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2 Original Article

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4 National Psoriasis Foundation COVID-19 Task Force Guidance for Management of Psoriatic Disease 5 During the Pandemic: Version 1

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137 Abstract:

138

Objective: To provide guidance about management of psoriatic disease during the COVID-19pandemic.

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142 Study Design: A task force (TF) of 18 physician voting members with expertise in dermatology,

143 rheumatology, epidemiology, infectious diseases, and critical care was convened. The TF was

144 supplemented by non-voting members which included fellows and National Psoriasis Foundation

145 (NPF) staff. Clinical questions relevant to the psoriatic disease community were informed by

146 questions received by the NPF. A Delphi process was conducted.

147

Results: The TF approved 22 guidance statements. The average of the votes was within the category
 of agreement for all statements. All guidance statements proposed were recommended, 9 with high
 consensus, 13 with moderate consensus.

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152 Limitations: The evidence behind many guidance statements is limited in quality.

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Conclusion: These statements provide guidance for the management of patients with psoriatic
 disease on topics ranging from how the disease and its treatments impact COVID-19 risk and

outcome, how medical care can be optimized during the pandemic, what patients should do to lower

157	their risk of getting infected with SARS-CoV-2 and what they should do if they develop COVID-19.
158	The guidance is intended to be a living document that will be updated by the TF as data emerge.
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161	Capsule Summary:
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163	The NPF COVID-19 Task Force produced 22 guidance statements to promote optimal
164	management of psoriatic disease during the pandemic.
165	Shared decision making is recommended as is adherence to evidence-based
166	recommendations when available. The guidance statements will be updated when necessary
167	in accordance with rapidly evolving science of COVID-19.

168 INTRODUCTION

169

Severe Acute Respiratory Coronavirus 2 (SARS-CoV-2), a single stranded RNA virus that binds to 170 the ACE-2 receptor and causes illness called COVID-19, has precipitated devastating personal, 171 economic and societal repercussions worldwide.¹⁻⁴ SARS-CoV-2 usually causes a mild, self-limited 172 illness, but about 15% of affected individuals have a more severe, sometimes life-threatening course, 173 with the risk of poor outcomes increasing with age and comorbidities.⁵⁻⁷ Diffuse alveolar damage and 174 acute respiratory distress syndrome are the most common presentations in severe COVID-19. 175 Additionally, thrombo-embolic events along with direct and indirect viral-induced injury may target the 176 skin, gastrointestinal tract, kidney, heart and brain with devastating consequences.⁸⁻¹⁰ 177

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The type 1 interferon response, required to clear the virus, is often insufficient in the early phase of SARS-CoV-2 infection but a delayed persistent elevation may develop as the illness progresses.¹¹ Profound dysregulation of innate and acquired immunity can occur with more severe COVID-19, including significant lymphopenia as a direct result of viral-induced apoptosis and necrosis of lymphocytes in the spleen and lymph nodes.¹² The persistent interferon response can result in systemic hyperinflammation (a.k.a., cytokine storm).^{13, 14} Several of the cytokines elevated in severe COVID-19 patients (TNF, IL-6, and IL-17) are also elevated in patients with psoriatic disease.¹⁵⁻¹⁷

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The current model of COVID-19 is that immune suppression in early infection may be harmful by
 allowing uncontrolled SARS-CoV-2 replication and dissemination, but may be helpful in severe illness
 by limiting organ damage from a dysregulated hyperimmune response¹⁸. Many treatments used for

psoriatic disease directly or indirectly impact immune pathways involved in COVID-19.¹⁹⁻²² Patients 190 and providers are concerned about the safety of immunomodulating agents in the setting of the 191 192 COVID-19 pandemic. These concerns are particularly relevant given that many of the comorbidities 193 associated with psoriasis and psoriatic arthritis (PsA), including obesity, diabetes and cardiovascular disease, are risk factors for the development of severe COVID-19.^{23, 24} To address the questions 194 posed by patients and providers, the National Psoriasis Foundation (NPF) commissioned a COVID-195 19 task force (TF) to develop scientifically-based guidance that promotes optimal management of 196 197 psoriatic disease during the pandemic.

