Technology Assessment and Feasibility Study of High-
Throughput Single Cell Force Spectroscopy

By

He Cheng

B.E. Bioengineering
Nanyang Technological University, 2009

SUBMITTED TO THE DEPARTMENT OF MATERIALS SCIENCE AND
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DEGREE OF

MASTER OF ENGINEERING IN MATERIALS SCIENCE AND ENGINEERING

AT THE

MASSACHUSETTS INSTITUTE OF TECHNOLOGY

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Signature of Author: ………………………………………………………………………
Department of Materials Science and Engineering

Certified by: …………………………………………………………………………………
Christine Ortiz,
Thesis Supervisor
Professor of Materials Science and Engineering

Accepted by: …………………………………………………………………………………
Christopher Schuh
Chair,
Departmental Committee on Graduate Students
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Abstract

In the last decade, the field of single cell mechanics has emerged with the development of high resolution experimental and computational methods, providing significant amount of information about individual cells instead of the averaged characteristics provided by classical assays from large populations of cells. These single cell mechanical properties correlate closely with the intracellular organelle arrangement and organization, which are determined by load bearing cytoskeleton network comprised of biomolecules.

This thesis will assess the feasibility of a high throughput single cell force spectroscopy using an atomic force microscopy (AFM)-based platform. A conventional AFM set-up employs a single cantilever probe for force measurement by using laser to detect the deflection of the cantilever structure, and usually can only handle one cell at a time. To improve the throughput of the device, a modified scheme to make use of cantilever based array is proposed and studied in this project. In addition, to complement the use of AFM array, a novel cell chip design is also presented for the fine positioning of cells in coordination with AFM cantilevers. The advantages and challenges of the system are analyzed too. To assess the feasibility of developing this technology, the commercialization possibility is discussed with intellectual property research, market analysis, cost modelling and supply chain positioning. Conclusion about this technology and its market prospect is drawn at the end of the thesis.
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Chapter 1 Introduction

1.1 Overview and Scope

The mechanical properties of single cells are important characteristics for identifying the stage of cell development, proliferation and health status. Single cell mechanics is the study of the mechanical properties of cells and their individual responses to mechanical stimuli. As shown in Figure 1, the field of single cell mechanics has gained increasing attention in past decades. It has become increasingly recognized in past 20 years that not only chemical substances from hormone secretion or signalling pathways affect the behaviours of cells, the mechanical interactions of cell and its environment, which is related to the cytoskeleton resides in cytoplasm [1, 2], can also dictate important changes in the life cycle of cells. For example, the transformation of mesenchymal stem cells (MSCs) to matured functional cells such as osteocytes and chondrocytes [3] during embryonic development involves the change of cytoskeleton as well as the mechanic properties of the differentiating cells [4].
The study of single cell mechanics involves the measurement of cell properties such as quasi-static indentation stiffness [5], shear modulus [6], loss, storage, and complex modulus in response to dynamic oscillatory loading [7] and relaxation time constants [8]. The detecting of single cell mechanics can sometimes measure force and displacement as small as a few pN and angstroms [9], these measurements impose high requirements in terms of resolution, accuracy, response time and throughput on measurement instruments. In this thesis, a novel technology of high-throughput cell force spectroscopy will be proposed and analyzed in details in terms of system components, fabrication process, technology limitations and potential future developments. In addition, analysis on commercialization is also carried out, including intellectual property research, market analysis, cost modelling and supply chain positioning, to assess the feasibility of developing this technology.

1.2 Single Cell Mechanics

1.2.1 Single Cell Mechanics for Cell Characterization

Recent studies suggest single cell mechanics can be used to characterize cells in terms of cell phenotype, origin and disease. In the research carried out by Darling et al [10], the viscoelastic properties of cells were used as biomarkers to differentiate three cell lines (chondrocytes, osteocytes and adipocytes) derived from mesenchymal stem cell (MSC), so that all four cell lines can be separated. The group used commercially available atomic force microscope (AFM) to perform indentation on cells, and subsequently measured the stress relaxation time constant to estimate the elastic modulus and viscoelastic modulus. They found in the experimental configuration, when cells were adhered to substrate, there was significant difference (p<0.0001) in elastic modulus and viscoelastic modulus among all four kinds of cells.

The logic of using single cell mechanics as a biomarker lies in the connection of single cell mechanics and cytoskeleton; single cell mechanics manifest cytoskeleton structure thus could be an indication of cell status [11]. Cytoskeleton is a network composed of filamentous proteins of actin, tubulin or one of several classes of intermediate filament [1], which are major contributors to the mechanical properties of cells and are also responsible for intra cellular transportation as well as cell
locomotion [12, 13]. Cytoskeleton mechanically connects intra cellular structures to extra cellular environment through the docking by transmembrane proteins such as integrin and Neural Cell Adhesion Molecule (NCAM) on extra cellular matrix (ECM) [14, 15].

It is found that cell deformability is related to cytoskeleton too, reduced assemble of cytoskeleton increases the deformability [16]. When cells age, their cytoskeleton also changes as there is a reduction in the amount of constituent biopolymers of cytoskeleton [17, 18]; as a result, their elastic modulus and shear modulus decrease, making them less resistive to external loading such as tension and shear [19]. In another previously cited work [20], the author concluded single cell biomechanical responses change due to cytoskeleton rearrangement.

Another obvious example demonstrating the correlation between single cell mechanics and cytoskeleton is the phenotype of a single cell could be altered by the decreased formation of actin, resulting in a morphological change which transforms the cell into spherical shape with a chondrocytes-like morphology [21].

1.2.2 Single Cell Mechanics for Diseases Indication

Cytoskeleton’s material properties are able to affect the response of cells or tissue to external loading. Cell or tissue’s abnormal responses to mechanical loadings can induce various diseases, such as asthma, osteoporosis, atherosclerosis and diabetes; and they contribute to the clinical presentation of these important diseases [22]. Since single cell mechanics reflects the status of cytoskeleton, it is possible that the measurement of mechanical properties of cell can indicate abnormalities in cytoskeleton, thus may be potentially used for diagnostic purposes.

As a highway for signal transduction, cytoskeleton provides various localization mechanisms for enzymes like glycolytic enzymes (aldolase, glyceralde hyde-3-phosphate dehydrogenase, lactate dehydrogenase, and phosphofructokinase-1 [23]), protein kinases (Tyrosine kinases [24], protein kinase C [25]), phospholipases (phospholipase A$_2$ [26]) and GTPases [27]. Upon the conformational change of cytoskeleton under mechanical stimuli, the enzymes linked with the network have to undergo architecture change correspondingly. If their conformations are altered, the
thermodynamic and kinetics of the chemical reactions will change inevitably. Cytoskeleton also affects the transportation of ions or intracellular compartments; for example, F-actin can affect the ion channels of Na\(^+\) and Ca\(^{2+}\). When the assembly of F-actin is prohibited by drugs like cytochalasins [28] in culture medium, the measurements of ion flow from patch clamp experiment showed the conductivity of these ions increased dramatically, however, upon stabilization of F-actin by adding drugs like phalloidin [29], the conductivity of these channels dropped in comparison with previous case. Thus, if an abnormal cell mechanics property is detected, it will likely to manifest the mechanotransduction is impaired, either the ion channels or the enzymes linked on cytoskeleton may not function properly.

As such, many diseases can be correlated with the dysfunction of cytoskeleton system, and can be recognized with altered single cell mechanics. It has been recognized 20 years ago that cytoskeleton may be related liver illness [30]. Zatloukal and coworker [31] have reviewed that in diseases like Alcoholic and non-alcoholic steatohepatitis (ASH & NASH), Copper toxicosis, Cholestasis as well liver cancer, the degradation of keratin filament and the disassemble of the intermediate filaments usually happen. Besides, there is also imbalanced production of keratin subunits, so that the assembly of keratin filaments is greatly hindered [32]. As a result, the cells loss the mechanical support to physically maintain proper cell morphology and mechanical integrity, resulting in the ballooning of the cell [33].

Cell mechanics can also be a marker for apoptosis, it has been shown that cells are less resistive to internal stress due to degradation and restructuring of all three cytoskeletal filaments. The change in mechanical properties of the cell may even produce blebbing of cytoplasm membrane, which is the hallmark of programmed cell death [34-37]. In experiment, cell death can be induced directly by applying drugs like taxol and vinblastine to purposely disrupt of microtubule [22].

In cancer cells, deformability has been shown to be positively correlated with the degradation of cytoskeleton and the metastatic rate. In the research carried out by Ochalek et al [16], four variants of the same cell line with different metastatic rates are tested for their mechanical deformability. In the experiment, the variant with the least metastatic rate had the lowest deformability while the most metastatically unstable variant had the highest stability. When all variants were treated with drugs
preventing the assembly of cytoskeleton, their deformability increased, strongly suggesting a correlation between the involvement of the cytoskeleton and their carcinogenic potential.

In another work by Suresh et al [20], single cell mechanics was used to characterize two distinct human diseases: gastrointestinal cancer and malaria. During their experiment, they found the elastic modulus of treated cancer cell to be significantly different from that of untreated ones, and they suggested it could be due to the reorganization of cytoskeleton near nucleus. In their test of red blood cell (RBC) in vitro with optical tweezers, the shear modulus of RBC increased about 10 fold because of parasite infection, resulting in the easy corruption of RBC when experiencing high shear fields.

Suresh’s experiment results were further confirmed by the study of RBC mechanics when RBCs from patients with sepsis and leukostasis were tested in microfluidic devices [38]. The experiment subjected RBC to narrow fluidic channels where cells had to deform in order to flow with fluids in the channel. It was shown RBCs from patients suffering these hematologic diseases had a decreased deformability compared to normal RBCs. The authors also concluded these diseases affected the status of cytoskeleton, as when drugs affecting cytoskeleton (cytochalasin D and pentoxifylline) were applied to RBCs, their deformability increased significantly.

Because of the important role cytoskeleton playing during the differentiation, proliferation and other events in cell life cycles, the abnormality in cytoskeleton is the source of many diseases. As previous examples showed, the measurement of single cell mechanics reflects the status of cytoskeleton, it is thus important for the understanding of cell biology as well as disease etiology to obtain this information [39].

1.2.3 Modelling of Single Cell Mechanics

There are a few models relating the mechanical properties of cells to the measurement done with various tools, thus bridging the macroscopic measurement with deeper understanding of cytoskeleton.
Besides the mostly used model where cells are treated like homogenous viscoelastic entities, the other widely used model to relate cell’s mechanical behaviours to cytoskeleton is the tensegrity model. The stability of the system depends on tensional integration, rather than compressional continuity. In this model, the cytoskeleton of a cell is represented by a group of interconnected bars and viscoelastic cables. And it is viewed that these structures are interconnected between cells in a tissue through the mechanical linkage of ECM [40]. Upon loading, tensegrity structure tends to deform globally rather than locally, that all its mechanical component can rearrange in order to respond to the applied force [41].

