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Modeling Calcium Dynamics in T Cells

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Acknowledgement

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1. Introduction

As part of the bodies' defence against viruses t cells can undergo activation in their lifetime. Whether and when a t cell activates is an interesting topic when studying immunology. However, measuring activation is often done indirectly by using the correlation between calcium concentration within the cell and activation. Measuring the calcium concentration leads to a time series that can be analysed by experts for activation.

This work aims to provide an algorithm for automatic detection of activation. By using approximation algorithms the time series is fitted with sigmoid functions. This reduces the data from hundreds of values in a time series to few parameters of the approximation function. Additionally, the parameters are chosen to be valuable for interpretation by experts. This parameter representation of the data is then filtered and used for clustering the data into activated and unactivated cells.

The provided algorithm can be used on any data set provided a positive control and negative control is provided. The most simple use case of finding the number of activated cells in the data set is described in more detail.

The proposed algorithm is tested with two of the most common types of t cells, human Jurkat cells and mouse 5c.c7 cells.

The main question this work aims to answer is which criteria can distinguish between unactivated and activated cells. Additionally, a criterion for detecting cells which activated before the experiment began will be investigated.

When only looking at activated cells some interesting questions are whether there are different types of activated cells and how they are different. A typical pattern observed in activated cells is that they show oscillations of the calcium concentration. Analysing the frequencies of these oscillations might be interesting.

Lastly differences between mouse and human cells show whether the proposed algorithm is applicable to the two most common types of t cells studied.

To summarize the questions are

- Which criteria can distinguish between unactivated, activated and pre-activated cells?
- Do different types of activated cells exists? How are they different?
- With which frequencies does the Calcium concentration repeat after activation?
- Is there a difference between mouse and human cells?

with the first question being the most relevant.

1. Introduction

This work starts with a chapter on optimization algorithms. In chapter 2 the relevant algorithm, Trust Region Reflective Algorithm, is attained from other algorithms, which are described as well.

Following is chapter 3 focusing on the biology of t cells. All relevant components of t cells for changes in calcium concentration are described. Their interconnections are described.

Next chapter 4 describes the structure and experimental setup for retrieving the data. Some processing steps performed on the data are outlined.

The main focus of this work, approximating the calcium concentration, are described in chapter 5. Here the approximation function is inferred and described. Parameter descriptions are provided. Pseudocode for the approximation algorithms are given and described. The parameters found from the approximation of the data sets used in this work are analysed. Oscillations are approximated in this chapter as well.

In chapter 6 the clustering algorithm Gaussian Mixture Model is described and applied to the output of the approximation. Visual representation of the clustering is shown.

Chapter 7 aims to answer the questions posed above by using the approximation and clustering described in the other chapters.

A final discussion of the results provided in this work is provided in chapter 8. The outlook is featured here as well.

2. Optimization Algorithm

An optimization problem is any problem where a function $f : X \rightarrow Y$ is given, and we search for the point $x \in X$ such that $f(x)$ is minimal or maximal. Obviously the minimum or maximum must not exist, as the example $f : (0, 1) \rightarrow \mathbb{R}, x \mapsto x$ demonstrates by not having either. Investigating conditions on X, Y and f such that a minimum or maximum exists is mathematically interesting. However, when implementing an optimization algorithm the true minimum or maximum can sometimes not be found even if it exists and is instead replaced by a sufficiently good approximation.

2.1. Gradient Descent

An iterative algorithm for finding the minimum of a differentiable function $f : \mathbb{R}^n \rightarrow \mathbb{R}$ is gradient descent. As the name suggests it uses information of the gradient ∇f . Locally the negative gradient always points into the direction of greatest descent. The idea is to follow this direction for the next guess of the minimum. The pseudocode of this approach is given below.

Algorithm 1: Gradient Descent

```
input :  $f : \mathbb{R}^n \rightarrow \mathbb{R}$  ... differentiable,  $x_0 \in \mathbb{R}^n$ 
output:  $x \in \mathbb{R}^n$ 

1 begin
2   for  $n = 0$  to  $max\_iterations$  do
3     if improvement is smaller than threshold then
4       | break
5     end
6     set or calculate step size  $\gamma_n$ 
7      $x_{n+1} = x_n - \gamma_n \nabla f(x_n)$ 
8   end
9    $x = x_n$ 
10 end
```

If we consider a function with a local minimum, that is not a global minimum, gradient descent might not converge to the optimum. An example of such a function can be seen in figure 2.1 along with the first few values x_n of gradient descent. The starting value was chosen to not have convergence to the global minimum. For a different starting value the global minimum can be reached.

Improvements can be made by choosing good step sizes, starting value or by starting

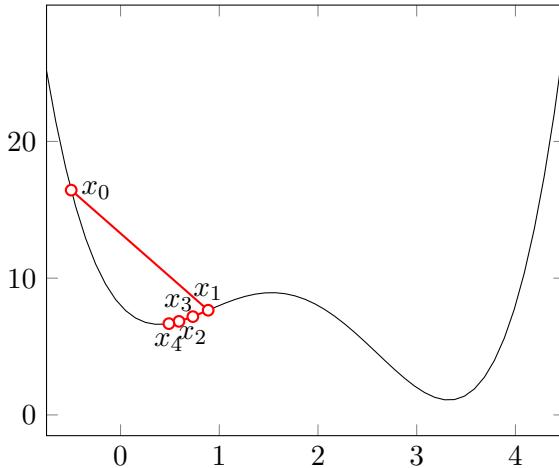


Figure 2.1.: The function has two local minima. For this starting value and step size gradient descent approaches the local, but not global minimum.

with different values and comparing the results.

2.2. Least Square Problem Algorithms

We now focus on the Least Square Problem and give an introduction into various algorithms solving this problem.

In the example dealt with in this work we are given some data points $((x_k, y_k))_{k \in \{1, 2, \dots, n\}}$ and want to find a close approximation in the form of a function $g(x, a_1, a_2, \dots, a_m)$ where for every list of function parameters $a = (a_1, \dots, a_m)$ we have the function $g_a(x) : \mathbb{R} \rightarrow \mathbb{R}, x \mapsto g(x, a_1, \dots, a_m)$. Searching for a good approximation can be reformulated as searching for the minimum of $r(a) := \sum_{k=1}^n |g_a(x_k) - y_k|^2$ or any other error function. This form of optimization problem is called the Least Square Problem.

First we want to discuss some variations of the problem. Easiest to solve are linear problems. These can be formulated as the minimization of $\|Ax - b\|^2$, and solved using calculus by $x = (A^T A)^{-1} A^T b$ provided the rank of A is full.

Often we want to constrain the search for a minimum under some properties. For linear problems we can find a formulation as

$$\text{minimize } \|Ax - b\|^2 \text{ subject to } Cx = d.$$

Finding a solution can be done by minimizing $\|Ax - b\|^2 + \lambda \|Cx - d\|^2$ for very large λ .

General least square problems are formally given as a residual function $r_f(x)$ which tells us whether a function f is a good approximation at the point x . We therefore want to find a way to minimize $\|r_f(x)\|^2$.

As $\|r_f(x)\|^2 \geq 0$ we can turn to the simpler problem of finding a root. However, a root must not exist, in which case we want to find the value closest to zero. This is then the minimum of the function.

Some algorithm for minimization are now discussed below.

2.2.1. Gauss–Newton Algorithm

The idea behind this algorithm is that it is easy to find the intersection with zero of a linear function. If we linearize $r : \mathbb{R}^n \rightarrow \mathbb{R}^m$ locally, we can approximate the root by finding it of the linear approximating function. This is demonstrated in figure 2.2.

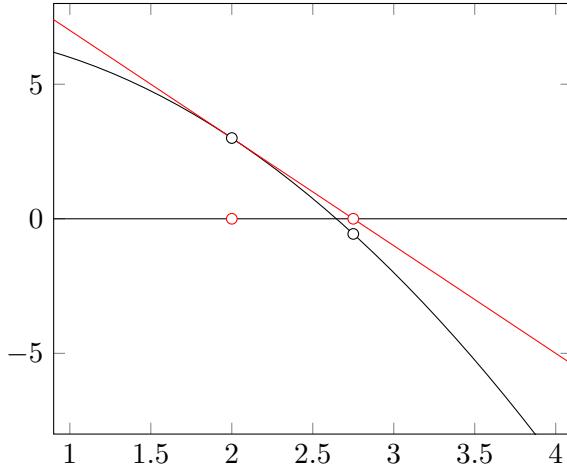


Figure 2.2.: By approximating the black function by a line an approximation of the root has been found.

Iterating this step of linear approximating gives us the Gauss-Newton Method. In figure 2.3 we can see that indeed x_n seems to converge towards the root of the function.

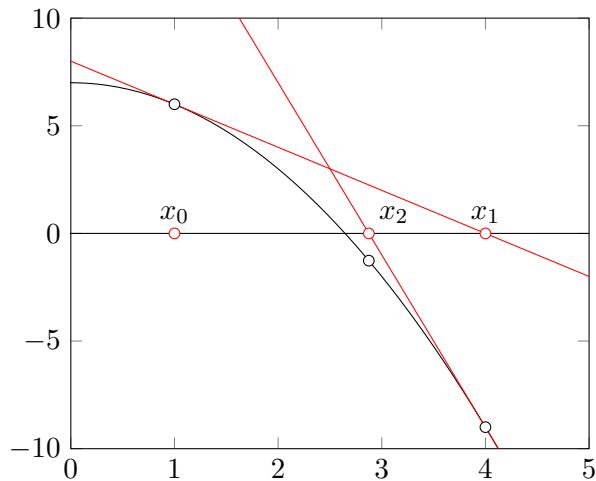


Figure 2.3.: Iteratively applying linear approximation gives the Gauss-Newton Method for approximating the root.

2. Optimization Algorithm

Define Dr as the Jacobian matrix $\left(\frac{\partial r_i}{\partial x_j}\right)_{ij}$. Using Taylor's theorem we get the linear approximation

$$r(x) = r(a) + Dr(a)(x - a) + h(x)(x - a) \approx r(a) + Dr(a)(x - a) \text{ with } \lim_{x \rightarrow a} h(x) = 0.$$

Rewriting this as $r(x) \approx Ax - b$ where $A := Dr(a)$ and $b := Dr(a)a - r(a)$ gives us the algorithm for this method. As $Dr \in \mathbb{R}^{n \times m}$ we solve $Dr^T Dr x = Dr^T b$ in order to get a system with square matrix. If $n = m$ we can skip this step and get the so-called Newton algorithm as a variant.

Algorithm 2: Gauss-Newton

```

input :  $r : \mathbb{R}^n \rightarrow \mathbb{R}^m$  ... differentiable,  $x_0 \in \mathbb{R}^n$ 
output:  $x \in \mathbb{R}^n$ 

1 begin
2   | for  $n = 0$  to  $max\_iterations$  do
3   |   | if  $\|r(x_n)\|^2$  close enough to zero or  $\|x_n - x_{n-1}\|$  is too small then
4   |   |   | break
5   |   | end
6   |   | Calculate  $A_n := Dr(x_n)$ 
7   |   | Calculate  $b_n := A_n x_n - r(x_n)$ 
8   |   | Solve  $A_n^T A_n x_{n+1} = A_n^T b_n$ 
9   | end
10  |  $x := x_n$ 
11 end
```

Gauss-Newton is guaranteed to find a local minimum x if r is twice continuously differentiable in an open convex set including x , Dr has a full rank and the initial value is close enough to x .

For the example demonstrated in figure 2.4 we can see that choosing a particular starting value leads to a loop in which only two points are explored as possible roots. More extreme examples exists in which Gauss-Newton gets increasingly further away from the root, due to an increasingly flat incline the further we get from the root. One example of such a function can be seen in figure 2.5.

Gauss-Newton has two problems, the starting value being too far from the root and Dr not having full rank. This can be combated using the technique of dampening. Instead of moving the new guess all the way to the root of the linear approximation we only move part of the way. How far to move can be determined by a dampening factor λ_n or a constant λ .

2.2.2. Levenberg-Marquardt Algorithm

This section is following section 18.3 of the book Introduction to Applied Linear Algebra by Stephen Boyd and Lieven Vandenberghe[BV18].

