

I. Use Case Description	
Use Case Name	<i>Personalized Depression Treatment Ontology</i>
Use Case Identifier	
Source	
Point of Contact	<i>Cole Feuer, Nancy Zhang, Gunnar Eastman</i>
Creation / Revision Date	<i>10/4/24</i>
Associated Documents	

II. Use Case Summary	
Goal	<i>To create an ontology that can improve the personalization of depression treatment by mapping relationships between patient demographics, treatment types, and outcomes.</i>
Requirements	<i>The system must integrate data from clinical trials, genetic studies, and patient-reported outcomes. It must have practical use in supporting clinicians in creating the best treatment plan for patients.</i>
Scope	<i>Focus on integrating genetic, demographic, and treatment outcome data to help clinicians find better treatment strategies for depression patients faster and with less trial-and-error.</i>
Priority	<i>High, as improving the speed and accuracy of mental health treatments has important social and medical implications.</i>
Stakeholders	<i>Primary stakeholders: Clinicians, mental health researchers, and patients. Secondary stakeholders: Hospitals, mental health organizations, and pharmaceutical companies.</i>
Description	<p><i>This use case focuses on the creation of a personalized depression treatment ontology aimed at improving the efficiency and accuracy of mental health treatment. The ontology will integrate diverse data sources, including patient demographics, genetic data, clinical trials, and patient-reported outcomes. The system will assist clinicians in formulating personalized treatment plans, reducing the trial-and-error approach often associated with mental health care.</i></p> <p><i>Principal Actors:</i></p> <p><i>Clinicians: Use the system to recommend personalized depression treatments based on patient data.</i></p> <p><i>Patients: Provide data on treatment outcomes, allowing the ontology to evolve with real-world experiences.</i></p> <p><i>Researchers: Input clinical trial and genetic data into the system to enhance its accuracy.</i></p> <p><i>Genetic Counselors: Utilize genetic data to offer tailored treatment recommendations for patients.</i></p> <p><i>Restated Goals:</i></p> <p><i>Provide clinicians with personalized treatment recommendations for depression based on patient-specific data.</i></p> <p><i>Minimize the trial-and-error approach in mental health treatment, leading to faster and more accurate care.</i></p> <p><i>Integrate ongoing patient-reported outcomes and clinical trial data to continuously improve treatment recommendations.</i></p>

Actors / Interfaces	<i>Primary Actors:</i> <ul style="list-style-type: none"> ● Clinicians: Use the ontology for recommending treatments. ● Patients: Provide patient-reported outcome data. ● Researchers: Input and analyze clinical and genetic data. <i>Systems:</i> <ul style="list-style-type: none"> ● Clinical trials databases ● Genetic databases ● Ontology management system ● Electronic medical records systems.
Pre-conditions	<i>A clinician or researcher is logged into the system.</i> <i>The system has access to clinical, genetic, and patient-reported outcome datasets.</i>
Post-conditions	<i>A treatment plan with a predicted success rate is recommended for the patient.</i> <i>Data from the current treatment experience is fed back into the ontology to improve future predictions.</i>
Triggers	<i>The clinician inputs a patient's demographic and genetic information into the system.</i> <i>The need for treatment recommendations triggers the use of the ontology.</i>
Performance Requirements	<i>Response time: needs to provide treatment recommendations reasonably quickly</i> <i>Scalability: needs to scale with data imputed and new research</i> <i>Concurrency: needs to be accessible to multiple clinicians and researchers at a time.</i> <i>Updatibility: needs to adapt to new research and update suggestions accordingly</i>
Assumptions	
Open Issues	

III. Usage Scenarios

Scenario 1: Clinician Seeking Treatment for a New Patient

A clinician is meeting with a new patient who has been diagnosed with depression. The patient has not previously received treatment for depression, and the clinician wants to recommend the most effective treatment option while minimizing the trial-and-error approach that is commonly used in mental health care. The clinician uses the ontology-based system to provide personalized treatment recommendations based on the patient's demographics and genetic data.

Scenario 2: Genetic Counselor Providing Personalized Treatment Recommendations

A genetic counselor, specializing in the treatment of mental health conditions, is meeting with a patient who has struggled with depression and has not responded well to traditional antidepressants. The counselor uses the ontology-based system to recommend personalized treatment options based on the patient's genetic profile and demographic information.