Figure 2 shows one of this model, where 6 bars and 24 viscoelastic cables were used to construct the model. In loading scenario, numerical analysis has to be used to solve for loading response. In this particular example, four nodes are fixed in order to simulate the attachment of cell to substrate surface. External loading can be applied though any of the nodes. In most simulations, only first order deformation of bars is considered, because higher order deformations will only become significant with larger dimension, which is less relevant when modelling cells.

Tensegrity model has been shown to be able to predict the mechanical measurement from experiment data, in fact, tensegrity model was adopted because its prediction coincides with experiments data where measured cell’s elastic modulus increases linearly with deformation [42]. Moreover, modified tensegrity model predicts accurately that the global structural viscosity and global structural elasticity changes proportional to the square of the size of cell [43]. In addition, tensegrity represents the combined mechanical behaviours of cytoskeleton components. Like F-actin, though each single component of tensegrity does not exhibit linear increase of elastic modulus under loading, The combined network showed linear increase in elastic modulus just as combined F-actin network [44]. There may even be some connection between the structures of tensegrity network to physical cytoskeleton structures. In cytoskeleton systems, microfilament (MF) can produce internal stress through internal actomyosin sliding as the viscoelastic cable in tensegrity model, and the stiffness of MF can be altered with different concentration of crosslink proteins among molecules [45, 46].
There are also models correlate the mechanical properties and measurements on special cells (like RBC, which do not have nucleus when it is mature) [47, 48]. It must be pointed out that cell like RBC cannot be modelled with tensegrity, because their underlying structures are essentially different. In tensegrity modelling, the 3D network corresponds to the actin filament network inside cell, while in RBC the major contribution of mechanical properties comes from spectrin network under the double layer of lipids.

Figure 2. Tensegrity model of cell mechanics. A tensegrity model with 24 elements (up) and tensegrity model subjected to tension (bottom) [40, 43]
1.3 Tools for Mechanical Properties Measurement

In the past 20 years, single cell mechanics has gained increasing interest in the research community. Since physical measurement can be correlated to the status of cytoskeleton, it has been of great interest to develop measurement instruments for probing the mechanical properties of single cells. There have been numerous tools developed for this purpose, the earliest of which is the use of micropipette to perform aspiration [49, 50] on nontransformed and transformed cells. Micropipettes are generally made by pulling hot glass capillary tubes. Suction can be applied to cell surface once micropipette is attached to the target cell. The deformation of the cell measured with respect to time reveals the viscoelastic properties [51]. For example, in the measurement of viscosity of blood granulocytes, the experiments revealed that the viscosity of the cell changes with respect to temperature, from $2 \times 10^3$ poise at 27°C to $10^3$ poise at 37°C [52]. Micropipette aspiration has also been used in measuring the Poison’s ratio of chondrocytes and osteocytes in [51], which ranges from 0.36 to 0.38.

Magnetic bead microrheology is another technique used in measuring the mechanical properties of cells [53]. In this kind of set up, magnetic beads functionalized with ligands that can attach to cell surface are used. When magnetic field is applied, tangential force exerts onto the surface of cells by the magnetic beads. In [53], the shear modulus of fibroblasts is measured to range from $2 \times 10^{-3}$ Pa m to $4 \times 10^{-3}$ Pa m. The authors also found the bulk shear modulus to be $0.5 \times 10^{-4}$ Pa, membrane coefficient of friction to be $2 \times 10^9$ Pa s/m corresponding and cytoplasmic viscosity of $2 \times 10^3$ Pa s. By using this tool, it is also revealed that cells can be viewed as composed of materials near their glass transition temperature, and cytoskeleton modulate cell’s responses to mechanical stimuli by altering effective noise temperature of the matrix [54]. A scaling law of elastic and frictional properties and was observed between different kinds of cells [54]. Another similar tool is optical tweezers, which uses similar principles to measure local mechanical properties of cells by utilizing laser deflection particles [55, 56]. In [55], the authors estimated the local shear modulus of red blood cell membrane to be $2 \times 10^{-6}$ Nm$^{-1}$.

Osmotic loading subjects cells to different osmotic pressures and detects the change in cellular volume, which is an indication of whole cell mechanical properties [57]. Through the use of osmotic loading, it is found the normalized volume of cells change
linearly with respect to osmotic pressure, and the instantaneous and equilibrium elastic moduli as well as the apparent viscosity of the cells were significantly decreased by hypoosmotic loading, but were unchanged by hyperosmotic loading [58].

Other techniques quantifying whole cell mechanical property include rheology measurement by plating manipulation where the cell is attached and traction or uniaxial compression is applied through the deformation of the plate or the plate, the applied force and measured displacement are in Newton and mm range respectively. Microplating is also possible where the resolution of this technique can be increased. In the experiment reported in [60], microplating was used to measure the loss and storage modulus of single HeLa cells. The authors found the storage modulus to be around 660 Pa when stimulation was performed at a frequency of 1 Hz, which is quite close to the AFM measurement result at low frequency. Microfluidic devices can also be used for measuring the mechanical properties of single cells, like their resistance to shear or their ability to deform [61].
1.3.1 Mechanical Probing by AFM

Among the characterization techniques, one important probing method is the use of AFM to quantify single cell mechanics. Atomic Force Microscope (AFM) is a unique tool which enables high resolution measurement of the mechanical properties at sub nanometer level [62]. In addition, unlike other high resolution scanning probe microscopes, *in situ* force measurement on biosamples is also possible because AFM can operate in a wide range of environments [63, 64].

AFM is developed from scanning tunnelling microscope (STM); it was originally designed for the detection of the interactions between probe tip and substrate surface. It provides flexible measurement of surface topography, friction, surface adhesion and deformation [65]. Typical AFM probes consist of a deflectable cantilever and a tip at one end of the cantilever. The cantilever must be resistive to deflection in two directions while prone to deflection in the third direction. Typically, cantilevers are fabricated so that their resonance frequency is above 10 kHz [66]. The cantilever structure can be as large as a few micrometers, while the radius of curvature of the end of tip can fall below a few nano meters (10nm for Si and 20-50 nm for SiN [66]). The tip of AFM is usually fabricated from silicon or silicon nitride, which moves into close proximity to interact with sample surfaces. AFM tip can be coated with various materials for alteration of mechanical, thermal and electrical properties [67-71].

AFM probe interacts with sample with both short range and long range forces; van der Waal forces act between sample and probe tip, while other forces such as electrostatic force capillary force, chemical force as well as Casimir force can also play a part in measurement. Because measurement is done by force interaction, no fixation or staining of sample is required, thus the device is particularly suitable for the testing of living tissue. The device is also able to quantify both static and dynamic loading responses of cell by applying forces at different frequencies controlled by piezo force generator relayed to an external signal generator.

AFM measures samples with length scale from a single molecule to a whole cell, with a force range from several pico newton to nano newton. AFM obtains force-displacement curves, which reveals elastic modulus of cells [72, 73]. In addition, hysterereses between approach and retraction curves can be used to evaluate the time-dependent modulus and time constant [74].
AFM typically operates in three modes, namely, contact mode, noncontact mode and tapping mode. In contact mode, total force felt by the cantilever tip is repulsive. In noncontact mode, total force exerted on the tip of cantilever is often attractive, and atomic resolution is achievable [75]. And due to the shape of the potential well of van der Waal interactions, there will always be a distance region that the AFM probe is unable to work in with, as such, the measurement can only achieve a lower resolution [66].

Typically, biosamples are measured in noncontact/tapping mode, not only to achieve higher resolution, but also to prevent the destruction of sample by indentation or friction between tip and sample. In noncontact/tapping mode, the cantilever is usually driven at a preset frequency. A change in the frequency of vibration corresponds to the external force felt by the cantilever. The following relationship shows the estimation of force by detecting the modulation of frequency, where $k$ is the stiffness of a single leg cantilever beam.

\[
\frac{\Delta f}{f} = -\frac{1}{2k} \frac{\partial F}{\partial x}
\]
AFM has been used to probe cell surface elasticity. For example, one of the pioneering studies of using AFM to probe the mechanics of single cell was carried out in 1994 by Hoh and Schoenenberge [62, 76], in their experiment, they measured the elastic modulus of MDCK and R5 cell and cellular dynamics with time lapse AFM. AFM indentation on cells revealed the loss and storage moduli follows power law, where the complex modulus at any frequency can be expressed as a base complex modulus times the power of that frequency over the base frequency. For example, in the measurement of complex modulus of human lung epithelial cells [77], the group found both storage and loss modulus increased with indentation frequency up to 100 Hz. The exponent for storage modulus was measured to be around 0.2, and the exponent for the loss modulus was measured to be around 0.067.

![Figure 5. Complex modulus measurement by AFM indentation [77]](image)

Because of the high resolution from AFM, there has been experiments using AFM for cell regional force mapping, for example, in [6], the group used AFM to detect the loss and storage modulus at nucleus region and peripheral regions respectively. The group found the storage and loss modulus to be around 300 Pa and 95 Pa respectively when indentation is performed at 0.1 Hz at nuclear region. However, the peripheral regions were found to be more rigid than nuclear region at all indentation frequencies.
It is also possible to directly connect the measured mechanical properties with cytoskeleton structures in living cells thanks to the high resolution of AFM. In the work done by Christian Rotsch and Manfred Radmacher [78], it has been shown that AFM measurement directly correlates with the cytoskeleton structure; and can reflect the change of cytoskeleton upon the addition of drugs. In their experiment, NRK fibroblasts were measured in contact mode with an AFM from Veeco ® and a force map was generated from AFM measurement. Elastic modulus was then obtained from force displacement curve of the AFM measurement. The cells were subsequently stained with dyes that were selective to F-actin. As the following figure shows, the features on AFM force map correspond to the stained structure by the dye, confirming that AFM can directly probe cytoskeleton. In addition, when cytochilatin was used to prevent the assembling of cytoskeleton, AFM measurement showed less features and a decreased elastic modulus.

AFM can also accommodate to wide range of cells, not only eukaryotic cells can be probed, prokaryotic cells like bacteria are also suitable for AFM investigation. For example, the surface elasticity of bacteria M. gryphiswaldense was determined by applying compressional stress on cell wall [79].

Figure 6. AFM measurement directly correlates with cytoskeleton. NRK fibroblast (a), AFM image (b), magnified view of AFM image (c), cytoskeleton staining (d), magnified view of staining (e) force map (f) [78]
Chapter 2 High Throughput Single Cell Measurement Technology

Although it has been demonstrated that the single cell mechanics can be quantitatively assessed by a range of techniques mentioned previously, it is important to realize that most of these techniques are of extreme low throughput, i.e. only capable of handling a limited number of cells in a reasonable experiment time frame [80]. As a result, the data produced by these techniques may not be an accurate reflection of the actual status of a whole cell population.

