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Journal Pre-proof

Use of teledermatology by dermatology hospitalists is effective in the diagnosis and management of inpatient disease.

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74 forward; telemedicine; dermatology hospitalists

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79 CAPSULE SUMMARY

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81 - Inpatient access to dermatologists is limited, highlighting an opportunity to utilize

82 teledermatology within the inpatient setting.

83 - Teledermatology in the inpatient setting may be a clinically acceptable option for

84 diagnosis, evaluation, and management. This may represent a novel and effective option

85 for hospitals.

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120 ABSTRACT:

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122 *Background:* Patient outcomes are improved when dermatologists provide inpatient consults.

123 Inpatient access to dermatologists is limited, illustrating an opportunity to utilize

124 teledermatology. Little is known about the ability of dermatologists to accurately diagnose and

125 manage inpatients using teledermatology, particularly utilizing non-dermatologist generated

126 clinical data.

127 *Methods:* This prospective study assessed the ability of teledermatology to diagnose and manage

128 41 dermatology consults from a large urban tertiary care center utilizing internal medicine

129 referral documentation and photos. Twenty-seven dermatology hospitalists were surveyed.

130 Interrater agreement was assessed by the kappa statistic.

131 *Results:* There was substantial agreement between in-person and teledermatology assessment of

132 the diagnosis with differential diagnosis (median kappa = 0.83), substantial agreement in

133 laboratory work-up decisions (median kappa = 0.67), almost perfect agreement in imaging

134 decisions (median kappa = 1.0), and moderate agreement in biopsy decisions (median kappa =

135 0.43). There was almost perfect agreement in treatment (median kappa = 1.0), but no agreement

136 in follow-up planning (median kappa = 0.0). There was no association between raw photo quality

137 and the primary plus differential diagnosis or primary diagnosis alone.

138 *Limitations:* Selection bias and single-center nature.

139 *Conclusions:* Teledermatology may be effective in the inpatient setting, with concordant

140 diagnosis, evaluation, and management decisions.

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144 BACKGROUND:

145 Tele dermatology is the remote dermatologic assessment of patients, in real-time (“live
146 interactive”), by accessing stored data (“store-and-forward”), or a combination of the two
147 (“hybrid”), with worldwide applications.(1) Tele dermatology has been studied in general triage,
148 consultation in remote locations, and monitoring of chronic skin conditions.(1) In addition to
149 increased access to dermatologists, potential benefits of store-and-forward tele dermatology
150 include cost reduction due to fewer face-to-face (FTF) consultations,(2) reduced travel time and
151 opportunity cost due to missed work,(3-5) and reduced contagion spread amid infectious disease
152 outbreaks.

153 Significant clinical evidence supports the outpatient use of store-and-forward
154 tele dermatology.(2-10) In contrast, tele dermatology has been studied in the inpatient setting to a
155 limited degree. A significant practice gap exists between the demand for inpatient dermatology
156 services and access to dermatologists,(11, 12) often a source of frustration for inpatient providers
157 and patients. Dermatology hospitalists represent a clinical group with expertise in complex
158 medical dermatology and the diagnosis and management of skin diseases affecting hospitalized
159 patients. Involvement of dermatology hospitalists in the care of hospitalized patients has been
160 found to improve patient outcomes.(13) In a subset of cases, inpatient tele dermatology reduces
161 time for the primary medical team to receive a response for a dermatology consultation.(14)

162 Dermatologist interest in inpatient tele dermatology is high. A survey of attending dermatologists
163 demonstrated that 61.5% agreed or strongly agreed that tele dermatology helps inpatient care.(15)
164 Another study found that 95% of hospital and emergency department practitioners would utilize
165 a tele dermatology consult service if available, however only 5% believed that tele dermatology
166 would be equivalent to a face-to-face (FTF) consult.(16) This finding supports the need for

167 additional studies evaluating inpatient teledermatology, which may shift perception and
168 encourage adoption of inpatient teledermatology.