Scenario 3: Psychiatrist Adjusting Medication for a Patient with Anxiety Disorder

A psychiatrist is treating a patient with generalized anxiety disorder (GAD) who has experienced limited success with current medication and has reported side effects. The psychiatrist uses the ontology-based system to refine treatment options by considering the patient's medical history, genetic markers, and past medication responses. The system provides personalized recommendations for alternative medications or dosages that are more likely to be effective while minimizing side effects based on the patient's specific profile.

IV. Basic Flow of Events

Basic Flow I			
Step	Actor (Person)	Actor (System)	Description
1	Clinician	Treatment ontology recommendation system	The clinician logs into the treatment recommendation system.
2	Clinician	Treatment ontology recommendation system	The clinician enters patient information (age, gender, genetic data).
3	N/A	Treatment ontology recommendation system	The system searches through the ontology to find relevant treatments based on similar cases.
4	N/A	Treatment ontology recommendation system	The system suggests potential treatments with predicted effectiveness.
5	Clinician	Treatment ontology recommendation system	The clinician selects a treatment and provides it to the patient.
6	Clinician	Treatment ontology recommendation system	The clinician logs treatment outcomes into the system for future analysis.

Basic Flow II			
Step	Actor (Person)	Actor (System)	Description
1	Clinician	Treatment ontology recommendation system	The clinician logs into the treatment recommendation system.
2	Clinician	Treatment ontology recommendation system	The clinician enters incomplete patient data (age, gender) but also provides medical history (previous medications taken, etc.)
3	N/A	Treatment ontology recommendation system	The system searches through the ontology to find relevant treatments based on similar cases, avoiding the previously taken medications, and medications similar to those already taken by the patient.
4	N/A	Treatment ontology recommendation system	The system suggests potential treatments with predicted effectiveness.
5	Clinician	Treatment ontology recommendation system	The clinician selects a treatment and provides it to the patient.
6	Clinician	Treatment ontology recommendation system	The clinician logs treatment outcomes into the system for future analysis.

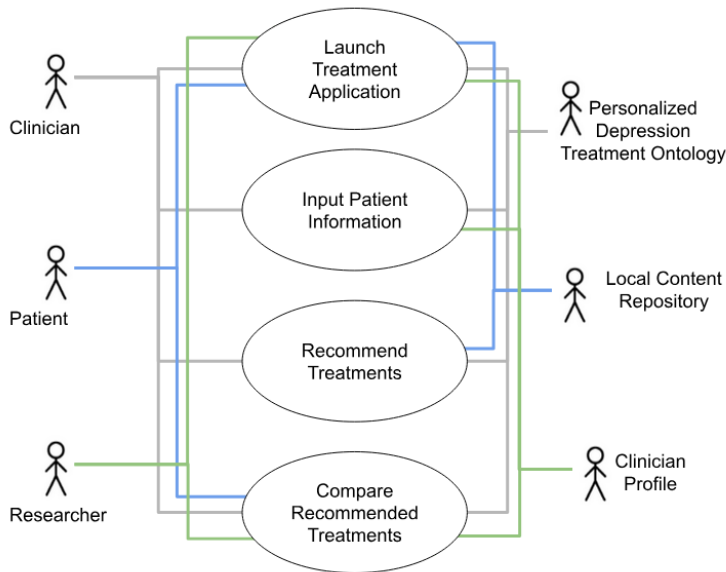
V. Alternate Flow of Events

Alternate Flow of Events			
Step	Actor (Person)	Actor (System)	Description
1	Clinician	Treatment ontology recommendation system	The clinician enters patient data, but the ontology finds insufficient data for an accurate prediction (e.g., for rare genetic conditions).
2	Clinician	Treatment ontology recommendation system	The system requests additional clinical trial or research data related to the case.

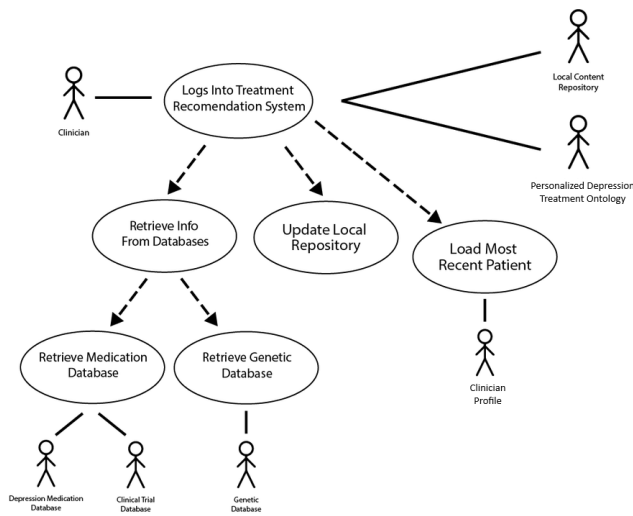
3	Clinician	Treatment ontology recommendation system	The clinician is provided with a generic recommendation based on broad population data rather than personalized data.
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VI. Use Case and Activity Diagram(s)