This represents certain limitation when it comes to the measurement of any heterogeneous cell population, which is often the case if the measurement is done on any sample derived from living organisms. As it is now known that in any biological events like growth, development and disease, there involves a coordinated reaction of a heterogeneous population of cells [81]. It is also noted that the distribution of the measured mechanical properties may not always be the same as assumed [38]. One example was given in the measurement of cell deformability through microfluidic devices. The passing time of RBC through a channel was recorded and used as a parameter to represent single cell mechanics. As shown in the following figure, contrary to common assumption that a Gaussian distribution should characterize the transit time, the cells derived from patients with hematologic diseases exhibited non-Gaussian distribution that the majority of the cell population either pass within 2 seconds or will not pass within 8 seconds.

Moreover, when tested with neutrophil, control group exhibited dramatic differences with group treated with drugs inducing inflammation, suggesting the distribution of cellular mechanical properties can also be related to the status of the cells, thus could be useful for characterization and diagnosis. It is also important to determine how the different parts of metabolism are coordinated and balanced through the study of heterogeneous cell population. The lack of distribution information could represent a flaw in measurement.
Furthermore, it must be stressed that although some techniques do measure the property of a large number of cells, it is quite often the average properties of the whole population are finally presented. For example, an increase in fluorescent measurement of a clone can indicate higher activities of bacteria in that clone, however, the measurement may not be an accurate indication of cell status as it could mean either each cell has an increase in their activity level, or more cells turns from “off” state to “on” state.
As a consequence, there is a need to produce new technology that would allow the measurement of multiple individual cells in a reasonable experiment time frame. Recent technology development has improved the throughput of flow-through optical microfluidic devices, so that the throughput could be reached at 1 cell/min [82, 83] to 50-100 cells/min [38].

AFM technology holds great potential for single cell mechanics measurement; it cannot be replaced by microfluidic devices in that it can produce direct and high resolution measurement on single cell in a controlled and more precisely manner; therefore it is of great interest to develop a high throughput platform based on AFM principles of operation. It is desirable that this technology should be capable of being multiplexed, and in the same time be friendly for mass production and cost reduction.

In the following section, a new technology platform is proposed to fulfil the goal of high throughput single cell operations. The proposed technology is composed of three major components, with a high throughput AFM system as a functional basis of the technology, which is able to combine the high resolution and high throughput advantages. A cell chip is the complementary component to the AFM system, which
is able to help the operation of AFM on biological samples. Lastly, a controller system is needed in order to coordinate the whole set up.

2.1 High-throughput AFM system

The AFM system should be able to operate in a high speed fashion so that the throughput can be increased. There are two ways to increase the operating speed of the system, namely, to accelerate each probe and to integrate multiple probes in the system. The first approach uses less stiff cantilever and elimination of cantilever resonance [84, 85]. However, this approach faces severe limitations that some applications require certain probe-sample interaction time [86-90], so that the probing speed cannot be increased over certain limit; besides, the magnitude of speed increase is limited.

Highly parallel array of AFM probes is more favourable because the throughput can be increased proportional to the number of probes used. The proposed measurement system is composed of a force measuring instrument with multiple cantilevers
functioning as AFM tip array and cell chip for the precise positioning of cells. In addition, a controlling system is necessary in order to coordinate the two components and to analyse the acquired data.

The proposed device employs an array of AFM tips individually controlled by picoforce piezos. There is a cell chip loaded on a platform and is controlled by three motors so that it could move in X, Y and Z directions independently. With this configuration, multiple cells can be tested simultaneously in one round, with each cell measured by one AFM tip, and the next round of readout can be obtained by moving platform to next designated coordinates.

The proposed device should have much higher throughput compared to conventional AFM, thus it can be used for high throughput cell mechanical property testing for cell biology study. Besides, this device should be able to test bioproducts such as engineered tissue, single cell mechanical testing is commonly used as a merit metrics of engineered tissue performance.

The number of cells tested per round is determined by the number of AFM tips in the array. Carrying out the testing for several rounds does not only generate averaged properties of the tested batch of cells, but also produces statistical information of this batch. Distributions, variations and deviations of mechanical properties can be obtained for better characterization of not only a single cell, but also of a group of cells. Generally speaking, two categories of such devices are mostly developed. The first category consists of device primarily developed by IBM (termed “millipede” system) [91].

2.1.1 Millipede Array

IBM Research Division, Zurich Research Laboratory presented a scheme using a square array of AFM cantilevers for patterning polymer by indentation [91], where 32×32 (1024) probes were used to create patterns on substrate surface. The following figure demonstrated the concept of IBM’s parallel array “millipede”.
The millipede device consists of a square array of cantilevers arranged in a row-column manner. Each cantilever is in charge of a field about 100µm × 100µm [93]. The cantilevers are entirely fabricated from silicon for good thermal and mechanical properties through surface micromachining by either plasma etching or wet etching [88]. Each cantilever consists of two legs acting as soft springs; legs are highly doped to reduce the resistance and energy dissipation, while tip is lightly doped. Metal wirings are avoided on leg region to avoid any electron migration or parasitic effect.

Legs are specifically produced with less mechanical stiffness to reduce the force required for deformation, so that the tip and substrate can be well protected [88].

The unique point about this millipede system is that whole array is controlled by only one z-scaner, so that the complexity of the system can be reduced [93]. However, several z-scanners can be used if more precise control of the vertical position of the array is needed [94]. As a result, the system requires uniform substrate surface to avoid the damage of the array. When approaching substrate surface, four cantilevers integrated at the corners of the array will detect the tilt of the substrate relative to the array [88].

The cantilevers in this array are not mechanically coupled and can be individually actuated by the application of voltage pulse [93]. When actuation is performed, each
individual cantilever receives two electrical pulses from the row and column it belongs to, if the sum of the pulses is nonzero, an indentation action will be performed [95].

A more complex array of 64×64 (4096) cantilevers has been fabricated [94], demonstrating the possibility of building high throughput devices by this approach. However, the application of millipede system to biological sample faces severe limitation. The millipede system is controlled by only one z-scanner [93]. Although this setup reduces the system complexity, it requires uniform substrate surface to avoid the damage of the array. Due to inherent surface roughness of biological samples, the variation in z direction can be as huge as several microns, which may disable the probing of the array to the surface. In addition, though the cantilevers are not mechanically coupled, their signals are coupled because the signal is transmitted in rows and columns, i.e., cantilevers in the same row will receive the same actuation signal (amplitude) transmitted by that row, so do cantilevers in the same column. As a result, the cantilever array lacks the ability to control each cantilever flexibly. Furthermore, each actuation made by the cantilever is static, rather than vibrational actuation used in tapping mode or non-contact mode AFM, which is more likely to destroy soft materials. Another restriction of the millipede array comes from the probing method and response time; IBM chose to use heat conduction to probe-sample distance, and with the measurement of resistance change to detect beam deflection. Both detection and actuation is too slow and too inefficient [96] for probing soft tissue because temperature change has to overcome the heat capacity of cantilever and heat dissipation to environment [94].

2.1.2 Individually Controlled Cantilever with Sensor and Actuator

The other kind of cantilever array is composed of individually controlled cantilevers with integrated sensing and actuation functionality. A High number of cantilevers form the self-actuated arrays, and these arrays can be further integrated into a large scale array for achieving even higher throughput [97]. For achieving actuation functionality, the cantilevers in this case mainly use piezoelectric materials, which can transfer the externally applied voltage into strain. This sort of array is generally faster in operation [98].
The piezoelectric material used could be materials from group III-V or group II-V compound materials, for example, ZnO is the most widely used piezoelectric material in MEMS, in addition, AlₓGa₁₋ₓAs and Pb(ZrTi)O [99] can also be used for higher piezo constant [100].

The pioneering work of the use of parallel scanning probes with individually controlled cantilevers was initially used by S. C. Minne in 1994 [101] to pattern amorphous silicon on single crystal silicon 100 surface as a lithography mask. Early developments from the group used one array of two piezoresistive cantilevers, with one in feedback loop set to constant force of 600 nN and the other in an open loop. Both cantilevers were operated under the condition of 20V bias voltage. They discovered that individual feedback of the cantilevers must be established in order to achieve high precision force control. The group also used arrays of four parallel cantilevers to image 100µm × 400µm surface with the same amount of the time usually only sufficient for 100µm × 100µm square area. Thus it corresponds to a 3 fold increase in throughput in the parallel cantilever system.

However, this group did not manage to integrate the sensing function onto each cantilever in the array, they used laser deflection detector to probe the cantilever bending. When larger number (4 cantilevers) of cantilevers was used, due to limited space, they were not able to use one laser deflector for each cantilever, so individual control was not achieved. As such, the author concluded further development on integration of sensing with AFM array is the key to the application of AFM array with large number of cantilevers for high precision operations.

The integration of the sensing unit in cantilever itself is necessary, because it greatly reduces the space needed for sensing modules. As illustrated in Figure 4, conventional AFM employs laser and photodiode set up for the purpose of cantilever beam deflection detection. For low level integrations of AFM like in previous case, separated sensing unit has already shown its limitations. As such, for the integration of a higher number of cantilevers, it would be impossible to continue to use the separated sensor and cantilever design.

There are a few categories of sensors that can be readily incorporated onto cantilevers, for example, capacitors [102, 103] can be built onto the cantilevers for sensing. Piezoelectric materials can also be incorporated onto the cantilevers for the detection
of beam deflection by measuring the voltage across the material. Another way to integrate the sensing functionality onto the cantilever is to use piezoresistive effect of silicon. It has been demonstrated that the resistivity of silicon could change upon deflection, although no charge is produced. One example of using piezoresistive sensing is demonstrated below in [98] and improved in 1996 [104], where the sensor and actuator were integrated onto cantilever together. The following figure shows one of the cantilevers in the array.

The design uses V shaped leg for achieving the sensing functionality, where the bending of the cantilevers can be transferred into the resistance change along the V shaped leg. Current is passed through the V shaped support from one leg to another. The change in the resistance will induce the change in current so that the deflection can be calculated.

![Side View](image)

**Figure 11. Schematic drawing shows cantilever with integrated sensor and actuator [98]**

The cantilever beam is deposited with a layer of 3.5 μm thick ZnO on top of the Si beam. If a field is applied in the direction of the beam, the cantilever will bend due to the expansion of ZnO. Below the silicon oxide insulator, the doped Si functions as a piezoresistor to detect the bending of the beam.
However, this design suffers inherent flaw that the piezoresistor sensor spans the entire range under ZnO actuator; it detects not only the bending caused by force on tip, but also the bending of ZnO actuator itself. The signal thus has to be electronically compensated for correct reading.

A modified design was then produced to compensate for the piezoelectric effect caused by ZnO. The following figure shows the modified design of cantilever beam.