199 METHODS

- 200 See online supplement for detailed methods.
- 201

202 Establishment of Task Force

203 The COVID-19 TF includes 18 physicians with a variety of expertise relevant to decision-making in

204 the pandemic from different geographical areas within the United States and Canada, many of whom

have frontline experience managing a surge of COVID-19 patients (E Table 1). The TF was

supplemented by non-voting members which include 4 trainees in dermatology, rheumatology, and

infectious diseases, 1 post-doctoral fellow in epidemiology, as well as senior staff from the NPF.

208

209 Evidence Synthesis

FDA, and NIH.

210 The TF co-chairs completed weekly literature searches for COVID-19 in relation to psoriatic disease.

211 TF members also recommended papers of broad importance to COVID-19 related to its basic

biology, epidemiology, and treatment. Additional sources of data were obtained from the CDC, WHO,

213

214

215 **Development of Clinical Questions**

The TF met every 2 weeks to discuss the developments in the literature and clinical experience.

217 Clinical questions relevant to the psoriatic disease community were iterated and informed by

questions received by the NPF from the broader patient and clinical community. The questions were

subdivided into 5 categories and work groups with balanced expertise were formed. Each work group

of the TF convened to draft responses to the clinical questions based on the available evidence.

221 These responses were reviewed and drafted into guidance statements.

222

223 Modified Delphi Process

The guidance statements were presented to the 18 TF members using a modified Delphi process including 2 rounds of voting with discussion in between. The Delphi approach was based on the RAND appropriateness method, which has been extensively validated.²⁵⁻³¹

227

228 TF members were asked to report their level of agreement anonymously with each guidance statement on a scale of 1-9. A rating of 1 corresponded to "complete disagreement," 5 corresponded 229 to "uncertain or neutral," and 9 corresponded to "complete agreement." The members were able to 230 231 provide anonymous written comments. Median vote ratings of 1-3, 4-6, and 7-9 were defined a priori as disagreement, uncertainty/neutral, and agreement, respectively. Panel consensus was 232 determined to be "low" when ≥ 5 votes fell into the 1–3 rating range with ≥ 5 votes concurrently falling 233 into the 7-9 rating range. Consensus was interpreted as "high" if all 18 votes fell within a single tertile, 234 with all other combinations considered as "moderate" levels of consensus. The results were analyzed 235 236 by the NPF with an independent analysis of the data by a non-voting member of the TF, which 237 vielded identical results.

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- 241

- 243 **RESULTS**:
- 244

The TF Delphi was completed over a 2 week period (E Table 2). Five categories of questions were explored (E Table 3) with 100% complete voting on 22 guidance statements (Table 4 & E-Table 4). The median was within the category of agreement for all statements, with the number of votes outside the range of agreement being only 1 or 2 for statements where agreement was not unanimous. All guidance statements were recommended, 9 with high consensus, and the remainder with moderate consensus.

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Category 1: What are the effects of psoriatic disease itself on SARS-CoV-2 infection and COVID-19
 illness?

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Patients with psoriatic disease appear to have similar rates of infection with SARS-CoV-2 and 255 COVID-19 outcomes³²⁻³⁶ as the general population (Guidance 1.1). However, uncertainty remains 256 regarding this question. First, a few reports suggest that psoriasis patients may be more prone to 257 infection with COVID-19 or have worse outcomes.³⁷⁻³⁹ For example, a United Kingdom study with 258 259 over 17 million patients found a small but statistically increased risk of death from COVID-19 (fully adjusted Hazards Ratio (HR) 1.19 [95% CI 1.11-1.27]) in individuals with either psoriasis, rheumatoid 260 arthritis or lupus.³⁸ It is unknown from this study the degree to which the observed finding is driven by 261 psoriasis, its severity, or treatment. Additionally, patients with psoriatic disease may be prone to 262

thrombotic complications that can also occur in COVID-19.⁴⁰⁻⁵⁴ There was unanimous agreement that
 severity of COVID-19 is driven by risk factors such as older age and comorbidities (Guidance 1.2). ^{32,}
 ^{35, 36, 38, 55-58} Psoriatic disease—particularly severe psoriasis— is associated with many of the
 comorbidities that drive COVID-19 mortality.^{44, 48, 59}

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Category 2: What are the effects of psoriasis or psoriatic arthritis treatment on SARS-CoV-2 infection and COVID-19 illness?

