SHORT REPORT

Healthcare Resource Use and Costs Among Individuals with Vitiligo and Psychosocial Comorbidities: Retrospective Analysis of an Insured US Population

Jennifer H Lofland¹, Samyuktha Darbha², Ahmad B Naim¹, David Rosmarin³

¹US Medical Affairs, Incyte Corporation, Wilmington, DE, USA; ²Health Economics and Outcomes Research, Datawave Solutions Corp, Cranbury, NJ, USA; ³Department of Dermatology, Indiana University School of Medicine, Indianapolis, IN, USA

Correspondence: Jennifer H Lofland, US Medical Affairs, Incyte Corporation, 1801 Augustine Cut-Off, Wilmington, DE, 19803, USA, Tel +1 302 498 5861, Email jlofland@incyte.com

Purpose: This study aimed to describe healthcare resource utilization and costs among individuals with vitiligo who were diagnosed with ≥ 1 psychosocial comorbidity, using data from US claims databases.

Patients and Methods: A retrospective, observational cohort analysis of the IBM MarketScan Commercial and Medicare supplemental claims databases for US individuals with vitiligo aged ≥ 12 years and a first vitiligo claim between January 1 and December 31, 2018, was undertaken to assess psychosocial burden, including mental and behavioral health comorbidities.

Results: Of the 12,427 individuals included in the analysis, nearly 1 in 4 (23.5%) who had vitiligo were also diagnosed with ≥ 1 psychosocial comorbidity. A greater percentage of these individuals versus those who were not diagnosed with a psychosocial comorbidity had a vitiligo-related prescription claim (50.2% vs 45.4%; *P*<0.0001), especially for oral corticosteroids (25.4% vs 16.6%; *P*<0.0001) and low-potency topical corticosteroids (9.0% vs 7.6%; *P*<0.05). Total vitiligo-related healthcare resource utilization and costs were consistent among individuals with and without psychosocial comorbidity despite significantly (*P*<0.05) higher vitiligo-related ER visit utilization and expenditure among those with psychosocial comorbidity. Furthermore, individuals diagnosed with vitiligo and ≥ 1 psychosocial comorbidity had significantly (*P*<0.0001) greater utilization of all-cause mean prescription claims (25.0 vs 12.8), outpatient services (other than physician and ER visits: 19.5 vs 11.3), outpatient physician visits (10.1 vs 6.4), inpatient stays (0.6 vs 0.1), and ER visits (0.4 vs 0.2) and incurred significantly higher mean (SD) direct medical expenditures (\$18,804 [\$46,621] vs \$9833 [\$29,094] per patient per year; *P*<0.0001). **Conclusion:** Individuals with vitiligo who were diagnosed with ≥ 1 psychosocial comorbidity incurred greater total all-cause but not vitiligo-related healthcare resource utilization and expenditures than those without diagnosis of psychosocial comorbidities. Identification of psychosocial comorbidities in individuals with vitiligo may be important for multidisciplinary management of vitiligo to reduce overall burden for individuals with vitiligo.

Keywords: vitiligo, mental health, behavioral health, healthcare resource utilization, HCRU, healthcare costs

Introduction

Vitiligo is a chronic autoimmune disease in which patches of skin are progressively depigmented due to melanocyte destruction.¹ The prevalence of vitiligo in the United States ranges from 0.1% to 1.5%, thus affecting up to 5 million people based on 2022 US population estimates.^{2–5} Vitiligo is associated with significant quality-of-life impairment in routine activities, employment, and psychosocial health.^{6,7} Notably, individuals with vitiligo are approximately 5 times more likely to have depression than healthy individuals.^{8,9} In a previous study, individuals with vitiligo were more likely to be hospitalized for a mental health condition, with higher costs of inpatient care and increased length of stay, than those without vitiligo.¹⁰ In this cohort study, we analyzed healthcare resource utilization and costs among individuals

© 2024 Lofland et al. This work is published and licensed by Dove Medical Press Limited. The full terms of this license are available at https://www.dovepress.com/terms work you hereby accept the Terms. Non-commercial uses of the work are permitted without any further permission from Dove Medical Press Limited, provided the work is properly attributed. For permission for commercial use of this work, please see paragraphs A2 and 5 of our Terms (https://www.dovepress.com/terms.php).