169 This study investigates the diagnostic and management agreement between inpatient FTF and
170 store-and-forward teledermatology evaluations utilizing remote digital evaluations for hospital-
171 based dermatology consultations.

172 METHODS

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174 Eligible patients for this study were admitted to Massachusetts General Hospital between July
175 and August 2013 and had a dermatology consultation staffed by a dermatology hospitalist with
176 more than six years of inpatient experience, defined as the Primary Dermatologist (PD). This
177 yielded a sample of 108 patients. Only those consultations with digital images and non-
178 dermatology evaluations involving the dermatologic complaint were included. Cases were
179 selected if the accuracy of the PD's diagnosis was able to be confirmed based on testing,
180 response to therapy and final diagnosis at discharge. Based on these inclusion criteria, a total of
181 42 patients were initially included (Figure 1). One case was excluded from analysis to preserve
182 the generalizability of study results,(17) as this patient presented with multiple concomitant
183 dermatologic complaints and the documentation did not specify the specific focus of the
184 dermatology consultation.

185 For teledermatology review, data abstractors not involved in the care of the included cases
186 packaged patient data into surveys by unique numerical patient identifiers. Each survey set
187 contained seven individual cases, randomly assigned to each survey set from the total case pool.
188 Each individual case contained the relevant history and physical exam notes generated by a non-
189 dermatologic internal medicine or emergency medicine provider. In addition, all data such as
190 laboratory studies, imaging, microbiology, pathology, and digital images up to the day of the

191 consult that would have been available to the PD were included. Finally, a
192 diagnosis/management questionnaire was included. The order of case examination within each
193 survey set was fixed across all TDs. Patient identifiers were uniquely created and stored safely.
194 This study was approved by Partners Institutional Review Board (IRB) #2018P002762.
195 Only non-dermatologic patient history and physical exam notes were included to mimic real-
196 world settings. Photographs were captured primarily by Dermatology Residents from the
197 Harvard Combined Dermatology Residency. Camera use was heterogenous and included Sony
198 NEX5N 12MP and 5MP iPad Mini. Images were obtained both by using the original digital
199 images and screengrabs from the electronic medical record. Study data were collected and
200 managed using Research Electronic Data Capture (REDCap) tools hosted at Partners.(18, 19)
201 The packaged cases were sent to 27 experienced dermatology hospitalists in order of response to
202 request for participation at various academic institutions across the U.S. Each remote
203 teledermatologist (TD) received six to seven cases within a secure REDCap survey
204 (Supplemental Figure 1). Each clinical case was evaluated by 4-5 unique TDs.
205 The surveys included the option to list a primary diagnosis as well as a maximum of three
206 differential diagnoses. The workup and management plans offered were as follows: (1) biopsy,
207 (2) topical therapy, (3) systemic/oral therapy, (4) microbiology, (5) labs, (6) transfer to the burn
208 unit, if not already there, (6) recommend continued patient monitoring as an inpatient, and (7)
209 recommend follow-up as outpatient for dermatologic condition. Once the TD selected a
210 treatment plan, s/he was prompted for free-text details. Both the correct mode and type of
211 therapy was assessed. If the selected treatment differed between the PD and the TD but both
212 options were within the accepted standard of care for that disease, these treatments were

213 considered concordant. This was to minimize the effect of stylistic practice differences in
214 grading appropriateness.

215 The follow-up plan options were: (1) sign-off and no need for future follow-up either inpatient or
216 outpatient, (2) outpatient follow-up, no need for additional inpatient dermatology evaluations
217 (“sign off”), (3) no need to see the patient tomorrow, but evaluate if the primary team requests
218 and ensure outpatient follow-up planned, and (4) see the patient tomorrow and follow closely.