![Modified Scheme for Integration of Sensing and Actuation with two-leg cantilever and Si functioning as piezoresistor](image)

**Figure 13. Modified Scheme for Integration of Sensing and Actuation with two-leg cantilever and Si functioning as piezoresistor [104]**

The modified design added another layer of heavily doped Si under ZnO actuator. this layer of Si has two benefits, firstly, by increasing the doping under ZnO region, the piezoresistive coefficient is decreased by 80% [104], thus reducing the coupling of piezoresistor sensor under ZnO and ZnO actuator. Another benefit is the increased doping reduced resistivity of the sensor, thus for a given stress level, the absolute change in resistivity increases. For actuation, ZnO is used to position the tip relative to substrate surface. Piezoresistor in this case is used as sensor as well as conductive path; it is thus necessary to separate the reading signal from signals for lithography.

An AC bridge can be used for the purpose of signal separation. Deflection sensing signals are usually of higher frequency (above 100 kHZ), while the electrical pulse for nanoindentation is typically of a frequency around several hundred HZ [105].
2.1.3 Cantilever Array Design

The proposed cantilever array will utilize piezoelectric materials for the purpose of sensing and actuation. Unlike in the case of capacitor sensor and piezoresistive sensor, where sensing and actuation are carried out by separate structures, the sensing and actuation function can be achieved by the same piezoelectric material, thus it is ideal for the purpose of integration. It has been demonstrated that perovskite structured materials usually have higher piezoelectric constants, and can be used for MEMS application with conventional planar processing techniques [106]. Although AlGaAs materials also offer high piezoelectric coefficient, the fabrication of such materials typically needs sputtering or MOCVD, which increases the cost of fabrication. On the other hand, the fabrication of perovskite materials can be achieved with simple method as spin coating, which does not add significant cost on the production. As such, perovskite structured material is chosen as actuator material for cantilevers.

Pb(Zr,Ti)O$_3$ is chosen for actuator fabrication. The fabrication process is modeled below. A single leg design of the cantilever beam is chosen; compared with the V shaped, single leg cantilever takes less space, so integration can achieve higher density. However, it cannot use the piezoresistive effect for sensing as it’s difficult to use single leg silicon as piezoresistor without short circuit. As such, an alternative is to measure the deflection of the PZT material itself by the electric potential it produces.

The proposed cantilever array must fulfill the requirement for cell testing and in the same time be compatible with fabrication technologies currently used in industries. The design of this device is inspired by the work of Itoh et al. [99].

For the adaption of this device to the testing of cells, the array must be able to accommodate the possible variations in cell size. As a result, a longer cantilever length is needed so that more deflection can happen over the whole cantilever range. And if the variation of cell size does occur at some position, DC voltage could be applied to specifically deflect the cantilevers at these positions. However, the use of long cantilever will produce a lower the first resonance frequency [107, 108], producing a lower sensing speed in tapping and non-contact mode. To counter this problem, short cantilevers could be used with actuators producing larger force, so that both large deflection and fast sensing can be achieved simultaneously.
The fabrication process of the device is modelled below:

A. A photoresist layer is deposited on Si substrate (yellow), and then holes corresponding to the base diameter of tip are made by exposing the resist. Cr deposition is carried out and followed by a lift off process, so that circular Cr patches (black) of the same diameter as the base of the tips are produced.

B. Reactive ion etching is carried out so that silicon columns are created at the positions of tips.

C. Thermal oxidation is carried out to change Si to SiO$_2$, and then the substrate is submerged with KOH which etch the columns into tips.

D. A layer of Pt (green) is deposited by sputtering on the desired location of substrate; the Pt layer will later be patterned into one set of electrodes.

E. The PZT layer (blue) is prepared by a Sol-gel process. To prepare Pb(Zr,Ti)O$_3$, zirconium-n-butoxide dissolved first in glacial acetic acid, 2-propanol, and 1-butano [109], subsequently, Ti(OC(CH$_3$)$_2$)$_4$ and Pb(CH$_3$COO)$_2$ are also dissolved in the solution. The mixture is hydrolyzed before cross link agent ethylene glycol is added. The solution is further dissolved by 2-propanol and 1-butanol and acetic acid before deposition. The method is derived from reference [110, 111].

F. To deposit the PZT material on substrate, spin coating is carried out followed by baking. Multiple spin coatings are necessary in order to achieve the desired thickness of the PZT material. After the last spin coating, the structure should be moved to an oven for annealing at 600 degrees to facilitate the formation of perovskite structure.

G. Another layer of Au (pink) is deposited on top of the PZT material, to function as the other set of electrodes.

H. The final step is to etch away undesired part by dry etching, and then a layer of SiO$_2$ is sputtered to passivate the device.

It is already shown this approach can be used for batch fabrication of up to 50 cantilevers. Batch fabrication of an array with higher number of cantilevers is possible [99, 112]. However, it is probably more favourable to use more such arrays to integrate into a higher level array, so that modular design can be achieved. In the case of cantilevers break down, it would not be necessary to replace the whole array, instead, only the sub array containing the broken cantilever needs to be replaced.
The device is operated by applying voltages between two electrodes (*green and pink layers*). DC voltage induces a static deflection while AC voltage induces dynamic actuation. A small amount of tip displacement can be applied on each cell and the corresponding force can be measured. To detect the responses of cells, the voltage generated by the bending of PZT between the two electrodes is monitored.

The group by Itoh showed cantilevers can be fabricated in this manner to achieve a resonance frequency about 64 kHz and actuation sensitivity of 150 nm/V. In addition, they also demonstrated their cantilevers can accommodate a sample variance at z direction as large as 1.5 μm [99] with the length of cantilever equal to 200 μm.
2.2 Cell Chip

Due to the employment of cantilever array, cells have to be precisely placed on substrate at corresponding positions to cantilever tips. In addition, substrate has to provide sufficient affinity so that cells will not be displaced by the force applied by cantilever tip. A cell chip is necessary to fulfil these requirements. It has been shown that small particles and cells can be precisely manipulated by vibration force (acoustic tweezers), hydrodynamic force (microfluidic tweezers) and optical force (laser tweezers) [113]. However, these methods require extensive labour if large quantity of cells are to be manipulated.

When coupled with other technology, recent advances in MEMS application provide simple methods for achieving precise positioning for a large quantity of cells. MEMS based positioning methods can be generally characterized into positioning by optical guiding, by mechanical fluidic trapping, by chemical modification and by the application of electric fields.

Optical guiding involves the use of laser, which can generate forces in piconewton range (up to 300 pN [114]) for manipulating small particles. The force is generated with a high power focused laser beam passing through the dielectric particle. When placed around the beam waist, the dielectric particle feels the greatest force due to the largest electric field gradient. It is also necessary to build an environment so that the light travels in a medium with refractory index different from that of the particle [115].

A high throughput laser positioning system involves the use of hollow optical fibre, where laser carries cells from source to the hollow optical fibre [116]. By the guiding of the optical fibre, cells can be deposited when they exit from the other end of the fibre. By moving the fibre, arbitrary patterns can be formed on the target. It has been demonstrated that this positioning technique can achieve a resolution under 1 µm at a very high particle fluxes up to 10 kHz [117]. However, some drawback of this technique hinders its application to biological samples, as moving cells require very high power, the concern of viability of single cells subjected to the laser arises [118].

Mechanical fluidic patterning uses patterned structures to disturb the flow of cells in the laminar flow of liquid and selectively trap cells at desired location. Examples of this kind of positioning include the use of dam/overflow structures [119] or sandbag
structures [120]. In dam structure, the flow is blocked by dam perpendicular to the flow direction, and cells are deposited at the foot of dam. The problem of this design is the difference of the hydrodynamic pressure across dam structure is too great, that multiple rows of cells can be deposited at some position, inducing undesirable cell aggregation at certain location. The modified overflow design uses two parallel channels with different hydrodynamic pressure; the hydrodynamic pressure difference can be controlled such that only a single row of cells is deposited.

Similarly, sandbag structures fabricated from photolithographic processes also partially block the flow in channel. Although majority of the liquid flows along the arrow (arrows shown in figure) direction, a small portion of fluid flow over the sandbag and bring cells to be deposited at the depicted positions (red circles). Other
designs uses similar principles may involve the use of microwells in channel. A two-step process has been reported by Langer [121] to use poly(ethylene glycol) (PEG) to fabricate wells within micro channels for accurate positioning.

Chemical modification is another widely used technique to confine cell in certain positions. Typical chemical modification involves the deposition of protein prior to the deposition of cells [122]. Many of these deposited proteins are extracted from ECM, like fibronectin (FN), collagen I, collagen II and Laminin [122, 123]. Proteins can be deposited through a photolithography process with lift off. Synthesized poly peptide can also be used for depositing. The common feature of this kind of poly peptide is that they contain a tail sequence of Arg-Gly-Asp (RGD) tri-amino acids [124]. It is believed RGD is one of the primary recognition sequences for integrin [125, 126], which is a transmembrane protein for regulating a host of cell responses including cell focal adhesion. Other materials, such as selectin, enzymes and antibody [127] can also be deposited on substrates to enhance cell adherence.

Another approach of chemical surface modification employs a technique named micro contact printing [128, 129]. Before printing, soft lithography is usually carried out to produce a stamp with polymers like PDMS. Self-assembled monolayers (SAMs) are printed with stamps onto substrate which is surface-modified by gold, silver or titanium deposition [128]. The stamp defines the positions for SAMs deposition. This printed pattern can form a ultra thin mask (10nm) for subsequent deposition of another material, or it could be used directly as created patterns.

\[
\begin{align*}
\text{CH}_3 & \quad (\text{EG})_n \\
\text{CH}_2=\text{CH}_2 & \quad \text{CONH}_2 \\
\text{CF}_3 & \quad \text{CO}_2\text{H} \\
\text{COCH}_3 & \quad \text{OH} \\
\text{CO}_2\text{CH}_3 & \quad \text{PO}_4^{2-} \\
\text{F}, \text{Cl}, \text{Br} & \quad \text{CN}
\end{align*}
\]

*Figure 15. Chemical compositions of SAM layer, X is the end group [124]*
Most widely used SAMs are alkanethiolates with different end groups. It is found SAMs terminated with methyl groups adsorb proteins while SAMs terminated with oligo (ethylene glycol) are extremely resistant for protein adsorption. Thus, SAMs can be used to link previously mentioned biomolecules with substrate. In addition, cells can directly attach to SAMs, it is found that the adherence strength of cells on SAMs layer is greatest with SAMs terminated with hydroxyl group, followed by SAMs terminated with carboxyl group and amino group, the adherence is the lowest with SAMs terminated with methyl group [124].