557

with vitiligo and psychosocial burden, including mental and behavioral health comorbidities, in the United States using data from a claims database.

Materials and Methods

A retrospective, observational cohort analysis of the IBM (now Merative) MarketScan Commercial (for employer health plans) and Medicare supplemental claims databases was performed to identify individuals with an International Classification of Diseases, 10th revision (ICD-10) diagnosis code for vitiligo (codes: A67.2 [vitiligo of pinta], L80 [vitiligo], H02.73X [vitiligo of eye/eyelids and periocular area]) from January 1 through December 31, 2018 (index date corresponds with the first claim in this period). The databases were linked by encrypted unique patient IDs. Data in these databases included healthcare resource utilization. Eligible individuals were ≥ 12 years old with a medical claim for vitiligo in 2018 (first claim was index date) and 12 continuous months of post-index enrollment in their healthcare plan. Comorbidities and prescription data were identified based on prespecified codes from ICD-10 and the National Drug Code, respectively, during the 12-month post-index period for each individual. Psychosocial comorbidities included depression, anxiety, adjustment disorder, bipolar disorder, schizophrenia, attention-deficit/hyperactivity disorder, alcohol abuse, and substance abuse.¹¹ Sleep disorder was also captured but was not used for group stratification purposes (ie, recorded for both groups). Prescription data included all paid costs captured in the claims data in the IBM MarketScan Commercial and Medicare supplemental claims databases. Demographics, healthcare resource utilization, and expenditures were aggregated at an individual level and summarized per patient per year (PPPY; defined as the average of total costs in 1 year per patient; expenditures were in 2018 US dollars [\$]). Out-of-pocket costs were not available in this dataset and not included in the analysis. Error claims (ie, those with missing age or gender and or negative total costs) were excluded from the analyses. Data from the study population were summarized using descriptive statistics, and the subgroups of individuals with and without ≥ 1 diagnosed psychosocial comorbidity were compared using the chi-square test for categorical variables and t test for continuous variables.

Results

Overall, 18,309 individuals \geq 12 years old were identified as being diagnosed with vitiligo in 2018; 12,427 who had \geq 12 months of continuous post-index coverage were included in the analyses (Table 1). Mean (SD) age was 43.7 (16.9) years, and 53.2% were female. Most individuals (93.5%) were identified from the MarketScan Commercial database, and 6.5% were identified from the Medicare supplemental claims database. Almost one-fourth (23.5%) had a diagnosis of \geq 1 psychosocial condition, with the most common psychosocial comorbidities among those individuals being depression (30.8%), adjustment disorder (20.3%), anxiety (20.3%), and sleep disorder (18.4%). Sleep disorder was recorded in a significantly lower percentage of individuals with vitiligo who did not have diagnosed psychosocial comorbidities (7.2%; *P*<0.0001). Approximately one-tenth (9.0%) of individuals with vitiligo had \geq 1 comorbid autoimmune disorder, which occurred at a similar rate in individuals with versus without \geq 1 psychosocial condition (Table 1).

During the 12-month post-index period, 52.8% of all 12,247 individuals diagnosed with vitiligo were seen by a dermatologist for any reason, and 46.5% had a pharmacy claim for prespecified vitiligo-related prescriptions. Half (50.2%) of 2923 individuals with ≥ 1 diagnosed psychosocial comorbidity had a pharmacy claim for vitiligo-related treatments (vs 45.4% of 9504 individuals without a diagnosed psychosocial comorbidity; *P*<0.0001). Overall, individuals with versus without ≥ 1 psychosocial comorbidity had greater utilization of vitiligo-related pharmacy claims (1.5 vs 1.2; *P*<0.01; Figure 1A); however, total mean (SD) PPPY vitiligo-related pharmacy costs were similar (\$238 [\$1592] vs \$217 [\$1205]; *P*=0.52; Figure 1B). Individuals with versus without ≥ 1 psychosocial comorbidity had similar rates of ≥ 1 vitiligo-related inpatient stay (0.6% vs 0.3%; *P*=0.07), outpatient visit (93.6% vs 94.1%; *P*=0.33), and emergency room (ER) visit (0.4% vs 0.2%; *P*=0.10); healthcare resource utilization and costs were consistent in both groups for vitiligorelated inpatient stays and outpatient physician visits. Significant differences were, however, seen for vitiligo-related ER visits healthcare resource utilization (0.01 vs 0.00; *P*<0.05) and mean (SD) PPPY healthcare expenditures (\$22 [\$535] vs \$3 [\$67]; *P*<0.05) among individuals with versus without ≥ 1 psychosocial comorbidity. Nevertheless, total annual vitiligo-related expenditure was similar for the 2 groups (mean [SD] PPPY, \$1187 [\$5012] vs \$1112 [\$4228]; *P*=0.47).

