

INTRODUCTION

Background:

Risk modeling for breast cancer screening is a standard of care in many areas that focus on cancer outcomes. Overall, an American woman's lifetime risk of breast cancer (BC) is 1-in-8, or 12.5%. However, due to different environmental, lifestyle, and genetic risks, women have differing relative risk profiles. Currently, breast experts define above-average risk as 15-20% lifetime risk, and "high risk" as >20%. Rurally, risk strategies for risk-adapted BC screening are often age-based due to lack of access to services that include risk models used in the existing population. Often, high risk women are not screened appropriately or offered risk reduction strategies. We have developed a manual process for risk assessment that combines lifestyle/biological factors with genetic risks and have instituted this model in our radiology screening program at TOBH since 2018, initially using a nurse navigator to help direct these patients. Women fill out a manual paper questionnaire at the time of the screen, given by the radiology technicians, which answers questions that are then used to calculate a lifetime risk of BC score. The model selected for this population is the Tyrer-Cuzick model version 8b, based on previous data examined at this institution. This calculation requires manual input of all questions answered by patient (15-20) into a website (Ibis) that estimates the patient's risk of breast cancer over their lifetime and gives an age-adjusted estimate for the normal woman as well for comparison. This calculation must be done quickly by the rad techs, then included to the reading radiologist in time to be included in the body of the report. How efficiently this is done, has not been examined, but due to the elements of human factors and busy schedules, we anticipate this is not the most efficient process to capture high risk women. We believe we are not capturing most women who qualify for high risk and referring them appropriately. We anticipate it is a problem with lack of resources, lack of education, and lack of efficient communication of this information to the patient. We also hypothesize a program that depends on multiple layers of human communication with the patient is inherently inefficient, as we plan to develop an EMR-based platform that automates this process for the provider and the patient.

