

### BACKGROUND

- Independently, opioid use disorder (OUD) and liver disease (LD) pose considerable clinical concerns.
- Naltrexone and acetaminophen have known hepatoxic properties when taken separately.
- Psychosocial factors that Black and Hispanic populations face in healthcare create disproportionate OUD and LD treatment.
- There are gaps in the literature for understanding how acetaminophen use among Black and Hispanic populations contribute to OUD and LD treatment outcomes.

## **METHOD AND CHARACTERISTICS**

- Examined descriptive data of N=631 individuals with OUD and LD, prescribed medication for OUD (MOUD), in electronic health record (EHR).
  - n=578 with acetaminophen; n=53 without acetaminophen
  - Propensity score matching used to examine demographic and clinical characteristic differences by acetaminophen use.
- Conducted literature review and consulted with industry experts to identify and prioritize future research needs related to acetaminophen use among historically marginalized populations with OUD and LD.

| Patient demographic characteristics by acetaminophen use after propensity score matching* |                             |                                |         |  |  |  |  |
|---|-----------------------------|--------------------------------|---------|--|--|--|--|
|   | Acetaminophen<br>Use (n=53) | No Acetaminophen<br>Use (n=53) |         |  |  |  |  |
|   | M (SD) / N (%)              | M (SD) / N (%)                 | p-value |  |  |  |  |
| Age   | 50.4 (13.9)                 | 50.2 (14.3)                    | 0.929   |  |  |  |  |
| Race  |                             |                                |         |  |  |  |  |
| White   | 39 (73.59%)                 | 51 (96.23%)                    | 0.001   |  |  |  |  |
| Black or African<br>American  | 14 (26.42%)                 | 10 (18.87%)                    | 0.353   |  |  |  |  |
| Ethnicity   |                             |                                |         |  |  |  |  |
| Not Hispanic or<br>Latino   | 53 (100%)                   | 51 (96.23%)                    | 0.153   |  |  |  |  |
| Unknown Ethnicity   | 0 (0%)                      | 10 (18.87%)                    | 0.001   |  |  |  |  |
| Sex   |                             |                                |         |  |  |  |  |
| Male  | 27 (50.94%)                 | 26 (49.06%)                    | 0.846   |  |  |  |  |
| Female  | 26 (49.06%)                 | 27 (50.94%)                    | 0.846   |  |  |  |  |

\*Propensity score matching balanced groups by age and sex. Indicator code and indicators with 0% after matching were excluded.

# Identifying High Priority Research Needs for Acetaminophen Use among Black/Hispanic Populations with Opioid Use Disorder and Liver Disease Autumn Barnes, B.A., Lissette Saavedra, Ph.D., Amber Jarnecke, Ph.D.

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# More research is needed to address medical and social disparities in prescribing practices and acetaminophen use among historically marginalized individuals with opioid use disorder and liver disease.

