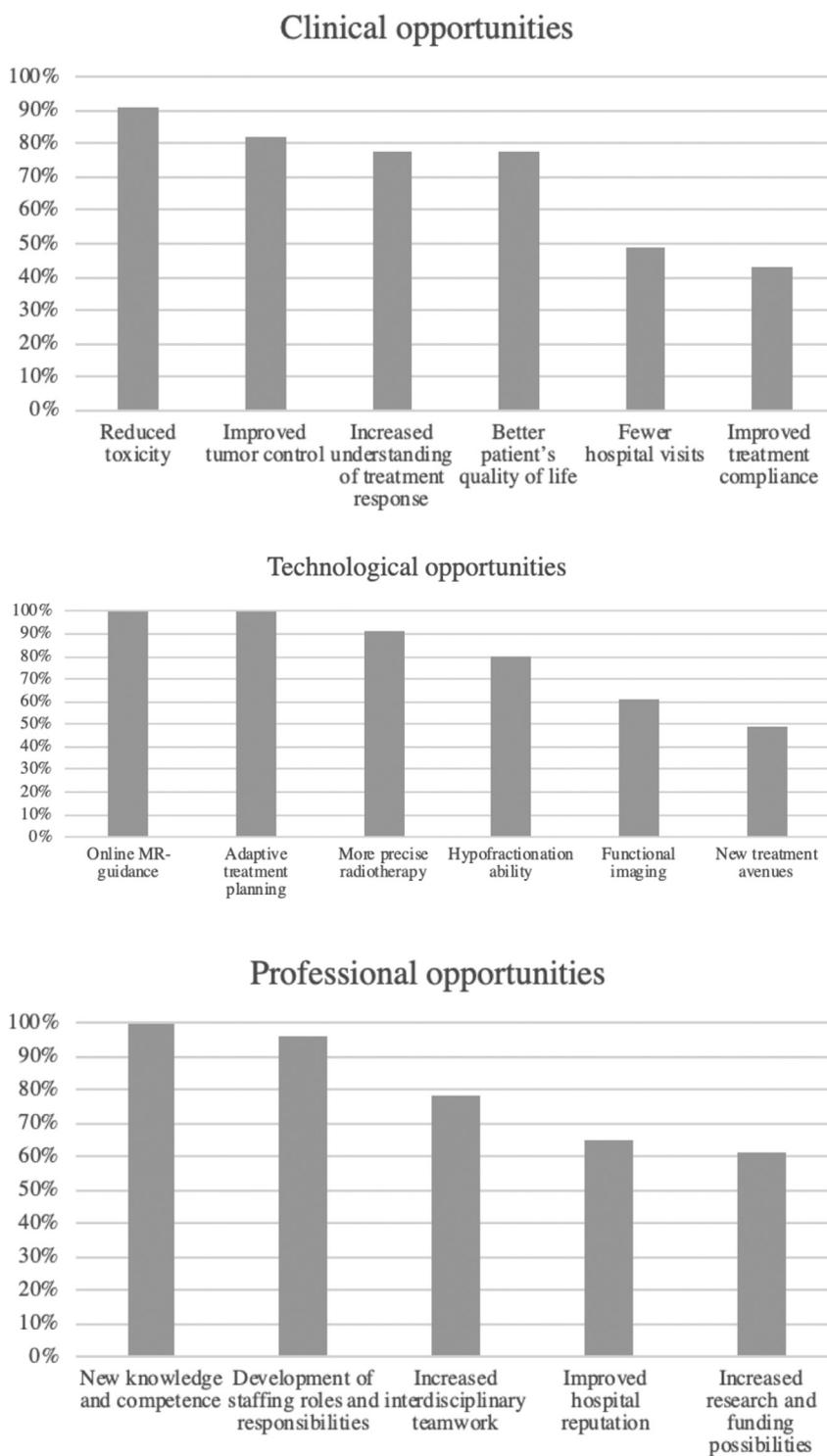


Fig. 1 Frequency of interviewees in percentages who discussed opportunities to MR-Linac implementation in US hospitals, by main theme and subtheme.

Given the ongoing technology development, interviewees felt they could also benefit from research opportunities with funding potentials, which positively affect their own careers. To elaborate, implementing the MR-Linac is perceived by technology adopters to

access prestigious technology which could allow delivery of higher-quality care. Being perceived as a “pioneer” in this area by providing this new technology could also be greatly beneficial for clinical expert and hospital reputation.