Table I	Demographics	and Baseline	Characteristics	of Individuals	Diagnosed with	Vitiligo
---------	--------------	--------------	-----------------	----------------	----------------	----------

Characteristics	All Individuals With Vitiligo (N=12.427)	No Psychosocial Comorbidities	≥I Psychosocial Comorbidity	P value
		(n=9504)	(n=2923)	
Age, mean (SD), y	43.7 (16.9)	43.7 (17.0)	43.5 (16.5)	0.619
Age groups, y, n (%)				0.061
12–17	1306 (10.5)	1026 (10.8)	280 (9.6)	
18–29	1493 (12.0)	35 (.9)	358 (12.2)	
30-49	4418 (35.6)	3325 (35.0)	1093 (37.4)	
50–64	4378 (35.2)	3366 (35.4)	1012 (34.6)	
≥65	832 (6.7)	652 (6.9)	180 (6.2)	
Sex, n (%)				<0.0001
Female	6609 (53.2)	4816 (50.7)	1793 (61.3)	
Male	5818 (46.8)	4688 (49.3)	1130 (38.7)	
Employment status, n (%)				0.056
Full-time	9519 (76.6)	7308 (76.9)	2211 (75.6)	
Part-time or seasonal	183 (1.5)	151 (1.6)	32 (1.1)	
Early retiree	544 (4.4)	413 (4.3)	131 (4.5)	
Unknown/other	2181 (17.6)	1632 (17.2)	549 (18.8)	
Psychosocial comorbidity, ^a n (%)				
Sleep disorder	1222 (9.8)	684 (7.2)	538 (18.4)	<0.0001
Depression	900 (7.2)	0	900 (30.8)	<0.0001
Adjustment disorder	594 (4.8)	0	594 (20.3)	<0.0001
Anxiety	594 (4.8)	0	594 (20.3)	<0.0001
ADD/ADHD	314 (2.5)	0	314 (10.7)	<0.0001
Alcohol abuse	89 (0.7)	0	89 (3.0)	<0.0001
Bipolar disorder	73 (0.6)	0	73 (2.5)	<0.0001
Schizophrenia	I (<0.I)	0	I (<0.I)	0.235
Substance abuse ^b				
Sedative, hypnotic, or anxiolytic	I (<0.I)	0	I (<0.I)	0.235
Other stimulant	I (<0.I)	0	I (<0.I)	0.235
Cocaine use disorder	I (<0.I)	0	I (<0.I)	0.235
Opioid	I (<0.I)	0	I (<0.I)	0.235
Autoimmune comorbidity, ^a n (%)				
Autoimmune thyroid disease	505 (4.1)	355 (3.7)	150 (5.1)	0.0008
Type I diabetes mellitus	250 (2.0)	189 (2.0)	61 (2.1)	0.741
Psoriasis	210 (1.7)	155 (1.6)	55 (1.9)	0.358
Alopecia areata	99 (0.8)	73 (0.8)	26 (0.9)	0.519
Pernicious anemia	92 (0.7)	63 (0.7)	29 (1.0)	0.069
Inflammatory bowel disease	61 (0.5)	45 (0.5)	16 (0.5)	0.617
Sjögren's syndrome	59 (0.5)	39 (0.4)	20 (0.7)	0.060
Scleroderma	44 (0.4)	35 (0.4)	9 (0.3)	0.631
Systemic lupus erythematosus	29 (0.2)	18 (0.2)	(0.4)	0.067
Rheumatoid arthritis with rheumatoid factor	23 (0.2)	18 (0.2)	5 (0.2)	0.840
Adult dermatomyositis	6 (<0.1)	4 (<0.1)	2 (0.1)	0.631
Addison's disease	I (<0.I)	0	I (<0.1)	0.235

Notes: P-values <0.05 are in bold. ^aIndividuals could be diagnosed with \geq I comorbidity. ^bSubstance abuse other than cocaine use disorder is uncomplicated. **Abbreviations**: ADD, attention deficit disorder; ADHD, attention-deficit/hyperactivity disorder.