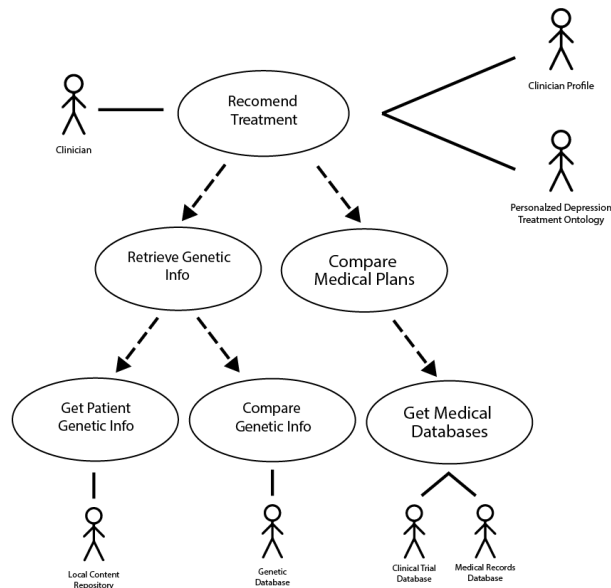
Overview Diagram



Launch Treatment Recommendation System Diagram



Recommend Treatment Diagram



VII. Competency Questions

What is the most effective treatment for patients with specific genetic markers associated with depression?

Example Answer: The system identifies that patients with a particular genetic marker (e.g., serotonin transporter gene polymorphisms) have shown a higher response rate to selective serotonin reuptake inhibitors (SSRIs). Based on this, the ontology suggests SSRIs as the recommended treatment with a success prediction of 75%, supported by clinical trial data and patient outcomes by clinical trial data and patient outcomes from similar cases.

Which treatment options have the highest success rates for patients aged 30-45 with treatment-resistant depression? In this case, patients demographic includes age 30-45, genetic marker is chromosome 3p25-26 that is often exhibited by families with multiple members that are diagnosed with Major Depressive Disorder. Treatment type will be medication, specifically SSRIs, with the treatment outcome hopefully being remission.

How Ontology Was Used: The ontology utilizes relationships between entities such as "patient demographics," "demographics," "genetic markers," "treatment types," "treatment types" and "treatment outcomes," "treatment outcomes." These relationships allow the system to infer connections between a patient's genetic data and the most relevant treatment options, even when some information may be missing or incomplete. By organizing knowledge in a structured and interconnected way, the ontology enables more accurate predictions than simple rule-based systems. For example, if we only had simple rule-based systems, we would not be personalizing treatments based on patients and only fitting them in "if... else" statements basically. By utilizing ontologies and an inference model, it can help go through a database that has a bunch of medications and with each relationship it has connected from "patient demographics," "genetic markers," "treatment types," and "treatment outcomes", it will be able to give a couple personalized treatment plans with high accuracy success outcomes.

Commented [1]: The highlighted informations need to be linked with the concepts from the revised terminology to determine the degree of completeness of the terminology.

Commented [2]: Including an example would greatly enhance the explanation. Simply stating that knowledge is organized in a structured and interconnected way isn't sufficient to justify the need for an ontology or to illustrate how semantics can be beneficial. An illustrative example would make these concepts clearer and more convincing.

How does age impact the effectiveness of certain treatments for depression?

Example Answer: For patients aged 30-45, the ontology suggests that a combination of cognitive behavioral therapy (CBT) and ketamine has shown a 60% improvement rate in cases of treatment-resistant depression. This recommendation is supported by aggregated data from clinical trials and patient-reported outcomes.

How Ontology Was Used: The ontology integrates clinical trial data and patient-reported outcomes categorized by demographic factors, such as age. It mapped the success rates of various treatments for specific age groups, identifying that for patients aged 30-45, a combination of cognitive behavioral therapy (CBT) and ketamine resulted in a 60% improvement rate in treatment-resistant cases. This information was drawn from an aggregated analysis of clinical trials that tested these treatments on patients within this age range, allowing the ontology to infer the most effective strategies for similar future patients.