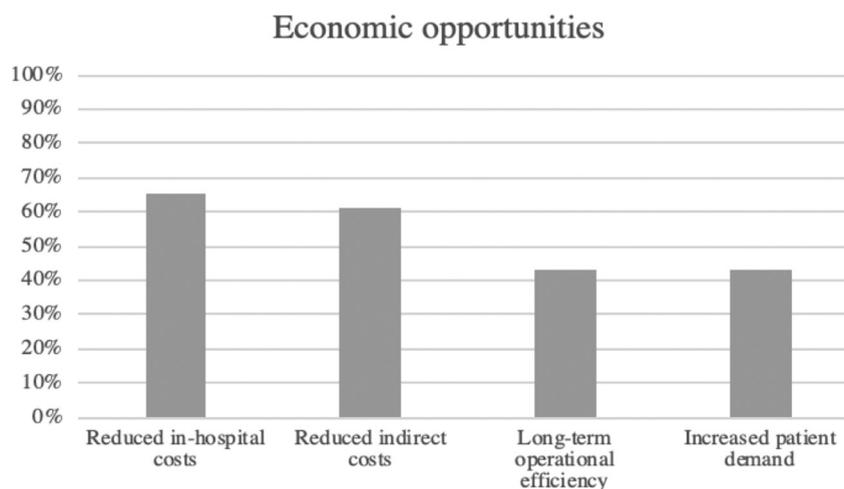


Fig. 1 Continued.

Economic opportunities

Most interviewees expected long-term efficiency and operational benefits from implementing the MR-Linac. Automation of elements in the clinical workflow, such as planning and contouring, were perceived to increase efficiency. Furthermore, the possible absence of pretreatment planning within an MR-only online workflow could decrease the duration of the total care pathway. Another example, the increased potential for hypofractionation could allow operational efficiencies for both the provider and the patient, such as decreased treatment sessions as well as interruptions and hospital admissions.

Operational efficiencies were also expected to evolve over time as technological development continued. Interviewees reported communication efficiencies, for example by the automation of tasks together with more consistent multidisciplinary communication and reporting. Furthermore, interviewees emphasized that the clinical benefit of decreased radiation related side effects and less treatment fractions could result in improved overall efficiency of hospital systems (eg, lower burden on anesthesia provision).

To elaborate, a possible reduction in the workforce per treatment of technology use is expected. On the one hand, the ongoing software development is expected to enable an automated process with faster quantification of the dose, which may lead to faster treatment delivery with reduced staff time and requirement (eg, less presence of the radiation oncologist and physicist during actual treatment delivery). On the other hand, professional experience and learning curves can reduce the education time needed to use the technology effectively.

Implementation of the MR-Linac was also thought to potentially improve economic outcomes of radiation oncology departments by providing state-of-the-art

treatment, particularly in the US health care system where health services are privatized and hospitals often compete against one another for patient referrals. Hence, interviewees mentioned that implementing the MR-Linac technology could possibly attract more patients with cancer to the providing center. One interviewee stated: “Especially well insured patients have this belief that that as Americans, we should have the latest and greatest technologies, almost regardless of cost.”

Barriers

We identified 5 barriers for implementation of the MR-Linac technology in US hospitals. These barriers were technology, professional, organizational, market, and regulatory related. [Figure 2](#) shows the percentages of the interview cohort who discussed the barriers, by main theme and subtheme.

Technological barriers

Current radiation therapy centers often lack MR-imaging facilities. Technology users therefore face substantial investments for implementing the MR-Linac in the construction of technical facilities, maintenance, information technology, safety assurance, human resource policy, and personnel training. To elaborate, the combined functionality of both the MRI device and the radiation delivery device raises technical complexity and necessitates the acquirement of new skills and additional understanding of MRI sequences.

Furthermore, the current lack of comparative effectiveness data and the ongoing technical development complicates a clear identification of potential benefits and return