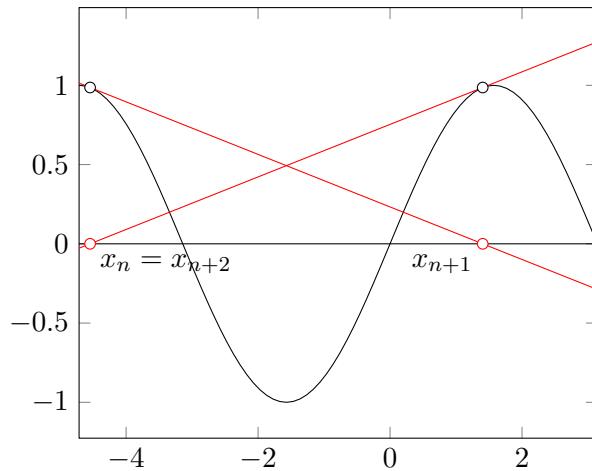


Figure 2.4.: For a poor choice of starting values Gauss-Newton can never find the root of the function $\sin(x)$.

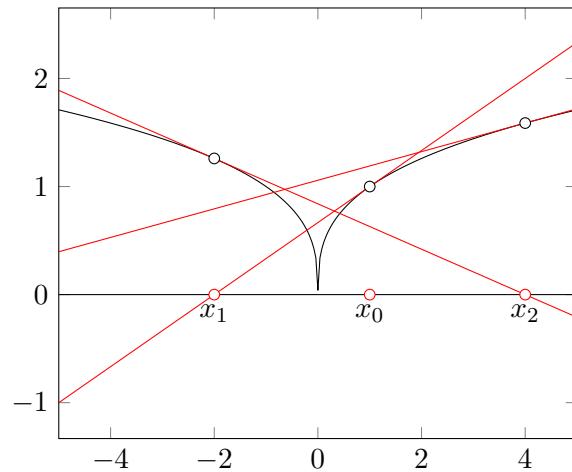


Figure 2.5.: Finding the root of the function $\sqrt[3]{|x|}$ using Gauss-Newton is only possible if the starting value x_0 is chosen as 0, which is the root. For any other value we have that the guess gets further and further away. Indeed for any x_n we have $x_{n+1} = -2x_n$.

2. Optimization Algorithm

As stated above, a shortcoming of Gauss-Newton is that for x far from x_n we must not have that $r(x) \approx r(x_n) + Dr(x_n)(x - x_n) =: \hat{r}(x, x_n)$. Levenberg-Marquardt addresses this by minimizing $\|\hat{r}(x, x_n)\|^2 + \lambda_n \|x - x_n\|^2$. The first part is the same as above, while the second objective expresses our desire to not stray away too much from the region where we trust the linear approximation. The parameter λ_n is a positive parameter specifying how far the trusted region extends.

Writing the above idea as a single squared norm to minimize gives us the problem

$$\text{minimize } \left\| \begin{pmatrix} Dr(x_n) \\ \sqrt{\lambda_n} I \end{pmatrix} x - \begin{pmatrix} Dr(x_n)x_n - r(x_n) \\ \sqrt{\lambda_n} x_n \end{pmatrix} \right\|^2.$$

We observe that as λ_n is positive the left matrix has full rank. From this it follows that a unique solution exists.

The change of including λ_n translates into the algorithm as replacing solving $A_n^T A_n x_{n+1} = A_n^T b_n$ in Gauss-Newton by solving $A_n^T A_n z + \lambda_n z = A_n^T b_n + \lambda_n x_n$.

The question of how to choose λ_n arises. If too small x_{n+1} can be too far from x_n to trust the approximation. If too big the convergence will be slow. If in the previous step the objective $\|r(x_n)\|^2$ decreased we decrease λ_{n+1} slightly. If the last step was not successful λ_n was too small. Therefore we increase λ_{n+1} .

Pseudo code of the resulting Levenberg-Marquardt algorithm is shown below. The stopping criteria of $\|2Dr(x_n)^T r(x_n)\|$ being too small is known as the optimality condition. It is derived from the fact that $2Dr(x)^T r(x) = \nabla \|r(x)\|^2 = 0$ holds for any x minimizing $\|r(x)\|^2$. Note that this condition can be met for points other than the minimum.

Algorithm 3: Levenberg-Marquardt

```

input :  $r : \mathbb{R}^n \rightarrow \mathbb{R}^m$  ... differentiable,  $x_0 \in \mathbb{R}^n$ ,  $\lambda_0 > 0$ 
output:  $x \in \mathbb{R}^n$ 

1 begin
2   for  $n = 0$  to  $max\_iterations$  do
3     Calculate  $A_n := Dr(x_n)$ 
4     Calculate  $b_n := A_n x_n - r(x_n)$ 
5     if  $\|r(x_n)\|^2$  close enough to zero or  $\|2A^T r(x_n)\|$  is too small then
6       | break
7     end
8     Solve  $(A_n^T A_n + \lambda_n)z = A_n^T b_n + \lambda_n x_n$ 
9     if  $\|r(z)\|^2 < \|r(x_n)\|^2$  then
10      |  $x_{n+1} := z$ 
11      |  $\lambda_{n+1} := 0.8\lambda_n$ 
12    else
13      |  $x_{n+1} := x_n$ 
14      |  $\lambda_{n+1} := 2\lambda_n$ 
15    end
16  end
17   $x := x_n$ 
18 end

```

Coming back to the example where Gauss-Newton failed we once again consider the function from figure 2.5. In comparison this time we are able to find a good approximation of the root using Levenberg-Marquardt. A few steps are demonstrated in figure 2.6.

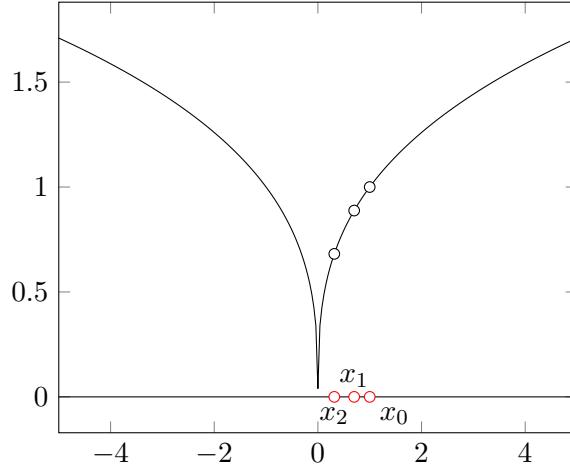


Figure 2.6.: In comparison to Gauss-Newton, Levenberg-Marquardt is able to find the root of the function $\sqrt[3]{|x|}$. From the starting value $x_0 := 1$ and using $\lambda_0 := 1$ the guesses x_n move towards the root $x = 0$.

Further improvements to this algorithm can be made using good starting values perhaps from the output of other algorithms or by letting the algorithm run multiple times with different starting values and comparing the results.

2.2.3. Algorithms for Bounded Least Square Problems

The algorithms described above do not consider bounds. For bounded problems two algorithms known as Trust Region Reflective Algorithm and Dogleg Algorithm with Rectangular Trust Regions can be used.

Trust Region Reflective gets its name from the use of trust regions as in Levenberg-Marquardt as well as reflecting along the bounds [BCL99]. If an iterative x_n lands outside the bounds set, it is replaced by a reflected value within the bounds. This ensures each iteration is feasible as a solution.

As the name suggests the Dogleg Algorithm with Rectangular Trust Regions uses rectangular trust regions as opposed to ellipsoids. [VL04] As the bounds are specified as a rectangle to stay within, this results in the intersection of trust region and bounds to be rectangular. The resulting minimizing problem is solved with an adequate algorithm [NW99].

In Python the library SciPy provides the three methods Levenberg-Marquardt, Trust Region Reflective and Dogleg with Rectangular Trust Regions for solving Least Square Problems.

3. Background Information on T Cells and Calcium Concentration

Lymphocytes form a key component of the immune system. T cells are a type of lymphocyte and are responsible for responding to viruses, fungi, allergens and tumours. Different subtypes of t cells exist, that perform various responsibilities. They are transported throughout the body via the lymphatic system and blood.[KCF18]

Precursor cells are formed in the bone marrow. Once they are transported to the thymus they undergo maturation and selection to become t cells. Each cell forms receptors, called t cell receptors (TCR), that respond to one particular out of many (10^6 – 10^9) possible short pieces of proteins, called peptides. These peptides are attached to the major histocompatibility complex (MHC) present on antigens and antigen presenting cells (APC). Important aspects of the selection are ensuring that the t cells react to foreign peptides, but not to those present on the body's own cells.[AH24]

In positive selection, cells in the thymus present peptides on their MHC. If a t cell is unable to bind, it will undergo apoptosis, a type of cell death. T cells which were able to bind receive survival signals. Negative selection verifies that t cells will not attack the body's own cells. This is done by only selecting t cells which only bind moderately to the peptides presented, as a strong bond suggests that these t cells would have a high likelihood of being reactive to own cells.[Hag18] If a t cell passed both the positive and negative selection it is transported to the periphery.

There are multiple types of peripheral t cells. Native t cells respond to new antigens. Cytotoxic t cells kill cells which present peptides on their MHC compatible with the t cells TCR. Helper T cells activate other parts of the immune response. Memory t cells shorten the reaction time when the same antigen is encountered again at a later point in time. Suppressor t cells moderate the immune response.[Gan97]

3.1. Components of a T Cell

T cell components relevant in activation and subsequent changes in intracellular Ca^{2+} are listed below and schematically shown in figure 3.1.

- **T cell receptor (TCR):** Receptor on the cell surface that can recognize peptides. By the simultaneous triggering of the TCR and co-stimulator, signalling is induced that leads to activation.
- **Co-stimulator:** A stimulation of co-stimulatory molecules is necessary in order for signalling to occur as part of activation.

3. Background Information on T Cells and Calcium Concentration

- **Endoplasmic reticulum (ER):** A series of connected sacs in the cytoplasm that is attached to the nucleus. Important functions are folding, modification and transportation of proteins.[Rog24]
- **Ca²⁺ permeable ion channel on the ER:** There are several Ca²⁺ channels present on the ER. Some receptors are responsible for releasing Ca²⁺ into the cytoplasm, when the intracellular Ca²⁺ concentration is low. [SB16]
- **Ca²⁺ storage in the ER:** Ca²⁺ is stored in the ER and can be released by Ca²⁺ permeable ion channels on the ER.
- **Cytoplasm:** The semi-fluid substance enclosed in the plasm membrane. It contains organelles, ions, proteins and molecules.
- **Stromal interaction molecule (STIM):** If the Ca²⁺ storage in the ER is depleted STIM proteins cluster where the ER is in the vicinity of the plasm membrane and assembles CRAC, which then leads to uptake in extracellular Ca²⁺. [SB16]
- **Plasm membrane:** A semipermeable structure forming the wall of the cell made up of lipids and proteins. Ion channels and transport proteins allow certain substances to move through.[Gan12]
- **Ca²⁺ release activated Ca²⁺ channel (CRAC):** Opened after a decrease in ER stored Ca²⁺ is sensed by STIM, these channels intake Ca²⁺ from outside the cell.[SI13]
- **Cytoskeleton:** A system of fibres within the cell, that allows it to change shape and move.[Gan12]
- **Nucleus:** An organelle that stores most of the DNA, controls cell growth and cell division. A double membrane separates it from the cytoplasm.[CA22]

Relevant components of APC are the

- **Major histocompatibility complex (MHC),** which can present peptides, and the
- **Co-stimulator,** which can form a bond with the co-stimulator on a t cell.

Both are present on the surface of the APC.

3.2. Activation of T Cells

Activation is necessary for t cells to divide and perform their functions.[Gan97]

When a native t cell encounters a peptide on an APC that is compatible, a bond is formed between the TCR on the t cell and the peptide-MHC complex on the APC. This recognition can be triggered by less than ten molecules of foreign substance and is therefore described as near perfect. Sufficiently long contact is necessary between the APC and the t cell in order for the t cell to activate. The role of contact time in t cell activation is modelled by Morgan et al.[ML23].

