



# FRONTOTEMPORAL DEMENTIA (FTD)

By: Basiru-Lee-Leigh

Nicholas Gonzalez

Mentor: Dr. Karen Bell




# What is FTD

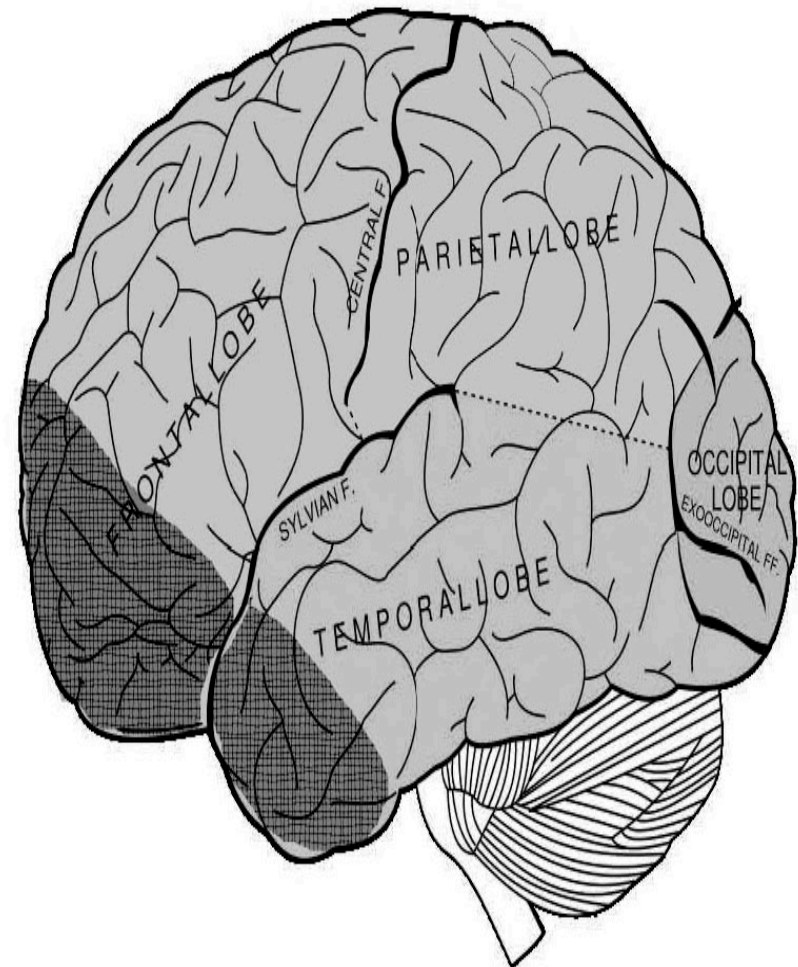
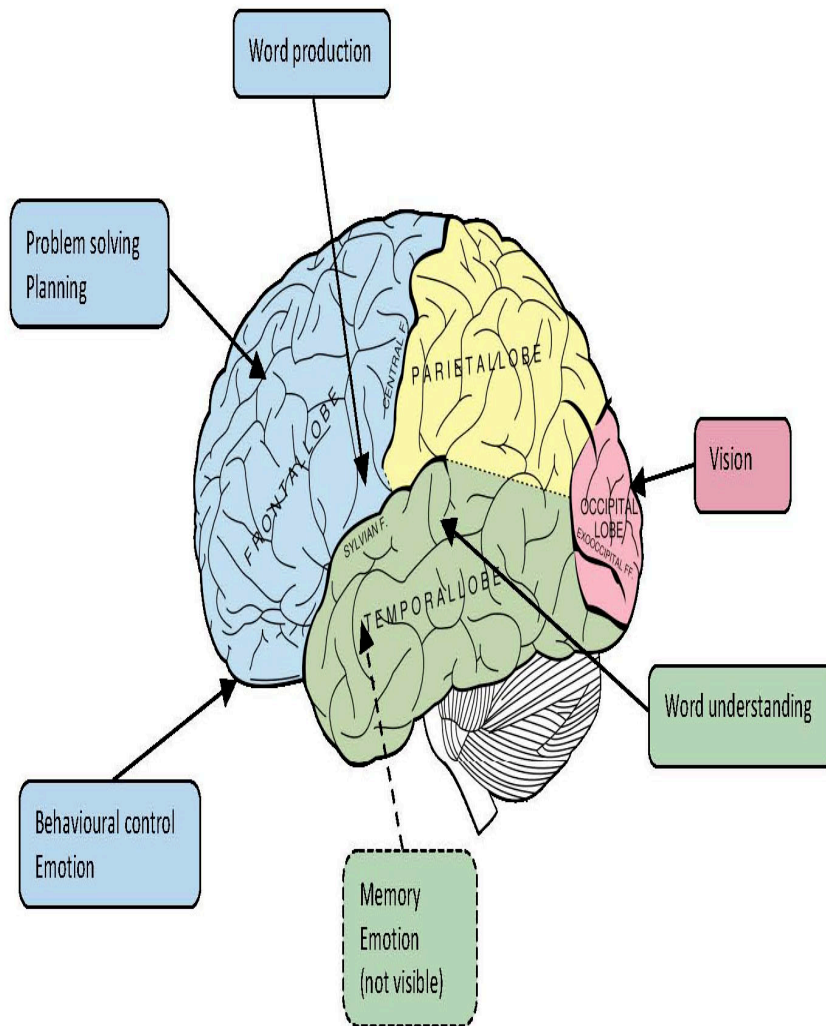
- A subtype of Frontotemporal Lobar Degeneration.
- A degenerative condition of the anterior part of the brain.
- FTD involves a disturbance of behavior and personality such that patients suffer a change in their character from previous selves.
- FTD differs from other degenerative conditions like Alzheimer's and CJD.
- FTD is marked by dramatic changes in personality, behavior and some thought processes.
  - Insidious onset, slow progression
  - Deficits in behavior, judgment, language, social conduct