219 TDs rated their degree of comfort in managing the case as a dermatologist, as well as the quality
220 of each image.

221 Outcomes measured were concordance between the PD and the TDs for the following: primary
222 diagnosis, primary diagnosis plus differential diagnosis, decision to biopsy, laboratory work-up,
223 imaging, treatment, and follow-up plan. Primary outcomes were defined as primary plus
224 differential diagnostic concordance as well as management plan concordance, the rationale of
225 which was to assess whether teledermatology could result in an appropriate work-up and
226 management leading to an effective outcome for the patient. Secondary outcomes were primary
227 diagnostic concordance alone, as well as concordance in work-up.

228 Primary diagnostic concordance was defined as agreement between the primary diagnosis
229 provided by the PD and the TD. Primary diagnostic plus differential diagnostic concordance was
230 defined as the PD’s diagnosis being among the differential diagnosis of the TDs in cases when
231 the primary diagnosis was discordant. The diagnoses themselves, and not diagnostic family, were
232 used in calculating diagnostic concordance.

233 *Statistical Analysis*

234 We calculated the prevalence-adjusted bias-adjusted kappa (20) to quantify the concordance
235 between a) the TDs’ and PD’s primary diagnosis, b) TDs’ primary diagnosis plus differential

236 diagnosis and PD's primary diagnosis, and c) TDs' and PD's management plan (separately for
237 each of the five domains: biopsy, work-up, imaging, treatment, and follow-up). The following
238 criteria were used to assess significance: values ≤ 0 as indicating no agreement, 0.01–0.20 as
239 none to slight, 0.21–0.40 as fair, 0.41–0.60 as moderate, 0.61–0.80 as substantial, and 0.81–1.00
240 as almost perfect agreement.⁽²¹⁾ We evaluated the associations of the calculated concordance a)
241 and b) with TDs' years of experience and the reported photo quality rating, and the associations
242 of the calculated concordance c) with photo quality using the Pearson correlation coefficient. We
243 also evaluated the associations of TDs' level of comfort managing patients (with photos and
244 story alone) with photo quality and TDs' years of experience using the Wilcoxon rank sum test.
245 All were conducted using R version 3.6.1 (<https://www.r-project.org/>).

246 RESULTS

247 Table 1 depicts the characteristics of the patients included in the study surveys. The mean age
248 was 54.1 years (standard deviation (SD) 23.7), 43.9% were female, 75.6% identified as
249 Caucasian, and 68.3% as Non-Hispanic or Latino. The final diagnoses are provided that were
250 used to evaluate diagnostic concordance. Diagnoses fell under a diverse set of diagnostic
251 families, consisting of hypersensitivity reactions (29.3%), vascular (19.5%), infectious (17.1%),
252 inflammatory (17.1%), neoplastic (7.3%), iatrogenic (4.9%) and traumatic (4.9%).

253 The TDs were 40.7% female and practiced in diverse academic institutions from all geographic
254 regions of the United States. The mean number of years' experience of each of the TDs was 7.0
255 (SD 1.2) (Table 2). Out of all cases, 45.1% of TDs felt comfortable managing the case as a
256 teledermatologist. The mean number of differential diagnoses per TD per individual case was 2.6
257 (SD 0.4).

258 There was fair concordance between PD and TD primary diagnosis alone (median concordance
259 66.7%, interquartile range (IQR) 57.1% to 78.6%; median kappa=0.33, interquartile range (IQR)
260 0.14 to 0.57), with substantial agreement between PD and TD primary plus differential diagnosis
261 (median concordance 91.7%, IQR 85.7% to 92.9%; median kappa=0.83, IQR 0.71 to 0.86).

262 There was substantial agreement in pursuing additional laboratory work-up (median concordance
263 85.7%, IQR 85.7% to 92.9%; median kappa=0.67, IQR 0.43 to 0.79), and almost perfect
264 agreement in imaging decisions (median concordance 100%, IQR 50.0% to 100.0%; kappa=1.0,
265 IQR, 0.0-1.0). There was moderate agreement in the decision to biopsy (median concordance
266 71.4%, IQR 53.6% to 85.7%; median kappa=0.43, IQR 0.07 to 0.71). There was almost perfect
267 agreement in treatment plans (median concordance 100%, IQR 85.7% to 100.0%; median
268 kappa=1.0, IQR 0.67 to 1.0). There was no agreement in the follow-up plan (median
269 concordance 50.0%, IQR 42.9% to 66.7%; median kappa=0.0, IQR -0.14 to 0.14). Figure 2 is a
270 pair of histograms depicting the distribution of kappa values for agreement between the TDs' and
271 the PD's primary diagnosis (Figure 2A), and primary plus differential diagnosis (Figure 2B).

