

Sun. Feb 5th

- The purpose of this experiment is to get familiar with the STM by attempting to take atomic resolution images of Graphite and Gold

- Preliminary Reading: Manual for Burleigh STM
 - Chapters 1, 2, 4

- Notes on Chapter 1: History of microscopy

Optical Microscopy

- Magnifying lens w/ resolution given by
- $$\frac{\lambda}{2n}$$
- $\lambda \leftarrow$ wavelength of light
 $n \leftarrow$ index of refraction of lens
 $\lambda \rightarrow$ ~0.1 microns

Electron Microscopy

- De Broglie wavelength of electron
- $$\lambda = \frac{h}{p}$$
- $h \leftarrow$ plank's constant
 $p \leftarrow$ electron momentum

- Electron can be accelerated to momentum

$$p = \sqrt{2mV}$$

through potential difference V

• For $V = 10^3$ Volts, $\lambda < 1\text{ \AA} = 0.1\text{ nm}$

- Shorter wavelengths of light encounter problems?

I don't really understand this part (p. 13)

- Transmission electron microscope

- Useful for study of regular lattices

- TEM requires a vacuum to avoid electron scattering from air

- TEM analogous to optical microscope

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- Scanning Electron Microscope (SEM) images surfaces by analyzing intensity of reflected electron beam and secondary electrons produced by incident beam
- An electron beam is scanned across surface
- Resolution 50-100 Å
- Non-conducting materials must be coated in gold
- Also requires vacuum

Acoustic Microscopy

- Can be used to image the inner structures of opaque materials

Scanning Probe Microscopy

- Prior are examples of wave microscopy
- Wave interaction w/ medium which affects index of refraction (wave speed)
- There is also probe microscopy. A point source interacts w/ sample to gather information
- Probe microscopy images 3D surface instead of inner structure of sample
- Different kind of probe tip interactions for different kind of microscopes
- The first was a scanning profilometer which measured deviations of a stylus as it scanned the surface

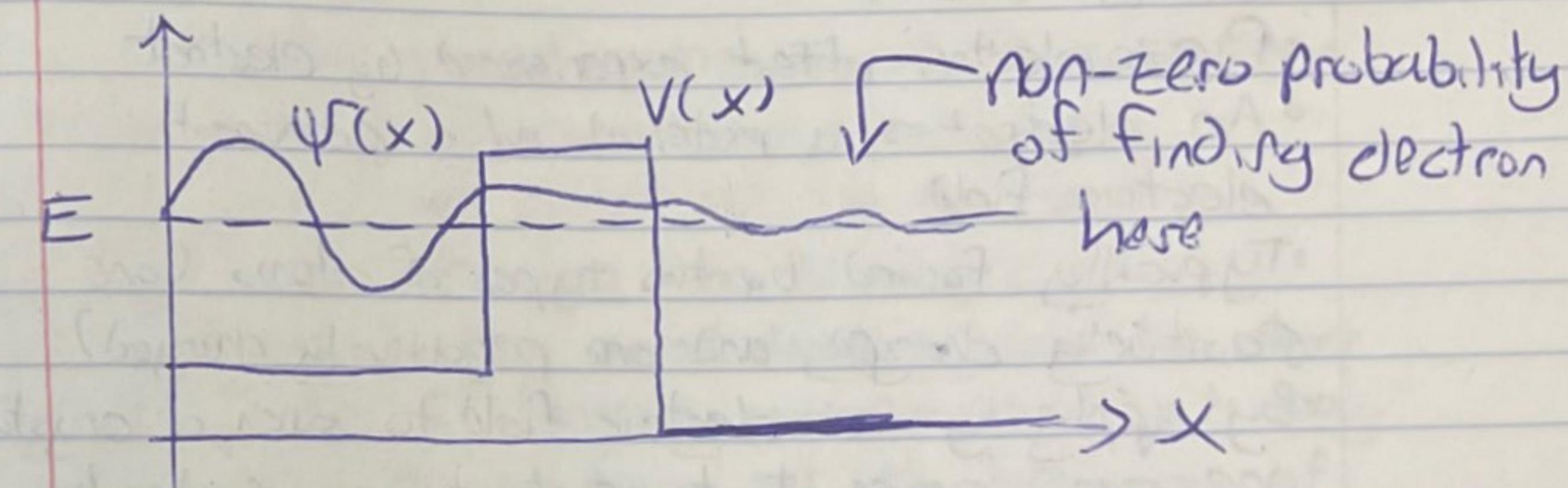
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Notes on Chapter 2: Operating Principles of STM

- 5 key concepts:
 - 1) Quantum Tunnelling
 - 2) Controlled motion over small distances using PZT piezoelectrics
 - 3) Negative feedback
 - 4) Vibration Isolation
 - 5) Electronic Data collection

Quantum Tunnelling

- An electron may be found across a potential barrier it does not have the energy to overcome



- The probability of transmission across a potential barrier of height V_0 and length L is

$$T(E) = \frac{(4\kappa d)^2}{(1+\kappa d)^2} e^{2L/\delta} \propto \exp(-L\sqrt{2m\varphi}/\hbar^2)$$

$$\text{w/ } \kappa^2 = \frac{2mE}{\hbar^2} \quad (\kappa d)^2 = \frac{E}{V_0 - E} = \frac{E}{\varphi}$$

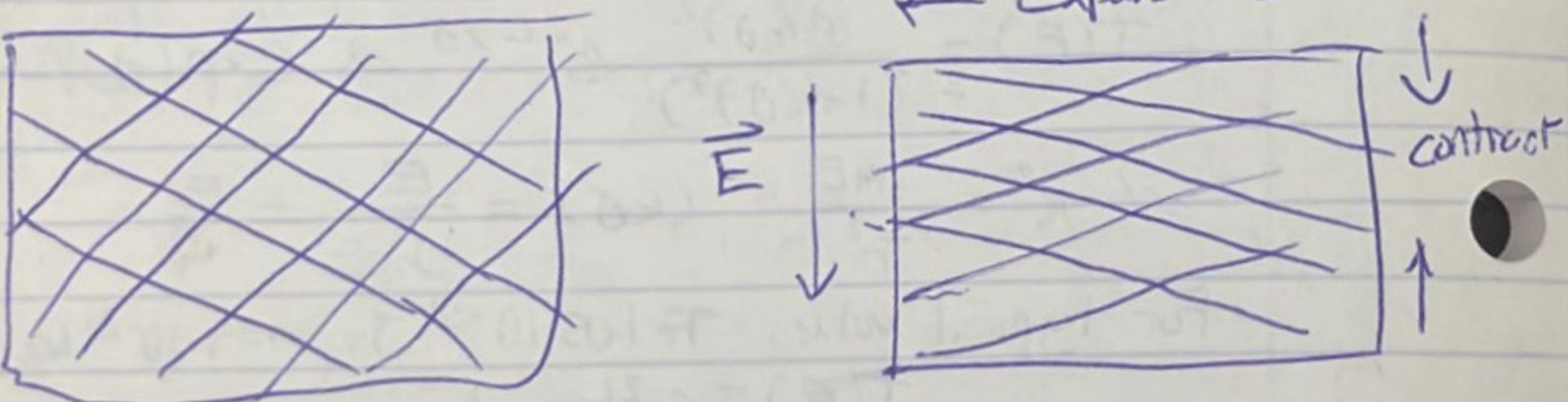
- For typical values, $T \approx 1.05 \cdot 10^{-34} \text{ Js}$, $m = 9 \cdot 10^{-31} \text{ kg}$, $\varphi = 5 \cdot 10^{-4} \text{ J}$, $T(E) = e^{-2L}$, L in angstroms