# Cause

- A variety of mutations on several different genes have been linked to specific subtypes of FTD. But more than half the people who develop FTD have no family history of dementia.
  - In some cases of FTD, the affected parts of the brain contain microscopic Pick bodies — abnormal protein-filled structures that develop within brain cells.
- 

# Areas of brain affected







# Symptoms

- Impairment in social skill
- Change in activity level
- Decreased Judgment
- Changes in personal habits
- Alterations in personality and mood
- Akinesia
- Failure or inability to make motor responses to verbal commands

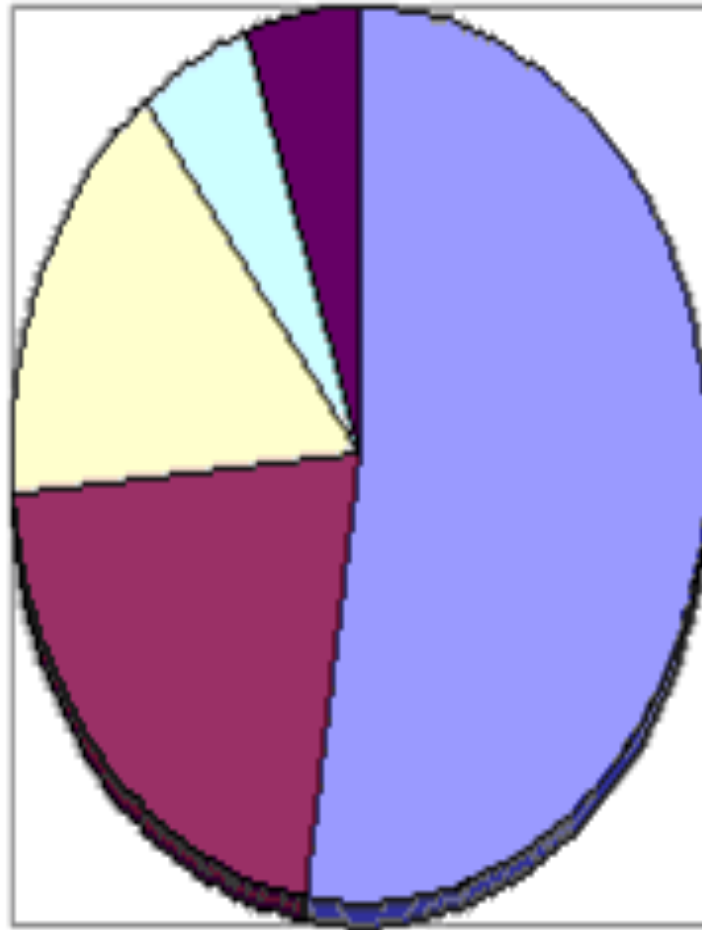


© ELSEVIER, INC. – NETTERIMAGES.COM



# Statistics

- Seven million Americans may be affiliated with a form of dementia
- FTD may account for 2-5% or 140,000 – 350,000 cases of dementia
- Occurs predominantly after age 40 and usually before age 65
- FTD have an equal chance of striking both men and women