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The existing literature suggests that treatments for psoriasis and/or PsA do not meaningfully alter the 271 risk of acquiring SARS-CoV-2 infection or having worse COVID-19 outcomes (Guidance 2.1)^{36, 60-86}. 272 Cyclosporine, the most broadly immunosuppressive of psoriasis treatments, was not found to alter 273 risk of COVID-19 in 130 patients in Italy with psoriasis or atopic dermatitis (2 became infected with 274 SARS-CoV-2 and recovered without hospitalization).⁷⁰ This study lacked a comparison group and is 275 too small to reach definitive conclusions. One study suggested that psoriasis patients on biologics 276 were more likely to be hospitalized for COVID-19 but did not adjust for risk factors known to drive 277 poor COVID19 outcomes.87 278

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The rheumatology literature also suggests that treatments used for psoriatic disease (such as TNF inhibitors and methotrexate) do not negatively impact COVID-19⁸⁸⁻⁹¹ with one large registry (600 case reports from 40 countries) finding that TNF inhibitors are associated with a reduced adjusted odds of COVID-19 hospitalization compared to patients with rheumatic conditions not treated with TNF inhibitors.⁹¹ Similarly, adverse effects of TNF inhibitors on COVID-19 were not observed in large registries of IBD patients.^{92, 93} Small case series have reported poor COVID-19 outcomes in patients on JAK inhibitors for PsA⁹⁴ and secukinumab for ankylosing spondylitis; ⁹⁰ however, these isolated reports could be due to selection bias, chance, or underlying comorbidity. By contrast, an analysis of about 1400 patients from the rheumatology, gastroenterology, and dermatology literature concluded that biologic or targeted synthetic disease-modifying antirheumatic drug therapy has not been associated with more severe COVID-19 outcomes.³²

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292 Given these data, patients who are not infected with SARS-CoV-2 should continue their biologic or oral therapies for psoriasis and/or psoriatic arthritis in most cases (Guidance 2.2). Nevertheless, the 293 294 existing literature is largely based on small case series or large registries of spontaneous reports and therefore shared decision-making between clinician and patient is recommended (Guidance 2.2, 2.4, 295 2.5). By contrast, studies in the rheumatology and gastroenterology literature have observed that 296 chronic use of oral corticosteroids is associated with worse COVID-19 outcomes (i.e., hospitalization, 297 or a composite outcome of ICU admission, ventilator use, and/or death).^{32, 91, 92} Chronic systemic 298 corticosteroids should be avoided, if possible, for the management of psoriatic arthritis (guidance 299 2.3)95. 300

301

302 **Category 3:** How should medical care be delivered to patients with psoriatic disease to lower their 303 risk of infection with SARS-CoV-2 while still ensuring quality of care?

304

The pandemic has disrupted the ability of patients and providers ability to meet in person due to personal protective equipment shortages, measures implemented to lower risk of SARS-CoV-2

transmission, and patients' personal and economic hardships.⁹⁶⁻⁹⁹ Patients express concern about 307 being exposed to SARS-CoV-2 in the clinical setting either directly or indirectly (i.e., on public 308 transportation). Telemedicine can achieve similar outcomes for psoriasis patients compared to in 309 person care with a dermatologist;¹⁰⁰⁻¹⁰² however, limited information available on management of 310 psoriatic arthritis with telemedicine.^{103, 104} Telemedicine should be considered when pandemic 311 conditions limit in-person visits (Guidance 3.1).¹⁰⁵ However, there are limitations of telemedicine, and 312 therefore some patients should be evaluated in person (Guidance 3.2). Office-based phototherapy 313 remains an important option for patients with psoriasis (Guidance 3.3, Table 5).^{106, 107} 314

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Category 4: What should patients with psoriatic disease do to protect themselves from becoming
 infected with SARS-CoV-2?

