

Project plan

Project name	Client / Sponsor	Project manager
Assisted Function Exoskeleton (AFE)	Mälardalens University	Mikael Ekström & Pontus Sundkvist

1 Executive Summary

The Assisted Function Exoskeleton (AFE) project aims to create a prototype of a practical upper-body exoskeleton designed to aid individuals in diverse sectors, including agriculture (e.g., fruit and berry picking), industrial applications (e.g., assisting in heavy lifting), and healthcare technology (e.g., providing support during surgeries or heavy lifting tasks and decrease muscle fatigue). This effort will involve comprehensive background research and embody the latest advancements in exoskeleton technology to gain an understanding of the concept. This knowledge will guide the project's development process, ensuring agreement with all relevant regulations and safety standards during the exoskeleton's construction. The project encompasses the following generalized tasks: The exoskeleton design and structure, determining the materials used in the construction of the exoskeleton to ensure reliability, functionality, and durability. The project will incorporate electronics to control the exoskeleton, utilizing Electromyography (EMG) sensors, force sensors, temperature sensors and motors as the actuators of the device. Enabling a logical fusion of information through data processing and developing a safe system that will control the torque on the exoskeleton motors and assist the user.

In an effort to enhance project completion and organization, specific roles and responsibilities have been assigned to individuals. The Exoskeleton project team includes Albin Gustafsson as the Hardware Team Leader, with Sebastian Ahlström and Moritz Schmidt serving as Hardware Developers. Furthermore, Irini Provatidis has taken on the role of Software Team Leader, while Jalal Taleb is responsible for software development. During this time, a collaboration with students from the Technological University of Panama is held, to gain additional ideas.

The hardware team is responsible for integrating electronics and mechanical components into the Exoskeleton. They are tasked with designing the Exoskeleton, selecting suitable materials and components to ensure its robustness and safety, and ensuring compliance with the project manager's requirements.

The software team handles data processing, analysis, and the development of the Exoskeleton's control system. Their responsibilities include accurately acquiring sensor data, filtering and detecting motion patterns using Artificial Intelligence and Machine Learning. They focus on designing a stable control system that not only collects critical feedback but also effectively estimates system errors to provide precise force commands to the Exoskeleton's motors.

2 Background

Human-robot collaboration systems have a wide range of applications, and currently they are entering the daily lives of humans. The development of sensor technology is establishing an interaction between humans and robots, which makes it possible to communicate, predict, and understand the current state of each partner of the system in a shared environment. Human-robot research is applied in diverse areas such as entertainment and education, robot-assisted surgery, intelligent vehicles and aircraft, assisted and rehabilitation technology, etc. Some of their basic aims are to enhance the quality of tasks, improve productivity, reduce the workload of humans, and help with the rehabilitation process after trauma.

To facilitate communication between humans and robots, predict intentions, categorize data, and develop control systems using sensor data input, researchers have used various signals that are non-biological and/or biological over the years. Biological signals are electrical signals that travel between our brain, skin, organs, glands, and muscles and are generated by the nervous system. The human body generates various biological signals: Electrooculogram (EOG), Electrocorticogram (ECoG), Electroencephalogram (EEG), Magnetoencephalography (MEG) and EMG. On the other hand, examples of non-biological signals are kinematic parameters such as force/torque sensor data, velocity, or signals

generated from temperature sensors, etc[1].

Exoskeleton control systems are applications reflecting assistive robotic technologies. The research and state of the art of this project are limited to the assistive technology of exoskeletons. In general, exoskeletons are categorized as upper-limb exoskeletons, lower-limb exoskeletons, or full-body exoskeletons. The majority of designed and developed exoskeletons are still in the experimental stage of clinical testing, and more research and effort are essential to bring them out of the laboratory.

Controlling any exoskeleton requires sophisticated technologies and methods. The main requirements of accuracy, long-term reliability, and safety are vital for the control system of exoskeletons. EMG is frequently used in the controlling methods of exoskeletons and prosthetics since it reflects the motion intention or muscle activity of the user. Furthermore, sensory data from temperature and force/torque sensors is also utilized as feedback control information in EMG-based control systems for exoskeleton applications. Even though EMG is one of the most efficient signals to utilize in the control method of an exoskeleton, muscle fatigue introduces variations in the EMG amplitude and frequency, which influence the input data to the control system. Senarath et al. researched hybrid EMG-EEG-based control approaches aimed at exoskeleton applications. Their paper includes a review of EMG-based methods proposed for assistive robot control. One of the aspects they aim to explore is related to muscle fatigue in EMG-based control systems. Their proposed method included training multiple fuzzy-neuro modifiers to adapt to the muscle fatigue conditions to compensate for the effects of muscle fatigue on EMG-based control, and the effectiveness of the results was experimentally validated. The results of their experiments indicated that an EMG amplitude feature such as EMG Root Mean Square (RMS) alone is not adequate as an input signal for an effective EMG-based control during muscle fatigue conditions. It is essential to use frequency domain EMG features as additional input features to identify muscle fatigue conditions and use them in EMG-based control systems[2].