3.2. Activation of T Cells

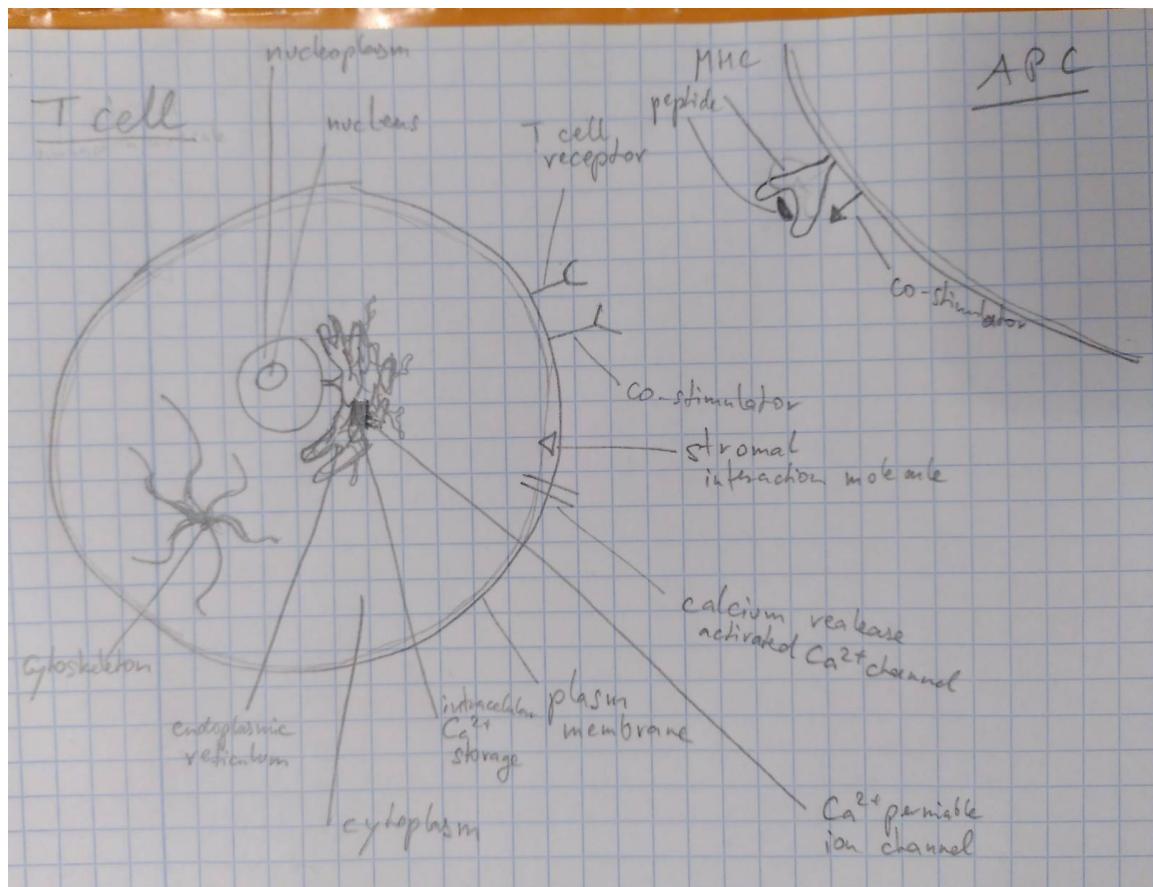


Figure 3.1.: Schematic view of a t cell and antigen presenting cell, with all relevant components. [TODO draw image]

3. Background Information on T Cells and Calcium Concentration

The presence of co-stimulatory molecules is needed for activation. The bond between the co-stimulatory molecules on the t cell and the one on the APC plays a role in signalling. Especially Ca^{2+} signals play a vital part in t cell activation.

An increase of Ca^{2+} in t cells during activation is caused by the stimulation of Ca^{2+} permeable ion channel receptors on the ER membrane. Ca^{2+} is released from the ER into the cytoplasm. Additionally, this decrease in Ca^{2+} is sensed by STIM, which leads to an influx of Ca^{2+} through plasma membrane CRAC channels.[\[SKJ09\]](#)

As the intracellular Ca^{2+} concentration is dependent on the interaction between Ca^{2+} sources and sinks, a variety of different forms in Ca^{2+} concentration have been observed. Examples are infrequent spikes, sustained oscillations and plateaus.[\[Lew01\]](#)

Intercellular Ca^{2+} increase together with other signals lead to a redistribution of receptors, signalling molecules and organelles.[\[JRB14\]](#)

4. Calcium Data of T Cells

From section 3.2, we gather that analysing the intracellular Ca^{2+} concentration gives us good insight in whether and when a cell activates. Additionally, it can be measured relatively easily by the method described in this chapter.

4.1. Structure of the Data

First we describe the structure of the data this work uses.

The data matrix has one row for each combination of tracked particle and frame number. In this context cells are called particles as the recording might feature non-cells that are detected as a cell and recorded in the data set. The information stored for each particle and frame combination is described in detail in 4.1.

Name	Data Type	Description
x	float64	Position of particle in pixels along the horizontal axis
y	float64	Position of particle in pixels along the vertical axis
frame	int32	Number of frame, with frame rate of 1 frame per second
mass short	float64	Brightness of cell in 340nm channel
mass long	float64	Brightness of cell in 380nm channel
ratio	float64	Calculated as mass short divided by mass long
particle	int32	Identification for each particle

Table 4.1.: Description and data type of all columns present in the data matrix.

One recording can have between 500 and 10000 particles and is between 700 and 1000 frames long, which corresponds to between about 11 and 17 minutes. The ratio recorded is typically between 0 and 5.

Four recordings were generated, with two each from human and mouse cells. For each cell type a positive and negative control was measured. In a positive control the conditions are such, that in theory every cell should activate, while in negative control the conditions are such, that none should activate. Due to stress on the cells caused by the movement or changes in temperature and other factors a few cells will activate before the recording starts, during the recording in the negative control or not activate at all in the positive control, regardless of the conditions.

4.2. Jurkat Cells, 5c.c7 Primary Mouse T Cells and Fura-2

The prototypical cell line to study T cell signalling is the Jurkat cell line.[\[ML23\]](#) It was obtained from the blood of a boy with T cell leukaemia.[\[SSB77\]](#) Different cell lines within

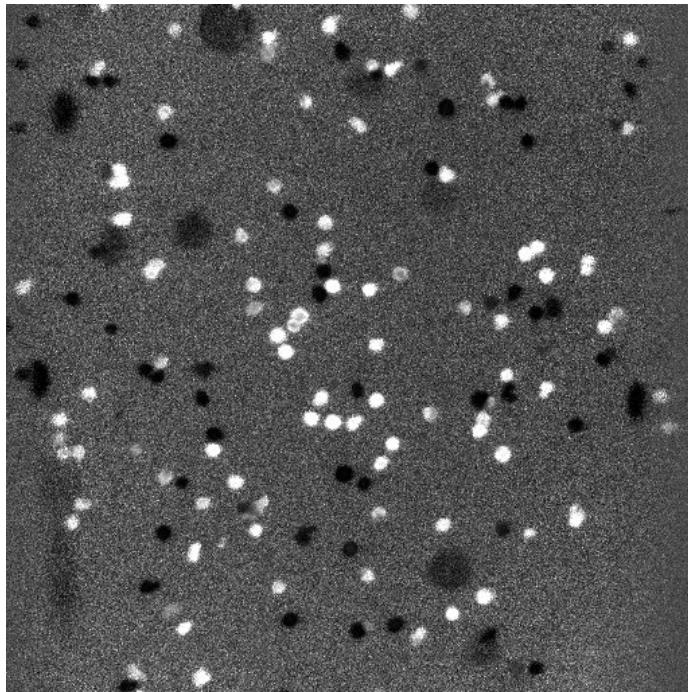


Figure 4.1.: Single frame showing the ratio of the 340nm and 380nm images from a recording of human Jurkat cells. Activated cells appear lighter, unactivated cells darker than the background. Big dark circles are out of focus cells that have not yet settled on to the plate.

the Jurkat family are described by Abraham and Weiss.[AW04] They provide a timeline of discoveries linked to Jurkat cells and t cell receptor signalling. Another type of t cells used in signalling studies are gathered from mice.

In order to be able to measure the intracellular Ca^{2+} concentration of cells they can be labelled with Fura-2. This method provides a way to record the Ca^{2+} concentration of multiple cells over a time period.[MMS17] Challenges encountered when using Fura-2 on certain cell types are described by Roe, Lemasters and Herman along with their respective solutions.[RLH90]

4.3. Measuring the Calcium Concentration of T Cells

After the cells have been labelled with Fura-2, a recording of up to 15 to 20 minute can be generated. To achieve this the cells and stimulant are photographed at both 340nm and 380nm wavelength once per second. The resolution of the images are 1.6um per pixel. By calculating the ratio of the two images at each pixel the Ca^{2+} concentration can be observed. An exemplary resulting image showing the ratio is shown in figure 4.1. The t cells appear a lighter shade than the background when activated and darker when not activated.

To activate the cells in the duration of the recording they are transferred to a plate

covered with replicas of the MHC-peptide complex normally present on APCs. This plate is then recorded as described above. For a negative control the plate is not covered with peptides, while for the positive control the peptide covering on the plate is very dense. Recordings of different densities in peptides lead to activation of a percentage of t cells.

4.4. Processing the Data

To track single t cells moving around during the video the sum of the 340nm and 380nm image of each second is calculated. This image provides the basis for separating t cells from the background. On this image all t cells will appear similarly light in colour. Therefore, it is used to track the movement of cells. Each cell is numbered, such that the same cell will have the same number during the video. For some cells the trajectory tracking is not perfect, resulting in a split of the numbering into multiple numbers for the same cell. The position and shade during both 340nm and 380nm as well as the ratio of each particle and each frame is then recorded into the data structure used in this work. The first roughly 50 frames at the start of the recording are discarded due to the video being out of focus. Additionally, cells only appearing in fewer than 300 frames are discarded as they most likely represent trajectories incorrectly tracked or split. The resulting data is then stored in a matrix structured as described in table 4.1.

5. Approximating the Calcium Concentration

If we have a look at the typical trajectory of the calcium concentration in activated and unactivated cells, shown in figure 5.1, we can see differences emerging. For one the maximum concentration value reached by most activated cells is higher. Another distinguishing feature is the presence of a steep incline at the moment of activation.

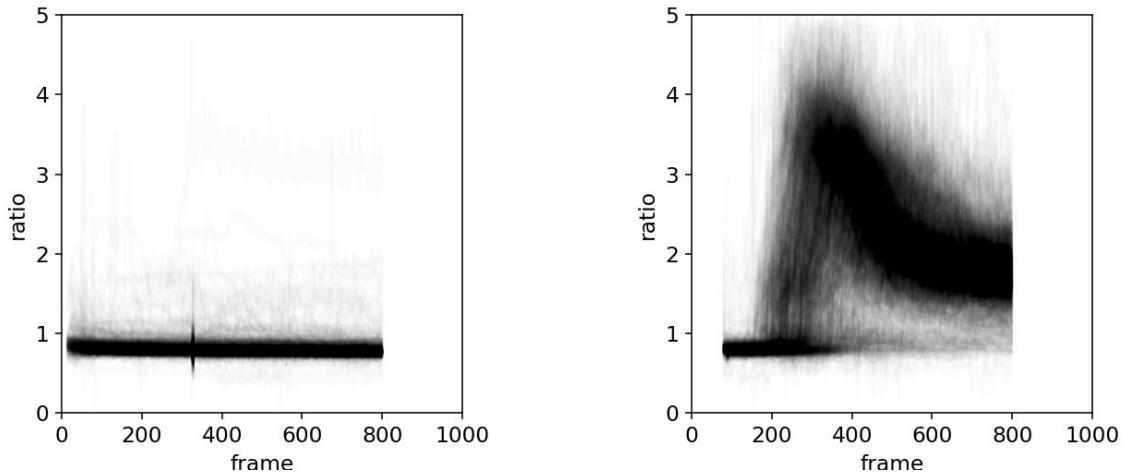


Figure 5.1.: Two plots of the overlapping calcium concentration time series of cells. On the left a negative control and on the right a positive control of mouse cells.

By modelling the time series with a function incorporating features such as the increase, maximum value and oscillations present in the decrease afterwards, we can extract these features more easily. By doing this, using approximation methods from chapter 2, we hope to have an easier method to answer the questions from the introduction.