The most prescribed vitiligo-related therapies among all 12,427 individuals in the analysis set were topical calcineurin inhibitors (TCI; 21.1%), oral corticosteroids (OCS; 18.7%), and medium-potency topical corticosteroids (TCS; 9.8%; Figure 2A). There were significant differences in categories of therapies being prescribed for vitiligo among individuals with versus without ≥ 1 diagnosed psychosocial comorbidity, including prescribed OCS (25.4% vs 16.6%; *P*<0.0001),



Figure I Vitiligo-related healthcare resource (**A**) utilization and (**B**) expenditure costs grouped by psychosocial comorbidity status in individuals with vitiligo. ^aDoes not include ER and physician visits. *P<0.05; **P<0.01. **Abbreviations:** ER, emergency room; PPPY, per patient per year.

TCI (19.3% vs 21.7%; P<0.01), low-potency TCS (9.0% vs 7.6%; P<0.05), and laser therapy (5.1% vs 6.1%; P<0.05; Figure 2B). Prescription patterns for other therapies were comparable between the groups.

Among all 12,427 individuals in the analysis set, 99.7% had ≥ 1 all-cause outpatient medical visit during the 12-month post-index follow-up, and 91.4% had post-index prescription claims for any reason. All-cause ER visits were made by



Figure 2 Percentage of individuals with prescription claims for vitiligo (**A**) among all individuals and (**B**) grouped by psychosocial comorbidity status. ^aSystemic nonsteroidal immunosuppressants include cyclosporine and methotrexate. Statistical analysis was performed using the chi-square test: *P<0.01; ****P<0.001. Abbreviations: OCS, oral corticosteroids; TCI, topical calcineurin inhibitor; TCS, topical corticosteroids; UV, ultraviolet.

15.1% of individuals, and 4.0% had a hospital admission, with a mean (SD) inpatient stay of 5.3 (14.2) days. Mean (SD) cost of all-cause and vitiligo-related healthcare PPPY for all individuals with vitiligo was \$11,943 (\$34,248) and \$1130 (\$4425), respectively (Table 2). More individuals with versus without \geq 1 psychosocial comorbidity had an all-cause inpatient stay (6.7% vs 3.1%; *P*<0.0001), visited the ER (23.8% vs 12.5%; *P*<0.0001), and had prescription claims (96.4% vs 89.8%; *P*<0.0001). Individuals with versus without \geq 1 psychosocial comorbidity had significantly (*P*<0.0001) greater utilization of all-cause mean prescription claims (25.0 vs 12.8), outpatient services (other than physician and ER visits: 19.5 vs 11.3), outpatient physician visits (10.1 vs 6.4), inpatient stays (0.6 vs 0.1), and ER visits (0.4 vs 0.2; Figure 3A). Mean (SD) all-cause PPPY total healthcare expenditures (\$18,804 [\$46,621] vs \$9833 [\$29,094]) and costs related to outpatient services (\$8942 [\$31,580] vs \$4518 [\$18,310]), prescriptions (\$4229 [\$14,760] vs \$2872 [\$14,399]),

Healthcare Resource	Mean (SD) Use PPPY ^a , n	Mean (SD) Cost PPPY ^a , \$	
All-Cause			
Inpatient stays/services	0.2 (3.1)	1680 (14,005)	
Outpatient services ^b	13.3 (18.3)	5559 (22,236)	
ER visits	0.2 (0.7)	616 (2830)	
Prescription claims	15.7 (18.6)	3191 (14,496)	
Outpatient physician visits	7.3 (6.1)	898 (872)	
Average all-cause annual co	11,943 (34,248)		
Vitiligo-related			
Inpatient stays/services	0.03 (2.5)	89 (2216)	
Outpatient services ^b	3.7 (13.4)	628 (3522)	
ER visits	0.00 (0.1)	7 (266)	
Prescription claims	1.3 (3.2)	222 (1307)	
Outpatient physician visits	1.7 (2.2)	183 (190)	

 Table 2
 Healthcare Resource Utilization for All Individuals with

 Vitiligo

Notes: ^aDefined as the average of total use in I year or total costs in I year per patient. ^bDoes not include ER and physician visits.

