#### Source:

# 1 | Preliminary Research

### 1.1 | Sources

https://www.frontiersin.org/articles/10.3389/fchem.2019.00540/full

#### 1.2 | **Notes**

### 1.2.1 | Target Processes

Enzyme catalysis
catalyzing reactions with actions

2. Protein-ligand binding

neurotransmitters (dopamine), protien is dopamine receptor how does the ligand bind the proper site to open the channel?

- · ligand: how to pronounce?
- 3. signal transduction

bind to other protein to trigger chain of actions

- · release calcium from intercellular stores
- 4. allosteric regulation
  - · a reason why knowing the structure and pockets is important
  - · predict allosteric cites
    - similar to non-competative inhibiton?
    - · but for dna binding protiens, like the dna transcription inhibitor
  - · ligand binds allosteric site and activates the protien

# 1.2.2 | Folding Simulation Methods

- 1. all-atom molecular dynamics (MD)
  - · Obtains all desired information regarding the kinetics and thermodynamics
  - (a) Time scale bottleneck
    - very slow (supercomputers -> microseconds of simulation)
    - · require microsecond to milisecond time scales
    - i. optimizations
      - A. conformational sampling?
        - · retains atomistic representation of the system
      - B. overcome kinetic trapping and thourough sampling of conformational space techniques
        - · umbrella sampling
        - · multicanonical algorithms
        - · simulated tempering

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- · transition path sampling
- · targeted molecular dynamics
- replica exchange method molecular dynamics (REMD)
- · accelerated molecular dynamics (AMD)
  - · see below
- 2. Accelerated molecular dynamics (AMD)

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## 1.3 | Meetings

### 1.3.1 | **12 oct 2020**

- · computational prediction modeling
  - · trying to predict the crystal structure
    - · why?
      - to analyze would this fit?
      - · does it work with this target
- · solving the structure
  - xray cristolography
    - · gold standard
    - · now got the structure
      - · what does that mean?
      - · can we simulate how it interacts?
      - · can you then do modeling on that to see if drug molecules work? are useful
- look at some concrete examples?
- · tell a biological story alongside with computational relevance piece
- 1. protien synthase

not as much simulation stuff

2. neurotransmitters

dopamine sodium rushes in, electrochemical and concentration gradient recharge gradient by releasing potassium

- (a) nerst equation electrochemical gradient as battery
- (b) goldman-katz equation
  - · applied to neuro
  - · ions?
- Case study
  - · why do we care? why is this useful
  - · knowing the structure can lead to some useful information
  - how did it lead to some sort of accelerated understanding?

4. prions

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- · how to pronounce?
- (a) CJD
  - is it inheritable?
  - · one case per million population
  - i. Casues
    - the gene that causes CJD in 5-10% of cases is PRNP
    - 87% of cases are sporatic
- (b) isoform
  - · a different set of intons and exons
  - · splicosome takes pre-RNA and cuts out intons
    - even if the pre-RNA had 10 exons, the splicosome might take a subset of those exons and remove the others
  - An isoform is a variant of that subset, an abnormal isoform is one that is "bad" and causes problems

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