The broad range and diversity of approaches available for the development and control of robots using surface Electromyography (sEMG) poses a significant challenge for researchers, prompting them to explore the optimal ways to design such systems. Song et al. present and provide a comprehensive overview of techniques and methods for controlling robots using sEMG. The article provides an overview of sEMG-based robot control concluding two important aspects:

1. sEMG signal processing and classification methods.
2. Robot control strategies and methods based on sEMG.

To detect limb movement or to control the system of an exoskeleton, raw sEMG signals cannot be used directly. This is to obtain a higher signal-to-noise ratio (SNR). The signals need to be processed through correct data acquisition, pre-processing, feature extraction, dimensionality reduction (if necessary), and pattern recognition. The selection of feature sets extracted by the processed signal is an important factor for the accuracy of the classifiers, which is the core of pattern recognition. They present a variety of sources related to sEMG and data processing methods as well as a statistical analysis to know the level of success of each attempt depending on the purpose and goal of the project. Numerous approaches have been explored and developed over the years. The filtering of raw EMG signals remains relatively consistent across many applications. However, when it comes to extracting features from EMG signals and implementing Artificial Intelligence (AI) algorithms, a wide array of approaches has emerged, each offering distinct solutions. Extracted features can be represented in the time domain (TD), frequency domain (FD), and time-frequency domain (TFD) of the EMG signal. The choice of features is heavily dependent on the specific application's objectives, and the same holds true for selecting the appropriate AI algorithm. Researchers have explored and used diverse approaches to classify data and achieve accurate predictions[3].

Most research on exoskeletons primarily focuses on rehabilitation purposes, aiming to support individuals with conditions such as spinal cord injuries, strokes, or other neurological issues. The control system of an exoskeleton application is one of the most important aspects of its development since it must be designed with respect to the needs of the patient. Delgado et al. have achieved a significant milestone by developing a simulated model that demonstrates effective human-exoskeleton synergy, particularly for individuals with weakened muscle activation. Their proposed strategy centers around an adaptive Fuzzy Sliding Mode Control, designed to manage the inverse dynamics of a nonlinear 4-degrees-of-freedom (DOF) exoskeleton, allowing for the estimation of muscle activation and effort. The system utilizes signals from the biceps brachii muscle, and the error in exoskeleton position serves as input for a fuzzy inference system. This system generates an output to adjust the sliding mode control law parameters, thus providing the correct assistive motor force to the actuators. The simulation model has successfully equipped users with the necessary support they require[4].

3 Purpose

The AFE project aims to assist healthy individuals with high workloads, demonstrating its feasibility and serving as a foundation for future research at Mälardalens University (MDU). By investigating its potential, we hope to inspire future students and researchers to further develop this concept and maximize its impact.

4 Goal

The project's objective is to establish the foundation for designing and developing a prototype single-arm AFE. This involves comprehensive research, including a thorough review of the current state-of-the-art, to validate the project's relevance in today's context. The exoskeleton will harness electrical impulses generated by muscular activity, which will be measured by sEMG sensors. These electrical signals, produced through muscle conduction, will undergo processing to generate commands that will activate the appropriate motors. Further on, the development of a stable control system and the utilization of the commands in combination with feedback sensor data will provide the necessary assistive force to facilitate the lifting of objects.

5 Scope

The scope of the project provides a general overview of the project's composition, outlining the responsibilities and tasks of each team. Each of these sections contains sub-tasks related to specific aspects of the project