- Alzheimer's Dementia
- Vascular Dementia
- Lewy Body Dementia
- Frontotemporal Dementia
- Other types of Dementia



# Comparison

Both FTD and Alzheimer disease (AD) are characterized by atrophy of the brain, and a gradual, progressive loss of brain function

## FTD

- FTD is distinguished by cerebral atrophy in the frontal and anterior temporal lobes of the brain
- Patients exhibit memory disturbances remain oriented to time and place and recall information about the present and past.
- Even in late stages of the disease, patients retain visuo-spatial orientation, and they negotiate and locate their surroundings accurately
- Life expectancy is slightly longer for FTD.
- Onset after 40, but mainly before 65
- It has no amyloid plaques or tau tangles that infect the brain.

## Alzheimer's Disease (AD)

- Alzheimer's affects the hippocampal, posterior temporal and parietal regions.
- patients experience severe memory loss and have an inability to learn new information.
- Onset at about 65.
- Has loss of nerve cells and contains tau tangles and amyloid plaques.



## 3 Pathological Subtypes

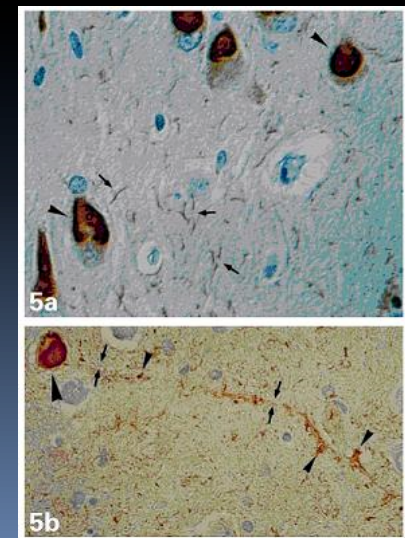
**bvFTD**

**Progressive nonfluent aphasia (PNFA)**

**Semantic dementia**

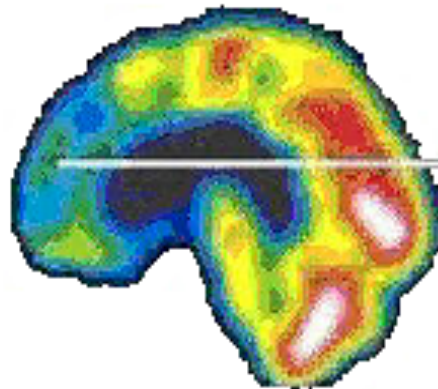
# bvFTD (Behavioral Variant FTD)

- Also referred to as Pick's disease.
- Patients with Pick's disease have an abnormal protein, pick bodies, inside nerve cells in the damaged areas of the brain.
- characterized by early and progressive changes in personality, emotional blunting and/or loss of empathy.
- Impairment of language may also occur, but is less prominent and would appear as a word-finding problem.
- It is the third most common dementia, Alzheimer's being the first.
- It is most commonly found in people of Scandinavian descent.
- Symptoms:
  - Hyperoral
  - Stereotyped and/or repetitive
  - Hypersexual
  - Impulsive acts
  - Apathy
  - Mood changes

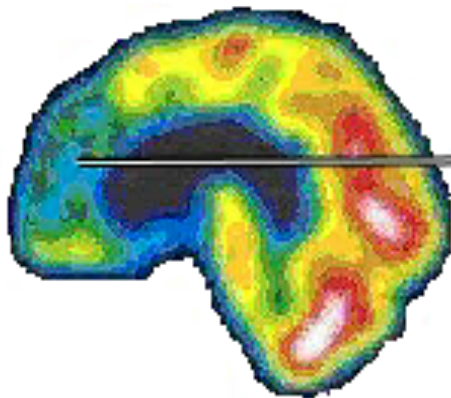


Pick bodies

## PET Scan - Pick's Disease



- Severely decreased brain activity in the Frontal Lobes resulting in significant cognitive, emotional, and behavioral disturbances.