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- For each angstrom change in separation, tunnelling probability decreases by an order of magnitude
⇒ Tunnelling current sensitively measures distance between tip and sample
- In SFP STM, atoms tunnel between sample and probe. Tip of probe is one atom
- Atoms are ~3 Å apart. So the probability of an electron tunneling from the second layer of atoms (not the closest one) is $e^{-2(3)} \approx 0.002$. i.e. The tip will mostly get electrons from outer surface!

Piezoelectrics

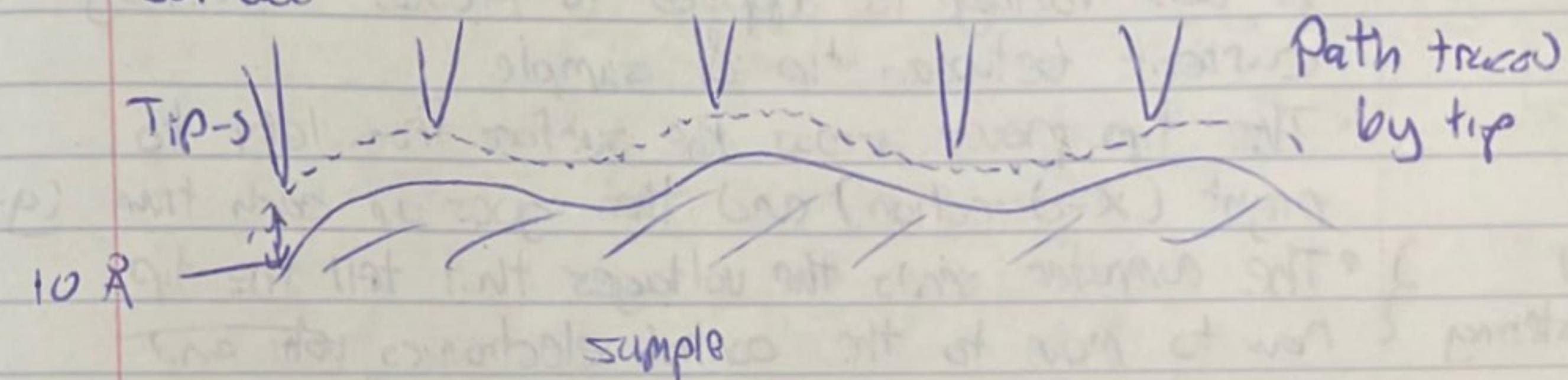
- Tip and sample separated by 10 Å
- To move tip this close, must use piezoelectric ceramics (PZT)
- Piezoelectric effect experienced by electret
- An electret is a material w/ a permanent electric field
- Typically formed by two types of atoms (one positively charged and one negatively charged)
- By applying an electric field to such a crystal, one can cause it to contract in one direction and expand in the other due to realignment of those atoms

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- and spaces
- Interactions in previous figure represent each kind of atom, one positive, one negative
- The piezoelectrics in the STM have a sensitivity of ~150 angstroms/volt
- The computer can control the voltage to mV accuracy ⇒ precision of less than an angstrom
- The motion of the STM tip is achieved using a piezoelectric cylinder using two electrodes
- By induction
- By changing the voltage of each electrode, one can extend, retract, and bend the tip

Negative Feedback

- Tunnelling current is measure of sample distance
- No current ⇒ tip too far
- Some current ⇒ tip near sample
- Electronics of obey the following rule:
"IF the tunnelling current decreases, move tip closer to sample. If tunnelling increases, move tip * away."
- This way, the tip traces out the topography of surface



Sun Feb 3rdVibration Isolation

- Everything is vibrating
- Walking around room causes vibrations in tables of $\approx 1 \mu\text{m} = 10^4 \text{ \AA}$
- We need to hold tip 10 \AA from sample, so the STM must be isolated from vibrations. 3 methods

a) Super-Conducting Levitation: A magnet above a super-conductor will float above its surface

b) Spring Damping: If the microscope is hung from a fixed surface, the springs, the springs will deform by

in response to vibrations of the fixed surface

c) Stacked plates w/ elastomer spacers: Place microscope on a series of metal plates separated by elastomer 'O-rings'

- The O-rings act like springs to dampen vibrations
- Each plate vibrates w/ $\approx 1/5$ the amplitude of the plate below

Electronic Data Collection

- STM image is built one data point at a time
- A bias voltage is applied to produce tunnelling current between tip & sample
- The tip moves across the surface from left to right (X -direction) and then goes up each time (Y -direction)
- The computer sends the voltages that tell the tip how to move to the control electronics, which then uses negative feedback to move the sample towards and away from sample (Z -direction)

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The negative feedback
Using the PZT scanner