5.1. Approximation Function

From studying the data in the two control groups we find to expect a function close to

$$f_{unac}(x) := u \quad (5.1)$$

for unactivated cells and

$$f_{ac}(x) := \begin{cases} \frac{a-u}{1+e^{-k_1(x-w_1)}} + u & \text{if } x \leq t \\ \frac{a-d}{1+e^{-k_2(x-w_2)}} + d & \text{else} \end{cases} \quad (5.2)$$

5. Approximating the Calcium Concentration

u ...	average value before activation,
a ...	value reached at the peak of activation,
d ...	average value after activation,
k_1 ...	steepness of increase,
k_2 ...	steepness of decrease,
w_1 ...	time point at which the increase happens,
w_2 ...	time point at which the decrease happens,
t ...	time point at which the increase ends, and the decrease starts,

Table 5.1.: List of parameters and their interpretation.

for activated cells. The parameters can be understood as described in table 5.1.

Figure 5.2 shows the above functions 5.1 and 5.2, and shows the relations to the parameters in unactivated and activated cells. The similarity between these functions and figure 5.1 can be observed.

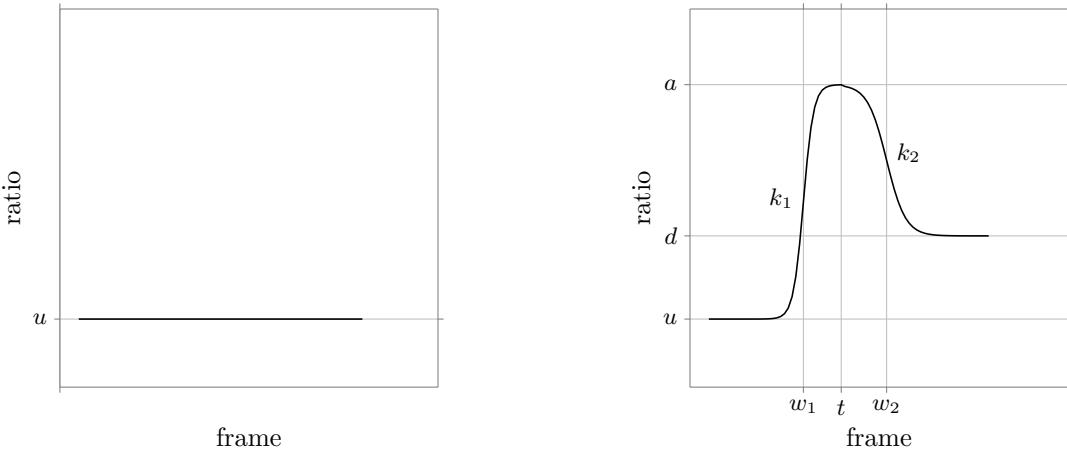


Figure 5.2.: Left shows the function f_{unac} defined in 5.1 with the parameter u . The right shows the function f_{ac} defined in 5.2 with the parameters $u, d, a, w_1, t, w_2, k_1$ and k_2 .

For our model to make sense we have to impose some conditions onto the parameters. We expect

$$0 \leq u \leq d \leq a, \quad w_1 \leq t \leq w_2, \quad k_1 > 0 \quad \text{and} \quad k_2 < 0.$$

There are multiple ways in which the parameters of f_{ac} can be chosen to get a function similar to f_{unac} . If w_1 is very large or $u \approx d \approx a$ then f_{ac} approaches a constant value of u , thus approximating f_{unac} . If the approximation of a cell has parameters with w_1 very large or $u \approx d \approx a$ we can therefore expect it to be of an unactivated cell. Otherwise, it is more probable to be activated.

5.2. Implementation of the Approximation

Now that we have defined our model functions we will implement a routine that fits such a f_{ac} -function through the data points of a particle recording.

First we give the pseudocode for approximating a single particles time series with the approximation function described above. It takes a (frame, ratio)-matrix of a single particle as input and returns the corresponding parameter list of the approximation.

Algorithm 4: Approximate

```

input : particle data as (frame, ratio) matrix
output: parameters describing the approximation

1 begin
2   set boundaries for parameters
3   set start values for parameters
4   use Trust Region Reflective Algorithm with boundaries and start values to get
      parameters
5   calculate corresponding approximation and add as fit_sigmoid columns to data
      matrix
6   return parameters
7 end
```

The parameters of f_{ac} used in the approximation are not independent of each other as we want to choose t to be the point at which the increasing part of the function, $(a - u)/(1 + e^{-k_1(x - w_1)})$, almost reaches the value a . We choose

$$t := w_1 - \log(1/0.99 - 1)/k_1 = w_1 - \log(1/99)/k_1$$

as the function has had 99% of the increase of the sigmoid curve up to this point.

Setting the boundaries in line 2 is non-trivial. We have noted that a condition such as $0 \leq u \leq d \leq a$, $w_1 \leq t \leq w_2$, $k_1 > 0$ and $k_2 < 0$ are expected. We want to impose them using boundaries in which the parameters must lie. However, boundaries for each parameter must not depend on other parameters. We can circumvent this by changing the parameters to be relative to each other. As $u \leq d \leq a$ we choose to use the three parameters u , $d - u$ and $a - d$. We can then set the lower boundary to be 0 which ensures

$$\begin{aligned} 0 \leq u \quad & \wedge \quad 0 \leq d - u \implies d \geq u \quad & \wedge \quad 0 \leq a - d \implies a \geq d \\ & \implies 0 \leq u \leq d \leq a. \end{aligned}$$

Using the same method, we choose the parameters $w_1 - start$ and $w_2 - w_1$, where $start$ is the first frame in which the particle was tracked. The resulting boundaries are described in table 5.2, where we set min val, max val and median val as the minimum, maximum and median of the particles' ratio data respectively while start and end is the first and last frame where data was recorded for this particle.

5. Approximating the Calcium Concentration

The condition $t \leq w_2$ can be violated, but it is ensured that at least $w_1 \leq w_2$.

The other conditions are met as $k_1 \in [0.05, 10] \implies k_1 > 0$ while $k_2 \in [-1, -0.01] \implies k_2 < 0$ and

$$t = w_1 - \underbrace{\log(1/99)/k_1}_{<0} \geq w_1.$$

parameter	lower bound	upper bound	starting value
u	min val	max val	min val
$d - u$	0	max val	median val - min val
$a - d$	0	max val	max val - median val
$w_1 - start$	0	end - start	0
$w_2 - w_1$	0	end - start	(end - start) / 2
k_1	0.05	10	0.1
k_2	-1	-0.01	-0.03
d	min val	2 max val	median val
a	min val	3 max val	max val
w_1	start	end	start
w_2	start	2 end - 2 start	(start + end)/2

Table 5.2.: Upper and lower bounds as well as starting value for each of the parameters.

The boundaries and starting values of d, a, w_1 and w_2 are derived from the parameters used in the implementation of the approximation, shown above the double line.

Starting values can have a big impact on the approximation reached by the algorithm. We want to choose starting values close to the expected resulting parameters. By choosing the starting value of $w_1 - start$ as 0, which corresponds to choosing $w_1 = start$, we favour the first increase in the data to be the point of activation. Otherwise, we are more likely to mistake an oscillation later in the data as the activation point. As we do not know when the activation happens when setting the boundaries we guess that w_2 will lie somewhere in the middle. Therefore we choose $(end - start)/2$ as the starting value for $w_2 - w_1$. The other starting values are chosen as we expect u to be low, a to be high, d to lie somewhere in the middle. Experimenting showed that k_1 often has a value around 0.1 while k_2 lies around -0.03.

Using algorithm 4 we now describe a routine which handles reading the data, some necessary preprocessing steps and saving of the resulting parameter lists.

Algorithm 5: Approximation Loop

```

input : file containing data matrix as described in section 4.1
output: parameters of the approximation of all particles as a matrix

1 begin
2   |  read data
3   |  filter data
4   |  for each single particle do
5   |    |  particle data := (frame, ratio) columns of this particle
6   |    |  if length of particle data is too short then
7   |    |    |  skip
8   |    |  end
9   |    |  parameters := approximate(particle data)
10  |    |  optionally show ratio data and approximation
11  |    |  save parameters
12  |  end
13  |  return matrix of all parameters
14 end

```

Filtering the data is necessary as the ratio can be very large if the denominator is small. Values are therefore bounded to lie within the interval $[0, 5]$. Any values higher than 5 are almost certainly caused by measurement errors. Values below 0 are definitely incorrect, as both the denominator and enumerator are measured as the brightness of a pixel, which can not be negative.

The visualization in line 10 of algorithm 5 generates images such as figure 5.3.

This work uses the python scipy function `scipy.optimize.curve_fit(function, xdata, ydata, p0=starting_values, method='trf', bounds=(lower_bounds, upper_bounds))` as it provides all the necessary functionality. The method parameter `trf` stands for Trust Region Reflective, as described in section 2.2.3.

5. Approximating the Calcium Concentration

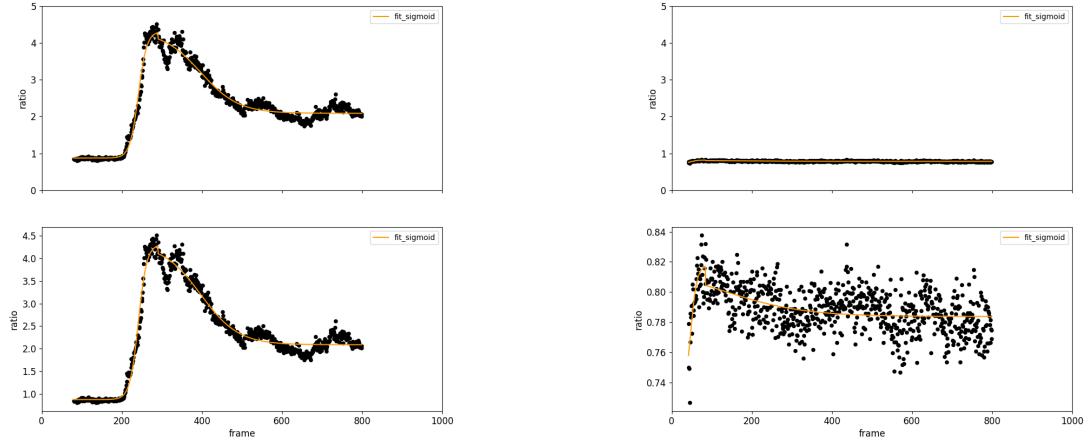


Figure 5.3.: The left plot shows the data in black and approximation in orange of an activated cell. The first plot is scaled from 0 to 5, the second one is scaled to fit the data. The right plot shows the same of an unactivated cell.

5.3. Analysis of the Approximation

We now give data on the parameters found from the above approximation. Some statistics are found in table 5.3. Figure 5.4 shows the distribution of the resulting parameters of the approximation. From the figure it seems the differences between activated and unactivated cells is biggest in the parameters activated value a , decreased value d as well as the steepness of increase k_1 .

As the datasets are not perfectly labelled, meaning there are activated cells in the negative control and vice versa, we have relatively high standard deviation.

We can use the mean and standard deviation of each of the parameters to find data points that can be considered outliers. We expect wrongly-labelled data, e.g. activated cells in the negative control, to be an outlier in the parameters a and d . However, activated cells in the positive control might have a decreased value d that is very low, around u . This makes it difficult to distinguish activated from unactivated cells when looking at the parameter d . Therefore, we choose a as the only parameter when filtering for these kinds of outliers.

A particle from the positive control dataset that has a value in parameter a higher than the median should still be classified as activated. Only a value lower than some threshold indicates an unactivated cell. The same holds for values of a lower than the median in the negative control dataset. In short, we want to filter out particles with a high value of a in the negative control and those with a low value of a in the positive control dataset. The question of how to choose the threshold will be discussed next.

As we do not have information on what percentage of cells behaved correctly in the positive and negative control we do not have enough information to choose threshold values without guessing. Instead, we can manipulate the threshold as a multiple of the standard

		Positive Control		Negative Control		
	Parameter	Average	Standard Deviation	Average	Standard Deviation	Difference
human cells	a	2.808	0.461	0.923	0.669	1.885
	u	0.663	0.521	0.613	0.311	0.05
	d	1.937	0.491	0.685	0.412	1.252
	k_1	0.263	0.428	0.524	0.963	-0.261
	k_2	-0.059	0.164	-0.163	0.292	0.104
	w_1	142.228	124.012	171.062	131.563	-28.834
	w_2	445.386	185.971	478.843	190.792	-33.457
mouse cells	a	2.9	0.907	0.876	0.186	2.024
	u	0.889	0.27	0.79	0.093	0.099
	d	1.749	0.407	0.804	0.129	0.945
	k_1	0.15	0.409	1.161	1.235	-1.011
	k_2	-0.1	0.195	-0.133	0.267	0.033
	w_1	295.809	77.207	100.712	112.352	195.097
	w_2	469.952	105.375	304.283	179.834	165.669

Table 5.3.: Statistics of the parameters retrieved from approximating the human cell data.

deviation until we filter out incorrectly labelled data, but would filter out correctly labelled data points if we increase the value. This trial and error approach led to different values for each of the four control datasets, which can be seen in table 5.4.

dataset	lower bound	upper bound
human positive	mean -3 std = 1.582	∞
human negative	$-\infty$	mean $+0.5$ std = 1.306
mouse positive	mean -2 std = 1.41	∞
mouse negative	$-\infty$	mean $+3$ std = 1.445

Table 5.4.: Thresholds in outlier detection in the different datasets.