5.1 Work-Breakdown-Structure (WBS)

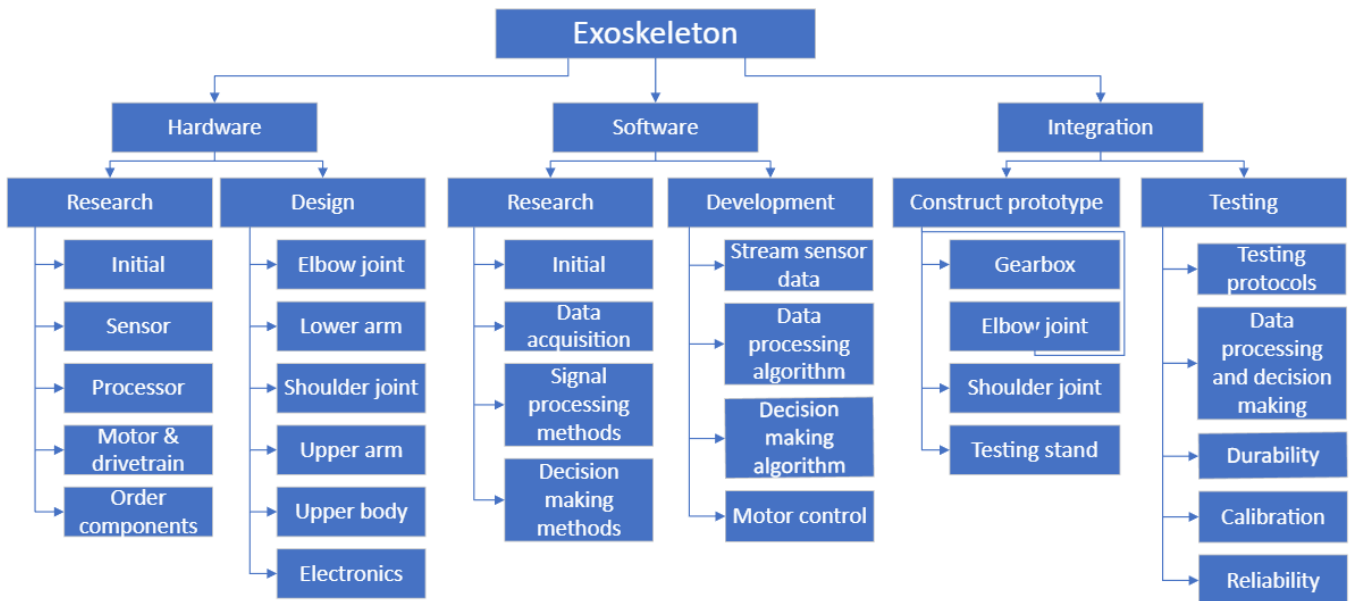


Figure 1: Work-Breakdown-Structure for each team. The basic responsibilities and tasks are illustrated.

6 Project Schedule

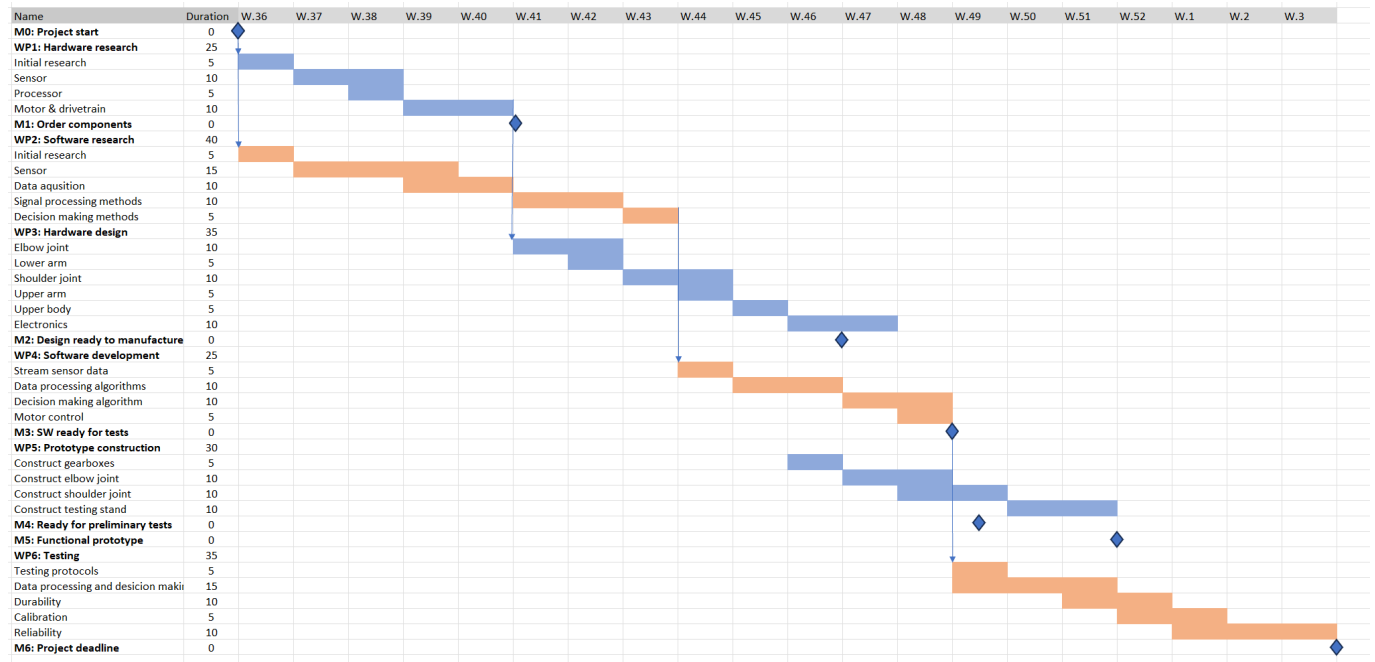


Figure 2: A GANTT chart showing the current project plan, including work packages and milestones that will be completed before the end of the project.

The project runs from the 4th of September 2023 until the 12th of January 2024, the project has several stages and assignments that need to be done in order to finish the project. The first stage is research and will last for 5 weeks, this period is used to find current research to serve as a basis to guide the design and development of the project. During this period two assignments also have to get done, this project plan is one and a preliminary poster to advertise the project is the second.

The next stage is the development stage, this is the stage where the design is made, and the software needed for the project is developed. It overlaps with the research stage and the construction stage and lasts for 8 weeks, during this time two progress reports are written as a reminder of what has been done and what needs to be done.

The third stage is construction, during this period the exoskeleton is being manufactured and at the end there will be a functional prototype. This stage overlaps not only with the development stage but also the testing stage and will last for 6 weeks. During this period the second progress report is being written and the final project poster will be made to advertise the project.

The final stage is testing, during this period of 3-4 weeks the prototype will be tested and tweaked to measure and improve the performance of the system. The last milestone to be done is the final project report which will be a summary of all of the work on the project and will contain the results and conclusions of the project.

7 Project and Product Requirements

This section provides a closer look at the project and product requirements. These requirements serve as the fundamental building blocks of successful project management and development, underpinning the entire project planning process, and ultimately guiding the project towards the successful attainment of its objectives.