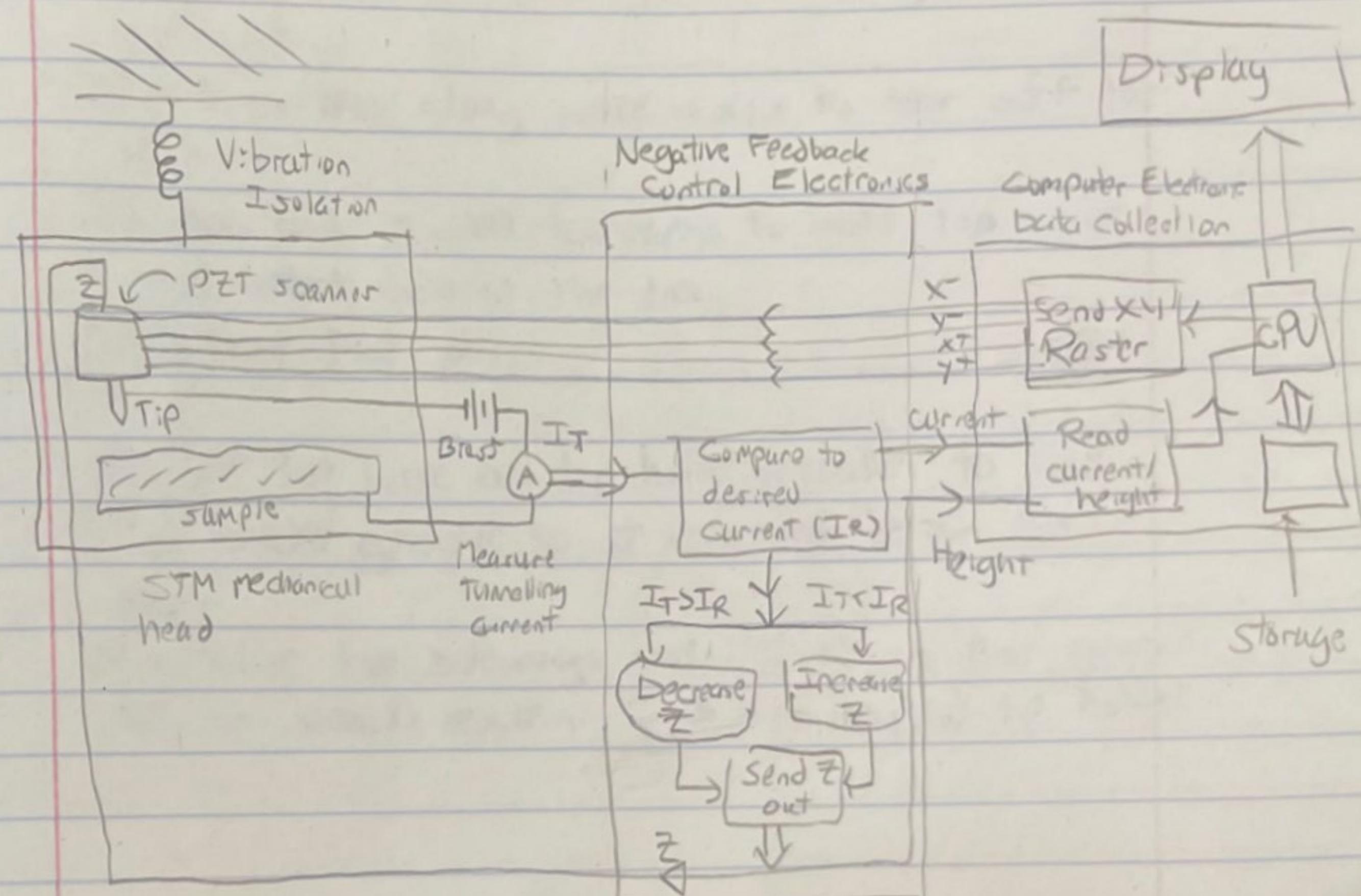
- The voltage required to maintain a constant tunnelling current (Z -voltage) is sent back to the computer

Computer's POV:

- a set of X and Y voltages corresponding to the tip's position are sent out
- a Z voltage corresponding to ~~to~~ the tip's height (the sample height is returned)

- On the display, the (X, Y) position are the Cartesian coordinates of image

- The image is in grayscale. The gray value of each point is proportional to the returned Z voltage



Tues, Feb 7th

- Showed up to lab and identified tools
- Here I will write a description of the microscope and available tools. NanoSTM PC
- Note: Previous STM student, Christian, came in and mentioned that the samples are mislabelled
- One sample is just sitting on table w/ no box?

Tues, Feb 7th

Preparing for Measurement procedure (from NanoSTM manual)

Initializing System

- 1) Connect power, & USB to computer
- 2) Turn on STM power
- 3) Start Nano control software on computer

Preparing and installing STM Tip

- 1) Clean wire cutters, flat nose pliers, and pointed tweezers w/ ethanol. Only touch P+ / IR wire w/ these
- 2) Remove old tip from STM using pointed tweezers
- 3) Hold end of wire firmly w/ ^{flat nose} pliers
- 4) move cutters at 4mm as obliquely as possible
- 5) Pull close cutters until wire is felt, but do not cut
- 6) Pull cutters along wire axis to tear off the tip
- 7) Use the pointed tweezers to hold tip wire just behind behind the tip
- 8) Release flat pliers
- 9) Put tip wire on tip holder, parallel to the tip holder groove so it crosses below tip clamp
- 10) Move tip sideways until it is in the groove
The tip should extend 2-3 mm beyond tip holder

Tues, Feb 7th

- * Not different samples require different preparation
The required preparation will be noted later

†

†

Mounting the sample

- 1) Unpack sample holder from plastic storage vial, touching only black plastic handle
- 2) Put prepared sample on magnetic end of sample holder using tweezers
- 3) Lower front end of sample holder onto sample holder guide spheres, then gently settle back end onto approach motor support
Place sample holder into scan head in such a way that it does not touch the tip and so the sample is not pulled by the magnet that pulls the sample holder down onto the approach motor

Tues, Feb 7th

- 10:03 AM, office prof. Tahir arrived. Office 915B
We discussed the experiment
- I must image these different surfaces and then use physics to interpret my results, measure size of atoms etc.
 - Print images and stick them in lab book

PreparingMounting Gold Sample

- Sample No. BT00623. Labelled Graphite but it is gold
- Requires no special preparation

Preparing Graphite (HOPG) Sample

- 1) Put sample on table w/ tweezers
- 2) Stick tape gently to top of HOPG and adhesive pull it off to remove top layer
- 3) Remove any loose flakes w/ tweezers

Tues, Feb 7th

First Measurements w/ Graphite

- Sample No. BT06351
- Got Ethanol and scotch tape from lab tech
- Initialized system (p. 9), blue  = LED activated at 10:28 AM
- Message saying 'No controller detected'?
 - Unplug and replug USB, and restarting system fixed this
 - Attempted to cut a wire but I dropped it
 - Having steady hands is really hard
 - Second attempt was successful but I dropped the tip on the paper at some point so ~~I~~ it's possible the end funny the sample holder + is the dull end
 - I picked the sharper looking end for the tip
 - I used a STAEDLER triangle to get ~~it~~ it
 - I accidentally used rounded tweezers instead of pliers to do this
 - Prepared HOPG as per p. 11
 - Mounted sample as per p. 10
- Load default parameters from File > Parameters
 - >> Load parameter settings and load C:\Program Files (x86)\NanoSurf NanoConfig\Default-STM.par

Tues, Feb 7th

Manual Course Approach

- 1) Push sample holder to ~1mm of tip
 - 2) Rotate sample so tip points to a flat area
 - 3) Place magnifying cover over scan head so you can see reflection of tip in sample
- Did this as instructed. Mirror may be hard to see in Graphite due to distortions and lack of light. The tip looks sharp!