Figure 16. General approach for micro contact printing. Molding of PDMS soft stamp from patterned master (a), Peel off PDMS stamp (b), stamp in contact with SAM ink (c), PDMS print on substrate (d) SAM layer produced as mask (e), Deposition of a secondary SAM (f) [128]
Electric fields can also be used in active cell positioning. The technique called dielectrophoresis (DEP) has already been applied through decades in sorting and separating particles [130]. DEP force is a result of the interactions between induced dipole in dielectric particles and the non-uniform electric field in the medium [131]. The DEP force can be calculated as

\[ F_{\text{DEP}} = 2\pi r^3 \varepsilon_m \text{Re}[f_{cm}(\omega)]\nabla E_{\text{rms}}^2 \]

where \( f_{cm}(\omega) = \frac{\varepsilon_p - \varepsilon_m^*}{\varepsilon_p^* + 2\varepsilon_m^*} \)

In the equation, \( \varepsilon_p \) is the permittivity of particle, \( \varepsilon_m \) is the permittivity medium. And \( \varepsilon^* \) is the complex conjugate of permittivity. Both negative and positive DEP are possible, negative DEP occurs when \( f_{cm}(\omega) \) is less than 0, which means medium is more polarisable than particle. In this case, DEP can repulse particles or cells away to the region of local minimum of gradient of the electric field. When \( f_{cm}(\omega) \) is greater than 0, particle is more polarisable than medium, positive DEP occurs which tends to attract cells to the local maximum gradient of electric field. It has been demonstrated that positive DEP is more suitable for cell positioning because it has the capability to separate individual cells and is useful for cell registration [132].

The following figure shows one of such DEP device combined with light electrode for massive cell manipulation. A photodiode coupled with and digital micromirror display is used to generate illumination pattern, which is projected onto a highly photo sensitive substrate made of heavily doped hydrogenated silicon. The projected pattern on the photo sensitive material produces electron hole pair to counter the applied AC field thus produces a local minimum in electric field. Correspondingly, the particles will be repulsed by negative DEPs to local minimum positions as defined by the lighting patterns. Figure 15 shows an advantage of light electrode DEP system, which is the pattern for particle positioning, can be altered arbitrarily by choosing the desired lighting pattern without the need of any hardware change. Figure 15 up demonstrates a circular light electrode, and figure 15 below demonstrates a square electrode array.
Figure 17. Mechanism of DEP patterning with light electrode (up), light electrode pattern for from cell array (bottom) [131]
2.2.1 Cell Chip Design

Because of the application of AFM arrays for measuring single cell mechanics, it is desirable for the cell chip to allow the approaching and retrieval of AFM cantilevers without tedious alignment operations. As a consequence, microfluidic device trapping cells with complex geometries should be excluded from the design. In addition, complex geometry means that more structure is needed for trapping, and the trapping efficiency, i.e., trapped cells per unit area, will decrease, which is unfavourable since it contradicts with the aim of increasing throughput.

Moreover, positioning resolution is important because the use of AFM places high requirement on this criterion. AFM tip should be in close proximity with sample before experiments start since there is only limited range of adjustment from piezo controller on AFM. In all afore mentioned techniques, the mechanical fluidic positioning has the lowest resolution [118], in addition, in this kind of patterning, cell agglomeration usually happens [124], which can further complicate the single cell force measurement.

Chemical positioning is able to achieve very high resolution with photolithography and micro contact printing. Although this group of techniques are suitable for low cost laboratory experiment, they are not suitable for commercial products to be used in the high throughput device. One very important feature in chemical surface modification is to enable cell attachment through focal adhesion. This process is time consuming and may take up to 24 hours for cells to firmly attach on substrate [128]. Another problem with chemical surface modification is to immobilize ligand or biomolecules onto surface will inevitably increase the complication of device handling. For example, there may come to be a very limited device shelf life and the cryo-storage of the device may be necessary. In addition, due to the specific interactions between ligands and cells, each cell chip fabricated with chemical surface modification may only be suitable for a limited range of cells, thus multiple chips must be purchased by the end user if different cells are to be tested.

It is desirable to build the device based on the electric field approach from the considerations mentioned above. Because electric field generating structures (like wires and vias) are well suited for miniaturization and have been studied extensively in electronics and semiconductor industries. Tools for generating these structures are
readily available. Moreover, due to the fact most control interface uses electrical signals; chips operating based on electrical fields are more likely to be easily controlled and automated.

It has been demonstrated most cells have similar electrophoresis mobilities [124]; as such, DC electrophoresis is able to exert force on a wide range of cells. However, DC electrophoresis lacks the ability to separate different kind of cells. As mentioned before, DEP has been successfully applied in the fine positioning of cells [133, 134], and different cells have a wide variety of responses to DEP forces, thus it is more favourable to use DEP for cell chip design.

Some concern rises about the viability of cells subjected to electrical field. It has been confirmed by various reports [130-133, 135] that the viability of cells can be maintained after DEP operations under controlled field strength. When the electric field does not exceed $10^4$ V/m, no loss of viability could be observed even under prolonged exposure [136, 137], when the electric field reaches $5 \times 10^4$ V/m, temporary membrane electroporation occurs, but cells will reseal after dielectrophoresis [118]. When the field strength exceeds $2 \times 10^5$ V/m, irreversible damage may happen to cells. Another feature of DEP is high frequency operation typically alleviates the electric-mechanical stress on single cell, so DEP is usually operated at high frequencies above 1 MHz [118]. As such, it is possible to define a region for safe operation for maintaining cell viability.

It should be noted in DEP operation, low conductivity medium should be used for avoiding heat generation and maximize the permittivity difference between medium and cell. Furthermore, by using low special medium is also able to prevent electrolysis of substance usually presents in normal culture medium [118].

The design of the chip is inspired by the work of Ho and colleagues [138, 139], where DEP was used to pattern liver cells. The schematic drawing of the design is shown below in Figure 16.

The chip can be represented as composed of three functional units. Component A represents terminals for external wiring to power source. Component B represents the surface patterning for assisting cell positioning. Component C represents the wire structure for producing desired DEP pattern, the red and green lines represent metal...
wires connected to an AC power source. D represents the chip structure after fabrication, only one unit of wiring is shown here, multiple units of wiring with same structure can present on the chip by simply repeating the basic wiring unit. All the wiring units are connected to two terminals respectively as shown in D, so that when in operation, all red wires connected to the red terminal will have the same potential and all green wires connected to the green terminal will have the same potential. The design of the wiring pattern must simultaneously provide electric field gradient in both longitudinal and transverse directions, so that the local maximum of the electric field gradient can be confined to a point. As a result, wires running in both longitudinal and transverse directions are needed. So the design of the wiring structure employs a branching configuration, where wires of sublevel are branched off from the main wire trunk. When this chip is operated, cells will be attracted towards the local maximum of electric fields, which are located at the branching point coming out of each wire trunk [118, 138-141]. A simulation was carried out later to confirm that the local maximum of the gradient field was indeed at the desired positions (See Appendix A).

Component B is used to assist the trapping of the cell mechanically by providing pits at the local field maximum, the pit pattern is needed to minimize the exposure of cells to DEP field, so that when DEP field is turned off after cell positioning is finished, cells can still be retained at the decided positions. Another use of the pit pattern is to hold cells steady after positioning, when excess cells must be washed away by increasing fluid flow and tuning down DEP strength. The dimensions of the pits are controlled so that it can only contain one single cell. Paired with positive DEP’s ability to separate cells, it is possible that the design can achieve high resolution, single cell positioning with a high throughput. Because most cells have average size around 10 µm, the diameter of the pit should not exceed 10 µm for common use.

Special pit may be designed to facilitate the trapping of cells with extreme dimensions. Also, since the mechanical properties of cell changes when focal adhesion happens [81], different experiment protocols should be used if the cell mechanical properties under focal adhesion is to be measured.
It must also be noted in the designing of this chip, it is assumed that the chip will be used with compatible AFM cantilever array, so that the positions of pits and the spacing between the pits corresponds to the specific AFM cantilever design.

Figure 18. Schematic of cell chip design. Wire bonding for power supply (A), SU-8 layer with pits (B), wire pattern on glass (C), Assemble of the chip (D)

Figure 19 illustrates a possible approach to fabricate such a chip. Transparent materials are preferred for fabrication because they allow the examining of cell patterning result under optical microscope. Another consideration is materials in contact with cells must at least be biocompatible. For the purpose of allowing optical
examination, glass is chosen as a preferred substrate. Commercially available glass wafer can be purchased with thickness ranging from 300 µm to 1 mm. for subsequent fabrication, glass surface is cleaned with oxygen plasma first and a photo resist layer is deposited by spin coating (B). After the photoresist has been cured, a photo mask can be used to generate designed pattern for metal line deposition (C). Metal like copper, gold and titanium can be deposited on glass by chemical vapour deposition [142] (D), then a lift off process is used to produce the final pattern of metal wiring (E). For creating the pit pattern, different biocompatible materials can be used. The most commonly used biocompatible polymer in MEMS is PDMS, it is also transparent. However, PDMS has a elastic modulus [143] that is very close to the elastic modulus of cells, which can be as high as 200 kPa [59, 144]. Thus, probing cell mechanical properties on PDMS may introduce error to the measurement. Other materials can be used include PMMA and PS, However, both PMMA and PS need contact printing to form the pits pattern, and additional adhesive (like epoxy) may need to be applied between PMMA/PS and glass.

Figure 19. Fabrication process of cell chip
For the purpose of easy fabrication, SU-8 is chosen to be used to form the pit pattern. SU-8 is biocompatible polymer that is commonly used as negative photoresist in UV range [145-147]. SU-8 offers high elastic modulus after being cured up to 5 GPa [148, 149], sufficient for the purpose of cell mechanics testing. SU-8 is spin-coated on glass substrate after the glass substrate is cleaned and dehydrated (F). The resist is baked and then pressed under heating conditions for bonding to glass wafer. The final step is to expose SU-8 layer to a pit pattern printed on photo mask film to form the pit pattern.

2.3 Integration and Control

With the design of AFM array and cell chip, the system can be integrated together. As mentioned before, a modular design is preferred in order to lower down the maintenance cost. So even if one module fails, it is possible to change that module only without the need of replacing the whole set. A schematic component diagram of the system is shown below. An array of four AFM sub arrays are shown on the graph, they are integrated into one higher level array. The blue platform represents cell chip mentioned in the previous part, while the area encircled under each cantilever array shows the area probed by that array. The cell chip can be mounted on a piezo actuator which can move the chip in 3 dimensions.