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Patients should be advised to follow measures that prevent infection with SARS-CoV-2 (Guidance 319 4.1, E-Table 6)¹⁰⁸. These prevention measures should be followed at work (Guidance 4.2) and school 320 (Guidance 4.3). In cases where measures to prevent transmission of SARS-CoV-2 at work or school 321 cannot be maintained, shared decision-making is recommended to determine if specific 322 accommodations are medically necessary (Guidance 4.2 and 4.3). Psoriasis, even when involving 323 the face or hands, is not a contraindication to face coverings and hand washing respectively, and a 324 variety of approaches can be applied to mitigate skin irritation (E-Table 7).¹⁰⁹⁻¹¹¹ Patients with 325 psoriatic disease should receive the seasonal inactivated (e.g. killed) influenza vaccine, which is of 326 special importance to individual and public health during the COVID-19 pandemic (Guidance 4.4). 327

Providers may consider temporary discontinuation of methotrexate for 2 weeks after flu immunization in order to improve the immunogenicity of seasonal influenza vaccine.¹¹²

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Category 5: What should patients with psoriatic disease do if they become infected with SARS-CoV 2?

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334 Patients with psoriatic disease who become infected with SARS-CoV-2 should monitor their symptoms (E-Table 8), discuss management of their psoriatic disease treatments with their health 335 care providers, and should be prescribed and adhere to evidence-based COVID-19 treatments, if 336 available (Guidance 5.1, 5.2).^{86 113-115} The mortality benefit of initiation of corticosteroids in patients 337 with severe COVID-19 outweighs the risks of potentially precipitating a psoriasis flare, and therefore, 338 acute systemic corticosteroids are not contraindicated for the management of COVID-19 in patients 339 with psoriatic disease (Guidance 5.3).¹¹⁵⁻¹¹⁷ Based on limited available data, and to be consistent with 340 prescribing information, it may be prudent to hold treatments that target the immune system in the 341 setting of suspected or confirmed SARS-CoV-2 infection, but the final decision needs to be 342 determined on a case by case basis. Consistent with guidance from the FDA and the American 343 College of Physicians, the use of hydroxychloroguine or chloroguine is not recommended to prevent 344 or treat COVID-19 in patients with psoriatic disease outside of a clinical trial (guidance 5.4). ¹¹⁸⁻¹³⁰ 345 Patients with psoriatic disease should be aware that infection with SARS-CoV-2 may result in a flare 346 of psoriasis, which may occur due to discontinuation of psoriasis treatments, treatment of COVID-19 347 with anti-malarial drugs, or due to triggering of inflammation as part of COVID-19 illness (Guidance 348 5.6)^{129, 131-133} 349

351	Patients with psoriatic disease who become infected with SARS-CoV-2 should follow CDC guidance
352	¹³⁴⁻¹³⁷ on home isolation and discuss with their healthcare providers when they can end home
353	quarantine (Guidance 5.7, E-Table 9). ^{116, 134, 138 139} In the event someone with psoriatic disease has
354	close contact (E-Table 10) with an individual with suspected or confirmed SARS-CoV-2 infection, they
355	should quarantine for 14 days after the last contact, as per CDC guidelines (guidance 5.8) ¹⁴⁰ . The
356	decision regarding continuing or holding psoriasis treatments during a period of quarantine should be
357	individualized on a case-by-case basis between patient and provider.
358	
359	Resumption of psoriasis and/or psoriatic arthritis treatments held during SARS-CoV-2 infection should
360	be decided on a case-by-case basis (guidance 5.5). The persistence of one or more symptoms of
361	COVID-19, such as fatigue or joint pain, beyond the acute phase of the illness can occur ¹⁴¹ , and may
362	complicate the decision to restart psoriasis or PsA medications. Therefore shared decision-making is
363	recommended (Guidance 2.5).
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372 **DISCUSSION**