Fine approach motor

- 1) Watch distance between tip & sample w/ help from magnifier
- 2) Click Advance button in approach group of acquisition tab until gap between tip & reflection is barely visible
- 3) IF motor not

- Motor moved as expected, but reflection of tip is hard to see
- I moved it as close as I could

Tues, Feb 7th

Automatic Final Approach

- 1) Preparation Group: Wizards \rightarrow Imaging
- 2) Give expected feature size & height to the best of your ability. Set scan quality

Next: Name Measurement Series

- 3) Activate Gallery panel in Info pane
- 4) Check History tab
- 5) Enter name in top entry box

Approaching:

- 6) Click Approach on Approach group & in Acquisition tab

- For approaching HOPG, left default vals.
- Feature size 50 nm
- Feature height 5 nm
- Medium speed / good quality
- Followed instructions
- Approach did not end after 20 seconds. Go back to repeat manual approach
- Rotated - Removed sample & rotated microscope so that reflection was more visible
- Remounted, and approached sample
- Automatic approach still not working
- Opened SPM parameters \rightarrow Approach
- Set move speed from 70% \rightarrow 100%
- Still taking way too long

Tues, Feb 7th

- Decreased gap using fine approach even more
- I repeated this process until the motor stopped moving in fine adjustment
- Tip may have crashed
- I may need to clean the motors next time

- Unmounted and packaged sample
- Turned off software & power
- Before Friday, figure out surface required feature size & height for graphite
- I left the tip in the clamp for next time

Fri, Feb 10th

- Tip is still positioned & sharp!
- Turned on STM as on pg. 9
- Prepared HOPG (sample BT063 + BT06351) as on 11
- Mounted sample as on 10
- Reflection visible!
- Fine approach as on 13
- Loaded default parameters but set approach speed to 200%
- System does not let me do that, set to 100%

- Automatic final approach at 9:36 AM
- Let that run while reading about ω_0 for approach too long in ~~the manual~~ operating instructions
- I think I can get a closer fine approach
- Put my eye right to magnifying lens and got a much closer fine approach
- Retry at 9:37 AM
- Quit at 9:44 AM
- Maintenance: Clean surfaces w/ acetone + cotton swap → sample support
- Turned off microscope & repackaged sample
- Sample fell on table

Fri, Feb 10th

Cleaning:

- Moisten tissue ~~skin~~ (KIMTECH Science Wipes) and clean interior surface of scan head
- Moisten cotton swap w/ acetone and clean sample holder support surfaces

- Moisten KIMTECH wipes and move along sample holder in axial direction. Put sample holder back in container
- Allow all parts to dry

- Reprepared sample BT06351 and remounted as before
- Redid page 13
- Still not working. I forgot to set appropriate parameters
- Loaded STM parameters file
- Set approach speed 100%
- Left default feature size & height

- It still is not working :C
- Automatic approach just goes on and on

- Prof Tahir walked in at 10:15 AM
- Tip was too close and not sharp enough
- Also tungsten tip that I am working with is okay, PI/IR not necessary

Fr, Feb 10th

Repeated the process on p. 9. This time I made sure to hold the wire w/ pliers instead of tweezers, and tugged on the cutters & in the wire's axial & ~~axial~~ direction for a much sharper tip.

- Remounted sample and reapproached
- Set values back to params to default STM w/ 70% speed
- Attempt at final approach, 10:36 AM
- Still nothing
- Replayed w/ manual approach until reflection area tip were almost indiscernible
- Approach done INSTANTLY
- Image size is $(250\text{nm})^2$
- I want to decrease this range
- I am just getting a flat colour map, even at $50\text{nm} \times 50\text{nm}$
- TA walked in. Said to play w/ different params
- Something was wrong w/ software. Probe light was red
- Restarted software and changed move speed to 100%
- Told me to start w/ big image and try to zoom later
- Mess w/ setpoint voltage and PID gain to get approach
- Set point 1nA, P-Gain 1000, I-gain 200, D-gain 0, scanning worked

Fr, Feb 10th

but image stopped imaging properly after a few lines

- Increased tip voltage from 50 mV \rightarrow 250 mV
- No change
- Inclined surface possibly
- Tip is crashing. Tip could be crooked or getting too far
- Line graph of Z-axis varying wildly

- Also, watch probe light on computer. Red = too close
- System froze as I played w/ values
- Restarted everything. Replugged USB
- Fine approach not moving at all!
- Restarted software!
- ~~Reapproached~~ Reproduced sample and a fresh new tip
- After setting speed 100% and tip voltage 250 mV, this worked
- TA explained that tip voltage is meant to decrease the potential barrier of air. So higher tip voltage means more tunnelling

Params:

Image size: 200 nm	Set point: 1nA	Tip Voltage: 250 mV
Time / line: 0.2 s	P-Gain: 1000	
Points / line: 128	I-Gain: 2000	
Rotation: 0°	D-Gain: 0	
• Produced image Image00044		
• Time / line Very sharp change		
• Time / line \rightarrow 0.4s produced Image00045		

Fri, Feb 10th

- 20
- Fri, Feb 10th
- Probe stuck after
 - Probe status orange. Images corrupted. Rec approach
 - System crashed
 - Restarted & tried again. Images corrupt again
 - Probably surface imperfections. I will cleave w/ tape again
 - The surface of this sample is extremely flaky and noisy
 (last time I mentioned there was a sample w/ no box)
 - Looks like graphite. It has a much smoother surface so I'll use that
 - During scan, red light. Tip crashed
 - Played w/ params trying to get approach to work
 - Tip seems to keep crashing
 - Decreased move speed to 55% (other params default)
 - That worked well enough to see details but images suddenly became white
- hypothesis
- Increased set point to 2nA
 - Probe light flashing orange-green
 - Set ~~Tip~~ voltage
 - Tip keeps losing contact w/ sample after a few lines
 - Cleaved sample w/ tape to remove bumpy outer layer and reapproached (Tip voltage 250mV, set point 1nA)

Params:

- The tip voltage lowers the potential barrier created by air. Higher voltage \Rightarrow more tunnelling
- Setpoint: the desired tunnelling current. Higher current \Rightarrow closer tip required
- PID \Rightarrow related to negative feedback cycle.
 Learn more about this

9PM

Fri Feb 9th

What is PID? From Wikipedia

- Define the error function

$$E(t) = J_{set}(t) - J_{true}(t)$$

where J_{set} is the current set point and J_{true} is the actual actual current

- PID defines a control function

$$U(t) = K_p E(t) + K_i \int_0^t E(t') dt' + K_d \frac{dE(t)}{dt}$$

- The PID values in the system must refer to $K_p, K_i, K_d \geq 0$

P-Term: related to the error right now

I-Term: accounts for the history of errors

D-Term: Accounts for future trend of errors

- In my case, the control variable should be the height distance between the tip and the sample?