The signals pathways are shown in single line (input) and double line (output). The signal processing system is modeled after commercially available controlling systems from National Instrument®. Analogue signals can be read out by data acquisition board (DAQ), which can also convert analogue input into digital signals. DAQ board then send signals to I/O boards for interfacing with computer. Desired stimulation form can be designed in computer and output to I/O board, which in turn control the power source for actuating cantilever arrays. The data acquired by this control system can be stored in the computer system for further analysis.
For data analysis, fast Fourier transform can be carried out based on sampled force data to generate complex modulus and phase angle [105]. Cells are modeled as viscoelastic homogenous spherical balls indented with small and infinite hard indenter [150]. Hertzian contact model and Maxwell-Weichert are coupled together for dynamic modeling. Most AFM based systems use Hertzian contact model based on a spherical tip to estimate the complex modulus of single cells. Revised Hertzian model can be applied to account for different tip geometry [151]. The real part of the complex modulus represents the storage of elastic energy, while the imaginary part represents the loss of the energy. The relationship between complex modulus, force and frequency are given by [105],

\[
|E_{\text{spherical}}^*(f)|= \frac{1-\nu^2}{2\sqrt{Rd_0}} \frac{F(f)}{d(f)}
\]

where \(d_0\) is the indentation depth at 0 frequency, \(\nu\) is the Poisson’s ratio and is assumed to be 0.4 for chondrocytes [152], and is specific to cell phenotype. \(R\) is the effective radius of contact.
A more accurate modeling of the force measurement takes into account of the geometry of the tip, for a pyramidal tip,

\[
|E^*_{pyramidal}(f)| = \frac{1 - \nu^2}{2\pi d_\rho \tan \alpha} \frac{F(f)}{d(f)}
\]

Where \( \tan \alpha \) defines the half angle of tip apex. The real part of the complex modulus signifies the energy storage part of dynamic responses, while the imaginary part signifies the energy loss part of dynamic responses corresponding to viscous flow.

Because of the utilization of continuum mechanics, the modeling of single cell mechanics is relatively simplified. It ignored the inherited heterogeneous nature of cell, which is composed of many microstructures. There have been experiments showing that the stress distribution on a single cell is far from homogenous, and tends to extend through cells [153] rather than to be localized as predicted by the models based on homogenous models [53, 154, 155].

In the model of cell mechanics of tensegrity, the discrete nature has already been taken into account. It has been demonstrated that the modelling of single cells as tensegrity mimic a number of mechanical behaviors or properties of cells. For example, this model predicts that the focal adhesion of cells will be accompanied by the malleable deformation of cytoplasm and the nucleus will polarize and move basally. This has been proved experimentally by the studying of living cells [156]. In addition, it has been shown by time lapse video microscope observation that the cytoskeleton network rearrange globally just as the tensegrity predicted [157]. The experiments on the focal adhesion of cell to ECM also suggest cytoskeleton is a tensegrity network because the compressional continuity was not observed in either ECM or cytoskeleton [156].

It is thus interesting to seek further improvement on the accuracy of measurement interpretation by incorporating tensegrity model in the mathematical modelling of single cell mechanics. A few models [158, 159] have been devised to take into account of tensegrity to model the dynamic responses of cells. By using these models, the internal mechanical loadings on tensegrity elements like bars, which correspond to microfilaments, and cables, which correspond to cytoplasm, can be derived from external loadings; it may be useful to provide a way to directly estimate the
cytoskeleton responses to external loadings. And furthering development of the software algorithm to interpret the mechanical measurements may lead improved performance of the system.

2.4 Challenge and Other Applications

There may be still some technological challenges faced by this technology for the realization of its full potential. One of which is to ensure the uniformity of the substrate to be probed. Due to the application of AFM platform, it is extremely important to make sure that the sample variation in Z direction can be accommodated by the cantilever system.

As a result, cells must be preselected by their size; a size variation range within the tolerance limits of the cantilever arrays must be specified for the purpose of cantilever array protection. There are some existing technologies like flow cytometry for cell size selection. Flow cytometry is a high throughput device capable of cell selection based on cell type, cell size or specific markers attached on cells. Although the selection of cell size is achievable with flow cytometry, it requires preparation of cell staining by fluorescent probes, thus increasing the experiment time frame [160-162]. Another problem of using flow cytometry is this could increase the cost of the system.

Another solution is to use microfluidic MEMS device for cell size sorting. The sorting mechanism can be based on mechanical size selection by forcing cells passing pores in the device [163]; or can be based on purely fluidic mechanics principles [164, 165]; or can be based on the same principle of flow cytometry [166, 167]. Although these cell sorting chips partially overcome the cost problem of conventional flow cytometry, they have their own troubles. For example, cell sorting devices based on pore size does not consider the deformability of cells, though it may not be a major concern for normal cells, it could potentially fail to sort cells with high deformability. Microfluidic devices on the other hand, do not have the same high sorting yield in conventional flow cytometry. Cell sorting based on light scattering principle like conventional flow cytometry can produce very high sorting yield, however, they still require the pre-stain of cell population.
Although the concern of cell size variation can be alleviated by the use of cell sorting devices, there is still deeper concern about the small size variation tolerance of AFM array. Even if cell sorting does not require time consuming staining process, it may still be a problem as only a fraction of cells can be selected from the total cell population; this could cause huge waste, not only of the materials needed for cell culture, but also of the cell source itself. This problem is more severe especially when precious cells such as stem cells are used.

Another problem associated with the selection of the fraction of cells bears the question whether this sub population could be a good enough sample to represent the whole population of cells. Single cell mechanics may vary with cell size; so that multiple batches of measurements have to be carried out for characterize the whole population.

A fundamental solution to this problem is to fabricate AFM arrays with higher tolerance. Less stiff materials [85, 168] could be used for the fabrication of array. Future research would be needed to solve the problem.

The proposed system has good application potentials to a variety of tasks. Although this technology platform is developed with the primary aim for high throughput probing of single cell mechanics by the nanoindentation of AFM tips, it is possible to modify this set up for other researches. With normal cell chip, cell response to dynamic compression can be tested, with AFM tips functionalized with particles that are large in size, whole cell mechanical property measurement can be obtained. If chemical agents are used to functionalize cell chip for cell binding, the tensile properties of single cells can also be tested.

AFM has found wide range of applications in pharmaceutical industries on molecular basis. For example, by chemically modifying the tip of AFM, it is possible to detect the binding forces between adhesive protein and the ligand of this protein on cell, thus providing a quantitative assessment of protein-cell interaction. The same principle can be applied in studying drug-cell interactions too [64]. In addition, due to the high resolution provided by AFM, force measurement on molecular level is possible, there are experiments performed on force measurement between single protein-ligand pair [169], DNA double strands [170] and single molecule domains [171]. With current design of the high throughput system, these molecules can be immobilized on
cantilever tips as well as in the pits, thus high throughput of single molecule mechanics is possible.

Beyond the measurement of mechanical properties, there are other applications of AFM in biological sciences; the flowing figure summarizes the application fields, and many of them can be benefited with the development of the high-throughput AFM.

![Figure 21. Possible applications of AFM technology in biology](image)

2.4.1 Beyond Biology

More applications of AFM can be found beyond the boundary of biological researches, and the high throughput platform is also desirable in those applications. Active AFMs with self actuations have been intensively researched in material science community for the use on surface patterning. For example, AFM can be used to mechanically patterning the polymer surface by indentation or plowing [96]. This was also the original idea of IBM’s millipede, which performs thermal mechanical indentation for
data storage [173]. Another patterning technique biases cantilever tip with regards to substrate surface. The oxidation is confined to the proximity of AFM tip, and it has been demonstrated on silicon[174], III-V compound semiconductors, silicon carbide and metal [175]. Furthermore, AFM arrays can be used to transfer molecules as ink onto substrate surfaces [176], which is known as dip pen nanolithography (DPN) [177]. This technology can help to directly modify substrate surfaces and to fabricate ultra thin mask on substrate, and is compatible with a wide variety of substrates, including organic and biological materials [112].

However, the probe based surface patterning faces similar dilemma as probe based single cell probing, as the low throughput of the device hinders the further adaption of this technology. This high-throughput AFM cantilever array may find potential of being applied in surface patterning as well. In fact, many researchers have already begun to seek the application of array technology. For example, C.F. Quate in Stanford University integrated four cantilever arrays, each consisting 50 cantilevers for with independent control for each beam in local oxidation lithography [112].
Chapter 3 Commercialization

3.1 Intellectual Property

Intellectual property (IP) assessment is an important component to assess the marketability of a new born technology. It can help to protect the IP of the new technology and at the same time to avoid any confliction of IP interest with established patent holders. The first part of this chapter focuses on the studying of existing IPs.

Patent searches were conducted to analyze IPs with two primary sources, namely, Google Patent Search (www.google.com/patents) and the US Patent Office's (www.uspto.gov) Patent Search. There are numerous existing patents in the fabrication and application of cantilever array as well as the cell chip. Most relevant patents about the fabrication process have already expired so that no cost is incurred by applying these technologies in the fabrication process. For example, to fabricate the AFM array and the cell chip, Sol-gel deposition, sputtering, photolithography and CVD are needed, the patents about sol-gel deposition of PZT materials with spin coating were filed since 1989 [178], the earliest patent about photolithography using SU-8 dates back to 1974, and the most relevant one was filed by IBM in 1989 [179]. The sputtering deposition was filed before 1950 [180] and CVD of metal was filed back to 1980s [181].

There are about 390 patents regarding the technology field of arrayed AFM filed between 1988 and 2010, suggesting this is a rather interesting research territory. These patents mostly use different design of cantilever systems, for example, the patent for multiple probe measurement uses mechanically coupled probes for testing [182]. Furthermore, many patents of multiple cantilevers centre around concept such as data storage [183]. The most relevant patents in this category are Atomic force microscope for biological specimens [184] and Tapping atomic force microscope [185]. These patents are about the applications of a single probe. There are about 80 patents regarding the application of arrayed AFM to biological study, however, most
of these patents concern about biomolecules testing. No patent exists about the application of AFM array for single cell mechanics testing.

There are 127 patents about cell chip for cell positioning. There are patents about cell positioning with mechanical means [186] as well as with dielectrophoresis means [187]. However, no patent was found to combine both techniques.

The patterns of the filed patents in related fields indicate that this high throughput device is entering a technology territory that is less explored. This new technology introduces novel applications of AFM arrays testing single cell mechanics with a novel cell chip design; the salient features of this system are not yet patented. In summary, no severe barriers for the new technology were found and the technology should be patentable.

In summary, the technology platform represents a collection of novelties which should be patentable in the following areas:

a) Design and manufacturing procedures of piezo AFM arrays  
b) Design and manufacturing procedures of cell chip for cell positioning  
c) Application of piezo AFM array and cell chip for single cell mechanics testing  
d) Measurement and controlling algorithms

However, it must be noted that there are a range of patents and publications in area a) already, it is most advisable to improve on the current design and manufacturing processes before the patents are files.

3.2 Market

With the study of single cell mechanics gaining more attention, devices for quantify single cell mechanical properties will face a growing demand. In addition, pharmaceutical companies and administrative agencies involved in the developing tissue engineering products need a tool that can assess the mechanical properties of cells from engineered tissue. Other potential applications are also accessible by some slight modification of this platform. Presently, there is no commercially available solution which is of high efficiency and in the meantime is a potentially multitask platform. The cantilever array based cell force microscope is designed to fulfil this
void by targeting at this high demand, price inelastic market. Great market opportunity could be expected for this product.