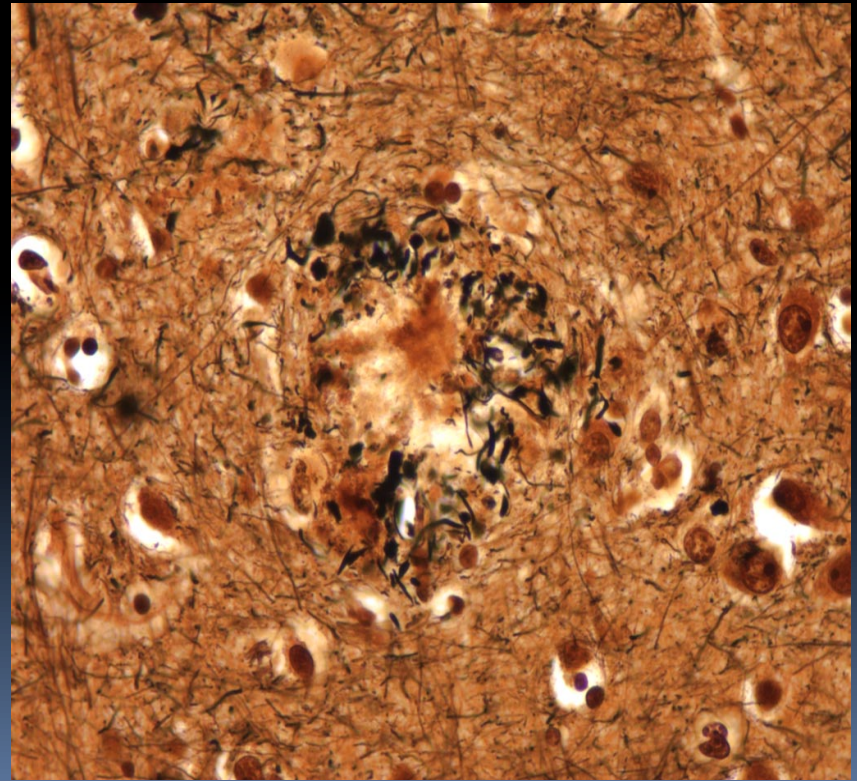
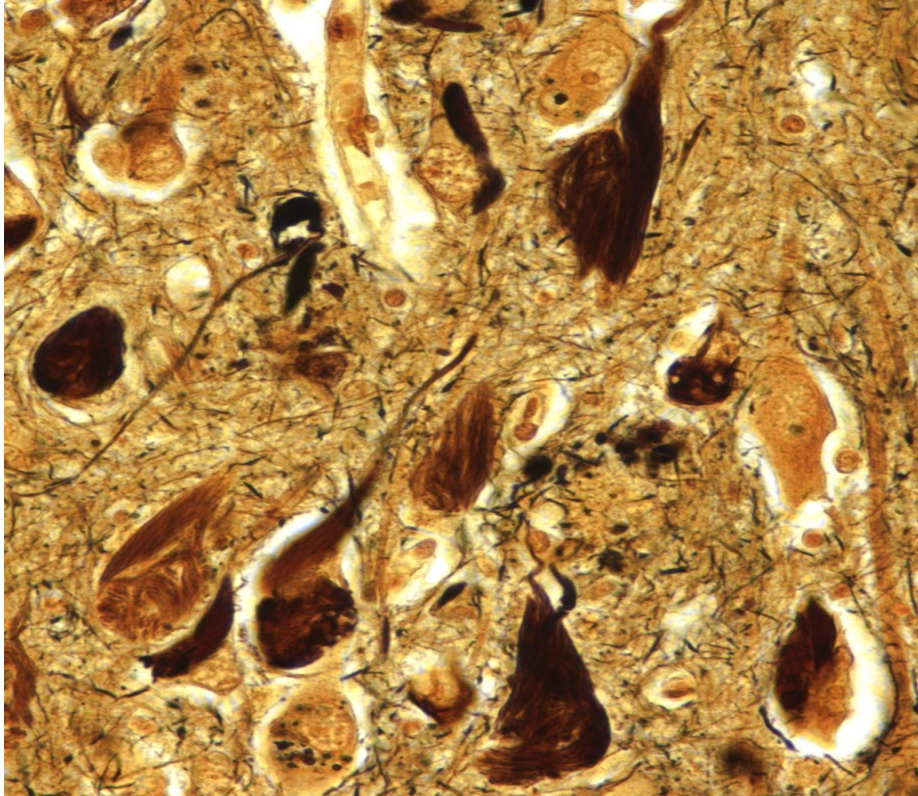
# Semantic dementia (SD)