Naturally we can use the same outlier detection with different parameters to find particles where the approximation failed to yield a good result.

These results will be used in section 7.1 to remove wrongly-labelled data from the datasets.

5.4. Adding Oscillation in the Decrease

In order to answer the questions from chapter 1 concerned with the oscillations happening in the decrease of the Ca^{2+} concentration we want to model them as well. We use a method often used when analysing oscillating data, called Fourier Transformation.

Fourier Transformation is used when an application is concerned with cyclic temporal data. Examples are sound waves, seismic data or oscillations of a skyscraper in strong wind. This data can be represented as a function of amplitude over time. Most of the time

5. Approximating the Calcium Concentration

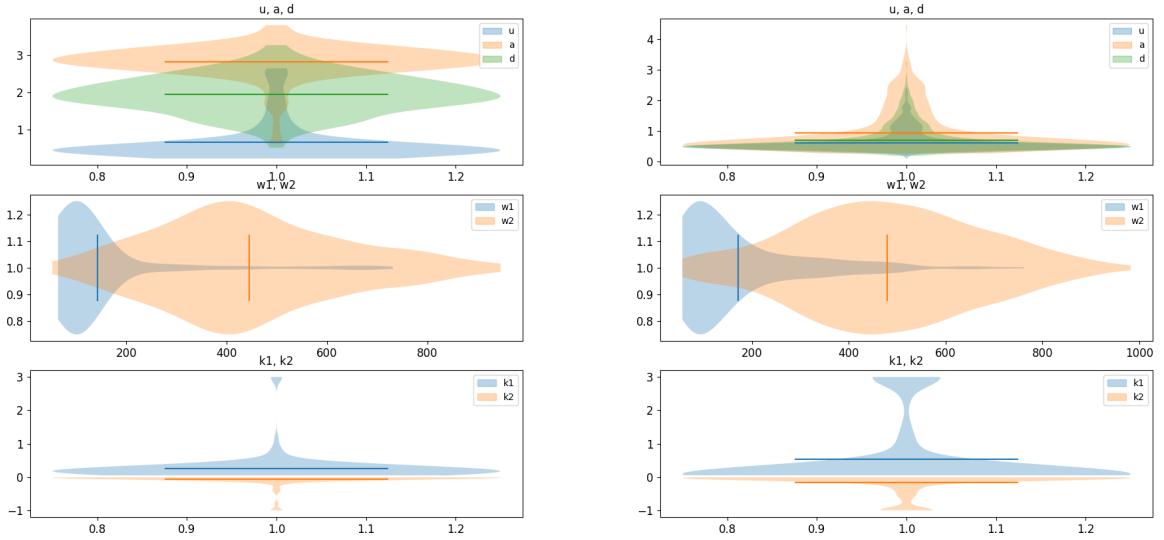


Figure 5.4.: Violin plots of parameters u, a, d, w_1, w_2, k_1 and k_2 from the approximations. The parameters of the positive control are on the left and those of the negative control are on the right. Both are of human cells.

we are not interested in the amplitude at a specific point in time, as a temporal shift would represent very similar information. Such a shift is demonstrated in figure 5.5.

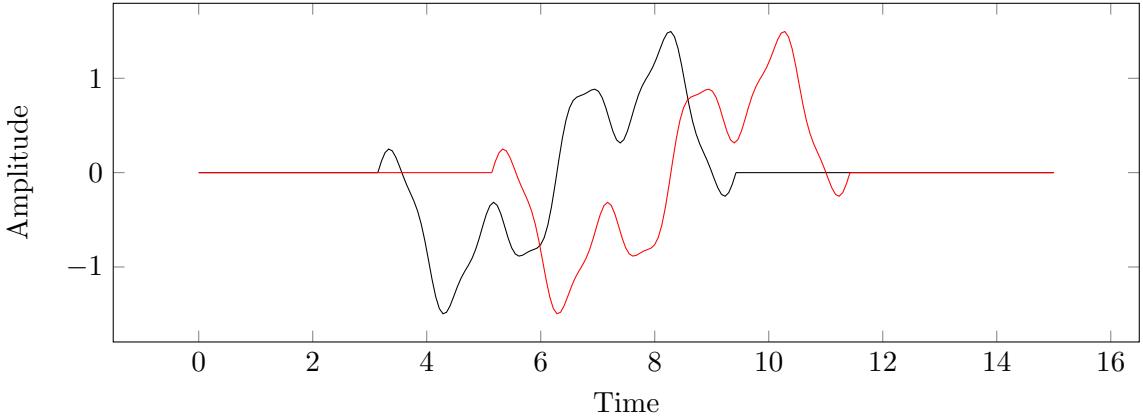


Figure 5.5.: Two signals that differ by a temporal shift.

As the function of sound waves or oscillations in the Ca^{2+} concentration in t cells is almost cyclic we might be interested in a decomposition into simple cyclic function, such as sine. We can then analyse the most prominent frequencies and their respective amplitudes. This gives a representation of the data, that can be easier to interpret. Fast Fourier Transformation (FFT) is an algorithm that transforms temporal data into such a representation of a weighted sum of sines.

As the oscillations happen in the decrease of the Ca^{2+} concentration we apply FFT to that part of the data. We can then filter out the 10 frequencies with the highest amplitudes

and use them to further analyse the oscillations. This gives an even better approximation of the data, which can be seen in figure 5.6.

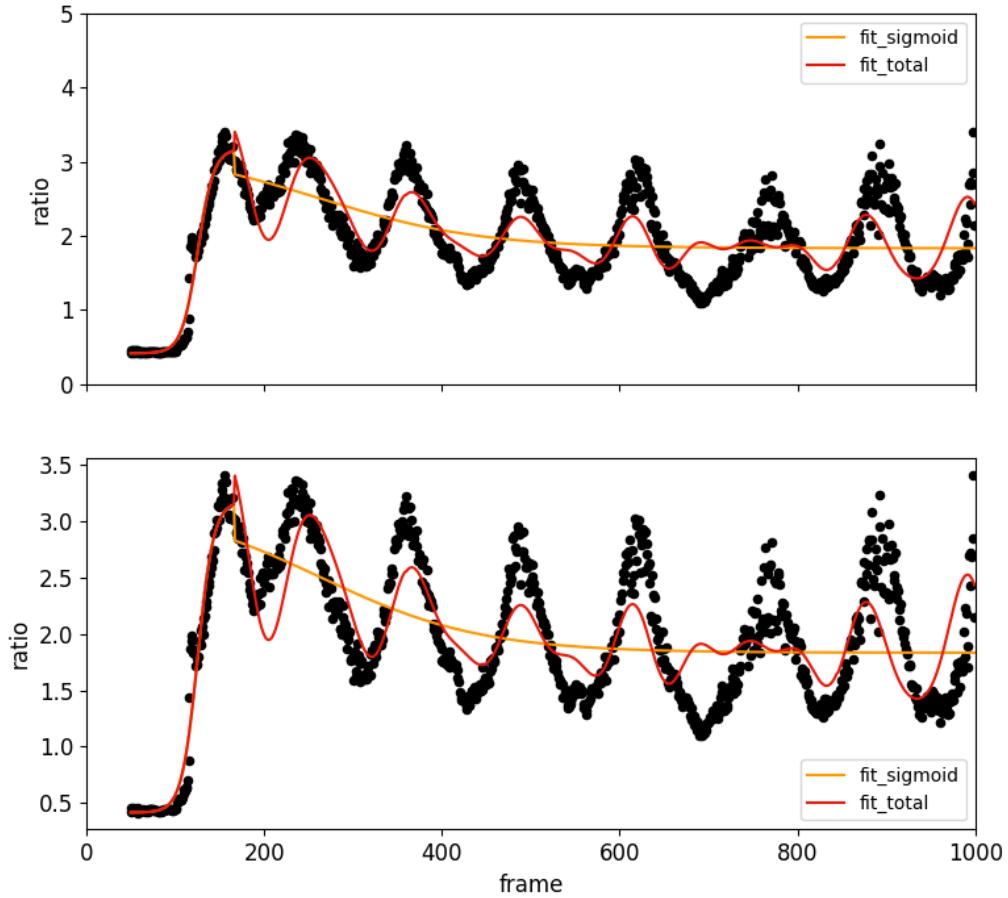


Figure 5.6.: The data of an activated cell with heavy oscillation is shown in black, simple approximation in orange and the approximation with FFT added in red. The first plot is scaled from 0 to 5, the second one is scaled to fit the data.

We can store the data gathered using the FFT as a list of frequencies and corresponding amplitudes.

[TODO analyse frequencies and amplitudes]

6. Clustering Algorithm and Application

The objective of classification is to find assignments between data points and categories. For some applications this can be done by taking correctly labelled data and comparing a new data point to the data points in different categories to see which category best fits. One such algorithm is k-nearest-neighbour. In our context the issue with this approach is that the data is only labelled as to which experiment it came from, e.g. positive control in human cells, negative control in mouse cells. However, as noted before not all cells from these experiments behaved as we expected them to, e.g. some cells activated in the negative control or did not activate in the positive control. Therefore, we choose to use a clustering algorithm which does not have the need for classified training data.

6.1. Gaussian Mixture Model

This section follows the article Gaussian Mixture Model by Reynolds [Rey+09].

Often gathered observations are distributed as a normal distribution. These distributions have a density function

$$g(x|\mu, \Sigma) = \frac{1}{(2\pi)^{D/2} |\Sigma|^{1/2}} \exp\left(-\frac{1}{2}(x - \mu)^T \Sigma^{-1} (x - \mu)\right)$$

with parameters D called the dimension, μ called the mean vector and Σ called the covariance matrix.

As we are concerned with clustering data points we expect the observed data points from different clusters to have different parameters in the normal distribution they come from. Assuming we have n different data sources gives us normal distributions $g_i(x|\mu_i, \Sigma_i)$ where $i = 1, \dots, n$. Additionally, we might have more data points being generated from some normal distributions while less from others. We can express this using another weight parameter w_i with $i = 1, \dots, n$. To normalize the weights we set the constraint $\sum w_i = 1$.

The distribution describing the entire dataset now can be described with the distribution

$$p(x) = \sum_{i=1}^n w_i g(x|\mu_i, \Sigma_i). \quad (6.1)$$

Gaussian Mixture Model is a method to retrieve these parameters w_i , μ_i and Σ_i for some D dimensional data points generated from n normal distributions.

From these parameters it is easy to cluster the data as we know where data points from the different clusters are expected to lie.

From equation 6.1 we expect every Σ_i to be independent of each other. In the context of Gaussian Mixture Models this is called having a full covariance matrix. However, we can eliminate some of the variables in the covariance matrix if we choose a diagonal covariance

6. Clustering Algorithm and Application

matrix. Additionally, we might specify to use the same covariance matrix for all i , which is called tied in this context.

Choosing a full covariance matrix is not necessary even if the data is expected to have statistically independent features, as the overall density is compromised from multiple normal distributions with diagonal Σ_i . This enables us to model correlations between features.