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The NPF COVID19 TF guidance statements serve to promote optimal management of psoriatic 374 disease during the pandemic. There are several strengths to the approach taken. First, the TF 375 assembled is a geographically diverse team with expertise in adult and pediatric dermatology. 376 377 rheumatology, critical care, infectious diseases, epidemiology, and basic and translational immunology with experience managing surges in COVID-19. The TF also includes trainees in 378 dermatology, rheumatology and infectious disease who are on the frontlines managing COVID-19 379 patients as well as senior staff from the NPF who are in touch daily with patients and providers 380 worldwide whose questions are brought to the TF. Second, we have established a robust process for 381 staying up to date with the latest literature relevant to COVID-19 and the management of psoriatic 382 disease resulting in the dissemination and evaluation of hundreds of peer reviewed publications by 383 the TF. Third, a validated Delphi approach enabled transparency and reproducibility of our process 384 for evaluating consensus statements.²⁵⁻³¹ 385

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Several limitations are acknowledged. First, the TF did not formally grade the strength of our recommendations.¹⁴² With the exception of guidance statements 4.4, 5.2, and 5.4, which are based on large scale randomized controlled trials, the evidence behind many of the guidance statements was often limited in quality. For example, studies evaluating the safety of treatments for psoriasis and psoriatic arthritis in the setting of COVID-19 involve small case series or large collections of case reports and thus should be considered preliminary. Large scale, longer term, population-based studies, with appropriate comparator groups, adjustment for relevant confounding variables, and complete ascertainment of clinically important COVID-19 outcomes are urgently needed. Second, the guidance is not intended to be proscriptive nor comprehensive. The ultimate judgment regarding how these recommendations should be followed is best left with the treating clinician and the patient in light of the circumstances presented by the individual patient, and the variability and biological behavior of the disease and therapeutics. Third, the TF does not have global representation of experts or direct inclusion of patients.

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The guidance statements are intended to be part of a "living" document that will be updated and amended when necessary by the rapidly evolving science of COVID-19. Readers are encouraged to visit <u>https://www.psoriasis.org/covid-19-resource-center</u> regularly for the latest guidance from the TF in order to promote optimal care and outcomes for patients with psoriatic disease during the pandemic.

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Guidance #	Guidance Statement	Level of Consensus
1.1	It is not known with certainty if having psoriatic disease meaningfully alters the risks of contracting SARS-CoV-2 (the virus that causes COVID-19 illness) or having a worse course of COVID-19 illness. Existing data, with some exceptions, generally suggest that patients with psoriasis and/or psoriatic arthritis have similar rates of SARS-CoV-2 infection and COVID-19 outcomes as the general population.	Moderate
1.2	The likelihood of poor outcomes from COVID-19 is driven by risk factors such as older age and comorbidities such as chronic heart, lung, or kidney disease and metabolic disorders such as diabetes and obesity. Patients with psoriatic disease are more prone to these comorbidities, particularly in those with more severe disease.	High
2.1	It is not known with certainty if treatments for psoriasis and/or psoriatic arthritis meaningfully alter the risks of contracting SARS-CoV-2 (the virus that causes COVID-19 illness) or having a worse course of COVID-19 illness. Existing data generally suggest that treatments for psoriasis and/or psoriatic arthritis do not meaningfully alter the risk of acquiring SARS-CoV-2 infection or having worse COVID-19 outcomes.	Moderate
2.2	It is recommended that patients who are not infected with SARS-CoV-2 continue their biologic or oral therapies for psoriasis and/or psoriatic arthritis in most cases. Shared decision-making between clinician and patient is recommended to guide discussions about use of systemic therapies during the pandemic (see guidance 2.5 for definition of shared decision making).	High
2.3	Chronic systemic corticosteroids should be avoided if possible for the management of psoriatic arthritis. If patients require chronic systemic corticosteroids for management of psoriatic arthritis, the dose should be tapered to the lowest dose necessary to achieve the desired therapeutic effect. Chronic systemic corticosteroid use for the treatment of psoriatic disease at the time of acute infection with SARS-CoV-2 may be associated with worse outcomes from COVID19 illness. It is important to note, however, that corticosteroids may improve outcomes for COVID19 when initiated in hospitalized patients requiring oxygen treatment.	High
2.4	Individuals newly diagnosed with psoriasis and/or psoriatic arthritis or who are currently not receiving treatment should be aware that untreated psoriatic disease is associated with serious impact on physical and emotional health, and in the case of psoriatic arthritis, can lead to permanent joint damage and disability. Shared decision making between clinician and patient is recommended to guide discussions about use of systemic therapies during the pandemic (see guidance 2.5 for shared decision making).	High
2.5	Providers recommend shared decision making with patients. Shared decision making between clinician and patient should be guided by several factors, including the potential benefits of treatment, the activity of skin and/or joint disease and response to previous therapies, as well as the patient's underlying risk for poor COVID19 outcomes, and ability to maintain measures to prevent infection with SARS-CoV-2 such as hand hygiene, wearing of masks, and physical distancing as required by	Moderate