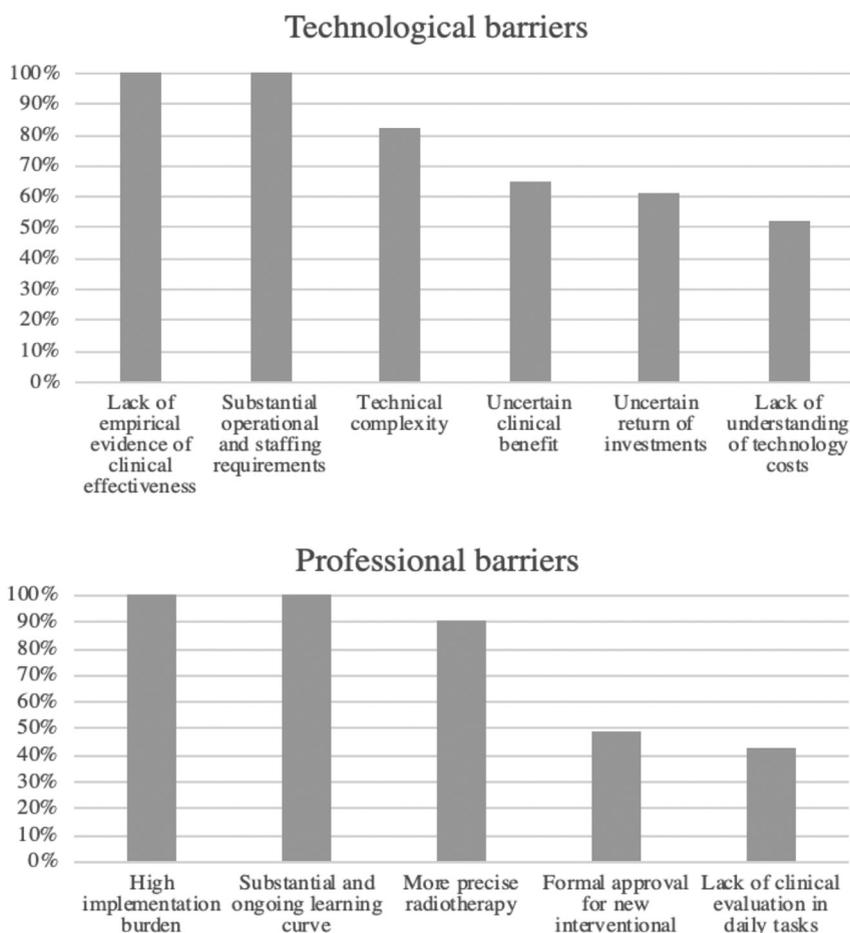


Fig. 2 Frequency of interviewees in percentages who discussed barriers to MR-Linac implementation in US hospitals, by main theme and subtheme.

of investment. Although the MR-Linac was expected to increase the efficiency and precision of existing radiation therapy treatment, interviewees were at the same time critical of the actual clinical effect given the current lack of comparative effectiveness data. The previously mentioned advantages, such as online high-resolution imaging during treatment and functional imaging have yet to be proven in clinical studies.

The interviewees also indicated a potential lack of understanding by technology users of the costs associated with MR-Linac treatment, especially as software development continues and the true treatment outcomes will evolve. One interviewee stated: “What would the cost be? What numbers are needed to actually make it a beneficial investment from a capital standpoint while the technological development still continues? I think that’s the main discussion.”

Professional barriers

Technology adopters may face hurdles from the burden of technology implementation in the radiation oncology

department to staff training of staff as well as treatment delivery to patients. As mentioned previously, several interviewees indicated a high investment burden and strategic decisions at a hospital level in early stages of technology implementation. To elaborate, technology users would need to obtain formal training to safely use the technology and to acquire MRI knowledge and skills given the routine use of MRI. Software developments inherent to the MR-Linac’s development require technology users to anticipate continuous learning. Hence, the implementation of technical advancements would need to be adopted into professional expertise before the actual use of the technology could flourish. One interviewee asked: “Do we have appropriate algorithms that are rooted in the science, that are also rooted in the clinical expertise of our medical staff?”

Another challenge is the formal approval for the introduction of new treatment workflows into clinical practice due to the lack of empirical evidence of effectiveness. Interviewees illustrated the importance of incorporating findings, as research data become available, into local clinical guidelines for appropriate technology usage in a timely manner. Therefore, interviewees also emphasized the importance of comparative effectiveness evaluation

