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Use of active comparator tirals in dermatology: A repeated crosssectional analysis

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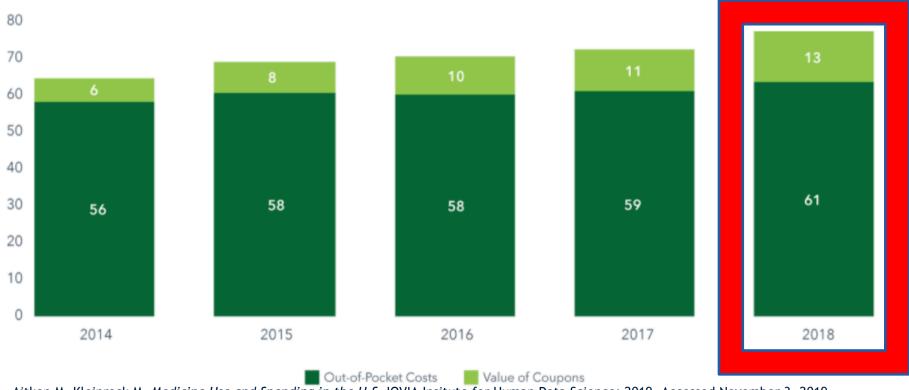
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Use of active comparator trials in dermatology: A repeated cross-sectional analysis

John Miller, Sophia Ly, Arash Mostaghimi MD, John Barbieri MD*

Introduction: Healthcare Spending Costs



Aitken M, Kleinrock M. Medicine Use and Spending in the U.S. IQVIA Insitute for Human Data Science; 2019. Accessed November 3, 2019.

• Overall out-of-pocket costs have risen to \$61 billion in 2018



Healthcare spending impact on patients?

Spending on prescription medications is expected to rise to \$420 billion by 2023.²



Nearly 30% of patients reported not taking prescription due to cost.



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The Role of Active Comparator Clinical Trials



While placebo-controlled trials are valuable to understand efficacy, active comparator trial designs provide data that can guide clinical decision making by allowing direct comparisons between similar drugs

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Active Comparators in Dermatology Topical Trials

In dermatology, subtle changes in concentration or combination of topical agents can be classified as a novel product.



While some new medications may improve patient outcomes, others may not offer meaningful benefit over existing less costly alternatives.

The Inquiry Question:

To evaluate trends in the use of active comparator trial designs for topical medications approved by the FDA between January 2002 and December 2020.

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Methods

Population

- Food and Drug Administration
- Physicians
- Patients

<u>Resources Used</u>

- Clinical Trials Database
 - Clinical Trial Information

- FDA Orange Book
 - Drug Approval Dates





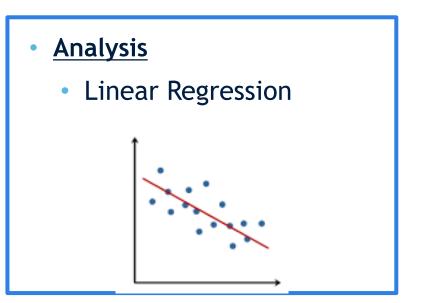


Outcomes & Analysis

<u>Outcomes</u>

- Frequency of Active
 Comparator over Time
- Frequency of Active
 Comparator by Trial Phase

256

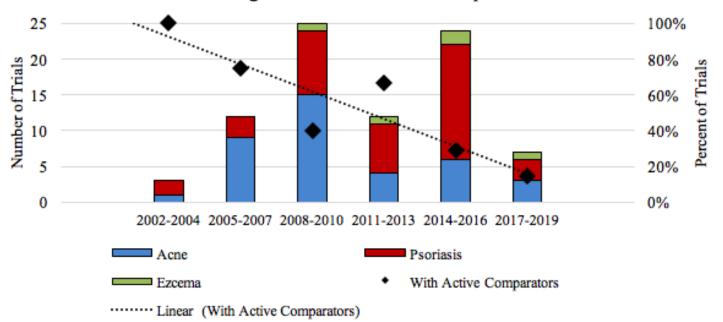




<u>Outcomes</u>

Between 2002 and 2020, the proportion of clinical trials for acne, psoriasis, and eczema with an active comparator has decreased.
 (-5.2% per year, 95% CI -1.7% to -8.6%)

Total Trials for Acne, Psoriasis, and Ezcema Per Year Compared to Percentage of Trials with Active Comparators



Outcomes

 Phase II studies were most likely to include and active comparator (71%), while phase III studies were least likely (32%).

% Trials with and Active Comparator by Clinical Indication

	Phase II	Phase III	Phase IV	Total
Psoriasis	83%	39%	29%	44%
	n=6	n=28	n=7	n=41
Acne	100%	35%	71%	50%
	n=1	n=23	n=14	n=38
Actinic Keratosis	75%	23%	80%	45%
	n=4	n=13	n=5	n=22
Antifungal	67%	15%	0%	30%
	n=6	n=13	n=1	n=20
Rosacea	0%	43%	0%	30%
	n=2	n=7	n=1	n=10
Eczema	50%	0%	0%	20%
	n=2	n=2	n=1	n=5
Other	77%	35%	0%	44%
	n=13	n=23	n=5	n=41
Total	71%	32%	47%	42%
	n=34	n=109	n=34	n=117

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Conclusions

 Although there is a greater need for comparative effectiveness data in the setting of a growing number of available treatments, our results highlight that the use of active comparator trials has decreased over time



Moving Forward (Impact):

It will be important for clinicians, patients, payers, and

Role of Patient Reported Outcomes

Absolute Lesion Reduction vs. Quality of Life Index

COSC-DENETICS ANALYSES

• FDA Regulated Guidelines

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Disclosures and Acknowledgements

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Citations

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