The market analysis of AFM with arrayed cantilevers is based on evaluations of current market for AFM and a case study. The data of AFM market research was drawn from ITRS, Frost & Sullivan [188, 189]. The material for case study was drawn from the financial report form 10K of the leading company (Veeco) in AFM market. It is also noted that the market research typically group AFM with closely related technology and products of Surface Tunnelling Microscope (STM) and Near Field Scanning Optical Microscope (NSOM).

The primary source for AFM market data is from World Microscopes Markets [189]. The AFM market was estimated about 572.3 million in 2008 [190], with a projected growth rate of 15.3% annually till 2013. The competition structure of the market comprises three tiers. The first tier is occupied by large companies like Veeco and JEOL, which offer whole range of product line and have a broad installation base. The second tier comprises smaller companies like Park System and JPK Instruments, which are typically privately owned, and exploit niche market. The third tier comprises companies produce various AFM accessories. The market strategy for our start up company is to enter second tier first, because small scale production can help the start up company to reduce the need for huge initial investment, and in the mean time to concentrate to the specialization it possesses. Further expansion to first tier is desired after the first phase of market entering is successful.

The key end user group of the market is segmented into general industry, including semiconductor industry, electronics industry and materials industry, biomedical companies and research institutions/universities. Among all customers, biomedical companies and research institutions hold about 50% of market share, which is the primary target of the proposed technology platform. The gross margin of AFM products is typically about 40% [191], which is also high enough to motivate large initial investment and subsequent R & D.

AFM market exists globally, the market with Asian pacific as the number one market which holds about 50% of market value. North America and EU hold the second and the third place.
From the global market division, it is necessary to set up the start up company in a way such that global operation is possible, so that the efficient capturing of the global market can be achieved.

3.3 Case Study

A case study on Veeco was carried out in order to better understand the environment faced by a strong company in AFM market. It is noted Veeco has several departments, only those related to AFM were used for the purpose of analysis.

Veeco offers a whole range of AFM lines, from simple low cost set up to more complicated systems for industrial solutions [44, 192]. In 2009, Veeco launched 5 new AFM models and placed 450 AFMs worldwide [191]. The operation of Veeco’s AFM unit has been greatly affected by the world economy downturn, but the projection is still optimistic as there is an increasing trend in requirement of devices capable of measuring material properties at nanoscale level. It may indicate that we may have relatively stable revenue regarding to the economy situations.
The sales of metrology product consists of about 30% of total net sales of Veeco, and the gross profit (40%) remains flat despite the economy situation. The majority of sales and procurement of Veeco relies primarily on several partners, which may impose risks on the supply chain.

Veeco spent about 15% of revenue in research and development; this may be due to the high speed of technological evolution. We recognize it is necessary for us to invest in R&D too if we want to keep up with competition.

In terms of revenue realization, AFM customers generally require no substantial provisional period and the installation is insignificant. As such, revenue is realized soon after the device is shipped. It may be beneficial for a start up company to minimize inventory cost and maximize cash flow.

3.4 Business Strategy

For the realization of the high-throughput AFM, two approaches may be preceded. The first approach is to align with existing market players, by utilizing the intellectual property to gain profit. The other approach would be set up a company to produce the technology platform. From the market analysis above, it is clear that the market is growth is rather strong in AFM sector, and the market capacity is large enough to hold both large and smaller companies. In this market, smaller companies may excel in certain areas by providing specific product to tailor the needs of specific customer, because the market provides a high profit margin for the device manufacturer. In addition, due to the protection of IP, these specific markets is not likely to be engulfed by the larger companies if the smaller companies are able to hold a comprehensive collection of patents. Meanwhile, profit generation from IP could not provide a gross margin as high as producing the product by one’s own, since in the profit chain model, a company producing the product will be able to reap margins from the bottom of the chain, which is R & D, until top of the chain, which are final product sales. Moreover, this technology break through has potentials to be applied in various fields in research and industry as illustrated in previous section; this represent a great opportunity for a company holding this technology to expand into various market. In summary of all above considerations, it is more likely that to build a start up company will eventually
provide more return on investment than solely relying on IP royalty; as such, the following section will discuss a few business aspects of starting a new company.

3.4.1 Supply Chain Positioning

To position a start-up company into supply chain is vital for defining its business model. The key of supply chain positioning is to define input and output of the business. From the market structure analysis above, it would be best if a start-up company can avoid direct competition with established market players. It is thus highly desirable if the start-up company sells only the core functional unit (AFM array and controller) as an expandable module to customer with existing AFM system, so as to minimize competition by helping the customers to avoid high capital investment and establishing partnership with current market players.

However, it is soon realized that this approach leads to the problem of over complicated product design, as there are more than 14 major AFM manufacturers and about 50 companies in total [193] in the market, it would be extremely difficult to provide compatible AFM arrays and controller modules across brands due to different interfaces used by various manufacturer.

As a result, it is more favourable to make the final product to be the whole AFM system with controller and software. The details of the product are summarized below.

Product components:

- Cantilever arrays: to probe single cell mechanics. A number of cantilever arrays can be integrated in order to achieve higher throughput.
- Cell chip: for precise positioning of cells before cantilever probing is carried out.
- Piezo platform: to adjust the relative position of cell chip and cantilever arrays
- Controllers: including DAQ board to sample the signals generated by cantilever arrays and I/O board convert those signals to digital input. Digital output is also received by I/O board from computer and converted to analogue output by the controller in order to control the power source.
- Software: software modulus for controlling the hardware and for interpreting the acquired data
- Power source

Companies producing AFM instruments seek to play in more than one sector for healthy business growth. The start up company will enter the rapidly developing industry as a device fabrication enterprise with strong technology and development background.

The supply chain for the start up company is illustrated in above figure. There are three subsectors in the supply chain, namely, the supplier side which in charge of providing raw materials and the controller modules; the start up company which is responsible for manufacturing of the cantilever array and cell chip, the design of the controlling algorithms and analysis software and R&D for product development; and distribution channel, which is responsible for providing customers with the final product. The fabrication of the controller circuits can be outsourced to more experienced companies who are in possess of required tools and are specialized in fabrication of chips the in order to decrease the investment and reduce cost. Several important factors have to be considered for the building of the supply chain:

- To secure the timely delivery of high quality raw materials
To ensure timely delivery of products to customers by partnering with established distributors
To build up a competitive price for product with technological edge

3.4.2 Cost Modelling

The cost assessment is conducted based on the estimations of fixed cost and variable cost. The cost figure could provide a guide on the continual optimization in cost, reliability and fabrication technique in the future. The actual cost analysis should be done in more details and consider more factors.

Two approaches were used for cost modelling. It is worth noticing that only crude estimation can be made before the actual process of production.

The fixed cost was estimated with a top-down approach with data gathered from Veeco’s financial report. A total of 16.6 million worth equipments, plants and properties has been acquired by Veeco for the production of AFM, and there annual production is 450 AFMs.

To estimate the fixed cost of the new start up, an evaluation of annual production target has to be devised first. Assuming we can produce 90 AFMs per year, a very rough approximation of the total fixed cost would be 3.32 million. The fixed cost is then spread out over the production period of AFM device, so if it is assumed that a linear relationship exists between production volume and the fixed cost, the average fixed cost can be calculated from the production lifetime.

The core fabrication process of this device is the production of AFM array and cell chip. The cost model is based on laboratory techniques for producing the system, the dimensions of materials used are estimated from literatures [99, 138-141]. However, it is worth noticing that this is also a rough estimation only since the cost in laboratory settings does not always reflect the cost in mass production.

The cost of materials and processes were obtained from various sources [194-207], CES Edupack ® was also used to facilitate the estimation of cost and processing. The variable cost was estimated to be 960 USD per array, with labour cost being the single
largest cost for device fabrication (52%). More detailed production cost breakdown can be referred to Appendix B.

![Variable cost in array fabrication](image)

![Variable cost in chip fabrication](image)

*Figure 24. Variable cost modelling for Piezo deposition and SU-8 coating*

The variable cost of the chip production is based on the estimation of chip size. An estimation of 1cm ×1cm is used; cost of producing chips with different size doesn’t vary dramatically with the size of chip, as size dependent component (material cost) only account for 27% of the total cost. Among the material cost, the cost of photoresist is almost negligible. For batch fabrication, labour cost can be averaged out
among the units produced; if 10 chips are fabricated per batch without the need of increasing the number of workers, one chip is estimated to cost below 500 dollars.

The cost of piezo scanner (USD 1563) was obtained from direct quotes, and the cost of controller (USD 5000) was estimated from the price of controller unit from National Instrument (NI-PXIE 8130). The variable cost of the software can be ignored as it does not require too much resource to produce once the algorithm is finished. In summary, the total cost of building this system may come to USD 11K (variable) plus fixed cost if 4 arrays are used with one chip, one pizeo scanner one DAQ and one I/O controller. The fixed cost can be averaged out through production time. If the fixed cost is comparable to other manufacturer, and minus the cost of piezo scanner which is common to all AFMs, the cost may be USD 9.5K (4 arrays + 1 chip + controller) higher than the cost of producing a normal AFM set.

For a start up company, to reduce risk, it is desirable to limit the investment to a small scale production first. If we assume a linear relationship between production volume and the fixed cost, and the production volume is 150 AFMs per year, from Veeco’s financial report, the unit cost variation with respect to production volume can be estimated. If a production time span of 5 years is assumed, the price of the high throughput device must be at least USD 18.37K for break even.

![Figure 25. Unit cost change over production volume](image)
3.4.3 Utility and Feasibility Analysis

From the product cost point of view, the device may cost up to USD 9.5K more than average AFM cost. Thus there could be some doubt whether this product would be accepted by the targeted market. Average AFM cost about USD 15K to 40K depends on the complexity, if the targeted market is the high end AFM users, the high throughput device may cost 25% more than conventional technology.

The major limitation of AFM is the low throughput problem. There is currently no competing technologies available in the market offering the same degree of automation, resolution and throughput in a single product. If four AFM cantilever arrays are supplied in one package, each array consists 50 cantilevers as used by Quate [112], the throughput will be 200 times of the throughput of a single probe AFM. Thus the benefit is quite likely to be more than the cost increase.

Through the case study, it is found AFM profit can be as high as 40%, and can remain flat even in economy down turn. It is possible for the start up company to lower down the gross margin in the first place to capture the market. Through the market analysis, it is clear that AFM market is expanding at a rate about 15% per year, and AFM for biomechanical research represents a major contribution of the market expansion. Moreover, this high-throughput device has the potential to be a multitask platform so its potential application is not just limited by the field biomechanical research. The market expandability for the start up company could be high due to both market expansion as well as technology branching out.

There are also various ways for lowering down the cost. Considering the major variable cost is down to the controller units, it is possible that the cost of the product can be lowered by establishing direct partnership with controller providers. Other cost reducing methods may include the optimization of fabrication processes and the renting of equipment, plant instead of purchasing.
Chapter 4 Conclusion

Single cell mechanics has emerged as an important branch in the studying cell biology. Although research has been focused on the characterization of single cell mechanical properties based on a small population of cells, it is recognized this approach cannot truly account the complexity of heterogeneous cell population in most living organisms. Thus for the purpose of quantitative assessment of a large population of cell, a force measurement device with high resolution and high throughput is necessary.