- SD accounts for 20% of all FTD cases.
- Difficulty with language is the key function problem experienced with patients with SD.
- A person with SD would substitute a complex word like “salmon” and use the word “thingy” many times since they cannot form the word.
  - It will eventually progress to a worsen state to where they lose the concept of a word and ask “ What is a fish?”
- Memory, however, tends to stay intact as it always has.
  - So they are fully aware of the time and place.
- Signs:
  - Early loss of word and object knowledge
  - SD patients eventually develop bvFTD behaviors such as:
    - Apathy and loss of empathy.




# Progressive nonfluent aphasia (PNFA)

- PNFA accounts for 20% of all cases of FTD.
  - Unlike SD, people with PNFA have problems pronouncing language rather than losing the meaning of words.
    - They would have problems in groups of friends or understanding complex words or sentences.
  - Some signs of PNFA are:
    - Slurred speech
    - Breathiness
    - Deficits in comprehension of syntactically complex sentences
    - Sound distortions or hesitations in initiation
    - And Grammatical deficits in language production
- 





# Treatments

- Currently, there are no specific treatment or cure for FTD, but there are treatments for symptomatic relief.
  - **Antidepressants**
  - **Antipsychotics**
  - **Anticonvulsants**
  - **Medroxyprogesterone**
- 




# Clinical Trials

- Conducted to allow safety and efficacy data to be collected for new drugs or devices
- Depending on the kind of product and the phase of its development, investigators register patients into small pilot studies
- Followed by bigger scale studies in patients that often compare to the new product with the presently prescribed treatment
- Safety and efficacy data are collected, the number of patients is typically increased.



# Clinical Trials On Memantine for FTD Treatment

- The purpose of the study is designed to determine whether Memantine is effective in slowing the rate of behavioral decline in FTD.
  - The study will also assess the safety and tolerability of long-term treatment with Memantine in patients with FTD or semantic dementia (SD);
    - whether Memantine is effective in slowing the rate of cognitive decline FTD;
    - and evaluate whether Memantine delays or decreases the emergence of parkinsonism in FTD.
- 

# References

- “CNS degenerative disease.” the internet pathology laboratory. July 21, 2009.  
<http://library.med.utah.edu/webpath/tutorail/cns/cnsdg..html>
- <<http://www.nlm.nih.gov/medlineplus/ency/article/000744.htm>>
- “frontotemporal dementia.” mayoclinic. July 18, 2009.  
<http://www.mayoclinic.com/health/forntotemporal-dementia/dsoo874>
- Bell, Dr.Karen. Personal Interview. 20 July, 2009
- <http://www.ftd-picks.org/frontotemporal-dementias/autopsy>
- Doraiswamy P, Murali and gwyther P, Lisa. The Alzheimer's action plan. 2008. pg. 28-31
- [http://memory.ucsf.edu/ftd/files/pdf/ftd/MAC\\_FTD\\_Primer.pdf](http://memory.ucsf.edu/ftd/files/pdf/ftd/MAC_FTD_Primer.pdf)
- Bioler F, and Traykov L. “Frontal lobes pathology and dementia. An apprasied of the contribution of the contribution of leanardo bianchi.” volume18, number 3, (march 1997): pg. 129-134.
- “ Forms of frontotemporal dementia.” UCSF. July 22, 2009. <[Http://memory.ucsf.edu/ftd/overview/forms/multiple](http://memory.ucsf.edu/ftd/overview/forms/multiple).>
- <http://www.ninds.nih.gov/disorders/picks/picks.htm>
- [http://nutmeg.conncoll.edu/Neurobiology-of-Disease/index.php/Picks\\_Disease](http://nutmeg.conncoll.edu/Neurobiology-of-Disease/index.php/Picks_Disease)



# Acknowledgements

- Dr. Karen Bell
- Dr. Evelyn Dominguez
- Dr. Ruth Tejeda
- Lynda Mules
- Columbia University Medical Center
- Dr. Sat Bhattacharya
- The HCS Staff
- Lee and Nick
- And everyone for listening!!