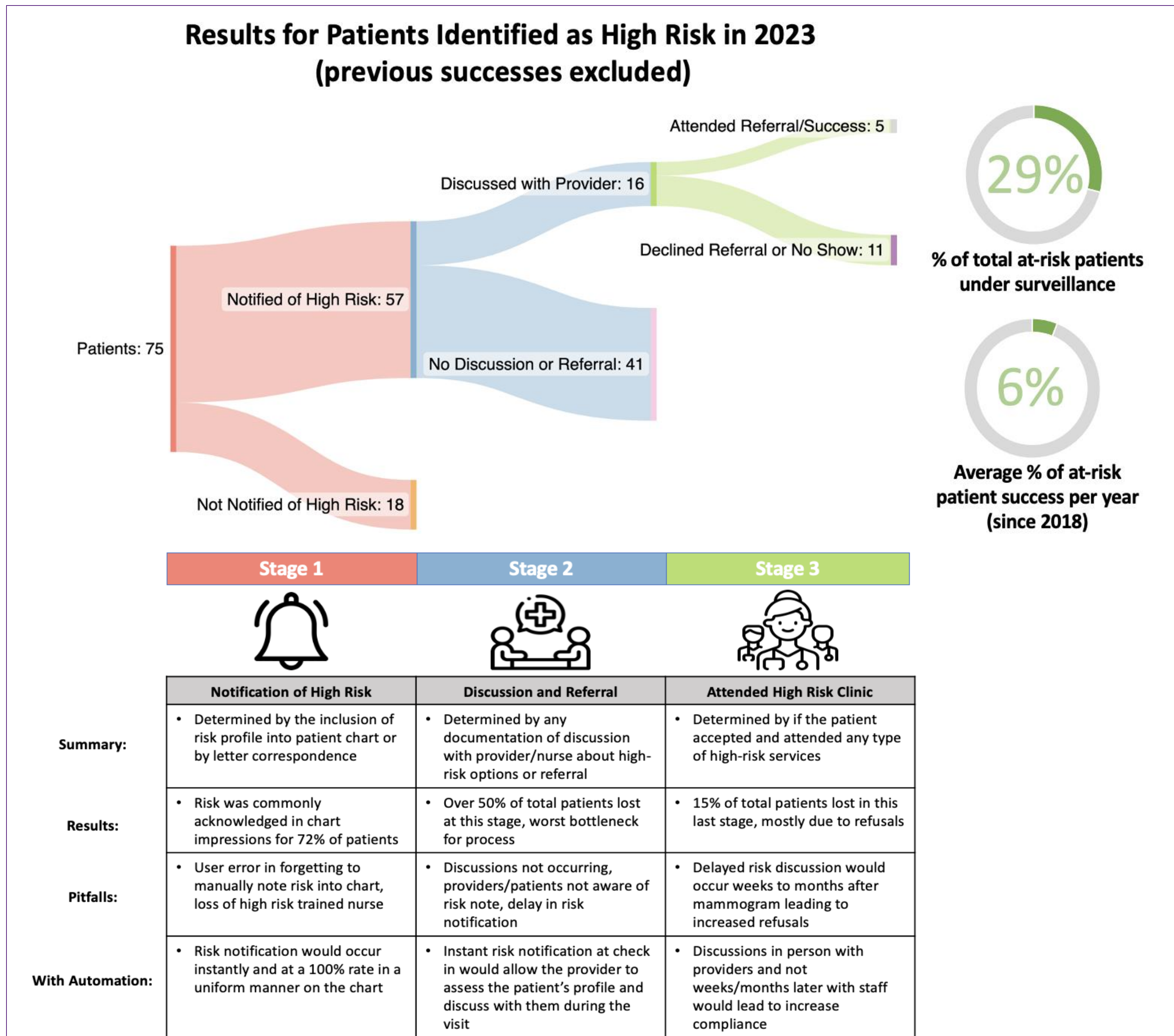
PURPOSE OF STUDY

To highlight the outcomes of patients identified as being at high-risk for breast cancer and identify the weaknesses in our process that an automated solution could address in the future

MATERIALS & METHODS

We looked at patients screened for BC at TOBH for 6 months for this project to obtain a recent estimate of our successes and failures in documenting high risk women. 2000 women underwent screening mammography from January 1 to June 1, 2023. Typically, TOBH screens 4500-5000 women a year on average. Average age for screening is 60 years, with ranges from 40 to 86 years of age. We obtained copies of the names of all women routinely screened for cancer. Patients undergoing diagnostic images were excluded for this study since we after the effects of this tool in screening risk, not in women presenting with a known mass or abnormality. N=2000 women met criteria for screening at TOBH, a critical access hospital. Once high-risk patients were identified, we tracked and documented every effort to get them referred to proper high-risk screening/treatment. The whole process from mammogram to attended referral was split into 3 stages: notification of high risk(1), discussion and offered referral(2), and attended high-risk clinic(3). Success and failure at each stage were recorded to determine where the bottlenecks were in our current manual process.

RESULTS



DISCUSSION

The total number of high-risk patients identified was 98. Of those 98, 23 had successfully been enrolled in high-risk care in the years prior since 2018. Among the remaining 75 patients, 57 advanced past Stage 1 because they received a notification detailing their risk profile within their chart or from a letter sent by the nurse navigator. However, 18 patients received absolutely no notice at all on their risk. In Stage 2, patients would advance in the process if they had a documented discussion with any staff about high-risk options or referrals. This stage was particularly unsuccessful, as we lost 41 patients and only 16 advanced. In Stage 3, patients advanced to a "success" if they accepted their referral and ultimately attended it. 11 patients either declined or failed to attend the appointment. Ultimately, in the first six months of 2023, only 5 high-risk patients of the possible 75 were captured into specialized care pipeline. The patients would go on to receive genetic testing, familial and lifestyle counseling, and more detailed and frequent imaging in order to monitor their risk profile. This personalized care will undoubtedly ease their worries and allows give them a better chance of detecting any signs of breast cancer earlier, as the Tyrer-Cuzick version 8b was demonstrated to be 6 times more accurate at predicting breast cancer than the current age-based model we use in North Carolina today(5).

Although the results look bleak, when it put into perspective of 2018 and onwards, the actual success rate of the total patients was around 29%, since 23 other high-risk patients were already received specialized care. Per year, the OBH averages capturing around 6% of their high-risk patients into these programs. In the first half of 2023, those 5 patients already account for 7% of the available people, which mean that if this trend continues then 2023 could be a very successful year. Additionally, the weaknesses in the current manual process are very visible and would all be improved with the incorporation of an automated tool.

CONCLUSION & FUTURE STUDIES

Because TOBH is an accredited breast program, it is one of the only hospitals in ECU Health that offers this type of risk assessment tool to all its screening population. TOBH is unique in that is piloting a risk assessment tool for women, stratifying the risk for BC over a woman's lifetime and comparing that risk to an age-adjusted control. We found 6-7% of women being screened met criteria for "high risk" as defined by TC model, with lifetime BC risk 20% or higher. Communication of these results to patients was variable. Some results were included by the radiologist in the body of the mammogram report (for the primary care to see), and some reports included the number as a % but no note as to the relevance. Some women had already been captured by previous iterations of the screening process, since mammography is often a repeated screen every 1-2 years. We noted various correlations with success, and some correlations with failures of communication of this information, which we help to use in developing pathways for incorporating this information into an EMR based model later this year. Ultimately, if we learn how to repeat the successes, and improve the failures, we anticipate we will have better participation in our risk clinics once we roll out the automated model in 2023/2024.

Specifically, at Stage 1, the most common pitfalls were human error and staff layoffs, which would result in no interpretation of the risk profile to the patient. Additionally, due to delays in getting this data into the chart, providers lacked the proper information to conduct a discussion (Stage 2) with the patient while they were still in their office. This would result in them having to backtrack to call or send a letter to the patient weeks or months after the original. In contrast, an automated process could instantly populate this value into the chart and notify the provider as soon as the patient enters that data into the questionnaire, which would vastly improve results for stages 1 and 2. Additionally, its reasonable to project that capture rates at the final stage 3 would improve as patients would have an immediate discussion in-person with relevant hospital staff. This would carry significantly more weight with patients in conveying the importance of their risk profile that a letter or call that arrives weeks after.

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