Abbreviations: ER, emergency room; PPPY, per patient per year.

inpatient services (\$3289 [\$21,211] vs \$1185 [\$10,821]), outpatient physician visits (\$1274 [\$1149] vs \$782 [\$729]), and ER visits (\$1069 [\$3799] vs \$476 [\$2440]) were significantly (P < 0.0001) higher among individuals with versus without ≥ 1 psychosocial comorbidity (Figure 3B).

Discussion

Our analysis of US insurance claims databases found that nearly 1 in 4 individuals with vitiligo was diagnosed with ≥ 1 psychosocial comorbidity. A higher percentage of individuals with ≥ 1 psychosocial comorbidity had a pharmacy claim for vitiligo-related treatments, especially for OCS and low-potency TCS. OCS may be prescribed during vitiligo flares,¹² which were reported to occur during periods of stress in nearly two-thirds of patients in a recent global survey.¹³ Thus, greater claims for OCS in this subpopulation could be an indicator of increased vitiligo activity during periods of increased psychosocial stress¹⁴ among individuals with mental health comorbidities.

Individuals with ≥ 1 psychosocial comorbidity also had greater overall prescription claims versus those without a diagnosed psychosocial comorbidity, which is not unexpected given the potential need for treatment of associated comorbidities. Although approximately 1 in 10 individuals with vitiligo also had ≥ 1 comorbid autoimmune disorder, individual rates were similar in those with and without ≥ 1 psychosocial comorbidity. This indicates that psychosocial comorbidity is likely related to vitiligo rather than comorbid autoimmune burden. Individuals diagnosed with ≥ 1 psychosocial comorbidity also had disproportionately greater utilization of all-cause ER, outpatient services, outpatient physician visits, inpatient stays, and prescription claims. These differences were further reflected in direct costs PPPY in healthcare expenditures, highlighting the greater financial burden incurred by individuals with vitiligo who were diagnosed with ≥ 1 psychosocial comorbidity versus those without a psychosocial comorbidity. Total vitiligo-related healthcare resource utilization and costs were similar for individuals with versus without ≥ 1 psychosocial comorbidity. There was, however, higher annual vitiligo-related utilization for ER visits and pharmacy claims as well as higher expenditure for ER visits among individuals with versus without ≥ 1 psychosocial comorbidity, suggesting increased vitiligo-related distress (and possibly flares) among individuals who have psychosocial comorbidities. Additionally, the imbalance in all-cause total utilization and costs despite consistent vitiligo-related total utilization and costs among individuals with versus without ≥ 1 psychosocial comorbidity suggests that psychosocial comorbidity suggests that psychosocial burden contributes largely to the disparity.



В



Figure 3 All-cause healthcare resource (A) utilization and (B) expenditure costs grouped by psychosocial comorbidity status in individuals with vitiligo. ^aDoes not include ER and physician visits. ****P<0.0001.

Abbreviations: ER, emergency room; PPPY, per patient per year.

The underlying comparability between the MarketScan Commercial and Medicare supplemental populations and the prevailing 2018 US census population, with only a slight overrepresentation of female patients (51.5% vs 50.5%) and an underrepresentation of patients aged 65 years and older (4% vs 9%), provides a strong platform for this