The question now is how we can derive the parameters w_i, μ_i and Σ_i . We choose the approach which chooses the parameters where the likelihood that the data was generated by these parameters is maximal. This is known as maximum likelihood estimation. The likelihood can be expressed as

$$L(w_i, \mu_i, \Sigma_i | X) = p(X | w_i, \mu_i, \Sigma_i) = \prod_{t=1}^n p(x_t | w_i, \mu_i, \Sigma_i)$$

with $X = (x_1, \dots, x_n)$ being the recorded data. As $L(w_i, \mu_i, \Sigma_i | X)$ is non-linear in the parameters deriving the maximum is not trivial. Instead, we use an iterative approach which approaches the solution. Define $\lambda = (w_i, \mu_i, \Sigma_i)$. Simplifying to a diagonal covariance matrix gives us the iterative algorithm where we define the successor values $\bar{\cdot}$ as

$$\begin{aligned} Pr(i|x_t, \lambda) &:= \frac{w_i g(x_t | \mu_i, \Sigma_i)}{\sum_{k=1}^n w_k g(x_t | \mu_k, \Sigma_k)} \\ \bar{w}_i &:= \frac{1}{n} \sum_{t=1}^n Pr(i|x_t, \lambda) \\ \bar{\mu}_i &:= \frac{\sum_{t=1}^n Pr(i|x_t, \lambda)x_t}{\sum_{t=1}^n Pr(i|x_t, \lambda)} \\ \bar{\sigma}_i^2 &:= \frac{\sum_{t=1}^n Pr(i|x_t, \lambda)x_t^2}{\sum_{t=1}^n Pr(i|x_t, \lambda)} - \bar{\mu}_i^2. \end{aligned}$$

for w_i , μ_i and σ_i^2 respectively. One can show that with this iteration rule we have $p(X|\bar{\lambda}) \geq p(X|\lambda)$. The value $Pr(i|x_t, \lambda)$ is known as the a posteriori probability for the i -th component.

6.2. Implementation of the Clustering Algorithm

Python offers an implementation of Gaussian Mixture Model with the `sklearn` package. The function with parameters relevant to us is `sklearn.mixture.GaussianMixture(n_components, covariance_type)`. The number of components `n_components` can be any positive integer. The `covariance_type` can be one of ‘full’, ‘tied’, ‘diag’ or ‘spherical’ and describes what type of covariance matrix is used.

As the input of the Gaussian Mixture Model we use data points of the form `[a, u, d, k1, k2, w1, w2]`, where `a`, `u`, `...`, `w2` are the parameters of the approximation from chapter 5. As our goal is to separate data points from the four data sets mouse cells and

human cells each with a negative and a positive control, we use all particles as input. The pseudo code below describes the steps performed to reach a clustering of the data.

Algorithm 6: Separate

```

input : parameters of the approximations of all particles in all data sets
output: assignments to different clusters for each particle, means and standard
         deviation of each cluster

1 begin
2   initialize Gaussian Mixture by specifying n_components and covariance_type
3   apply Gaussian Mixture to matrix of all parameters of the approximations of
      all particle data sets
4   assign particles to clusters according to Gaussian Mixture results
5   compare assignments from Gaussian Mixture to those of the data set the data
      stems from
6   return assignments to clusters, means and standard deviations of every
      cluster
7 end

```

When comparing different covariance types in the Gaussian Mixture we see that using ‘diag’ we have the lowest error rate. The details are shown in table ???. Why reducing the number of parameters in the covariance matrix can yield better results is described in section 6.1.

full: 13.23%	tied: 12.7%
diag: 7.17%	spherical: 31.52%

Table 6.1.: Error as a percentage of particles being assigned the wrong component.

Using a diagonal covariance matrix we can now try to separate the four data sets and visualize the results. As the data is 7 dimensional we show lower dimensional representations of the data both as it is assigned according to the data set it stems from as well as the assignment from algorithm 6. The results can be seen in figure ??.

By comparing which data points are from which data set and where they were assigned by the Gaussian Mixture we can find an assignment between the two.

6. Clustering Algorithm and Application

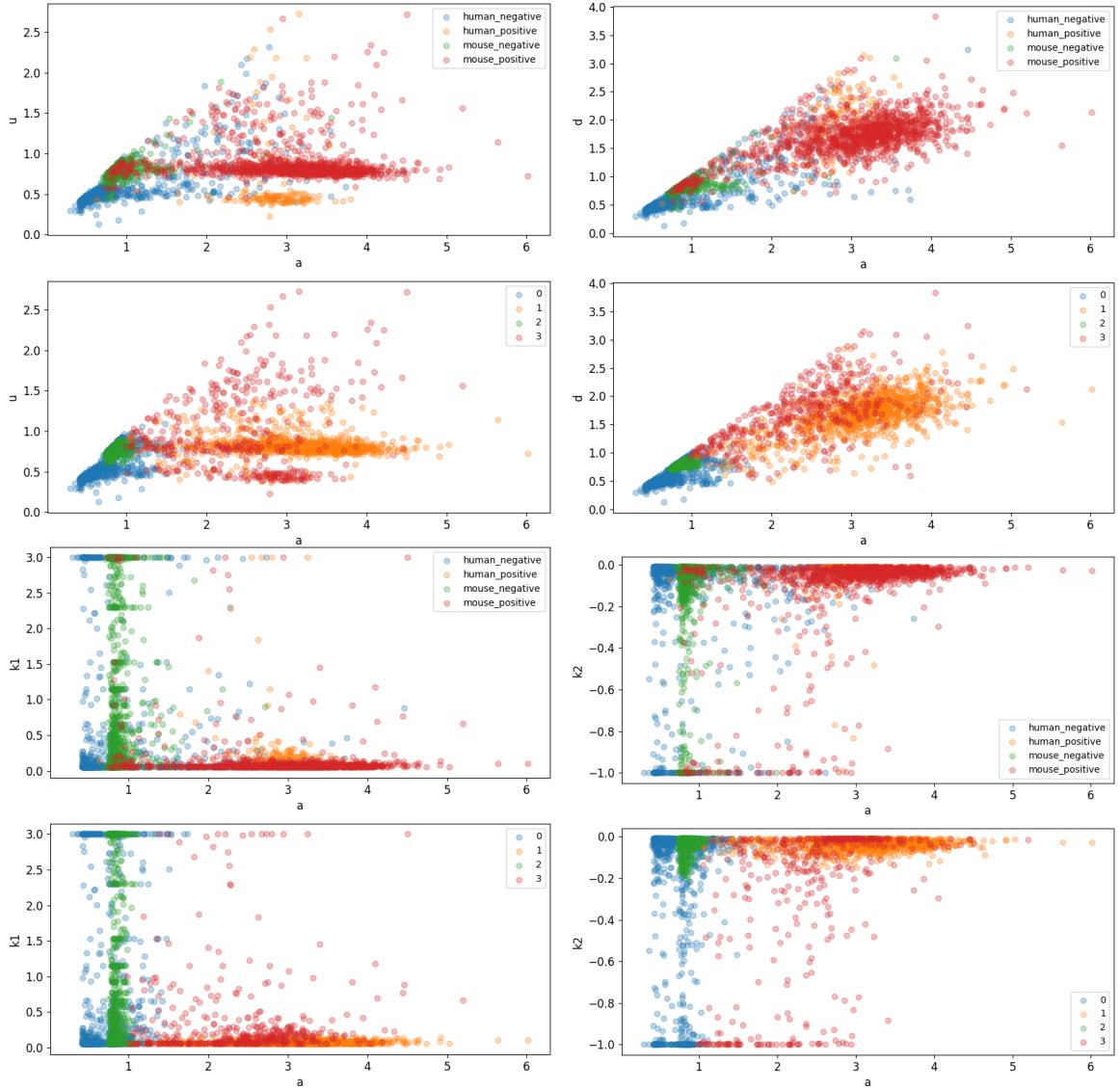


Figure 6.1.: Visualization of the clustered data points. Each pair of images depicts the clustering according to positive and negative control on either humans or mouse t cells in the upper image and the clustering according to the algorithm proposed in the lower image.

6.2. Implementation of the Clustering Algorithm

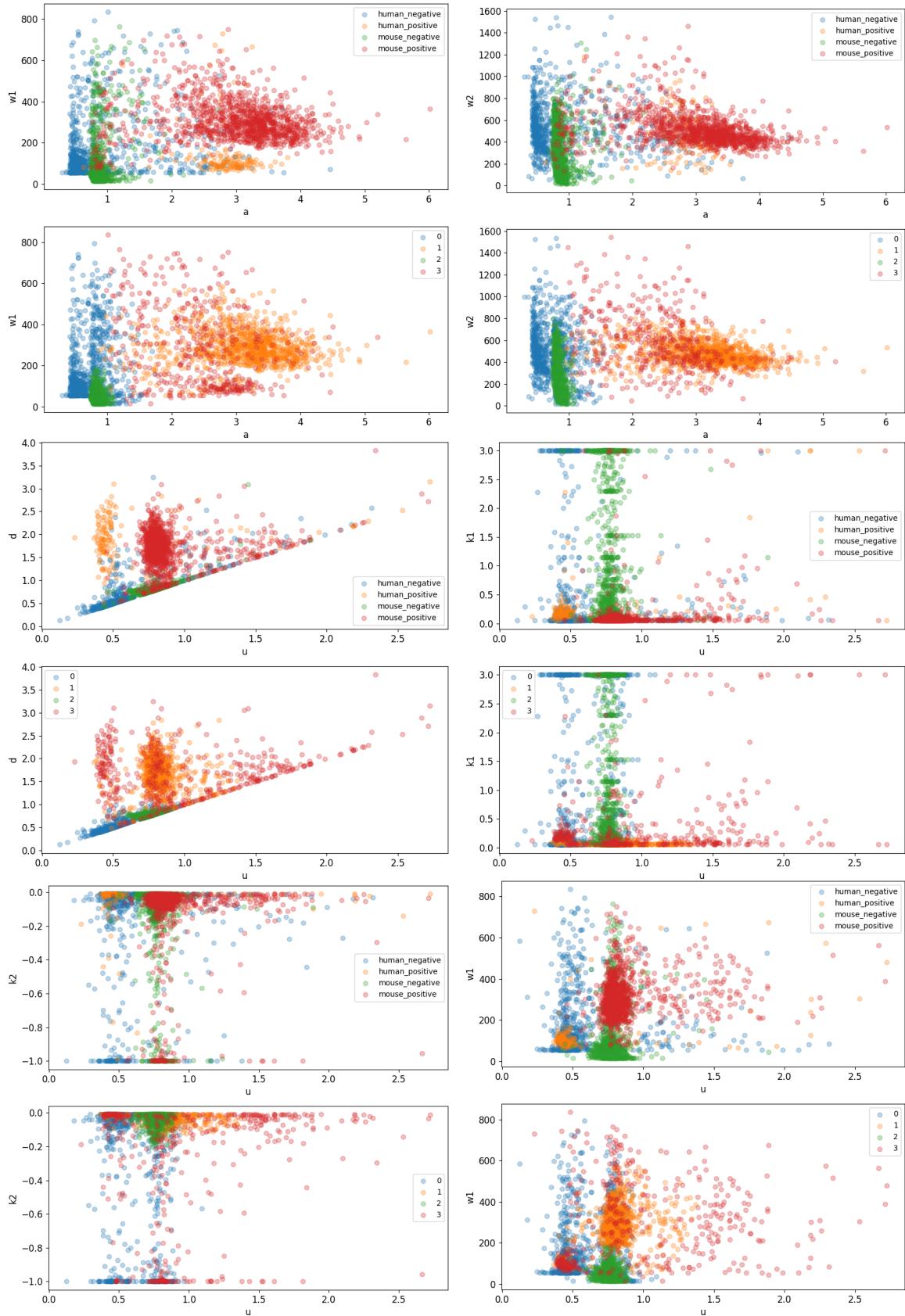


Figure 6.2.: Continuation of the visualization of the clustering results.