What alternative treatments are recommended for patients with genetic markers indicating poor response to SSRIs?

Example Answer: The system identifies norepinephrine-dopamine reuptake inhibitors (NDRIs) as more effective for patients with genetic markers that show low efficacy with SSRIs. This recommendation is based on genetic data and patient-reported outcomes from similar cases.

How Ontology Was Used: The ontology links genetic data with treatment efficacy outcomes. It identified that patients with genetic markers associated with poor response to selective serotonin reuptake inhibitors (SSRIs) responded better to norepinephrine-dopamine reuptake inhibitors (NDRIs). This connection was established through the ontology's integration of genetic markers with historical treatment success rates, gathered from both clinical trials and real-world patient-reported outcomes. The ontology used these correlations to provide an evidence-based recommendation for alternative treatments.

Commented [3]: Great! here we have a valuable opportunity to apply inference.

How can treatment recommendations be adapted for patients with comorbid conditions, such as anxiety or chronic pain, along with depression?

Example Answer: The ontology suggests treatments like mindfulness-based cognitive therapy (MBCT) for patients with both depression and anxiety. This recommendation is drawn from clinical trials showing that MBCT can effectively treat both conditions simultaneously, improving outcomes by 65%.

How Ontology Was Used: The ontology incorporates comorbidity relationships between conditions like anxiety, chronic pain, and depression. It used clinical trial data that measured the effectiveness of treatments addressing multiple conditions simultaneously, such as mindfulness-based cognitive therapy (MBCT), which showed improvements for patients with both depression and anxiety. By accessing data on dual-treatment effectiveness, the ontology identified MBCT as a recommended treatment, showing a 65% improvement rate, and proposed it as an effective strategy for managing both conditions in comorbid patients.

What is the most effective treatment option for patients with both depression and genetic markers linked to

bipolar disorder?

Example Answer: The ontology identifies mood stabilizers, such as lithium, combined with cognitive-behavioral therapy (CBT) as the most effective treatment for patients showing genetic markers for both depression and bipolar disorder. Clinical data indicates a 70% success rate in reducing depressive episodes in these patients.

How Ontology Was Used: The ontology cross-referenced genetic markers associated with bipolar disorder and their influence on depression treatment outcomes. It identified mood stabilizers, such as lithium, combined with cognitive-behavioral therapy (CBT), as effective based on clinical trials involving patients with both depression and bipolar disorder markers. The ontology was able to match patient genetic profiles to successful treatment protocols, providing a recommendation that reflects a 70% success rate in managing depressive symptoms for such patients.

Info Gotten From Each Source:

M. et al., Li, "Integrative functional genomic analysis of Human Brain Development and Neuropsychiatric Risks | Science"

Provides a comprehensive look at genetic markers and brain development, linking neuropsychiatric risks to specific treatment outcomes. This research can answer questions involving genetic predisposition and comorbid conditions. The study's functional genomic analysis would be highly relevant to understanding how genetic markers influence treatment efficacy, especially in complex conditions like depression with comorbid bipolar disorder or pharmacoresistance.

B. Gaynes and A. Rush, "Treatment Alternatives to Relieve Depression (STAR*D) study *The STAR*D study: Treating depression in the real world,*" CLEVELAND CLINIC JOURNAL OF MEDICINE

The STAR*D study is a large-scale investigation into treatment-resistant depression, focusing on various treatment strategies when initial options fail. It provides insights into the effectiveness of different treatments across demographics. The study can also offer data on treatment outcomes for specific genetic markers or age groups, helping answer questions related to treatment efficacy based on age or genetic factors.

GENOME-BASED THERAPEUTIC DRUGS FOR DEPRESSION (GENDEP)

The GENDEP project focuses on the relationship between genetic markers and antidepressant responses, which is crucial for understanding pharmacogenomics in depression treatment. This study links genetic data to treatment efficacy, providing an evidence base for personalized treatment strategies based on genetic variations and pharmacoresistance.

Kupfer, D. J., Frank, E., & Phillips, M. L. (2012). "Major depressive disorder: New clinical, neurobiological, and treatment perspectives." *The Lancet*.