analysis. Limitations of this study are those consistent with retrospective, administrative claims analyses, including the potential for sampling bias, incomplete records, improper data entry, and differences in actual versus physician-directed medication use. Data analyzed in this study were the latest available at the time of analysis, with the 2018 index period and 12-month post-index period (into 2019) predating the COVID-19 pandemic. Variables pertaining to individual and disease characteristics (eg, ethnicity/race, Fitzpatrick skin type, vitiligo extent, body surface area) were not available, so associations between these characteristics could not be analyzed. Furthermore, the cohorts in the study were not matched with respect to these variables, nor were data available for individuals without vitiligo who had mental and/or behavioral disorders, thus limiting the generalizability of the analysis. Healthcare resource utilization and costs directly related to mental and/or behavioral disorders were not collected. Data were not provided regarding whether diagnosis of psychosocial conditions occurred before or after the onset of vitiligo, and prespecified vitiligo-related treatments may not have been used for the treatment of vitiligo. No multivariate analyses were performed to determine relationships. Individuals who were not diagnosed with a psychosocial comorbidity may have undiagnosed conditions, potentially biasing the data in this group. Furthermore, out-of-pocket expenditures, which may be considerable due to payers not covering some vitiligo therapies,¹⁵ were not available within the dataset. Likewise, the direct, individual, and societal costs of vitiligo may be underreported in US claims databases.

Conclusions

In summary, nearly 1 in 4 individuals with vitiligo identified in a US claims database were diagnosed with ≥ 1 psychosocial comorbidity. While further investigation is required to better understand the burden of psychosocial comorbidities in individuals with vitiligo, we found that these individuals incurred greater all-cause but not vitiligo-related healthcare consumption and expenditures versus those with vitiligo but without diagnosed psychosocial conditions. These findings emphasize the need to identify psychosocial comorbidities among individuals with vitiligo and reduce the associated burden through multidisciplinary management of their vitiligo, including ensuring effective management of mental and behavioral health comorbidities.

Abbreviations

ER, emergency room; ICD-10, International Classification of Diseases, 10th revision; OCS, oral corticosteroids; PPPY, per patient per year; TCI, topical calcineurin inhibitors; TCS, topical corticosteroids.

Data Sharing Statement

Access to individual patient-level data is not available for this study.

Ethics Approval and Informed Consent

Ethics approval of the study protocol and informed consent were not required based on the use of deidentified insurance claims data and no direct patient involvement.

Acknowledgments

We thank Vijay N. Joish, MS, PhD (Incyte Corporation, at the time of the study) for leading the development of the analysis for this study, which was funded by Incyte (Wilmington, DE, USA), as well as Shreekant Parasuraman, PhD, and Jinan Liu, PhD, of Incyte for their critical review of the manuscript. Writing assistance was provided by Joshua Solomon, PhD, of ICON (Blue Bell, PA, USA) and was funded by Incyte.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis, and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Funding

Funding was provided by Incyte Corporation (Wilmington, DE, USA). The sponsor had roles in study design, data analysis, and data interpretation.

Disclosure

JHL and ABN are employees and shareholders of Incyte Corporation. SD is an employee of Datawave Solutions. DR has received honoraria as a consultant for AbbVie, Abcuro, Almirall, AltruBio, Arena, Astria, Boehringer Ingelheim, Bristol Myers Squibb, Celgene, Concert, CSL Behring, Dermavant Sciences, Dermira, Incyte, Janssen, Kymera, Kyowa Kirin, Lilly, Nektar, Novartis, Pfizer, RAPT, Regeneron Pharmaceuticals, Recludix, Revolo Biotherapeutics, Sanofi, Sun Pharmaceuticals, UCB, Viela Bio, and Zura Bio; has received research support from AbbVie, Amgen, Bristol Myers Squibb, Celgene, Dermira, Galderma, Incyte, Janssen, Lilly, Merck, Nektar, Novartis, Pfizer, RAPT, and Regeneron Pharmaceuticals; and has served as a paid speaker for AbbVie, Amgen, Bristol Myers Squibb, Celgene, Dermavant, Incyte, Janssen, Lilly, Novartis, Pfizer, Regeneron Pharmaceuticals, and Sanofi. The authors report no other conflicts of interest in this work.