7.1 Product requirements

The following section categorizes the given product requirements into two sections. That is non-functional and functional requirements. The non-functional requirements contain performance, scalability, security, usability and maintainability requirements, while functional contains input, output, data handling, processing, user interface and error handling requirements.

7.1.1 Non-functional requirements

1. The EMG sensors shall be Electromagnetic Compatibility (EMC) resistant.
2. The exoskeleton, and all the parts of it, shall share a common electrical ground.

3. Sensors shall be assigned a corresponding motor.
4. Sensors shall be assigned to a motor in such a way that they are not interchangeable.
5. A temperature sensor shall be placed on the battery pack.
6. Temperature sensors shall be present on each motor.
7. The Central Processing Unit (CPU) shall have a pre-coded emergency shutdown sequence.
8. The pre-coded shutdown of the medical device shall be executed in a controlled manner.
9. All maintenance-authorised investigators/engineers shall be provided with an individual "key" (access code) used to access the system.
10. All wireless communication shall be encrypted.
11. The medical device shall not be connected to any external wireless network (outside its own data processing network).
12. System code maintenance shall only be possible through physical connection to a system data port.
13. The system shall have a maximum of three (3) actuators per unit (arm).
14. The system shall have a maximum of three (3) motors per unit (arm).
15. The device shall shutdown in a controlled manner if the signal from the EMG sensors is zero or lost for a duration of more than 3 seconds.
16. The EMG sensors shall be fastened to the exoskeleton structure in such a way that they may only be placed correctly on the subject's body when wearing the medical device.
17. Structural parts of the system shall be capable of handling 40 kg without experiencing plastic or elastic deformation (breaking or deflecting).
18. The medical device shall weigh a total maximum of 20 kg.
19. The actuators shall be capable of handling a weight of 40 kg.
20. The subject shall be shielded from the battery pack.
21. The battery compartment/attachment point shall have vibration dampening rubber insulators.
22. The battery pack compartment shall be protected against inertia shock of 2000 J.
23. The system shall be able to operate normally between -20°C and 40°C.
24. The power supply shall consist of a Lithium ion battery pack.
25. The battery pack shall be a 28 Volt system
26. The structural parts of the medical device shall be of a non-conductive material (electricity) or completely electrically insulated.
27. The battery pack shall have a capacity equalling to four (4) hours of use at maximum power consumption.
28. The battery pack shall be modular.
29. The motors shall be axial motors.
30. The structural parts of the exoskeleton shall include a stop not allowing movement beyond the arm being straight.
31. The structure attaching the two modules shall not hinder the movement of the subject's shoulder blades.
32. The strap connecting from the back to the front shall not interfere with the female breast if present.
33. Power cable connectors shall be keyed in such a way that they only allow for correct connection between both cables and connectors.
34. Data cable connectors shall be keyed in such a way that they only allow for correct connection between both cables and connectors.
35. The structural parts of the exoskeleton shall be able to withstand a physical shock of 2000 Joule without compromising the structural integrity of the structural parts.

36. The medical device shall issue an auditorial warning when an controlled shutdown is initialised.
37. The medical device shall issue a visual warning when a controlled shutdown is initialised.
38. The exoskeleton device shall only operate if all motors of a device unit (arm) is functioning.
39. The battery compartment shall protect the subject from a explosion and/or fire caused by the battery.
40. The subject shall always use two modules (a pair) of the system (one on each arm).
41. The medical device shall be rated for 25 kg (15kg below max).
42. The structural parts of the medical device shall be replaced if the medical device has experienced a physical shock greater than 2000 Joule (e.g. drop from height).
43. The battery shall be charged separate from the medical device and the subject.
44. The battery pack shall be charged in a controlled environment.
45. The battery pack shall be handled by a qualified and authorised investigator during the charging process.
46. System alterations shall be performed by a trusted investigator/engineer.
47. If the subject has experienced injuries to the arm resulting in a bone fracture, sprain, dislocation or strain usage of the device shall be avoided.

7.1.2 Functional requirements

1. The device shall always have a non-zero signal from the EMG sensors for assistive force to be applied by the device.
2. The temperature sensor shall monitor the temperature of the battery pack.
3. The temperature sensors of the motors shall monitor the temperature of the motors.
4. The minimum range of the EMG sensors shall be 3-80 Hz.
5. The EMG sensors shall take five (5) measurements every second.
6. All system maintenance shall be logged automatically with date and time as well as location in the code.
7. The individual "key" (access code) used during the maintenance shall be logged.
8. System code back-ups shall be performed every 80 hours of use.
9. The CPU block shall have a maximum failure rate of 10^{-3} faults/hour.
10. The CPU block shall have a maximum error rate of 10^{-4} errors/hour.
11. Each actuator shall have a maximum failure rate of $6 * 10^{-6}$ faults/hour.
12. Each actuator shall have a maximum error rate of $6 * 10^{-4}$ errors/hour.
13. Each motor shall have a maximum failure rate of 10^{-3} faults/hour.
14. Each motor shall have a maximum error rate of 10^{-4} errors/hour.
15. Each EMG sensor block (one block per measuring point) shall have a maximum failure rate of 10^{-3} faults/hour.
16. Each EMG sensor block (one block per measuring point) shall have a maximum error rate of 10^{-4} errors/hour.
17. The force sensor shall measure the force output to the exoskeleton from the actuators.
18. The force sensor shall have a minimum resolution of 1-220N of force.
19. Each force sensor block (one block per actuator) shall have a maximum failure rate of $2 * 10^{-3}$ failures/hour.
20. Each force sensor block (one block per actuator) shall have a maximum error rate of $3 * 10^{-4}$ errors/hour.
21. Every component of the system shall be compatible with a 28 volt power input.
22. The power supply shall have a maximum failure rate of 10^{-3} failures/hour.
23. The power supply shall provide correct amount of power to the system with a maximum error rate of 10^{-4} errors/hour.