This comprehensive review focuses on new developments in the understanding and treatment of major depressive disorder (MDD), with an emphasis on the neurobiological mechanisms and treatment options, including newer antidepressants and personalized medicine. This source could provide updated insights into pharmacoresistance and the effectiveness of newer treatments like ketamine or novel antidepressants, especially in various age groups.

VIII. Resources

Knowledge Bases, Repositories, or other Data Sources

Data	Type	Characteristics	Description	Owner	Source	Access Policies & Usage
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<i>(dataset or repository name)</i>	<i>(remote, local/in situ, etc.)</i>	<i>e.g. – no cloud cover</i>	<i>Short description of the dataset, possibly including rationale of the usage characteristics</i>	<i>Source (possibly a system, or remote site) for discovery and access</i>
GENDEP (Genome-Based Therapeutic Drugs for Depression) Dataset	Genetic and Clinical Data	Combines genetic data with clinical outcomes from antidepressant treatment trials	The GENDEP project focuses on the genetic determinants of antidepressant response, making it highly relevant for an ontology that aims to personalize treatments based on genetic profiles.	Publicly available (European Genome-phenome Archive)
STAR D (Sequenced Treatment Alternatives to Relieve Depression) Dataset	Clinical Trial Data	Large-scale, longitudinal, real-world study of depression treatment	one of the largest and most comprehensive studies of depression treatment. It tracks patients across several levels of antidepressant treatment, detailing effectiveness, side effects, and patient-reported measures over time.	Publicly available (European Genome-phenome Archive)
UK Biobank	Genetic, Demographic, and Health Data	Large cohort with comprehensive genetic, demographic, and health-related data.	The UK Biobank includes both genetic and health data from over 500,000 participants. It provides great data on depression diagnoses, symptoms, and genetic factors, which can be integrated into the ontology to provide personalized treatment insights.	UK Biobank (need to apply)
PsychENCODE Consortium Data	Genetic and Epigenetic Data	Multi-omic data combining genetic, epigenetic, and transcriptomic information	The PsychENCODE project provides detailed genetic and epigenetic data, offering insights into how gene expression impacts depression and mental health. This is valuable for understanding the biological mechanisms that might influence treatment responses.	PsychENCODE Consortium, (NIH data portals)

External Ontologies, Vocabularies, or other Model Services

Resource	Language	Description	Owner	Source	Describes/Uses	Access Policies & Usage
<i>(ontology, vocabulary, or model name)</i>	<i>(ontology language and syntactic form, e.g., RDFS -</i>	<i>If the service is one that runs a given ontology or model-based application at</i>		<i>Source (link to the registry or directly to the ontology, vocabulary, or model where that model is</i>	<i>List of one or more data sources described by and/or used by the model</i>	

	N3)	a given frequency, state that in addition to the basic description		maintained, if available)		
POEM	OWL	Psychometric Ontology of Experiences and Measures	Kelsey Rook	https://tetherless-world.github.io/P-OEM/	Mental Health Terminology	open

Other Resources, Service, or Triggers (e.g., event notification services, application services, etc.)

Resource	Type	Description	Owner	Source	Access Policies & Usage
(sensor or external service name)		Include a description of the resource as well as availability, if applicable	Primary owner of the service	Application or service URL ; if subscription based, include subscription and any subscription owner	

IX. References and Bibliography

List all reference documents – policy documents, regulations, standards, de-facto standards, glossaries, dictionaries and thesauri, taxonomies, and any other reference materials considered relevant to the use case

[1] B. Gaynes and A. Rush, "Treatment Alternatives to Relieve Depression (STAR*D) study The STAR*D study: Treating depression in the real world," CLEVELAND CLINIC JOURNAL OF MEDICINE, vol. 75, no. 1, 2008, Available: <https://www.ccjm.org/content/ccjom/75/1/57.full.pdf>

[2] "GENOME-BASED THERAPEUTIC DRUGS FOR DEPRESSION (GENDEP)," CORDIS, <https://cordis.europa.eu/project/id/503428> (accessed Sep. 26, 2024).

[3] M., et al Li, "Integrative functional genomic analysis of Human Brain Development and Neuropsychiatric Risks | Science," Science, <https://www.science.org/doi/10.1126/science.aat7615> (accessed Sep. 26, 2024).

[4] Kupfer, D. J., Frank, E., & Phillips, M. L. (2012). "Major depressive disorder: New clinical, neurobiological, and treatment perspectives." The Lancet.

X. Notes

There is always some piece of information that is required that has no other place to go. This is the place for that information.