In Nano STM operating Manual:

- Increasing P-Gain: decreases error signal
- Increasing I gain: decreases error signal over time
- Increasing D-gain: decreases fast changes in error signal, but amplifies high frequency noise

Tues. Feb 14th

- Tip from last time \Rightarrow is still in STM

- Prepared graphite sample

- Manual approach, Tip may have crashed

- Attempt approach w/ pg F1 params

- But we move speed 55% \rightarrow 0.5 m/s or pg 20

- Approach didn't go through. Time to replace tip

- Accidentally crashed tip while approaching sample

- I will cut the end off the same wire but again

- It is too short now

- Success. Now try final approach approach w/ move speed $\times 60\%$

- It worked!

- Produced images 46 and 47

- Still fairly choppy images

- I will increase resolution. 256 points per line

- And time to 0.5 s per line

- About halfway through image, probe light turned orange. Too far. Must reapproach

- Probe light flashing orange/green ~~brief~~ during scan

- About halfway through scan, reapproach

- Each slice of the image doesn't quite line up. Some sort of drift of the tip perhaps?

- Product was ~~at~~ image 0048

- Perhaps the tip is still too far for a good image. I will try tuning voltage tip voltage to 150mV so it needs to be closer

Tues
Feb 14th

- Try PID (100, 200, 0)
- Interview w/ prof Majoribanks
- Try PID (100, 200, 0)
- Image 0053, less grainy, but drift is still apparent
- Image 0054 is down skin, looks very different
- Maybe I

- Try PID (100, 50, 0)
- Probe light red. Do not set PID too low. Causes this to happen? Why?
- Perhaps too slow of a response to changing terrain causes crash

Discussion w/ TA:

- Change set point & time per line. Worry about PID later. Explore how these affect quality
- Try P=1000. I want P greater than I?
- Start at 20nm, once a good image is seen, try 50nm. Then 10nm and so on

Tues
Feb 14th

- Stopped during image to reapproach w/ 60 nm left
 - Image 0049 a bit better
 - Reapproach, try at 50.1 nm now
 - It is taking way too long to approach
 - Go back to 150nm
 - Experiment w/ PID
 - What is the effect of keeping PID ratios but decreasing the total amplitude?
 PID from (1000, 2000, 0)
 to (100, 200, 0)
 - Probe light turned red? No crash apparent. Still red after retracting tip. Restart software
 - Res. Reset params and * try again
 - Probe light turns red at as soon as I hit approach approach
 - I changed PID back to (1000, 2000, 0)
 - It worked again. Image 0050
 - Image 0051 is the downscan of same region!
 - Okay I'm going to zoom in to a flat area. Pick a flat. Flat area is an area w/ a constant & uniform colour on image
 - Zoomed into $(664 \text{ nm})^2$ area
 - Zoom again into $(587 \text{ nm})^2$ region
 - Product is image 0052. Very grainy.
 - I will try decreasing PID gain. Perhaps probe is reacting too fast
 - PID \rightarrow (400, 800, 0)
 - No difference. Left setup for 20 mins for
- these were pretty clear

Fri, Feb 17th

- Cut new tungsten tip
- Prepared HOPG sample and mounted in scan head w/ probe tip pointing to a smooth area
- Set params to those required for Image 50 & Image 51 and approached

That is:

Image size: 200 nm

Time / line: 0.5 s

points / line: 256

Move speed: 60%

Setpoint: 1 nA

PID: (1000, 2000, 0)

Tip voltage: 150 mV

- Approach taking too long. I noticed there was still a very big visible gap between sample and its reflection so I returned to manual approach

- The approach was successful!
- The resulting Image 0055 seems kind of warped
- I will take another one w/ the same settings. Perhaps the system just needs to settle
- At. Thermal effects acting on scan head on nm scale. Maybe turn down time / line for this, to scan faster
- Image 0056 looks similar to 55
- Turning down time at this scale only made

Fri, Feb 17th

Image streaker. I think 0.5 s / line is optimal

- Streaky image still. Perhaps sample must be cleaned and better tip cut.
- Image 0057 is the best it'll be at this scale? Time to zoom in
- Find a flat area in downscan
- Try 50 nm x 50 nm

Contour plot of current looks like pure white noise. Play Playing w/ params until this is not the case

- Images 58 and 59 are the best so far at this scale. Currently,
- Time / line = 0.25,
- Setpoint = 504 pA PID = (700, 200, 0)
- Tip voltage = 50.1 mV

The line graph is changing a bit for each line through. Plenty of room for

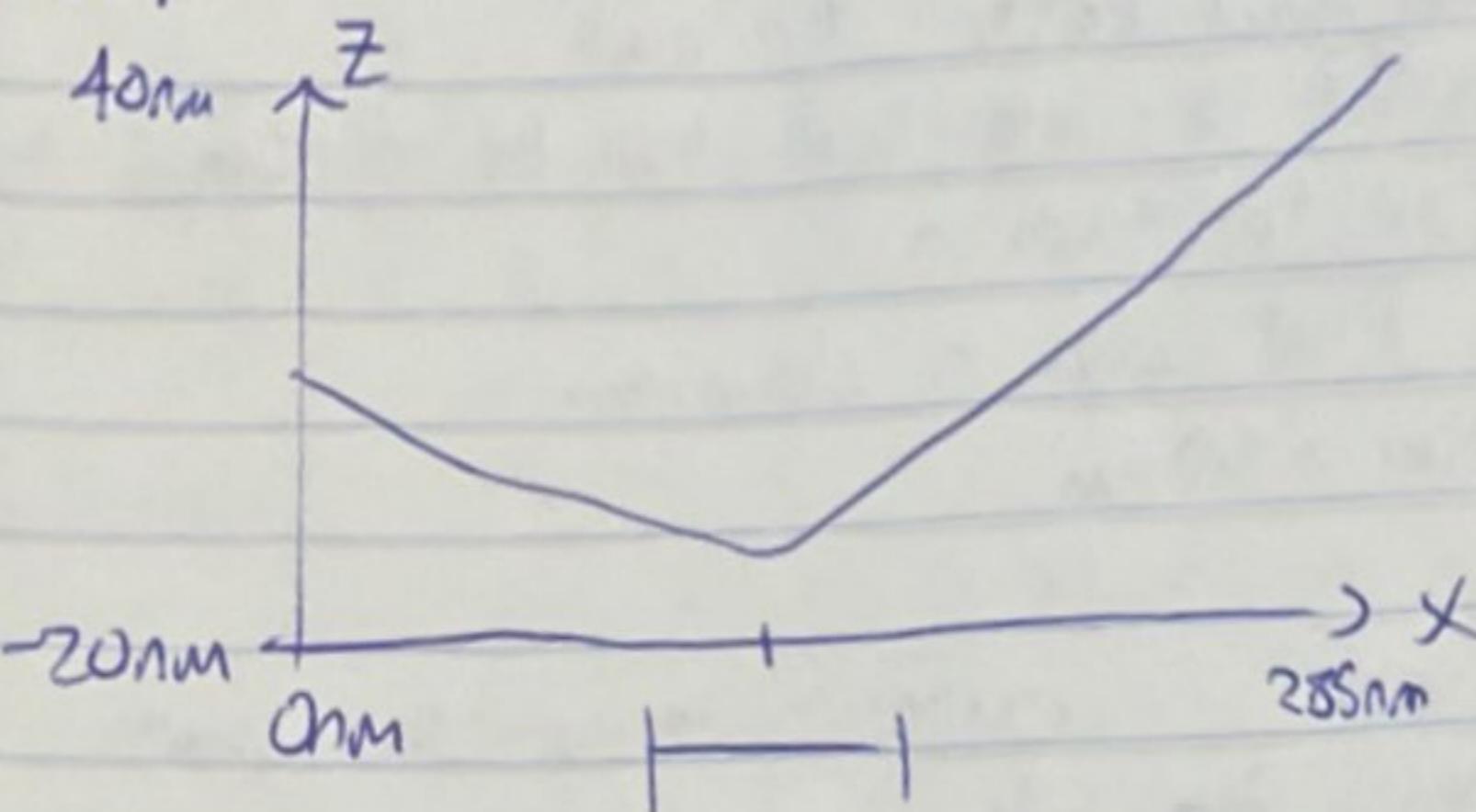
- Turning setpoint to 300 pA produced images 0060 and 0061 w/ time / line 0.01 s
- What am I looking at?
- Software crashed
- Restarted and reapproached sample
- I believe 60 & 61 are just noise.
- The picoamp scale for current is likely much