272 There was no association between experience of the TD and primary plus differential diagnostic
273 concordance (correlation=-0.27; 95% confidence interval (CI) -0.59 to 0.12, scatterplot in
274 Supplemental Figure 2, corresponding Supplemental Table 1) or primary diagnostic concordance
275 (correlation=-0.27; 95% CI, -0.59 to 0.12). There was also no association between years'
276 experience of the TD and decision to pursue laboratory evaluation (correlation=-0.19; 95% CI, -
277 0.53 to 0.21), biopsy (correlation=-0.32; 95% CI, -0.62 to 0.07), imaging (correlation=-0.19;
278 95% CI, -0.53 to 0.21), treatment decisions (correlation=-0.18; 95% CI, -0.53 to 0.21), and
279 follow-up planning (correlation=-0.06; 95% CI, -0.33 to 0.43).

280 There was no association between either raw photo quality and the primary plus differential
281 diagnosis (correlation=0.008; 95% CI, -0.18-0.19), or primary diagnostic concordance alone
282 (correlation=-0.07; 95% CI, -0.12-0.25). The Wilcoxon rank sum test of the TDs' comfort with
283 managing the case and years of experience indicated that TDs with fewer years of experience
284 were more likely to feel comfortable managing the patients as a teledermatologist (p=0.04).

285 DISCUSSION

286 This study illustrates that store-and-forward teledermatology may be reliable in the academic
287 inpatient setting, with strong agreement between PD and TD for diagnosis, work-up, and
288 management.

289 The high concordance of primary plus differential diagnosis is in-line with prior outpatient
290 literature,(8, 22) with studies demonstrating diagnostic concordance ranging from 41% to 100%
291 for store-and-forward cases.(2) This finding builds upon limited studies evaluating the use of
292 teledermatology in the inpatient setting.(12, 23, 24) As with prior study,(2) diagnostic
293 concordance improved when the differential diagnosis was taken into account.

294 The decision by TDs to pursue work-up in this study was highly concordant, with substantial
295 agreement in the laboratory work-up desired. However, there was only moderate agreement in
296 the decision to biopsy, which is in contrast with a prior inpatient teledermatology study finding a
297 >95% concordance in assessing need for biopsy.(12) This may be due to stylistic practice
298 differences or individual comfort level.

299 The treatment plans offered by the TDs were highly concordant with those of the PD, suggesting
300 that the outcomes of each patient may have been the same if managed by teledermatology, even
301 in cases where the primary diagnosis differed. This may be due to the high concordance of
302 primary plus differential diagnosis, leading to treatment plans applicable to multiple diagnoses.

303 The baseline inter-dermatologist variability that occurs even with face-to-face consultations must
304 also be taken into consideration, as a previous study of face-to-face, clinic-based dermatologists
305 has found diagnostic testing to be 85% concordant, medical-based therapy to be 85% concordant,
306 and clinic-based therapy 77% concordant, respectively.(22) Thus, some degree of discordance
307 may be expected.

308 The lack of concordance between TDs and the PD for follow-up plans suggests that in-person
309 evaluation may be needed prior to disposition planning. Stylistic differences also likely played a
310 role. Patient-specific factors may go into disposition planning, such as access to resources and
311 health literacy, which may contribute to the discordance between the PD and the TDs. Further
312 study of follow-up planning is needed to elucidate whether teledermatology may be reliable for
313 this use.

