



Impact of Dermoscopy Training for Primary Care Practitioners on Number Needed to Biopsy to Detect Melanoma

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Abstract

Introduction: The incidence of melanoma is on the rise. In trained hands, dermoscopy aids in the differentiation of melanoma from benign skin growths, including melanocytic nevi. This study evaluated the impact of dermoscopy training for primary care practitioners (PCPs) on the number of nevi needed to biopsy (NNB) to detect a melanoma.

Methods: We conducted an educational intervention that included a foundational dermoscopy training workshop and subsequent monthly telementoring video conferences. We performed a retrospective observational study to evaluate the impact of this intervention on the number of nevi needed to biopsy to detect a melanoma.

Results: The number of nevi biopsied to detect one melanoma decreased from 34.3 to 11.3 following the training intervention.

Conclusion: Dermoscopy training for primary care practitioners resulted in a significant reduction in the NNB to detect melanoma.

Introduction

The incidence of melanoma is rising across the United States, with Maine having the tenth highest incidence in the nation.¹ This, in combination with a national shortage of dermatologists, means that primary care practitioners (PCPs) must play a pivotal role in early skin cancer detection.² Studies have shown that PCPs desire more training in dermatology and the evaluation of cutaneous lesions.³⁻⁵ To address this need, we offered a dermatology training intervention targeted for PCPs that included a 90-minute Triage Amalgamated Dermoscopic Algorithm (TADA) workshop, subsequent monthly telementoring sessions via the Extension of Community Health Outcomes project (ECHO)⁶, and equipping participating clinics with dermatoscopes.⁷

Dermoscopy is a noninvasive tool that allows for visualization of colors and structures in the epidermis and dermis at 10x magnification. In trained hands, dermoscopy improves the user's ability to differentiate benign melanocytic nevi from melanoma.⁸ Despite the benefits of dermoscopy, fewer than 15% of PCPs are trained to use a dermoscope.³ The Triage Amalgamated Dermoscopic Algorithm (TADA) was developed to aid novice

dermoscopists in the evaluation of skin growths.^{9,10} TADA training equips learners with the skills needed to differentiate benign from malignant dermoscopic images with high sensitivity and specificity.¹⁰ While it is known that the use of dermoscopy reduces the number needed to biopsy (NNB) to detect melanoma, most of this literature is in the dermatology setting.¹¹ Our research aims to test if meaningful reductions in NNB are possible in the primary care setting as well.

Methods

This project was reviewed by the Maine Medical Center Institutional Review Board and was deemed exempt. To study the impact of this intervention on melanoma detection in the primary care setting, we developed a skin biopsy database. We then compared the NNB to detect melanoma before and after the educational intervention.

Educational Intervention

Participants were eligible if they were a primary care attending, resident, or advanced practice provider (APP) in family medicine, internal medicine, internal medicine/pediatrics, or pediatrics. Participation was voluntary and no compensation was provided. TADA workshops were taught in-person or virtually 11 times over the course of 8 months to 267 PCPs. The workshop included training on benign and malignant skin growths.⁸ Following workshop participation, learners were invited to join monthly ECHO telementoring sessions on dermoscopy. Seventeen primary care clinics were provided with dermatoscopes.

Data Extraction and Analysis

Using the current procedural terminology codes, we extracted all skin biopsies performed in our health system by PCPs between November 5, 2013 and November 6, 2021 from our electronic medical record. We conducted a retrospective chart review of the biopsy data. We manually reviewed pathology reports, and we entered results into Research Electronic Data Capture software (REDCap). In this analysis, only biopsy results that revealed nevi or melanoma were included. A total of 1,236 biopsies were in the preintervention group and 1,042 biopsies were in the postintervention group. NNB was used to assess differences in the ratio of biopsies that had nevi versus melanoma outcomes over time. The NNB ratio is calculated by dividing the total number of biopsies (nevi and melanoma) by the number of biopsied melanomas.¹² A lower NNB indicates a greater accuracy in differentiating benign and malignant lesions.¹³ We used a χ^2 test of independence to evaluate statistical significance by between biopsy outcome type (melanoma or nevus) and time period (pre- or postintervention).

Results

Preintervention, pathology revealed a total of 1,200 biopsies were nevi and 36 were melanomas. Postintervention, pathology revealed a total of 950 biopsies were nevi and 92 were melanomas. The NNB of nevi to detect a melanoma decreased from 34.3 preintervention to 11.3 postintervention. The χ^2 test of independence examining the relationship between biopsy outcome type and time period was significant (χ^2 [1, N=11,523]=36.21, P<.001). The average annual NNB results were lower in the postintervention time period than the preintervention time period (Figure 1).

Conclusions

Following dedicated training of PCPs in our health system on the clinical and dermoscopic findings in benign and malignant skin growths, the number of nevi biopsied to detect a melanoma was reduced. While educational interventions do not consistently result in practice change, dermoscopy training, ongoing telementorship, and easy access to a dermatoscope led to widespread dermoscopy use⁹ and a meaningful reduction in NNB to detect melanoma.

Following the intervention, there was an increase in the rate of biopsies performed and the total number of melanomas detected in the primary care setting. This could be due to multiple factors: improved detection of melanoma due to the use of dermoscopy, increased likelihood to perform biopsies of suspect skin cancers at the primary care clinic (rather than refer to dermatology), and/or expansion of the primary care workforce.

Reduced NNB has implications not only on patient experience, but also on health care costs. Due to climate change, rates of skin cancer in the United States are predicted to increase by 10% by 2050.¹⁴ With the rising skin cancer incidence, PCPs need to be equipped with the necessary skills to evaluate skin growths accurately. Reducing the NNB has the potential to improve quality, reduce cost, and foster provider engagement, all important aspects of the quadruple aim.¹⁵

However, we must be cautious when interpreting these results, as overdiagnosis of skin cancer is a legitimate concern.¹⁶ In the postintervention time period, there was an increase in the number of melanomas being detected in the primary care setting. The goal of early melanoma detection is to reduce melanoma morbidity and mortality. This does come with the potential for detection of slow-growing indolent types of melanomas. However, at present, clinicians do not have reliable bedside diagnostic or FDA-approved genetic tests to aid in determining the metastatic potential of a melanoma. Furthermore, early-stage melanoma is amenable to low-risk outpatient surgery at a reasonable cost.

This study has limitations. The sample size for melanomas was small and the study was conducted in a single health system. Without a control group, we were unable to determine how much of the improvement in biopsy ratios and NNB was due to additional clinical experience. We also have not accounted for changes in skin cancer incidence in our patient population.

Tables and Figures

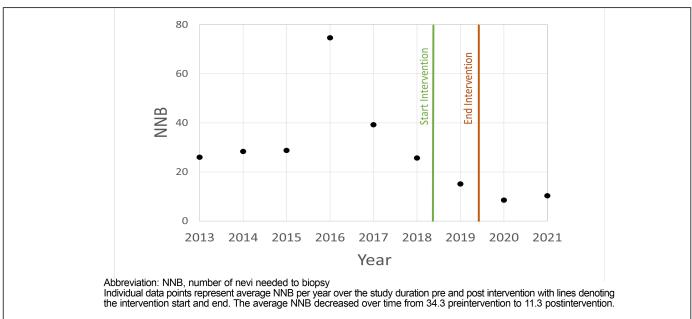


Figure 1: Average NNB to Detect a Melanoma per Year

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