78nm

Fr. Feb 17th

more noise sensitive than the nano amp scale

- To check if image is real, use line graph. Currently, for each slice, we have



zoom into this region
and check if pattern stays

- I got the same structure on the edge but it's hard to trust

- I'm cutting a new probe
- Image 063 w/ new tip and clean sample MUCH better, default settings w/
tip voltage 250mV

- Image 064 w/ setpoint 2nA, voltage 50.1mV
PID = (1000, 800, 0)

Zoom in for image 0065

- Zooming in beyond this point just gives noise
- Perhaps tip has picked up particles. Tried STM cleaning pulse and reapproached

Fr. Feb 17th

- Replaced tip & cleaned sample

- It is possible that tip got too close while zooming in. Try increasing voltage to prevent this.

- Evidence: After zooming back out, image was quite blurry

- My new strategy is to zoom in on well defined surface features and make them look more clear

- ~~Image 0072 for example~~

- Such as image 0072

- Setpoint $4 \rightarrow 6$ nA $\rightarrow 8$ nA gets image 73

- It is hard to say which image is best

- I will go w/ 74 or 73

- Zoom into flat region to the right of large feature. Image is still suffering from noise due to very small current variations

- Image 075. Noisy atoms?

- Image 076 and Image 077 are upscan and downscan respectively

- Line graph pattern is stable. Still changing a bit too quickly, but I think these are atoms

- Setpoint 7.51nA. Voltage 50.1mV

- Time 0.06s/line. 256 points

- PID (650, 350, 0)

- Zoomed in and features got bigger. These

Fri Feb 17th

are definitely atoms!! (Image 0078)

- Making tip voltage too big might increase noise
- It lowers the potential barrier due to air and may cause tunnelling from below surface, which causes noise

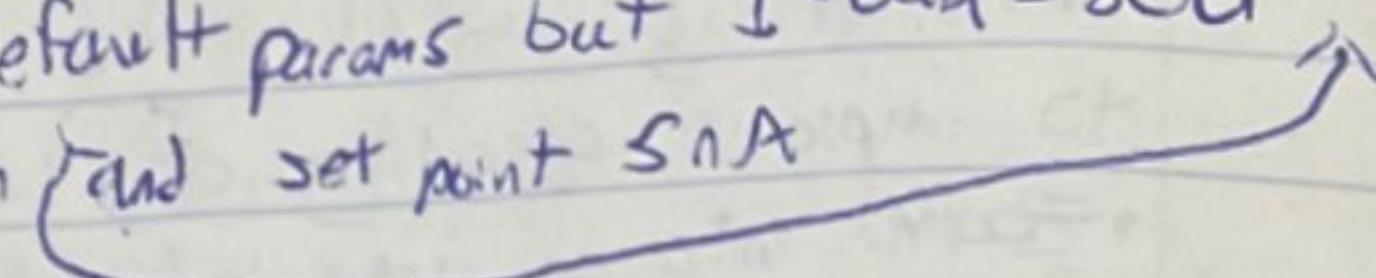
Outer

Tues, Feb 28th

- Turned on SIM & software
- Move speed 60%
- On pg 29, tip was replaced & did not crash so I will continue using it
- Cleaved graphite sample w/ tape
- Approached sample
- Tip crashed :C
- Cut a new one
- Tip may have gotten too close on approach
- The images produced are extremely (choppy) like image 44
 - Increasing time/line to 0.4, ^{from} 0.25 seems to improve quality at 200 nm scale
 - Zoomed into an area w/ well defined topography and saw nothing. Time to throw away tip
 - I trimmed the end of the same tip
 - Tip crashed
 - I keep trimming the tip, but the line graph is extremely choppy. I might just have to cut a new one entirely
 - New tip produced clear image w/ set point 2 nA and voltage 49.7 mV
 - Zoom into well defined topography
 - Feature in image 74. Continue zooming
 - Result is image 81. I threw out image 80 bcs I zoomed into the wrong area
 - Image 81 used set point 2 nA, voltage 49.7 mV
 $PID = (600, 500, 0)$, and time/line 0.12 s
 - Zoom in to the left of large peak
 - Image 82 time/line 26 ms

Tues, Feb 28th

Fri, March 3rd

- Looks like atoms
- Probe status went orange & I had to reapproach
- Zoomed back out to 50nm scale
- I have to basically start again
- Nice topography in Images 00083, 00084.
- I am just playing w/ params
- Image 0085 w/ default settings
- Zoom into center
- Image 0086 has a faint atomic lattice
- Again, I lost tip
- Get image 87 w/ default params but I-Gain = 800
- Zoom into flat region (and set point 5mA) 
- Result is Image 0088 (Time/line 52ms)
- I will let the system image for a while to reduce the effects of thermal drift
- Withdraw tip & ran STM cleaning pulse
- Unable to get a good image