24. The power supply shall not experience any errors that may impact its integrity with a maximum error rate of 10^{-7} errors/hour.
25. The structural parts of the system shall have a maximum failure rate of $4 * 10^{-6}$ failures/hour.
26. The structural parts of the system shall have a maximum error rate of $4 * 10^{-4}$ errors/hour.
27. The wires of the system shall have a maximum error rate of $2 * 10^{-3}$ errors/hour.
28. The fuses of the system shall have a maximum error rate of $7 * 10^{-4}$
29. Motors shall be able to provide a total maximum force of 200N.
30. The attachment points shall be capable of handling 400N of force in any direction without detaching and/or breaking.
31. Any maintenance shall be logged with a description of the change/-s made to the system.

7.2 Project requirements

1. To develop the prototype AFE
2. To follow stated Product requirements.
3. To respect budget constrain.
4. To respect time constrain.
5. To establish a collaborative partnership with Technological University of Panama

8 Limitations

Since the project is very complex, it will need to be limited to make it reasonable to achieve within the given time and budget. The limits can be categorized as both product limitations and project limitations; the limitations state what we aim to achieve and what we will not achieve at the end of the project.

8.1 Product functional limitations

The exoskeleton will offer a limited range of motion, specifically designed for safely lifting and raising boxes in compliance with safety regulations. It is important to note that this exoskeleton is intended as a proof of concept, featuring a single arm constructed from cost-effective materials. As a result, the final product will not be a fully functional upper-body exoskeleton for commercial use. The primary objective of this project is to assist in fruit picking and box handling by enhancing stamina, rather than solely increasing raw strength. Furthermore, our research aims to boost work efficiency, with a focus on productivity improvement rather than rehabilitation.

8.2 Project limitations

The project operates with a limited budget, which impacts the procurement of all the necessary components for constructing the device. Developing a prototype product under these constraints may affect the quality and final outcome of the exoskeleton. Additionally, the project faces time constraints. The design of a functional device may be affected by delays in component arrivals or the non-availability of components in manufacturers' stock. The project requirements, as provided by the project manager to ensure the safety of the medical device, pose a limitation. This is because the required components may not meet a specific requirement on the list, even though they fulfill all other specified requirements. At the end of the project, a prototype will have been constructed; it will consist of one arm that will respond to muscle activity. The motors of the arm will provide a force proportional to the muscle activity of the operator. The project will not provide a complete product, only a limited prototype to prove the concept. It will not be mounted to a person but to a stand so as not to endanger the person operating it.

9 Risk Analysis and Counteractions

As part of our project management, we carefully analyze risks. This involves identifying potential risks, rating their probability on a scale of 1 to 5 (from very unlikely to very likely), assessing their impact on a scale of 1 to 5 (from minimal to critical), and then calculating a risk score by multiplying the probability and impact ratings.

The risk score thresholds are interpreted as follows:

- 1-5: Low priority risks. These are monitored but may not require immediate action.
- 6-10: Medium priority risks. These require a mitigation plan and should be addressed as resources allow.
- 11-15: High priority risks. These require a detailed mitigation plan, and resources should be allocated to address these risks immediately.

- 16-25: Critical risks. These must be addressed immediately with a detailed and efficient mitigation plan to avoid severe project disruption.

Table 1 illustrates recognized risks, their evaluation, and planned responses. This includes mitigation techniques and reactions to these risks, ensuring that we are ready for unforeseen circumstances and can handle them successfully if they do occur.

Risk	Probability(1 to 5)	Impact(1 to 5)	Risk(P*I)	Risk Response
Team member absent during development	2	5	10	Develop a good communication plan, cross-train team members, have each development part be distributed in smaller groups.
Missed project deadlines	2	3	6	Multiple status meetings every week, one at the beginning to assign tasks, and one at the end to discuss progress and if needed relocate resources. Also, have bi-weekly meetings with stakeholders
Components breaking	2	4	8	Order extra components, Always be thorough in datasheets and do multiple checks before powering a newly build system.
Software bugs detected during tests	5	3	15	Thorough testing process, allocate resources for bug fixes.
Unexpected additional costs	2	5	10	Prepare a flexible budget that can withstand additional costs.
Incompatible hardware	1	5	5	Make proper research before hand, finding sources and guides supporting the current idea.
Scope creep & Communication issues	2	3	6	Multiple meetings every week to discuss current situation and plan progress according to the requirements.
Time and budget constrains	4	3	12	Allocate resources according to the budget plan.
Inadequate Testing	2	5	10	Follow a thorough predefined test plan, allocated resources and conduct peer reviews.
Health and safety risks of testing entire system	4	5	20	Follow safety tips [5], regular inspections, educate staff in protection and awareness.
System communication failure between components	3	5	15	If communication fails during use of device, immediate shot down and research of new communication protocols.