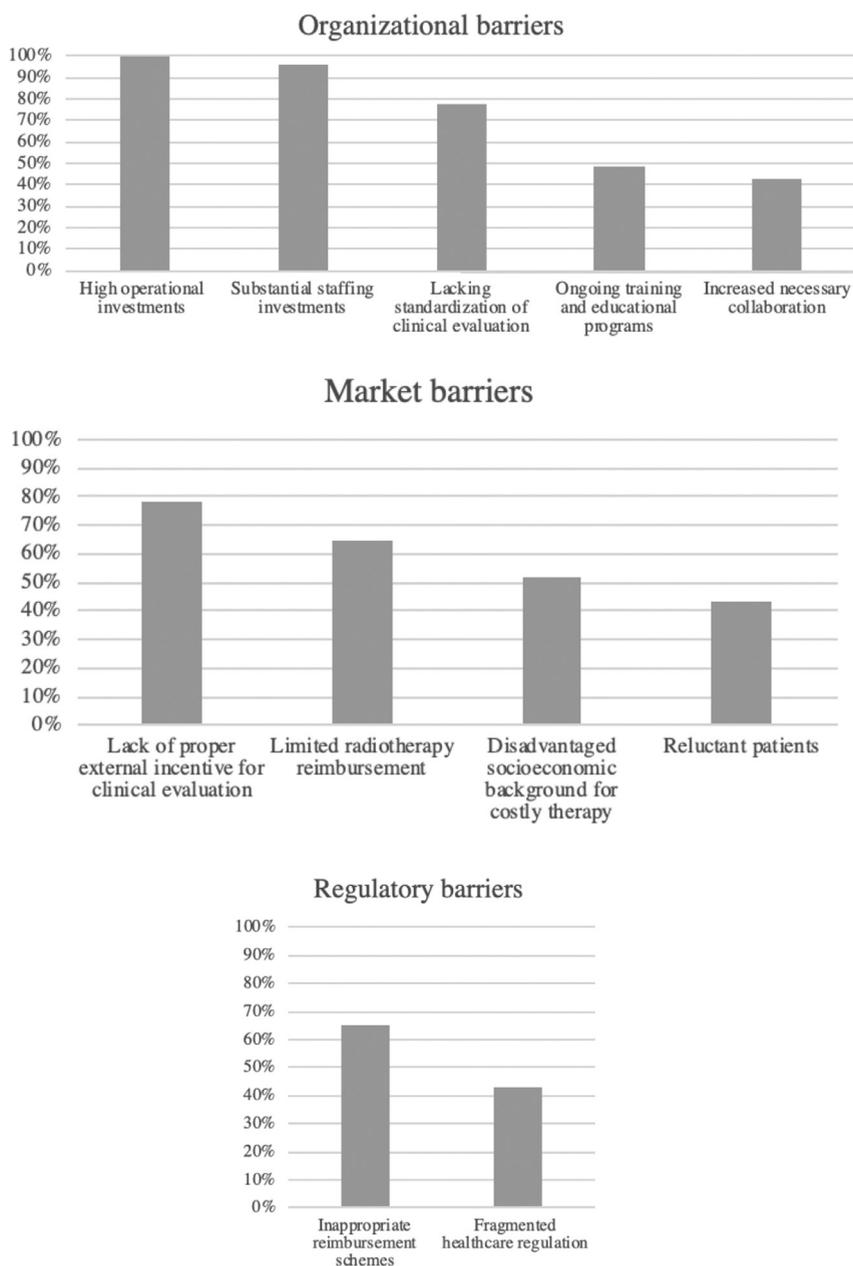


Fig. 2 Continued.

for technology implementation. Yet, there could be conflicts in prioritizing clinical treatment versus effectiveness research, given limited resources allocation at hospitals, due to the ability to have only one MR-Linac machine and the care burden because of the COVID-19 pandemic. Clinical evaluation was therefore not widely accepted as a main research objective among all interviewees.

Organizational barriers

As stated previously, the technology supply model requires substantial structural and staffing investments

for factors such as workflow, quality assurance, and the development of protocols. Alongside these implementation investments, the ongoing technology development requires radiation oncology departments to maintain training and educational programs. Furthermore, interviewees expected an increased necessary interaction between radiation oncology departments as well as supporting medical specialties such as radiology who would aid in target delineation.

Interviewees also emphasized that the COVID-19 pandemic was also an important factor leading to changes in resource allocation at the hospital; hence, for more commitment of human resources to direct care provision

rather than secondary tasks, such as expertise training and clinical evaluation of new technology. One interviewee noted: “Discussions of decreased reimbursements for radiation delivery are important. A lot of resources are being reallocated to fight the COVID-19 pandemic. So, it is actually a pretty uncertain time economically.”

Interviewees perceived the fact that value assessment and clinical trials (eg, demonstrating empirical evidence of [cost] effectiveness) not being integrated into departmental strategy and culture including vision, goals and key performance indicators, as a significant barrier to identifying the potential value offered by technology implementation.