314 Photo quality was not associated with primary diagnostic concordance or primary plus
315 differential diagnostic concordance. This suggests that even in cases in which image quality is
316 suboptimal, the reliability of teledermatology may not be impacted. However, while the authors
317 utilized images from heterogeneous sources, many photos utilized in the study surveys met the
318 minimum standards recommended for teledermatology.(25) Additionally, assessment of image
319 quality was not broken down into detailed components, such as lighting, focus, or capture of
320 clinically-relevant information. Photo quality and training in obtaining photos may be needed to
321 ensure good capture of the relevant areas when implementing teledermatology, as the study
322 photos were captured by dermatology resident physicians.

323 There was no association between experience of the teledermatologist and diagnostic
324 concordance, illustrating the generalizability of teledermatology across all ages of practicing
325 dermatologists.

326 There appeared to be a disconnect between concordance and the TDs' level of comfort in
327 managing each case as a teledermatologist. The TDs considered themselves comfortable less
328 than half of the time; however, their survey responses often aligned with the PD. This may be in
329 part due to the novelty of teledermatology. The TDs with fewer years of experience were more
330 likely to feel comfortable managing the case, aligning with prior literature,(26) reflecting an
331 opportunity to utilize teledermatology even in novice practice settings. Similarly,
332 teledermatology exposure in residency may correlate with comfort of use,(27) suggesting that
333 early incorporation of teledermatology in training may facilitate its implementation.

334 One of the greatest strengths of this study is the large sample size of TDs, mimicking the
335 heterogeneity of applying teledermatology to real-life practice settings. The distribution of
336 diagnoses included in this study reflects that of common dermatology consultations.(13)

337 Limitations of this study include its single-center nature and the fact that dermatology residents
338 captured the clinical photos. The dermatology residents may have had a more thorough
339 understanding of how to obtain a high-quality dermatology photo than non-dermatology staff,
340 who would be submitting the teledermatology consult in real-life. Training of non-dermatology
341 staff in obtaining high-quality images may be needed. On the other hand, camera technology has
342 likely improved today and may lead to heightened quality of photos in today's use of
343 teledermatology. Further study is needed to determine best practices for implementing an
344 inpatient teledermatology program.

345 In conclusion, teledermatology may be effective for managing dermatologic disease in the
346 inpatient setting and leads to highly concordant diagnostic, work-up, and management decisions
347 when performed by experienced inpatient dermatologists. This may represent a novel and
348 effective option for community hospitals and may be particularly applicable during times of

349 concern for spread of infectious disease, such as during the 2019-2020 outbreak of the severe
 350 acute respiratory syndrome coronavirus 2 (SARS-CoV-2).

351

352 **Table 1.** Demographic characteristics of patients included in this study.

Patient characteristic	Total (n=41)
Age in years, mean (SD)	54.1 (23.7)
Sex, n (%)	
Female	18 (43.9)
Race, n (%)	
Asian	2 (4.9)
Black or African American	4 (9.8)
Caucasian	31 (75.6)
Unknown	4 (9.8)
Ethnicity	
Hispanic or Latino	0 (0.0)
Not Hispanic or Latino	28 (68.3)
Unknown	13 (31.7)
Dermatologic consultation characteristics	
Chronology of skin findings, median (IQR) (days)	4.0 (2.0-14.0)
Medications, mean (SD)	7.0 (3.7)
Final diagnostic categories	
Hypersensitivity	12 (29.3)
Contact dermatitis (4)	
Drug hypersensitivity (6)	
Erythema nodosum	
Urticaria	
Vascular	8 (19.5)
Calciphylaxis	
Henoch-Schonlein purpura	
Leukocytoclastic vasculitis	
Lipodermatosclerosis	
Small vessel vasculitis	
Stasis dermatitis (3)	
Infectious	7 (17.1)
Atypical mycobacterial infection	
Bullous impetigo	
Eczema herpeticum	
Herpes simplex virus	
Erythema chronicum migrans (2)	
Varicella zoster virus	
Inflammatory	7 (17.1)
Atopic dermatitis	
Gout	

Granulomatous disease	
Hiradenitis suppurativa	
Miliaria rubra	
Pyoderma gangrenosum (2)	
Neoplastic	3 (7.3)
Carcinoma erysipeloides	
Kaposi sarcoma	
Nevus lipomatosus	
Iatrogenic	2 (4.9)
Steroid acne	
Warfarin skin necrosis	
Traumatic	2 (4.9)
Bateman's purpura	
Neurotic excoriations	

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Table 2. Characteristics of the surveyed teledermatologists.