Table 1: Risk Analysis

10 Development Plan

This Development Plan serves as an encompassing guide for the project, accurately outlining the development process, our chosen methodologies, and the unique roles and responsibilities of each team member. Its design prioritizes a smooth integration process for newcomers, regardless of their entry point in the project's lifecycle. By immersing yourself in this plan, you'll gain essential insights into our operational procedures and project-related information, enabling you to contribute effectively and align with the team's overarching goals.

10.1 Development Methodology

In this project, our team strictly complies to the Waterfall development method [6]. This method follows a linear approach, comprising distinct phases, including requirements, design, implementation, verification, and deployment. The fundamental principle of this methodology is that each phase must be completed in its entirety before progressing to the subsequent one.

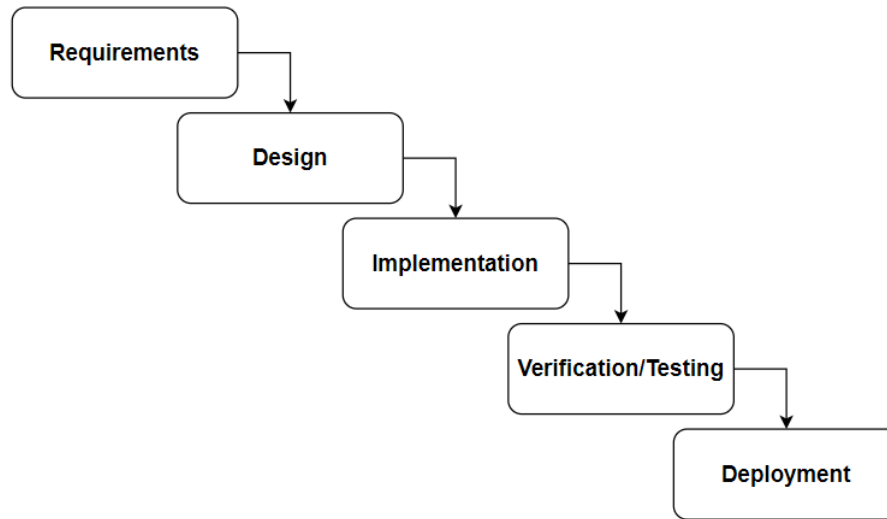


Figure 3: This image shows the Waterfall development method, which follows a step-by-step process where each phase must be completed before moving on to the next. For instance, you need to finish the requirements phase before starting the design phase. In simple terms, it is like water flowing down a set of stairs, taking one step at a time.

We chose this methodology due to its alignment with the initial project briefing, which provided a well-documented and clear list of requirements to be fulfilled. Furthermore, it suits our project exceptionally well, as this project demands thorough research and development. To provide a better understanding of this choice, we provide some of the advantages and disadvantages of this methodology below.

Advantages:

- Easy to manage and understand
- Clear objectives and requirements
- Greatly transfers information between sections

Disadvantages:

- Slow development
- Costly
- Hard to make changes
- Testing delayed to after completion

10.2 Team Structure and Roles

Our team comprises the following roles:

- **Hardware Team Leader:** Manages the hardware development process, coordinates with the software team, and ensures timely delivery while maintaining quality standards.
- **Software Team Leader:** Oversee the software development process, collaborate with the hardware team, and ensure the timely delivery of high-quality project milestones.
- **Hardware Developer:** Designs, develops, tests, and troubleshoots hardware components and collaborates with the software team.
- **Software Developer:** Develops, tests, and debugs software modules and coordinates with the hardware team.

10.3 Tools, Technologies, and Systems

We leverage the following tools, technologies, and systems in our project:

- **Project Management & Communication Tools:**

For successful communication and management, the platform Microsoft Teams 10.4.7 is used. This can be acquired from [7]. Additionally, the add on tool for teams Microsoft Planner 1.18.0 is also utilized. For general file management Github is used which can be obtained from [8].

- **Software Development Tools:**

The following software tools were applied for this development project, Arduino IDE 2.2.1 [9], Matlab R2023b [10], ConsensysBASIC v1.6.0 [11], Solidworks 2022 [12], Python 3 [13], Visual Studio Code [14].

- **Hardware Tools and Software:**

We utilize tools like a 3D printer Finder 3 [15] and the corresponding software for it FlashPrint 5.7.1 [16], Raspberry Pi 4 model B [17], PCB Design Software NI Ultiboard 14.2 [18].

Install the correct version numbers as specified for consistency and compatibility.

10.4 Coding Standards and Guidelines

To ensure high code quality and readability, we follow the google coding standard provided by *Styleguide* [19], which is a git directory. The site provides several guides for different languages, such as C#, C++, Python etc. Some of the basic universal guidelines can be viewed below:

- Code should be documented
- Each variable should have a descriptive and meaningful name
- Each identifier should be limited to one purpose if possible
- Proper indentation
- Limited use of global variables

For the rest of the guidelines see the given link.