- +8
- Today I will spend some time imaging gold
 - Time/line 0.35, setpoint 4mA, PID = (1000, 300, 0)
 - Tip voltage 500mV
 - Resulted in image 106 of gold surface
 - Rotate plane by 30°
 - Structures reproduced in image 107, not noise
 - Very strange in measurement direction
 - Image 108 very strange
 - Prepare new tip
 - Beautiful image 109 w/ new tip
 - Image 110 at 50nm scale
 - Image 111 at 20nm. Starting to look grainy
 - Image 112 at 5nm. Looks too homogeneous?
 - What happens if tip time/line $\rightarrow 0.065$? Horrible image 113. Back to 0.035 and wait
 - Images 114-116 at this setting
 - What if I increase time/line? 0.4s for Img 117, ~~118~~
 - Or 0.5s for Img 118
 - Gold is conductive \Rightarrow homogeneous electrons
 - This may mean I need to spend ~~+~~ a longer time imaging gold to see its features
 - But I need to balance this effect w/ thermal drift
 - Zoomed in to 2nm for Image 119, 120
 - Drifted probe too far from sample. Reapproach required
 - Img Run an STM cleaning pulse
 - I lost tip
 - Back to 100nm scale w/ Img 121

Fri, March 3rd

- I will trim the tip once more
- Img122 at 500nm
- Img123 at 1000nm
- Img124 at 5nm
- Reduce voltage to 0.0004mV
- Withdraw, cleaning pulse, reapproach
- Lost completely. Recut tip

- Could electrical interference be the culprit?
- If I don't know how to fix that
- In the meantime, mean time, I will rotate the surface sample and approach a new area

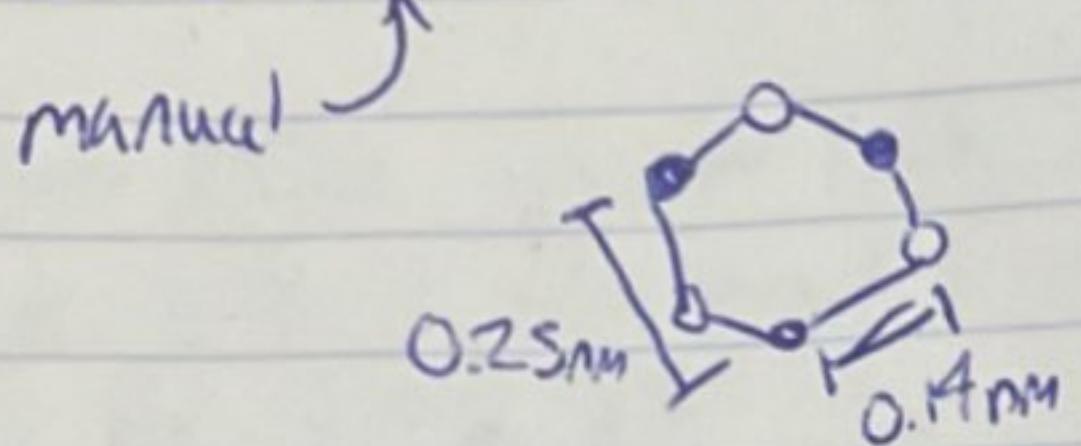
- Time/line 0.35, setpoint 1nA, PID = (1000, 800, 0),
voltage 500mV for img 126 at 200nm
- 50nm for Img 128
- 20nm for Img 129
- Img 130 w/ setpoint 5nA
- Lost the structure

Fri, March 3rd

- Back to imaging graphite
- Cut a new tip
- Cleaved graphite sample and mounted it
- ~~Image 125~~
- Graphite is steady in image
- Played w/ params and saw nothing
- Rotated surf. SIM cleaning pulse
- Image 131 default params, time/line 0.36s
- Zoom into ridge in ~~image~~, setpoint 1.5nA
- Time/line 0.16ms. Image 132
- Not seeing anything iC

Analysis

- Image00ff is my best photo of the graphite lattice
- Throughout the entire image there is a clear triangular lattice of black points; these are low areas/gaps between the atoms
- The actual atoms are a bit harder to make out due to distortion
- The image quality could be improved. There are white/blue streaks in the scan direction
- The NanoSTM describes a carbon lattice like so



- have neighbours in the plane below
 - do not
- Atoms w/o neighbours in the plane below appear higher than the others due to a higher electrical conductivity
- This structure is hard to make out in Image00ff. There are clearly white peaks and orange peaks throughout, but the lattice is unclear
- To get the atomic diameter, I will find a clear feature patch in my image and measure relatively the distance between the edges of two black spots

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- Picking a clear area to do this results in an atomic diameter of about 113.3pm, for carbon
 - The accepted value is 140pm.
 - My figure has a relative error of about 20%
 - This is an extremely rudimentary analysis based on an image of dubious quality
 - The true estimate of the diameter could be improved greatly by making use of a computer ~~algo~~ algorithm to find the average peak radius
 - I could improve my imaging by continuing to experiment w/ PID values and/or considering the effects of electrical interference or air currents in the environment
 - For the gold sample, I found it much easier to get an image at the 200nm scale than the graphite. Perhaps the conductive properties of gold made it easier to achieve this
 - I was not able to get a clear image at the atomic str level. For gold, you need to image for a longer time since electrons are distributed homogeneously through a conductor
 - However, the atomic lattice is sensitive and prone to changing due to thermal / environmental effects during this time, which makes getting a clear feature patch difficult

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- More time is required
 - Abstract included in STA summary and
STM-Summary-and-Abstract.pdf

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The following pages contain the images I took

Printing issue caused me to not get physical copies for images taken on pg. 34, 35
many