Characteristic	Total (n=27)
Sex, n(%)	
Female	11 (40.7)
Geographic distribution	
Northeast	13 (48.2)
Midwest	5 (18.5)
West	5 (18.5)
Southeast	3 (11.1)
Southwest	1 (3.7)
Years of experience, mean (SD)	7.0 (1.2)

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445 FIGURE LEGENDS

446 **Figure 1.** Selection criteria for cases to include in study.

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448 **Figure 2.** Distribution of Kappa values for agreement between the teledermatologists' and the
449 primary dermatologist's (A) primary diagnosis and (B) primary plus differential diagnosis.

450

451 SUPPLEMENTS

452 **Supplemental Figure 1.** Sample case within a survey set provided to the teledermatologists. The
453 original diagnosis provided by the primary dermatologist was an atypical mycobacterial
454 infection.

455

456 **Supplemental Figure 2.** Scatterplot of the correlation between the teledermatologists' (TDs')
457 primary plus differential diagnosis and the primary dermatologist's (PD's) primary diagnosis.
458 Each point represents a teledermatologist (TD), color-coded by which survey set the TD
459 participated in. The absence of clustering of points by color and the wide variation in TDs' years
460 of experience indicate that TDs' years of experience exhibit robust nonassociation with the
461 concordance between the TD's primary plus differential diagnosis and the primary dermatologist
462 (PD)'s primary diagnosis.