10.5 Version Control

We utilize Github [20] as a code hosting platform for version control and collaboration. To keep track of our work and help us to easily explore the changes we have made while coding. GitKraken [21] is also used, which makes Git commands and processes easy, fast, and intuitive. For detailed guidance on using this system, please refer to *Version Control with Git* [22].

10.6 Testing and Quality Assurance

We adopt robust testing and quality assurance practices that are integrated into our development plan. For comprehensive insights into our testing process, please consult the section 12, Testing Plan.

10.7 Integration and Deployment

Our integration and deployment strategies are aligned with the goals assigned to each team (hardware and software). This requires individuals within each team to independently integrate and deploy their solutions to address the problem. Our integration strategy involves Continuous Integration, while the deployment process includes preparation, testing, and ultimately, deployment.

11 Documentation Plan

To have a structure in what, how, when, who and where to document files during a project is essential for its success. Therefore, the following section will in detail present this projects Documentation Plan and answers what needs to be documented, followed by how and when to document. Finally it will be outlined who is responsible for documenting and where data shall be stored to assure data is easily accessible. This section will also discuss the review and approval process for these documents.

11.1 What to Document

- Daily progress report
- Meeting protocols
- Data sheets
- Computer Aided Design (CAD) models and drawings, including iterations
- Code, including iterations
- Testing
- Research

11.2 How and when to Document

The rules listed below should be followed when documenting in this project.

11.2.1 Daily progress report

Daily progress shall be updated daily at the end of the workday by every member of the project group and shall reflect what the respective group member has worked on during that day.

11.2.2 Meeting protocols

Meeting protocols may be informal but should always include the following: date, participants, topic of the meeting. They shall be created during or right after the meeting and the completed protocol shall be stored as soon as possible but no later than the day after the meeting.

11.2.3 Data sheets

Complete data sheets as provided by the manufacturer should be documented if the described hardware might or has been chosen for the project. Data sheets of hardware which was not selected shall never be deleted to assure a complete documentation and decision history.

11.2.4 CAD models and drawings

Every iteration of model and drawing shall be documented and include the following: name, iteration number, date, author, revision. All files shall be stored in the appropriate location at creation.

11.2.5 Code

Code should be commented as it is written. Finished functionalities should be reviewed and different iterations documented separately. GitHub is used to structure and document all code including all iterations.

11.2.6 Testing

All tests shall be documented including at least setup description, expected outcome, collected data, results and conclusion. Description and expected outcome shall be documented before the test by the responsible group member in form of a short testing plan. Data and results shall be added after the testing. The conclusion shall be revised by at least another group member.

11.2.7 Research

All relevant researched scientific papers shall be documented as soon as possible.

11.3 Who is Responsible

- **Hardware Team Leader:** Oversees and approves hardware-related documentation and assigns hardware-related documentation tasks.
- **Software Team Leader:** Oversees and approves, and assigns software-related documentation tasks.
- **Hardware Developer:** Creates and updates hardware-related documentation as assigned.
- **Software Developer:** Creates and updates software-related documentation as assigned.

11.4 Where to Store Documents

Documents shall be stored in Microsoft Teams according to the provided structure to ensure availability, accessibility and data security. All document that might be important for the continuation shall also be uploaded to Github as formal documentation for the client.

11.5 Document Review Process

All design documents shall be reviewed by at least one other member of the respective group.

12 Testing Plan

This Test Plan offers a comprehensive overview of our project's testing strategy, methodology, and procedures. It serves as a framework for the validation and verification of both software and hardware components to ensure compliance with specified requirements. The document is structured to provide clarity to team members at any stage of the project's lifecycle.

12.1 Testing Methodology

- **Compatibility Testing (CT):** CT is a software and hardware testing method that assesses how a software application performs in various environments and configurations. Our team will adopt this approach to ensure the seamless operation of components and to verify their compatibility. This proactive measure helps prevent issues from arising during the integration of all components into the final product [23].
- **Requirements-Based Testing:** This is a software testing approach that focuses on ensuring that a system or software application meets its specified requirements. This is critical for the software and hardware teams and must be followed to successfully develop the Exoskeleton.
- **Functional Testing (FT):** This approach focuses on verifying that hardware components or systems perform their intended functions correctly. Test cases are designed to validate each hardware function, such as input/output ports, sensors, or buttons, to ensure they work as expected. This approach is upheld to ensure the safety and reliability of the final hardware component composition [24].
- **Reliability Testing (RT):** RT assesses the consistent performance of both software and hardware over an extended duration, aiming to minimize failures. This encompasses tests measuring parameters like MTBF (Mean Time Between Failures) and assessing performance under challenging conditions. It is a critical testing method, particularly since project requirements specify a specific acceptable failure rate for the Exoskeleton [25].
- **Endurance Testing (ET):** This involves subjecting the hardware to continuous operation for an extended period to identify any performance degradation, overheating, or other issues that may occur with prolonged use. The aim is to develop a prototype what is robust and stable [26].

12.2 Testing Team Structure and Roles

Our testing team includes the following roles:

- **Testing Team Leader:** Software and Hardware Team leaders are in charge of the testing strategies, as well as organize and ensure that tasks are assigned to each team members.
- **Testing Engineer:** Manage tests with respect to the chosen testing methodology (e.g, CI, compatibility testing, and requirement testing) as well as report all issues and problems that can occur in the testing stage and mention possible solutions.