6. Clustering Algorithm and Application

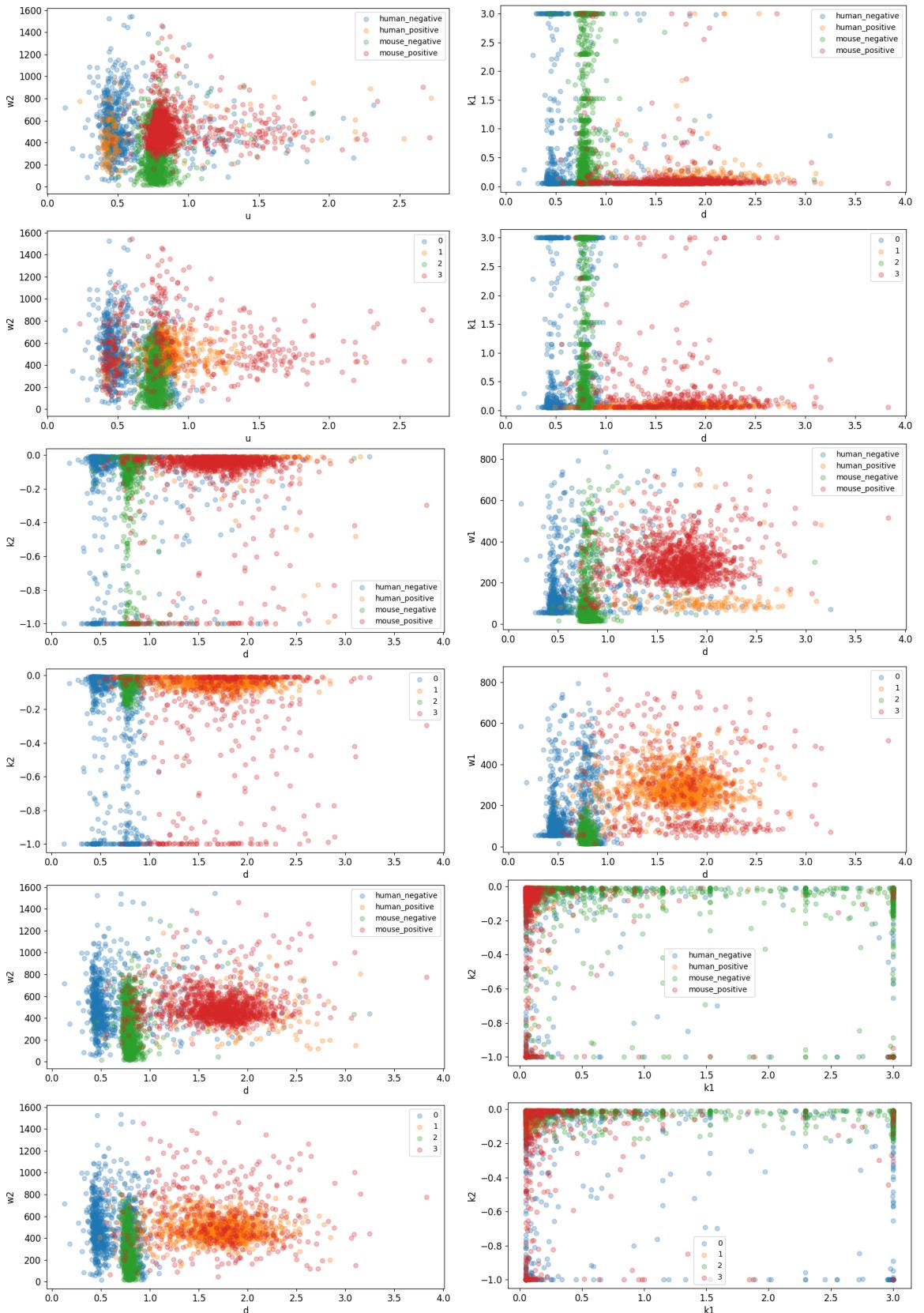


Figure 6.3.: Continuation of the visualization of the clustering results.

6.2. Implementation of the Clustering Algorithm

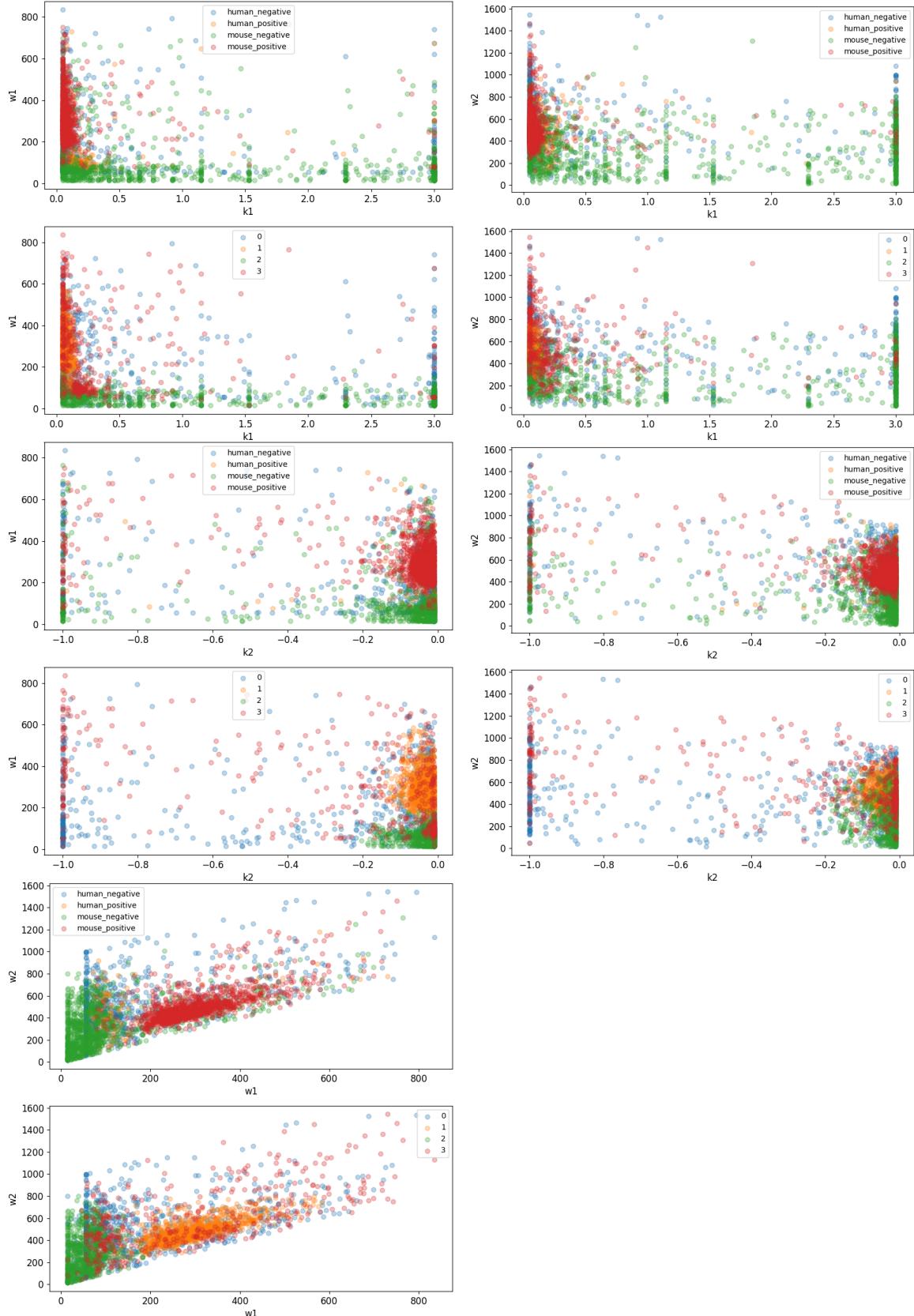


Figure 6.4.: Continuation of the visualization of the clustering results.

The relevant information derived from the Gaussian Mixture clustering is the means and covariances of the four components. From this we can decide which cluster a new data point belongs to. A use case might be to find percentages of activated cells in an experiment. Distinguishing between mouse and human cells does not have a clear use case. When specifying `n_components=2` we assume to have a cluster for activated and a second for unactivated cells.

Using the order of parameters a, u, d, k_1, k_2, w_1 and w_2 we get the mean and covariances for each of the data sets used in this work as

$$\begin{array}{lll}
\text{human positive} & w = 0.148 & \mu = \begin{pmatrix} 2.429 \\ 0.987 \\ 1.703 \\ 0.363 \\ -0.275 \\ 292.798 \\ 587.26 \end{pmatrix} \quad \Sigma = \text{diag} \begin{pmatrix} 0.541 \\ 0.245 \\ 0.276 \\ 0.395 \\ 0.131 \\ 37561.179 \\ 74216.954 \end{pmatrix}, \\
\text{human negative} & w = 0.278 & \mu = \begin{pmatrix} 0.735 \\ 0.586 \\ 0.612 \\ 0.83 \\ -0.267 \\ 179.044 \\ 491.68 \end{pmatrix} \quad \Sigma = \text{diag} \begin{pmatrix} 0.061 \\ 0.029 \\ 0.032 \\ 1.347 \\ 0.144 \\ 24314.336 \\ 59213.616 \end{pmatrix}, \\
\text{mouse positive} & w = 0.346 & \mu = \begin{pmatrix} 3.111 \\ 0.818 \\ 1.676 \\ 0.067 \\ -0.038 \\ 287.576 \\ 480.405 \end{pmatrix} \quad \Sigma = \text{diag} \begin{pmatrix} 0.478 \\ 0.016 \\ 0.128 \\ 0.0005 \\ 0.0006 \\ 7147.489 \\ 10182.191 \end{pmatrix}, \\
\text{mouse negative} & w = 0.228 & \mu = \begin{pmatrix} 0.842 \\ 0.762 \\ 0.78 \\ 1.335 \\ -0.037 \\ 56.052 \\ 285.085 \end{pmatrix} \quad \Sigma = \text{diag} \begin{pmatrix} 0.004 \\ 0.002 \\ 0.001 \\ 1.513 \\ 0.002 \\ 1149.598 \\ 29471.53 \end{pmatrix}.
\end{array}$$

When comparing these results to those of the average and standard deviation analysis done in table 5.3 we can see similarities. However in comparison, to the approach focusing on outlier detection in section 5.3 we now have an approach that is not only not dependent on parameters specified by a user, but can also be applied to a greater set of problems. A proposed way of answering the research questions from chapter 1 using these methods is described in chapter 7.

7. Results

7.1. Proposed algorithm for Detecting Activated T Cells

Question: Which criteria can distinguish between unactivated, activated and pre-activated cells?

Answer for activated and unactivated cells:

1. get positive control, negative control and experiment recordings
2. get parameters of each particle in all data sets using approximation described above
3. use outlier detection to filter out non-conforming cells from both the positive and negative control
4. use Gaussian Mixture Model with input parameters of filtered negative and positive control to get means and covariances of two clusters
5. predict the zugehörigkeit of the experiment particle parameters to the clusters to get a prediction of activation

For pre-activated cell detection:

Filter out those with too high u or to small w_1 using the outlier detection.

7.2. Types of Activated Cells

Question: Do different types of activated cells exist? How are they different?

Answer: Apply Gaussian Mixture Clustering to activated cells only. (Separate human and mouse cells, otherwise two clusters of eben das)

7.3. Oscillation in Decrease

Question: With which frequencies does the Calcium concentration repeat after activation?

Answer: Results from frequency analysis

7.4. Difference between Mouse and Human Cells

Question: Is there a difference in frequencies between mouse and human cells?

Answer: Compare mean and covariance between the two.

8. Discussion

can we give a value for accuracy of the proposed algorithms?

Discuss limitations

8.1. Outlook

better frequency analysis (look into what causes these oscillations)

apply discussed algorithm to enough data to give good approximations of average and standard deviation to hold for arbitrary data sets

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Bibliography

- [AH24] K Maude Ashby and Kristin A Hogquist. “A guide to thymic selection of T cells”. In: *Nature Reviews Immunology* 24.2 (2024), pp. 103–117.
- [AW04] Robert T Abraham and Arthur Weiss. “Jurkat T cells and development of the T-cell receptor signalling paradigm”. In: *Nature reviews immunology* 4.4 (2004), pp. 301–308.
- [BCL99] Mary Ann Branch, Thomas F Coleman, and Yuying Li. “A subspace, interior, and conjugate gradient method for large-scale bound-constrained minimization problems”. In: *SIAM Journal on Scientific Computing* 21.1 (1999), pp. 1–23.
- [BV18] Stephen Boyd and Lieven Vandenberghe. “Levenberg–Marquardt algorithm”. eng. In: *Introduction to Applied Linear Algebra*. Cambridge University Press, 2018, pp. 391–399. ISBN: 781316518960.
- [CA22] Geoffrey M Cooper and Kenneth Adams. “The Nucleus”. eng. In: *The cell: a molecular approach*. 19. edition. Oxford University Press, 2022, pp. 336–364. ISBN: 9780197583722.
- [Gan12] William F. Ganong. “Overview of Cellular Physiology in Medical Physiology”. eng. In: *Review of medical physiology*. 24. edition. Stamford, Conn: McGraw-Hill, 2012, pp. 35–66. ISBN: 9780071780032.
- [Gan97] William F. Ganong. “Circulating Body Fluids”. eng. In: *Review of medical physiology*. 18. ed. Stamford, Conn: Appleton & Lange, 1997, pp. 486–488. ISBN: 9780838584439.
- [Hag18] Kimberly Hagel. *Positive and Negative Selection of T Cells*. 2018. URL: <https://immunobites.com/2018/08/20/positive-and-negative-selection-of-t-cells/> (visited on 06/21/2024).
- [JRB14] Noah Joseph, Barak Reicher, and Mira Barda-Saad. “The calcium feedback loop and T cell activation: how cytoskeleton networks control intracellular calcium flux”. In: *Biochimica et Biophysica Acta (BBA)-Biomembranes* 1838.2 (2014), pp. 557–568.
- [KCF18] Brahma V Kumar, Thomas J Connors, and Donna L Farber. “Human T cell development, localization, and function throughout life”. In: *Immunity* 48.2 (2018), pp. 202–213.
- [Lew01] Richard S Lewis. “Calcium Signaling Mechanisms in T Lymphocytes”. In: *Annual Review of Immunology* 19. Volume 19, 2001 (2001), pp. 497–521. ISSN: 1545-3278. DOI: <https://doi.org/10.1146/annurev.immunol.19.1.497>. URL: <https://www.annualreviews.org/content/journals/10.1146/annurev.immunol.19.1.497>.