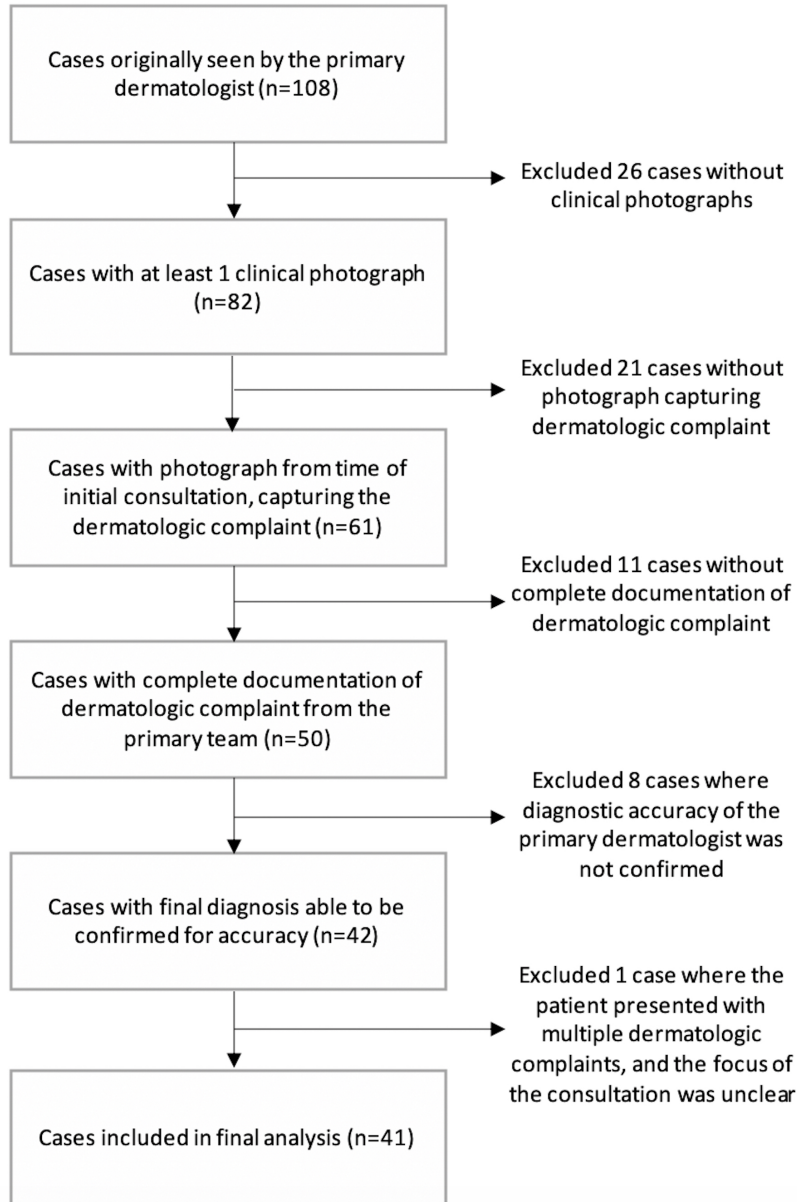
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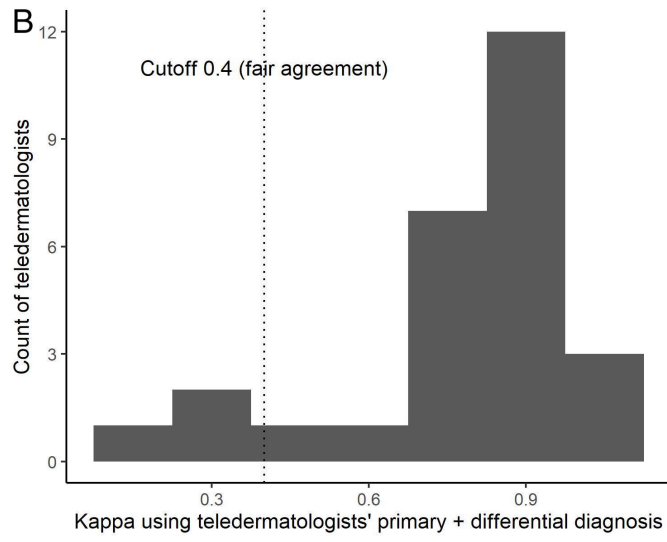
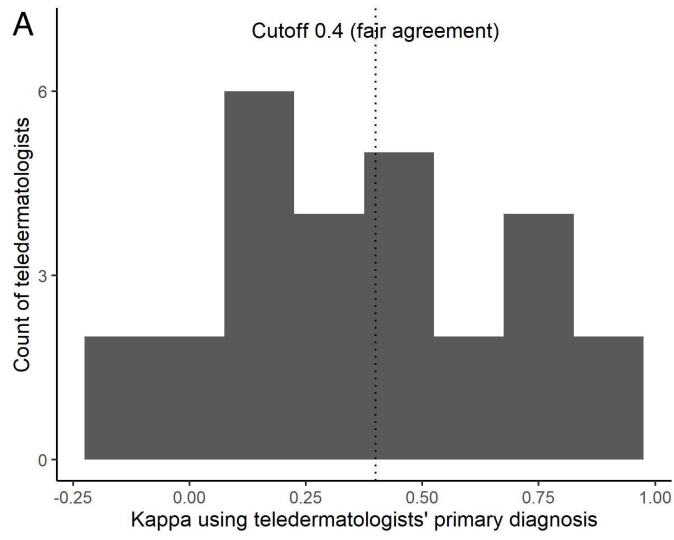
464 **Supplemental Table 1.** Tabular representation of the years' experience of the teledermatologists
465 (TDs) with corresponding kappa values for primary and primary plus differential diagnostic
466 concordance.

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Capsule summary:

- Inpatient access to dermatologists is limited, highlighting an opportunity to utilize teledermatology within the inpatient setting.
- Teledermatology in the inpatient setting may be a clinically acceptable option for diagnosis, evaluation, and management. This may represent a novel and effective option for hospitals.

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