	pandemic conditions. A review of known benefits of treatment accompanied by acknowledgment of the uncertainty related to the COVID19 pandemic and a discussion of a patient's individual circumstances and preferences should guide decision making.	
3.1	Telemedicine should be offered to manage patients wherever possible when local restrictions or pandemic conditions limit the ability for in-person visits. The following patients can be managed with telemedicine: Patients who are clinically stable and previously started on psoriatic disease treatment. Patients requiring a follow-up visit and refills for medication. New patients without timely access to in-person visits. Patients diagnosed with COVID-19 who are experiencing a significant flare. If telemedicine visits become inadequate to monitor patients' disease progress or manage new or evolving symptoms or signs of skin and joint disease, clinicians and patients should consider in-person visits.	Moderate
3.2	The following patients should be considered for in-person care if pandemic conditions allow (i.e., the clinical practice is open to see patients in person): Patients at risk for melanoma and non-melanoma skin cancer should be seen in person at a frequency consistent with standard of care for a full skin examination. New patients establishing care. Patients experiencing unstable psoriatic disease/flares. Patients requiring a thorough skin/or joint examination and a full physical examination for rheumatology patients.	Moderate
3.3	Providers recommend the recent guidelines published by Lim et al on how to optimize safety of office phototherapy for the patients and staff in the setting of the pandemic. See Table 5 for details.	High
4.1	Patients should be advised to follow measures that prevent infection with SARS-CoV- 2. These preventative measures include: To practice good hand hygiene, to maintain physical distancing from non-household members, and to wear a face covering of the nose and mouth when indoors (except in their own home), and when outdoors, but unable to maintain physical distancing. Face coverings should not be used in children under 2 years old due to risk of suffocation. See E Table 6 for details.	High
4.2	Patients with psoriatic disease should follow measures to prevent infection with SARS-CoV-2 in the workplace. If the work place environment does not allow for maintenance of prevention measures, a shared decision-making process between the patient and his/her clinician is recommended to determine if specific accommodations are medically necessary, especially for individuals whom, due to age or underlying health conditions, are at especially high risk for poor COVID19 outcomes.	Moderate
4.3	Youth with psoriatic disease should follow measures to prevent infection with SARS- CoV-2 while at school. These measures include maintaining 6 feet of physical distancing, consistently wearing masks if over the age of 2 years, and washing hands frequently. If the school environment is unable to ensure these prevention measures or families believe their child may not be able to adhere to these practices, we encourage discussion with the patient, caregivers, and his/her clinician to collectively develop a learning plan in the best interest and safety of the child.	High
4.4.	Patients with psoriatic disease should receive the seasonal inactivated (e.g. killed) influenza vaccine when it becomes available. While this vaccine will not protect against SARS-CoV-2, influenza vaccine lowers the risk of infection from seasonal influenza, which is of special importance to individual and public health during the	High