Bibliography

- [ML23] Jonathan Morgan and Alan E Lindsay. “Modulation of antigen discrimination by duration of immune contacts in a kinetic proofreading model of T cell activation with extreme statistics”. In: *PLOS Computational Biology* 19.8 (2023), e1011216.
- [MMS17] Magdiel Martínez, Namyr A Martínez, and Walter I Silva. “Measurement of the intracellular calcium concentration with Fura-2 AM using a fluorescence plate reader”. In: *Bio-protocol* 7.14 (2017), e2411–e2411.
- [NW99] Jorge Nocedal and Stephen J. Wright. “The Dogleg Method”. eng. In: *Numerical Optimization*. Springer, 1999, pp. 73–76.
- [Rey+09] Douglas A Reynolds et al. “Gaussian mixture models.” In: *Encyclopedia of biometrics* 741.659-663 (2009).
- [RLH90] MW Roe, JJ Lemasters, and B Herman. “Assessment of Fura-2 for measurements of cytosolic free calcium”. In: *Cell calcium* 11.2-3 (1990), pp. 63–73.
- [Rog24] Kara Rogers. *endoplasmic reticulum*. 2024. URL: <https://www.britannica.com/science/endoplasmic-reticulum> (visited on 06/23/2024).
- [SB16] Dianne S. Schwarz and Michael D. Blower. “The endoplasmic reticulum: structure, function and response to cellular signaling”. In: *Cellular and Molecular Life Sciences* 73 (2016), pp. 79–94. DOI: <https://doi.org/10.1007/s0018-015-2052-6>. URL: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4700099/>.
- [SI13] Peter B Stathopoulos and Mitsuhiro Ikura. “Structural aspects of calcium-release activated calcium channel function”. In: *Channels* 7.5 (2013). PMID: 24213636, pp. 344–353. DOI: [10.4161/chan.26734](https://doi.org/10.4161/chan.26734). eprint: <https://doi.org/10.4161/chan.26734>. URL: <https://doi.org/10.4161/chan.26734>.
- [SKJ09] Jennifer E Smith-Garvin, Gary A Koretzky, and Martha S Jordan. “T cell activation”. In: *Annual review of immunology* 27 (2009), pp. 591–619.
- [SSB77] Ulrich Schneider, Hans-Ulrich Schwenk, and Georg Bornkamm. “Characterization of EBV-genome negative “null” and “T” cell lines derived from children with acute lymphoblastic leukemia and leukemic transformed non-Hodgkin lymphoma”. In: *International journal of cancer* 19.5 (1977), pp. 621–626.
- [VL04] C Voglis and IE Lagaris. “A rectangular trust region dogleg approach for unconstrained and bound constrained nonlinear optimization”. In: *WSEAS International Conference on Applied Mathematics*. Vol. 7. 2004.

A. Python Implementation

We give a minimal version of the algorithm and code described in this work in Python.

```
1 import math
2 import scipy
3 import numpy as np
4 import pandas
5 from sklearn.mixture import GaussianMixture
6 from sklearn.cluster import KMeans
7 from sklearn.preprocessing import StandardScaler
8 import pandas as pd
9 import random
10
11
12 def approximate(dataframe):
13     def calc_t(w, k, alpha = 0.99):
14         tmp = 1 / alpha - 1
15         if tmp < 0.0001:
16             return None
17         try:
18             return w - math.log(tmp) / k
19         except ValueError:
20             return None
21
22     def approx_func_(x, w1, t, w2, a, d, u, k1, k2):
23         if t is None: # transition point lies outside datapoints
24             return u
25         elif x <= t: # logistic function before transition point
26             tmp = -k1 * (x - w1)
27             if tmp <= 32:
28                 res = (a - u) / (1 + math.exp(tmp))
29             else:
30                 res = 0
31             return res + u
32         else: # logistic function after transition point
33             tmp = -k2 * (x - w2)
34             if tmp <= 32:
35                 res = (a - d) / (1 + math.exp(tmp))
36             else:
37                 res = 0
38             return res + d
39
40     def approx_func(x_arr, w1_start, w2_w1, a_d, d_u, u, k1, k2):
41         w1 = w1_start + start
42         w2 = w2_w1 + w1
43         d = d_u + u
44         a = a_d + d
45         transition_point = calc_t(w1, k1)
```

A. Python Implementation

```
46     return (np.vectorize(approx_func_)
47             (x_arr, w1, transition_point, w2, a, d, u, k1, k2))
48
49 min_val = np.min(dataframe['ratio'])
50 median_val = np.median(dataframe['ratio'])
51 max_val = np.max(dataframe['ratio'])
52 start, end = min(dataframe['frame']), max(dataframe['frame'])
53
54 lower_bounds = (0, 0, 0, 0, min_val, 0.05, -1)
55 upper_bounds = (end-start, end-start, max_val, max_val,
56                  max_val, 3, -0.01)
57 p0 = (0, (end-start)/2, max_val-median_val, median_val-min_val,
58       min_val, 0.1, -0.03)
59
60 popt, *_ = scipy.optimize.curve_fit(approx_func, dataframe['frame'],
61                                     dataframe['ratio'], p0=p0,
62                                     method='trf',
63                                     bounds=(lower_bounds, upper_bounds))
64
65 w1_start, w2_w1, a_d, d_u, u, k1, k2 = popt
66 w1 = w1_start + start
67 w2 = w2_w1 + w1
68 d = d_u + u
69 a = a_d + d
70 t = calc_t(w1, k1)
71 return {"start": start, "end": end, 'w1': w1, 't': t, 'w2': w2,
72         'e': end, 'a': a, 'd': d, 'u': u, 'k1': k1, 'k2': k2}
73
74
75 def approximation_loop(file_name):
76
77     # read data
78     data = pandas.DataFrame(pd.read_hdf(f"../data/{file_name}"))
79
80     # filter data
81     data = data[np.isfinite(data["ratio"])]
82     data = data[np.less(data["ratio"], np.full((len(data["ratio"])), 5))]
83     data = data[np.greater(data["ratio"], np.full((len(data["ratio"])), 0))]
84
85     all_parameters = list()
86     parameters_saved = ["idx", "start", "end", 'w1', 't', 'w2',
87                          'a', 'd', 'u', 'k1', 'k2']
88
89     for particle_idx in set(data['particle']):
90         # get data of a single particle
91         single_particle_data = (
92             data.loc[data['particle'] == particle_idx][['frame', 'ratio']])
93
94         # skip if too few datapoints
95         if len(single_particle_data['frame']) < 300:
96             continue
97
98         try: # throws error if no best fit was found
99             parameters = approximate(single_particle_data)
100
```

```

101     parameters["idx"] = str(particle_idx) + file_name
102     all_parameters.append([parameters[e] for e in parameters_saved])
103
104     except Exception as e:
105         print(f"error in particle {particle_idx}: {e}")
106
107     return all_parameters
108
109
110 def separate(neg_par, pos_par, CLUSTERING_METHOD):
111     prediction_parameters = ["a", "u", "d", "k1", "k2", "w1", "w2"]
112
113     df_neg = pd.DataFrame(data=neg_par,
114                           columns=["idx", "start", "end", 'w1', 't', 'w2',
115                                     'a', 'd', 'u', 'k1', 'k2'])
116     df_pos = pd.DataFrame(data=pos_par,
117                           columns=["idx", "start", "end", 'w1', 't', 'w2',
118                                     'a', 'd', 'u', 'k1', 'k2'])
119     df_neg["activation"] = "negative"
120     df_pos["activation"] = "positive"
121     data = pd.concat([df_neg, df_pos])
122
123     # clustering
124     if CLUSTERING_METHOD == "gaussian_mixture":
125         clustering = GaussianMixture(n_components=2, covariance_type="diag",
126                                       n_init=10)
127     elif CLUSTERING_METHOD == "kmeans":
128         clustering = KMeans(n_clusters=2, n_init=10)
129     else:
130         raise RuntimeError(f"Value of CLUSTERING_METHOD set to "
131                             f"{CLUSTERING_METHOD}, which is not "
132                             f"one of [gaussian_mixture, kmeans].")
133     data["predicted_clusters"] = clustering.fit_predict(
134         data[prediction_parameters])
135
136     # find association between predicted clusters and files
137     neg_0 = len(data[(data['predicted_clusters'] == 0) &
138                       (data['activation'] == "negative")])
139     neg_1 = len(data[(data['predicted_clusters'] == 1) &
140                       (data['activation'] == "negative")])
141     pos_0 = len(data[(data['predicted_clusters'] == 0) &
142                       (data['activation'] == "positive")])
143     pos_1 = len(data[(data['predicted_clusters'] == 1) &
144                       (data['activation'] == "positive")])
145
146     permutation = (0, 1) if neg_0 + pos_1 > neg_1 + pos_0 else (1, 0)
147
148     return permutation, clustering
149
150
151 if __name__ == "__main__":
152     FILE_NAME_NEG_CONTROL = "human_negative/human_negative.h5"
153     FILE_NAME_POS_CONTROL = "human_positive/human_positive.h5"
154     FILE_NAME_EXPERIMENT = "human_positive/human_positive.h5"
155

```

A. Python Implementation

```
156     neg_con_par = approximation_loop(FILE_NAME_NEG_CONTROL)
157     pos_con_par = approximation_loop(FILE_NAME_POS_CONTROL)
158     experiment_par = pos_con_par # approximation_loop(FILE_NAME_EXPERIMENT)
159
160     # center all paramters
161     neg = pandas.DataFrame(neg_con_par, columns=["idx", "start", "end",
162                                         'w1', 't', 'w2', 'a',
163                                         'd', 'u', 'k1', 'k2'])
164     pos = pandas.DataFrame(pos_con_par, columns=["idx", "start", "end",
165                                         'w1', 't', 'w2', 'a',
166                                         'd', 'u', 'k1', 'k2'])
167     exp = pandas.DataFrame(experiment_par, columns=["idx", "start", "end",
168                                         'w1', 't', 'w2', 'a',
169                                         'd', 'u', 'k1', 'k2'])
170
171     all_data = pd.concat([neg, pos, exp])
172
173     scaler = StandardScaler()
174     all_data[["a", "u", "d", "k1", "k2", "w1", "w2"]] = (
175         scaler.fit_transform(all_data[["a", "u", "d", "k1", "k2", "w1", "w2"]]))
176
177     neg = all_data[all_data["idx"].isin(neg["idx"])].values.tolist()
178     pos = all_data[all_data["idx"].isin(pos["idx"])].values.tolist()
179     exp = all_data[all_data["idx"].isin(exp["idx"])].values.tolist()
180
181     # filter out outliers
182
183     n = min(len(neg_con_par), len(pos_con_par))
184     per, clustering = separate(random.sample(neg_con_par, n),
185                                random.sample(pos_con_par, n), "kmeans")
186
187     df_exp = pd.DataFrame(data=experiment_par,
188                           columns=["idx", "start", "end", 'w1', 't', 'w2',
189                                     'a', 'd', 'u', 'k1', 'k2'])
190     df_exp["predicted_clusters"] = clustering.predict(df_exp[["a", "u", "d",
191                                                               "k1", "k2",
192                                                               "w1", "w2"]])
193
194     print(f"positive: {len(df_exp[df_exp['predicted_clusters']] == per[1])}
195 / {len(df_exp)}")
```