| Themes from Literature and Experts   |  |  |  |   |   |  |
|--|--|--|--|---|---|--|
| Literature   | Psychiatrists/<br>Addiction<br>Researchers (2)   | Pharmacists (2)  | Substance Use<br>Therapist   | Senior Research<br>Psychiatrist   | Psychologist/<br>Addiction<br>Researcher  |  |
| the US, Hispanics have<br>ghest risk of developing<br>on-Alcoholic Fatty Liver<br>sease (1), and Blacks are<br>e most likely to develop<br>epatitis C. (2)<br>acks/Hispanics with OUD<br>as likely than White<br>unterparts to receive<br>JD treatment. (3)<br>altrexone and<br>etaminophen are<br>etabolized by the same<br>zymes; inhibition of these<br>zymes could cause drug<br>kicity. (4)<br>retaminophen and<br>altrexone may cause<br>hanced glutathione<br>pletion/hepatotoxicity. (5)<br>chnologies are being<br>veloped to measure<br>altrexone and<br>etaminophen in blood<br>asma. (6)<br>o published studies report<br>noracial differences in<br>edication management for<br>tients with OUD + LD. | <ul> <li>Careful consideration of<br/>Naltrexone for OUD + LD<br/>needed because<br/>Naltrexone can exacerbate<br/>liver damage.</li> <li>No known published<br/>guidelines on liver function<br/>levels, but generally if<br/>ALT/AST are 2x normal<br/>levels, proceed with<br/>caution.</li> <li>Important to consider when<br/>LD develops; Hepatitis C<br/>commonly co-occurs with<br/>OUD, this can lead to other<br/>liver conditions.</li> <li>Because there is concern<br/>with Naltrexone and liver<br/>function, important to<br/>examine acetaminophen<br/>use because of hepatoxic<br/>effects.</li> <li>Clinicians may not be<br/>aware of potential drug<br/>interactions if they do not<br/>ask about over the counter<br/>drug use – conducting<br/>research is one way to<br/>identify clinically significant<br/>interactions.</li> </ul> | <ul> <li>Pharmacists may consult<br/>with healthcare team on<br/>appropriate treatment.</li> <li>Software enables<br/>pharmacists to to search<br/>drug interactions.</li> <li>No formal database or<br/>system to find detailed<br/>information on drug<br/>interactions with pre-<br/>existing conditions (e.g.,<br/>LD).</li> <li>There is an interest in<br/>having a centralized<br/>database on OUD,<br/>MOUD, and other drug<br/>interactions.</li> </ul> | <ul> <li>Disproportionately low rate of treatment uptake/retention in Blacks/Hispanics compared to Whites.</li> <li>Hispanic community is high risk for LD; may lack access to healthcare knowledge/resources.</li> <li>Few providers speak and deliver interventions in Spanish.</li> <li>Politics/culture impact treatment uptake; acculturation, nation of origin impact outcomes.</li> <li>Other treatment barriers: job insecurity, lack of healthcare insurance, lack of access to treatment can lead to improper drug management.</li> <li>PCPs need to ensure patients with LD have citizenship/are on LD treatment before MOUDs.</li> <li>Most MOUD practitioners know potential risks, but little published research.</li> <li>Risk for developing LD after starting MOUDs and pain medications.</li> <li>Patients with OUD are not given much information on the risks of LD.</li> <li>Future studies need to incorporate lived experiences.</li> </ul> | <ul> <li>Naltrexone and<br/>acetaminophen are<br/>commonly taken together<br/>for pain.</li> <li>Often cannot replace<br/>acetaminophen with other<br/>medications (e.g.,<br/>nonsteroidals) because<br/>increased risk of bleeding;<br/>acetaminophen may be<br/>preferred for geriatric<br/>populations.</li> <li>No known interactions<br/>between Naltrexone and<br/>acetaminophen; both are<br/>hepatoxic but little<br/>evidence-based literature<br/>on the subject.</li> <li>No concrete, numerical<br/>guidelines for liver enzyme<br/>function; are guided by<br/>vague terms such as<br/>"elevated significantly."</li> <li>May be safe for patient to<br/>take continuous dose of<br/>Naltrexone and<br/>acetaminophen if AST and<br/>ALT are carefully<br/>monitored.</li> <li>Have to consider the costs<br/>and benefits of warning<br/>patients about<br/>acetaminophen and<br/>Naltrexone co-use.</li> </ul> | <ul> <li>Rate of opioid use among<br/>African Americans is high<br/>more research is needed<br/>to determine whether LD<br/>develops before or after<br/>opioid use initiation in<br/>African Americans.</li> <li>Need to consider<br/>social/cultural factors that<br/>contribute to interactions<br/>with medical professionals<br/>(e.g., African Americans<br/>less likely to go to the<br/>doctor due to historical<br/>harms).</li> </ul> |  |





DIVERSITY IN ADDICTION RESEARCH TRAINING

## RECOMMENDATIONS

# **Future Research Needs**

Hepatoxicity of Acetaminophen and Naltrexone Co-Use

Clinical Relevance, Training, and **Guidelines for Assessing Liver** Function with Acetaminophen and Naltrexone

Sociocultural Impacts on Dissemination of Information, Treatment and Outcomes for **Historically Marginalized People** with Opioid Use Disorder and Liver Disease

Understanding How Causation and Timing of Liver Disease are Associated with Opioid Use **Disorder and Treatment** Outcomes

## REFERENCES



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