References

- 1. Rodrigues M, Ezzedine K, Hamzavi I, Pandya AG, Harris JE. New discoveries in the pathogenesis and classification of vitiligo. J Am Acad Dermatol. 2017;77(1):1–13. doi:10.1016/j.jaad.2016.10.048
- 2. Kruger C, Schallreuter KU. A review of the worldwide prevalence of vitiligo in children/adolescents and adults. *Int J Dermatol.* 2012;51 (10):1206–1212. doi:10.1111/j.1365-4632.2011.05377.x
- Bibeau K, Pandya AG, Ezzedine K, et al. Vitiligo prevalence and quality of life among adults in Europe, Japan and the USA. J Eur Acad Dermatol Venereol. 2022;36(10):1831–1844. doi:10.1111/jdv.18257
- Gandhi K, Ezzedine K, Anastassopoulos KP, et al. Prevalence of vitiligo among adults in the United States. JAMA Dermatol. 2022;158(1):43–50. doi:10.1001/jamadermatol.2021.4724
- 5. US Census Bureau. U.S. Population Estimated at 332,403,650 on Jan. 1, 2022. Available from: https://www.census.gov/library/stories/2021/12/ happy-new-year-2022.html. Accessed May 25, 2023.
- Morrison B, Burden-Teh E, Batchelor JM, Mead E, Grindlay D, Ratib S. Quality of life in people with vitiligo: a systematic review and meta-analysis. Br J Dermatol. 2017;177(6):e338–e339. doi:10.1111/bjd.15933
- 7. Silverberg JI, Silverberg NB. Association between vitiligo extent and distribution and quality-of-life impairment. JAMA Dermatol. 2013;149 (2):159–164. doi:10.1001/jamadermatol.2013.927
- 8. Lai YC, Yew YW, Kennedy C, Schwartz RA. Vitiligo and depression: a systematic review and meta-analysis of observational studies. Br J Dermatol. 2017;177(3):708-718. doi:10.1111/bjd.15199
- 9. Wang G, Qiu D, Yang H, Liu W. The prevalence and odds of depression in patients with vitiligo: a meta-analysis. *J Eur Acad Dermatol Venereol*. 2018;32(8):1343–1351. doi:10.1111/jdv.14739
- Patel KR, Singam V, Rastogi S, Lee HH, Silverberg NB, Silverberg JI. Association of vitiligo with hospitalization for mental health disorders in US adults. J Eur Acad Dermatol Venereol. 2019;33(1):191–197 doi:10.1111/jdv.15255.
- 11. Ezzedine K, Eleftheriadou V, Jones H, et al. Psychosocial effects of vitiligo: a systematic literature review. Am J Clin Dermatol. 2021;22 (6):757-774. doi:10.1007/s40257-021-00631-6
- 12. Kim SM, Lee HS, Hann SK. The efficacy of low-dose oral corticosteroids in the treatment of vitiligo patients. Int J Dermatol. 1999;38(7):546–550. doi:10.1046/j.1365-4362.1999.00623.x
- 13. Hamzavi IH, Bibeau K, Grimes P, et al. Exploring the natural and treatment history of vitiligo: perceptions of patients and healthcare professionals from the global VALIANT study. *Br J Dermatol.* 2023;189(5):569–577. doi:10.1093/bjd/ljad245
- 14. Henning SW, Jaishankar D, Barse LW, et al. The relationship between stress and vitiligo: evaluating perceived stress and electronic medical record data. *PLoS One.* 2020;15(1):e0227909. doi:10.1371/journal.pone.0227909
- 15. Blundell A, Sachar M, Gabel CK, Bercovitch LG. The scope of health insurance coverage of vitiligo treatments in the United States: implications for health care outcomes and disparities in children of color. *Pediatr Dermatol*. 2021;38(2):79–85. doi:10.1111/pde.14714

ClinicoEconomics and Outcomes Research



DovePress

Publish your work in this journal

ClinicoEconomics and Outcomes Research is an international, peer-reviewed open-access journal focusing on Health Technology Assessment, Pharmacoeconomics and Outcomes Research in the areas of diagnosis, medical devices, and clinical, surgical and pharmacological intervention. The economic impact of health policy and health systems organization also constitute important areas of coverage. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit http://www.dovepress.com/testimonials.php to read real quotes from published authors.

Submit your manuscript here: https://www.dovepress.com/clinicoeconomics-and-outcomes-research-journal

🖬 🔰 in 🗖