 5.2 Patients with psoriat prescribed and addee therapies should be disease, as well as of treatment, if availabl care of the hospitaliz dermatologists, and/ 5.3 Systemic corticoster disease are not contreatment of CoVID-Cases of psoriasis fl but the clinical signif 5.4 Hydroxychloroquine treatment of COVID-Cases of psoriasis fl but the clinical signif 5.5 Resumption of psori infection should be of psoriasis and/or psoriasis a	c. Patients taking systemic medications for psoriasis or psoriatic uss the timing of influenza vaccination with respect to their edications with their health care provider in order to optimize the lenza vaccine.	
 prescribed and adher therapies should be disease, as well as or treatment, if availabl care of the hospitaliz dermatologists, and/ 5.3 Systemic corticoster disease are not contropotentially flaring particles and emonstrates the effective of the constrates of the constrates of the constrates the effective of the constrates of the constrate of the constrates of	ic disease who become infected with SARS-CoV-2 should one of their treatments with their s.	Moderate
 disease are not control potentially flaring parademonstrates the effective demonstrates t	ic disease who become infected with SARS-CoV-2 should be ere to evidence-based COVID-19 therapies. Evidence-based used, currently including supportive care for patients with mild dexamethasone (systemic corticosteroids) and remdesivir e, for hospitalized patients requiring supplemental oxygen. The zed patient should include consultation with rheumatologists, for infectious disease specialists as medically necessary.	Moderate
 treatment of COVID-Cases of psoriasis flbut the clinical signif 5.5 Resumption of psori infection should be of psoriasis and/or psoriasis and or a case-by-order of the solution of the cover and the provide the	oids for the management of COVID-19 in patients with psoriatic traindicated and should not be withheld due to the concern of oriasis upon withdrawal of corticosteroids when evidence fectiveness for treating COVID-19 illness.	Moderate
 infection should be of psoriasis and/or pso symptoms. In those made on a case-by- 5.6 Patients with psoriat result in a flare of ps of COVID-19 flaring 5.7 Patients with psoriat CDC guidance on he they can end home in COVID-19 symptom antipyretics and imp returning to work, as with severe cases of immunosuppressive the length of home is 	or chloroquine are not recommended for the prevention or -19 in patients with psoriatic disease outside of a clinical trial. are have been reported in patients on anti-malarial medications, icance is not well understood.	High
 result in a flare of ps of COVID-19 flaring 7 Patients with psoriat CDC guidance on he they can end home in COVID-19 symptom antipyretics and imp returning to work, as with severe cases of immunosuppressive the length of home is 	asis and/or psoriatic arthritis treatments held during SARS-CoV-2 decided on a case-by-case basis. Most patients can restart riatic arthritis treatments after complete resolution of COVID-19 who have had a severe hospital course, shared decision making case basis is recommended.	Moderate
CDC guidance on he they can end home in COVID-19 symptom antipyretics and imp returning to work, as with severe cases of immunosuppressive the length of home is	ic disease should be aware that infection with SARS-CoV-2 may oriasis based on case reports. The clinical significance of the risk psoriasis is not known.	Moderate
	ic disease who become infected with SARS-CoV-2 should follow ome isolation and discuss with their healthcare providers when isolation. We recommend waiting a minimum of 10 days after onset, along with fever resolution for 24 hours without rovement in other symptoms, before ending home isolation and a patients are unlikely to be infectious after this point. In patients f COVID-19 or when psoriasis patients are on medications with effects, we recommend a case-by-case approach to determining solation.	Moderate
themselves for 14 da	contact to someone with SARS-CoV-2 infection should quarantine ays after the last contact and discuss the management of their atment with their medical provider(s).	Moderate

- Table 5: Methods to Reduce Risk of SARS-CoV-2 Transmission During Delivery of Office-Based Phototherapy

	individual bag
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417 Adapted from Lim et al.¹⁰⁶

Abbre	eviations and Acronyms
•	SARS COV-2 : Severe Acute Respiratory Coronavirus 2
•	COVID-19: Coronavirus disease 2019
•	IL: InterLeukin
•	TNF: Tumor Necrosis Factor
•	PsA: Psoriatic Arthritis
•	TF: Task Force
•	NPF: National Psoriasis Foundation
•	IBD: Inflammatory Bowel Disease
•	CDC: Centers for Disease Control
•	WHO: World Health Organization
•	FDA: Food and Drug Administration
•	NIH: National Institutes of Health
•	HR: Hazards Ratio
•	JAK inhibitor: Janus Kinase Inhibitor
•	SD: Standard Deviation
•	IQR: Inter Quartile Range

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- 442 4.3
- 443
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445 **References**

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Capsule Summary:

- The NPF COVID-19 Task Force produced 22 guidance statements to promote optimal management of psoriatic disease during the pandemic.
- Shared decision making is recommended as is adherence to evidence-based recommendations when available. The guidance statements will be updated when necessary in accordance with rapidly evolving science of COVID-19.