Market barriers

Interviewees felt that they may not be appropriately motivated or financially compensated when it came to proving the effectiveness of the MR-Linac technology. For example, some felt that the external authorities focused more on safety aspects than on demonstrating clinical effectiveness or added-value for the patient. Furthermore, interviewees discussed that decreased financial reimbursement for new radiation therapy delivery techniques may not allow possible financial gains from implementing MRgRT (eg, a relative reduction in number of treatment fractions and hospital visits as a result of hypofractionation compared with traditional external beam radiation therapy).

The interviewees also addressed the implementation challenge of the presence of patient groups with disadvantaged socioeconomic background for new, costly therapy. As an example, the main hospital in which this study was conducted was identified as being socioeconomically diverse. This location included both wealthy, insured patients which will be beneficial for costly treatment implementation such as the MR-Linac, compared with relatively poor or uninsured patients, which may face hinder to afford costly therapy. Furthermore, while well-insured patients may hold the belief that Americans should have the “latest and greatest” medical therapy options available to them regardless of cost, other more historically marginalized communities (eg, African Americans) may be more reluctant to try innovative treatment such as the MRgRT.

One interviewee stated: “There are a lot of undertreated, underdiagnosed cancer patients here in this inner city. So, if you're using your resource dollars, you'll get the most benefit from a place like [city]. There is a negative to that. And part of that is patient's reluctance to come in and try new technology given historical health disparities with race and other socioeconomic status. It's both a huge opportunity, but also a challenge, especially here in [city].”

Regulatory barriers

Interviewees identified the lack of appropriate reimbursement arrangements as a barrier to implementing the MR-Linac. As stated previously, interviewees felt more motivated by federal regulators to demonstrate safety than clinical effectiveness for reimbursement of care. Interviewees also indicated a potential lack of consensus among clinical care providers and nonclinical reimbursement entities in regards to health care cost reimbursement. The definition of insured care costs would be more focused on how much the government or insurers were willing to pay to providers, rather than the costs incurred by providers to deliver treatment. The existence of such gaps between clinical care providers and reimbursement entities was experienced by interviewees to discourage the demonstration of value of new technology.

Furthermore, as health care and reimbursement are structured differently across states and individual hospitals in the US, some interviewees recognized that this fragmentation could hinder collaboration between MR-Linac providers when implementing the technology as well as having reproducible results across institutions.

Discussion

Our findings will help US hospitals to identify key points in their strategy when implementing the MR-Linac. MR-guidance in radiation therapy has been perceived a technical advancement with potential for therapy improvement and better patient outcomes, as well as scope for research and professional development.^{17,30,31} Yet, the technological complexity, the substantial operational and staffing investments, and the lack of empirical evidence of clinical effectiveness raise implementation uncertainties. Our findings also show that the lack of appropriate reimbursement and regulatory structures may complicate the actual deployment of the potential technology value.^{17,32} The convergence of diagnostic and therapeutic realms in the MR-Linac technology, confirms the essential task of solving the common gap between technical innovations in the field of radiation oncology and current national associated treatment guidelines as well as present reimbursement structures.³³⁻³⁵

The opportunities and barriers identified are fairly similar to those in literature such as the Dutch study on the implementation of the MR-Linac.¹⁹ Both in the US and Dutch cases, findings were consistent about the technology aspects, knowledge, and competence needed, and what it expects from radiation oncology departments to use new interventional procedures on the MR-Linac. Hence, the use of new interventional procedures on the MR-Linac with continued software developments require

ongoing staffing learning and workflow adaptations, approval from national regulatory authorities before clinical usage and necessary interactions with policy-makers and payers.

There is also consistency in the professional prospects, such as the development of staffing roles and required ongoing learning, and a higher hospital quality profile with potential economic growth. Interviewees in both cases were, however, also critical of the actual clinical added-value of the MR-Linac given the current lack of comparative effectiveness data. Defining the clinical value and operational profits of the MR-Linac at this time would be more of a challenge, given the ongoing technology development and indefinite treatment outcomes at present.^{21,36-39} The relative cost-effectiveness of the MR-Linac will vary across geographic settings due to specific treatment standards and patient populations in a particular health care system.

The inference of the different geographic contexts in the opportunities and barriers to the implementation of the MR-Linac shows similarities, but also differences. To illustrate, in the US case, interviewees mentioned potential challenges regarding certain patient groups with a reluctant attitude toward new therapy. In this geographic setting, interviewees also explicitly stated that they perceive few external stimuli to evaluate clinical effectiveness. The Dutch case, in contrast to the US case, reflected more explicitly on the intra- and interorganizational network required in technology implementation, such as the cooperation with the referring physician because of competitive specialties and payers to ensure treatment access and reimbursement. Thus, the implementation of the MR-Linac technology may encounter opportunities and challenges that are more relevant to one specific health care system.