The first part of thesis is concentrated on the biological back ground which connects single cell mechanics to various cellular activities. A deep correlation between single cell mechanic properties and cytoskeletal structures was explained, the application of single cell mechanics was shown including the characterization of cells and identification diseases.

In the second part, a system composited novel AFM cantilever array and cell is proposed and studied. Different AFM cantilever array technology and cell chip technology are compared and suitable technology was chosen for the fabrication of the device. It is found that the fabrication of the device does not contribute much difficulty for the realization of the technology. Although the technology may still face some challenge to realize its full potential, these challenges are not fatal for the technology and they can be overcome by the use of additional device or the advance of complementary technologies, such as cell sorting chip.

Feasibility study based on commercialization of the concepts was also carried out, through the analysis of intellectual property; it is shown that this technology would be patentable as it combines novel features of both AFM patents and cell chip patents. Market analysis was also carried out to assess environment faced by companies in this sector. Through the analysis of business strategy, the author found strong market growth and good IP protection could facilitate the start up and the expansion of the company, meanwhile to set up a start up company for producing such technology platform may generate more return for investment. As a result, it seems more
profitable to realize the technology through joining the device fabrication industry rather than through the collecting IP royalties. Although the fabrication cost was would be higher for such a system compared to common AFM, it is possible that the utilities provided by this technology overweighs its expenses. Further cost reduction may come from the expansion in production scale and advancing in manufacturing processes.

In conclusion, this technology represents advancement in the development force measuring devices, because no competing device with equivalent precision and throughput is offered in current market, this technology can receive positive market responses.
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"6.777J/2.751J Material Property Database."


[196] "ethylene glycol cost."
[197] MTI, "Cost of thermal oxide wafer."
[203] Y. E. K. S. A. Gridnev, A. V. Kalgin, E. S. Grigor, "Magnetoelctrical effect in layered composites PbZr0.53Ti0.47O3-Mn0.42Zn0.6Fe2O4," Protection of Metals and Physical Chemistry of Surfaces, vol. 45, 2009.
[205] "Acetic acid cost."
[206] "Butanol Cost."
[207] C. E. Package, "process coast."
Appendix A: Calculation of the Gradient of Electric Field in Cell Chip

The electric field strength is calculated by assuming the structure is assembled with infinite number of point charges.

For vertical wire trunk, if the length of the wire is l, the field strength can be calculated as:

\[
E_i = \int_{lower}^{upper} \frac{kQ}{x^2 + (l - y)^2} \, dl = \int_{lower}^{upper} \frac{kQ}{x^2 + (l - y)^2} \, d(l - y)
\]

\[
= kQ \frac{1}{x} \tan^{-1} \left( \frac{l - y}{x} \right)
\]

For each wiring unit, there are two vertical wire trunks, the electric field from the other one is calculated as:

\[
E_2 = \int_{lower}^{upper} \frac{-kQ}{(right - x)^2 + (l - y)^2} \, dl = \int_{lower}^{upper} \frac{-kQ}{(right - x)^2 + (l - y)^2} \, d(l - y)
\]

\[
= -kQ \frac{1}{right - x} \tan^{-1} \left( \frac{l - y}{right - x} \right)
\]

here upper represents the upper y-axis boundary of the wire trunk, lower represents the lower y-axis boundary of the wire trunk, left represent the x-axis position of first wire trunk, while right represents the position of the second wire trunk on x-axis.

For wire branches, if they are on the first wire trunk,

\[
E_{\text{branch}} = \int_{0}^{\text{length}} \frac{kQ}{[upper - (i - 1) \times \text{space} - y]^2 + (m - x)^2} \, dm
\]

\[
= kQ \frac{1}{[upper - (i - 1) \times \text{space} - y]} \tan^{-1} \left( \frac{m - x}{length - x} \right)_{b}^{a}
\]

\[
= kQ \frac{1}{[upper - (i - 1) \times \text{space} - y]} \left( \tan^{-1} \left( \frac{length - x}{upper - (i - 1) \times \text{space} - y} \right) - \tan^{-1} \left( \frac{-x}{upper - (i - 1) \times \text{space} - y} \right) \right)
\]
if they are on the second wire trunk,

\[
E_{2x} = \int_{right-length} -kQ \left[ upper - (i-1) \times space - y \right] \frac{1}{\tan^1 \left[ upper - (i-1) \times space - y \right]} d\left( m - x \right)
\]

Here space represents the interval between branching points; length represents the dimension of branching-out wire tips.

The gradient of electric field is calculated with Matlab.

A simulation is carried out by assuming two wire trunks present in one wiring unit, and each wire trunk has 11 branch-off points, each tip branching out has a length corresponding to 1/16 of the distance between two wire trunks. The simulation is preceded by assuming unit strength of the gradient of the electric field when the potentials of the wires reach the maximum points in AC voltage. The following figure shows a snapshot of the changing square of AC electric field gradient at certain moment. No significant change except the absolute values of the maximum will occur in other moments during device operation.
The simulation shows maximum gradient of electric fields can be achieved at the branching point, where it is desirable to build pits for assisting cell positioning.

The matlab codes used for simulation is listed below

```matlab
function A = Efield
upper = 100;
lower = -100;
left = 0;
right = 20;
number = 11;
space = (upper-lower)/(number-1);
i = (right - left -1)/1 + 1;
j = (upper - lower -1)/1 + 1;
A = zeros(j, i);
m =1; n =1;
for x = (left+0.5):1:(right-0.5);
    for y = (lower+0.5):1: (upper-0.5);
        A(m,n)= Ehorizontal(x,y, upper, lower, left, right, number) + Evertical(x,y, upper, lower, left, right);
        A(m,n) = A(m,n)^2;
        m = m + 1;
    end
end
```
m = 1;
n = n+1;
end
B = abs(gradient(A));
y = [(lower+0.5): 0.5 : (upper-0.5)];
x = [(left+0.5): 0.5 : (right-0.5)];
[Y X] = meshgrid(y, x);
mesh(B)

function E = Evertical(x, y, upper, lower, left, right)
E1 = 1/x*(atan((upper-y)/x)-atan((lower-y)/x));
E2 = -1/(right-x)*(atan((upper-y)/(right-x))-atan((lower-y)/(right-x)));
E = E1+E2;

function E = Ehorizontal(x, y, upper, lower, left, right, number)
space = (upper-lower)/(number-1);
length = (right-left)/16;
E1 = 0; E2 = 0;
for i = 1:1:number;
    E1 = E1 + 1/(upper-(i-1)*space-y)*(atan((length-x)/(upper-(i-1)*space-y)-atan((-x)/(upper-(i-1)*space-y))));
end
for i = 1:1:number;
    E2 = E2 - 1/(upper-(i-1)*space-y)*(atan((right-x)/(upper-(i-1)*space-y))-atan((right-length-x)/(upper-(i-1)*space-y)));
end
E = E1 + E2;
Appendix B: Data Used in Cost Modeling

It must be noted that the cost analysis here is just a rough estimation from various literature survey; actual process may have different cost. For fixed cost, the Form 10K from Veeco was used to assess the investment in plant, equipment as well as property. Veeco is the leading company in the field of manufacturing AFM. Estimation from Veeco gives an indication of the best performance one may be able to achieve with the invested resources because among all manufacturers, Veeco has the comparative advantage in obtaining the benefit of mass production. Veeco’s investment on plant, equipment and property is about 16.6 million in total, which gives a production volume of 450 AFMs a year. For the reason of simplicity, if a linear relationship between production volume and the fixed cost is assumed, and if the production life time is assumed to be 5 years, and all the equipment has to be upgraded after that, there will be an average fixed cost on each machine for USD 7377.

For variable cost, the cost of manufacturing the cell chip and the AFM array was separately calculated. The variable cost was estimated based on the processes used in the fabrication. For the reason of estimation, the processes were divided into common processes and unique processes, where common processes refer to the processes which do not have distinguishing characteristics related product produced, for example, etching, masking and MOCVD metal deposition are considered common processes because these processes have similar costs across the production of a range of devices. The cost of common processes can be conveniently estimated by modeling software such as Edupack© from GRANTA Intelligence. The unique processes refer to the processes specifically designed for the fabrication of the proposed technological platform; these processed include piezo material spin coating. Energy consumption is estimated based on process time and equipment power, labour cost is estimated with hourly rate set to USD 20 per hour. The following table and flow chart present data used in assessing the variable cost in the unique process.
Cr deposition as mask on Silicon wafer
13 um Cr, RIE for 7 um

KOH immersion
30 hrs CES
Thermal oxidation

Zirconium-n-butoxide dissolve

Dissolve Ti{OCH(CH\_3)\_2}\_4

Dissolve Pb(CH\_3COO)\_2

Dissolve ethylene glycol

Dissolve 2-propanol

Dissolve butanol

8 X Spin coating

110 5min drying
600 20 min baking 8 times

1.2 um thick layer
Anneal at 600 4 hours

Au/Cr deposition 0.2 um

SiO\_2 passivation, 0.2 um

KOH etching

Ar ion beam etching

<table>
<thead>
<tr>
<th>Materials</th>
<th>Unit cost</th>
<th>Amount</th>
<th>cost</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zr(C_4H_9O_4)_4</td>
<td>USD 22/kg</td>
<td>0.00065 kg</td>
<td>USD 0.014</td>
</tr>
<tr>
<td>Ti{OCH(CH_3)_2}_4</td>
<td>USD 618.6/kg</td>
<td>0.00028 kg</td>
<td>USD 0.175</td>
</tr>
<tr>
<td>Pb(CH_3COO)_2</td>
<td>USD 110/kg</td>
<td>0.00067 kg</td>
<td>USD 0.073</td>
</tr>
<tr>
<td>ethylene glycol</td>
<td>USD 12/L</td>
<td>&lt;1L</td>
<td>USD 12</td>
</tr>
<tr>
<td>acetic acid</td>
<td>USD 50</td>
<td>&lt;1L</td>
<td>USD 50</td>
</tr>
<tr>
<td>2-propanol</td>
<td>USD 45</td>
<td>&lt;1L</td>
<td>USD 45</td>
</tr>
<tr>
<td>butanol</td>
<td>USD 2</td>
<td>&lt;1L</td>
<td>USD 2</td>
</tr>
<tr>
<td>Wafer</td>
<td>USD 80</td>
<td>1</td>
<td>USD 80</td>
</tr>
<tr>
<td>SU-8</td>
<td>USD 600/L</td>
<td>0.002L</td>
<td>USD 1.2</td>
</tr>
<tr>
<td>Glass wafer</td>
<td>USD 80</td>
<td>1</td>
<td>USD 80</td>
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