12.3 Bug Reporting and Tracking

We use GitHub to log and track identified defects. Each testing team member is responsible for documenting and reporting bugs according to our bug reporting guidelines, outlined in GitHub repository [20].

12.4 Test Schedule

Our testing schedule aligns with the overall project timeline. The detailed test schedule can be found in this section 6.

13 Communication Plan

To facilitate the sharing of information among team members throughout the project lifecycle, a Communication Plan has been defined. Its purpose is to promote mutual understanding and foster cooperation among team members and stakeholders. The Exoskeleton Communication Plan is presented as follows.

- **Who (Target Audience):** Project team members such as the hardware and software developers, hardware and software team leaders, contribution team in Technological University of Panama , stakeholders, the examiner, the course responsible, and the supervisor.
- **Why (Purpose):** Project updates inform stakeholders about progress, task assignments, to manage deadline and potential issues and to share ideas with our contribution team in Technological University of Panama.
- **What (Type of Information):** Project updates, task assignments, meeting agendas, change requests, and risk alerts.
- **When (Timing):** Daily communication between team members, weekly communication with the stakeholders and biweekly meeting with our contribution team in Technological University of Panama.
- **How (Communication Channels):** Communication methods include email, direct communication, meetings and project management software and communication such as Microsoft Teams.
- **Responsible:** Hardware and Software Team Leaders.

Who	Why	What	When	How	Responsible
Project members	To discuss the progress of the ongoing plan and put goals for the coming days	Meeting agendas	Daily	Direct communication, meetings	HW,SW Team leaders
Project members	To address progress and issues, Plan for the next meeting with the Stakeholder	Meeting agendas	Weekly	Direct communication, meetings	HW,SW Team leaders
HW Team Leader	To keep updated on hardware status and needs, to know their responsibilities	New tasks, updates, changes	Everyday	Email, direct communication, meetings	Albin Gustafsson
HW Developer	To know their tasks and deadlines	Task Assignments	As needed	Email, direct communication, meetings, project management tools	Sebastian Ahlström and Moritz Schmidt
SW Team Leader	To keep updated on software status and needs, to know their responsibilities	New tasks, updates, changes	Everyday	Email, direct communication, meetings	Irini Provatidis
SW Developer	To know their tasks and deadlines	Task Assignments	As needed	Email, direct communication, meetings, project management tools	Jalal Taleb
Team members in Technological University of Panama	To share ideas and progress of the common project	Task Assignments	Biweekly	Email, project management tools	Project members
Stakeholder	To share our progress and to address any issues that may arise	Meeting agendas	Weekly	Email, direct communication, meetings, project management tools	Project members

14 Handover Plan

The following section describes this projects handover plan. It outlines the systematic process of transferring all documentation, results, reports and hardware to the project owner at the end of the project.

14.1 Handover Team Structure and Roles

Our handover team includes the following roles:

- **Handover Team Leader:** Coordinates the handover process, ensuring that all project deliverables and documentation are transferred wholly and accurately.
- **Handover Specialist Hardware:** Organises the handover of the hardware related documents.
- **Handover Specialist Software:** Organises the handover of the software related documents.

14.2 Handover Methodology

To ensure a complete and comprehensive handover process, the following methodology is employed.

- Handover specialists check inventory and itemize all deliverables
- Team members review itemization
- Deliverables are transferred to the project owner

These steps have been chosen to ensure a comprehensive and efficient handover process.

14.3 Handover Tools

We leverage the following tools, technologies, and systems during our handover process:

- **Inventory Management Tools:** We utilize Microsoft Teams 10.4.7 and GitHub to catalogue and track all project deliverables for handover.
- **Presentation Tools:** If necessary, we use PowerPoint 1808 to conduct presentations or demonstrations.

14.4 Handover Schedule

Our handover schedule aligns with the overall project timeline. See section 6 for the handover timeline.

14.5 Documentation

Comprehensive project documentation, including user manuals, technical documentation, and project reports, will be provided during the handover process. See section 11 for more information.

14.6 Presentation and Demonstration

Our team will conduct presentations and demonstrations where necessary to familiarize the stakeholders with the project's operation and results.

14.7 Final Sign-off

All deliverables will be handed over to the project owner at the 12.01.2024. The project owner will sign off, indicating successful handover and completion.

15 Individual Contributions

Chapter	Responsible	Contributor
1. Executive Summary	Irini Provatidis	Jalal Taleb Albin Gustafsson
2. Background	Irini Provatidis	
3. Purpose	Irini Provatidis	
4. Goal	Irini Provatidis	
5. Work-Breakdown-Structure (WBS)	Moritz Schmidt	Moritz Schmidt Albin Gustafsson & Irini Provatidis
6. Project Schedule	Albin Gustafsson	
7. Project and Product Requirements	Sebastian Ahlström	
8. Limitations	Sebastian Ahlström	
9. Risk Analysis and Counteractions	Sebastian Ahlström	Irini Provatidis
10. Development Plan	Sebastian Ahlström	
11. Documentation Plan	Moritz Schmidt	
12. Testing Plan	Jalal Taleb	
13. Communication Plan	Jalal Taleb	Irini Provatidis
14. Handover Plan	Moritz Schmidt	

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