The identified opportunities and barriers to implementing the MR-Linac could relate to each other. Market influences on technology implementation could be caused by regulation while also raising technological barriers.^{1,40} For instance, governmental decision-making informs reimbursement requirements and may therefore stimulate technology users to evaluate clinical effectiveness as well as to deliver new insured treatment regimes.⁴¹ Our findings also showed that technology users may not feel motivated to evaluate the clinical effectiveness by external bodies as well as face hinder in technology implementation due to the lack of appropriate reimbursement structures. Ultimately, securing proper reimbursement structures and regulatory approval for new interventional procedures, are essential for technology implementation.

Although market barriers could result from regulatory barriers, cultural barriers may determine these regulatory barriers.⁴² Our findings confirm that the lack of effectiveness evaluation in the radiation oncology department's strategy hinders to prove the technology's value as well as the steering and development of appropriate treatment

guidelines and staffing protocols. Regulation and policy such as financial incentives or cost sharing have an important albeit indirect effect on clinical decision making and staffing allocation.⁵ Thus, the implementation of the MR-Linac includes factors at clinical and technical level, but also at cultural, organizational, market, as well as governance level which can influence each other.

Our study provides the first early multifaceted assessment of opportunities and challenges to MR-Linac implementation in US hospitals, which inform local technology adopters to improve and facilitate treatment access.^{34,43,44} Technology users may face influences during implementation that are outside their ordinary scope of work. Because these influences are interrelated, the engagement with other health care decision-makers such as insurers, regulators, and policymakers could be relevant. Future efforts should generate empirical evidence of MR-Linac's effectiveness to prove expectations and justify return on investment concerns. The Multi-OutcoMe EvaluationN of radiation Therapy Using the MR-linac Study aims to generate empirical evidence of clinical effectiveness and safety, and to identify subgroups of patients who are most likely to benefit from the MR-Linac.⁴⁵

Approaches to assess the potential value of new medical technology, particularly those in early stages of development and implementation are gaining more attention.⁴⁶⁻⁴⁹ Early health economic analyses are useful for identifying areas in which new developing technologies could be cost-effective and conditions that must be met to achieve cost-effectiveness results.^{43,50} This analysis is particularly relevant for developing medical technologies such as the MR-Linac when both the costs and the effects of the innovation are still largely unknown, thereby guiding research and development and identifying potential meaningful treatment strategies.

The presented findings may be relevant for other convergent medical technology as the MR-Linac represents other technical trends in medicine, such as the application of artificial intelligence applications.³⁹ The converging characteristic and rapidly increasing role of digitization in medical technologies affects the user practices as well as the current organizational and regulatory settings, creating an additional dynamic context.⁵¹⁻⁵³ Our findings confirm this potential impact of MR-Linac implementation. In-depth knowledge about the functionality and configuration of convergent medical technology, and the redistribution of knowledge, responsibilities and care pathways as well as related regulation and market dynamics are important to further consider in future research for this type of technologies. Moreover, findings can be better interpreted across different health care systems when considering specific national dynamics, such as health care trends, socioeconomic, reimbursement, and regulatory structures.

Strengths of the present study include a thorough qualitative study design, which allows for maximal stakeholder

insight into questions asked. The study was conducted with insights from several US health care institutions and different types of stakeholders. Limitations of this study include difficulty contacting private and public insurance regulators, whose insights could have been useful regarding regulatory and policy factors affecting technology implementation. Furthermore, the COVID-19 pandemic during the study timeline limits the number of stakeholders interviewed and may influence the experiences and perspectives on MR-Linac implementation. Future research could include more implementation perspectives at another time point. Broader stakeholder perspectives across national borders can help to identify health care system-specific opportunities and barriers.

Conclusions

Our findings will help US radiation therapy oncology departments to identify main elements in their strategy when implementing the MR-Linac. Our study confirmed the current literature on MR-Linac implementation with technical and clinical prospects, but also reveals additional insights into professional, sociocultural, market, and regulatory challenges. We therefore address implementation factors that are overlooked in the current MR-Linac literature and which in particular are crucial in the US health care system. This leads to new connections to facilitate appropriate introduction of the MR-Linac in US hospitals and is a start to understanding its long-term effects in health care systems.

Supplementary materials

Supplementary material associated with this article can be found in the online version at [doi:10.1016/j.adro.2022.100953](https://doi.org/10.1016/j.adro.2022.100953).

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