Image 0044

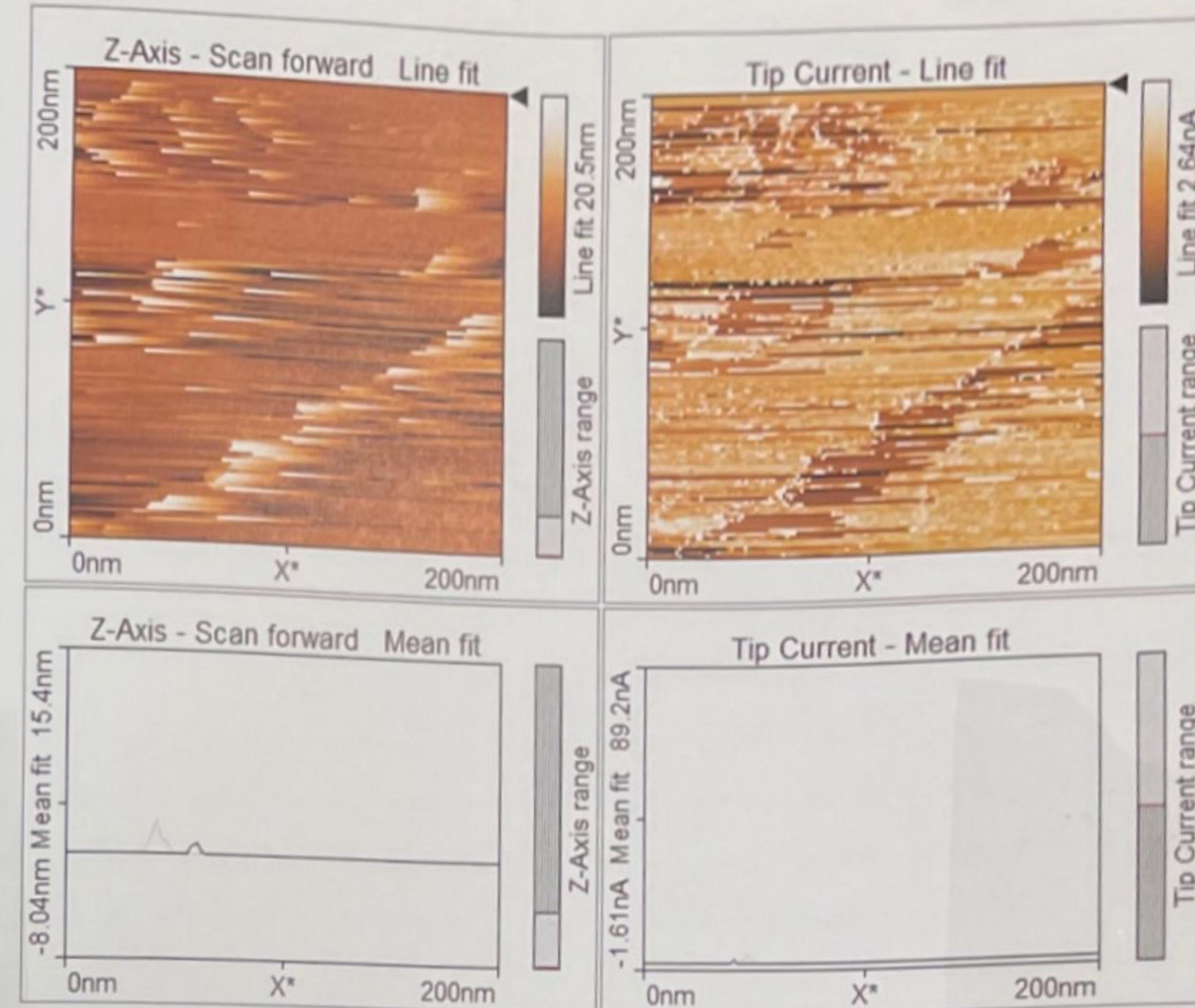
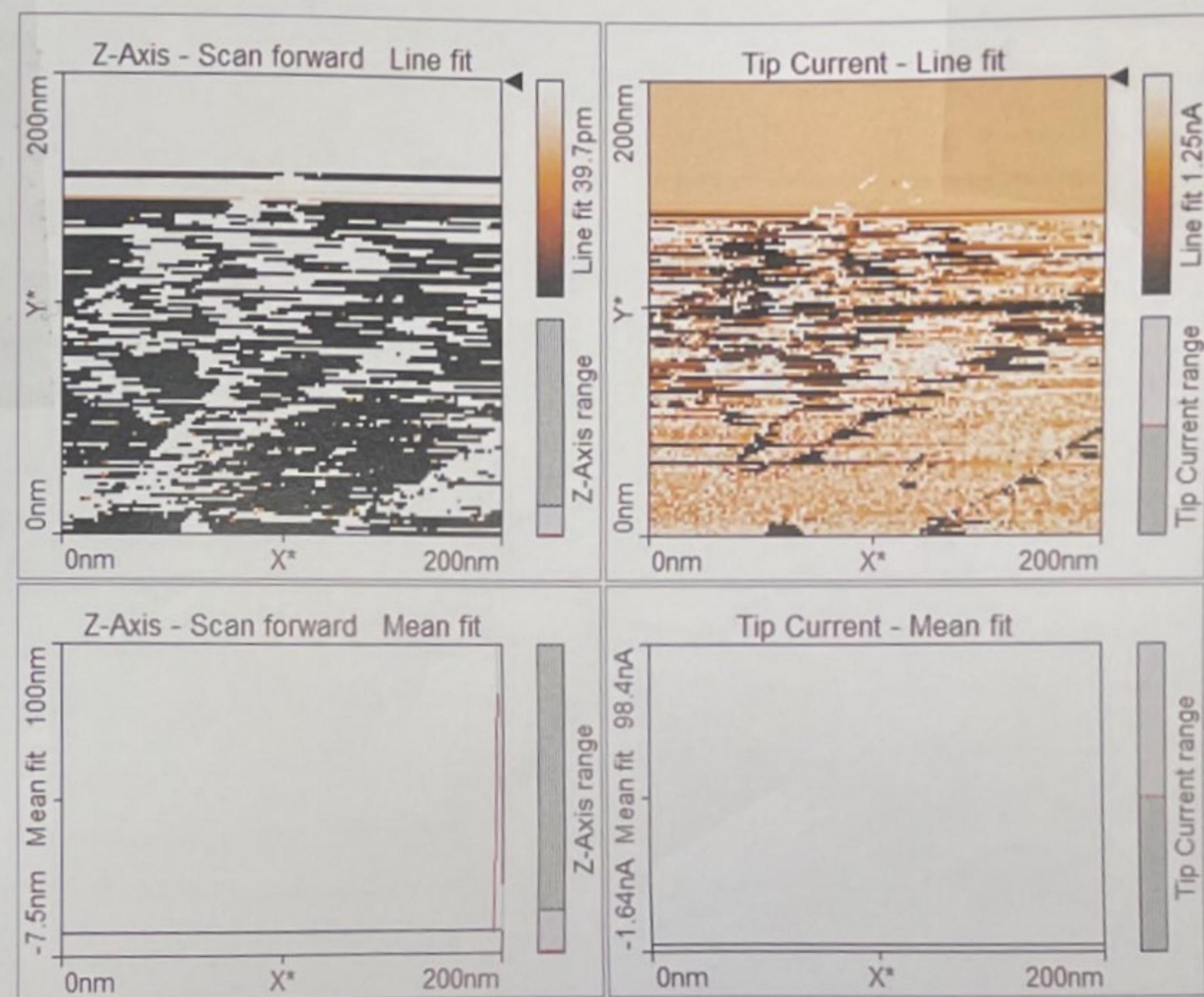


Image 0045



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Image 0046

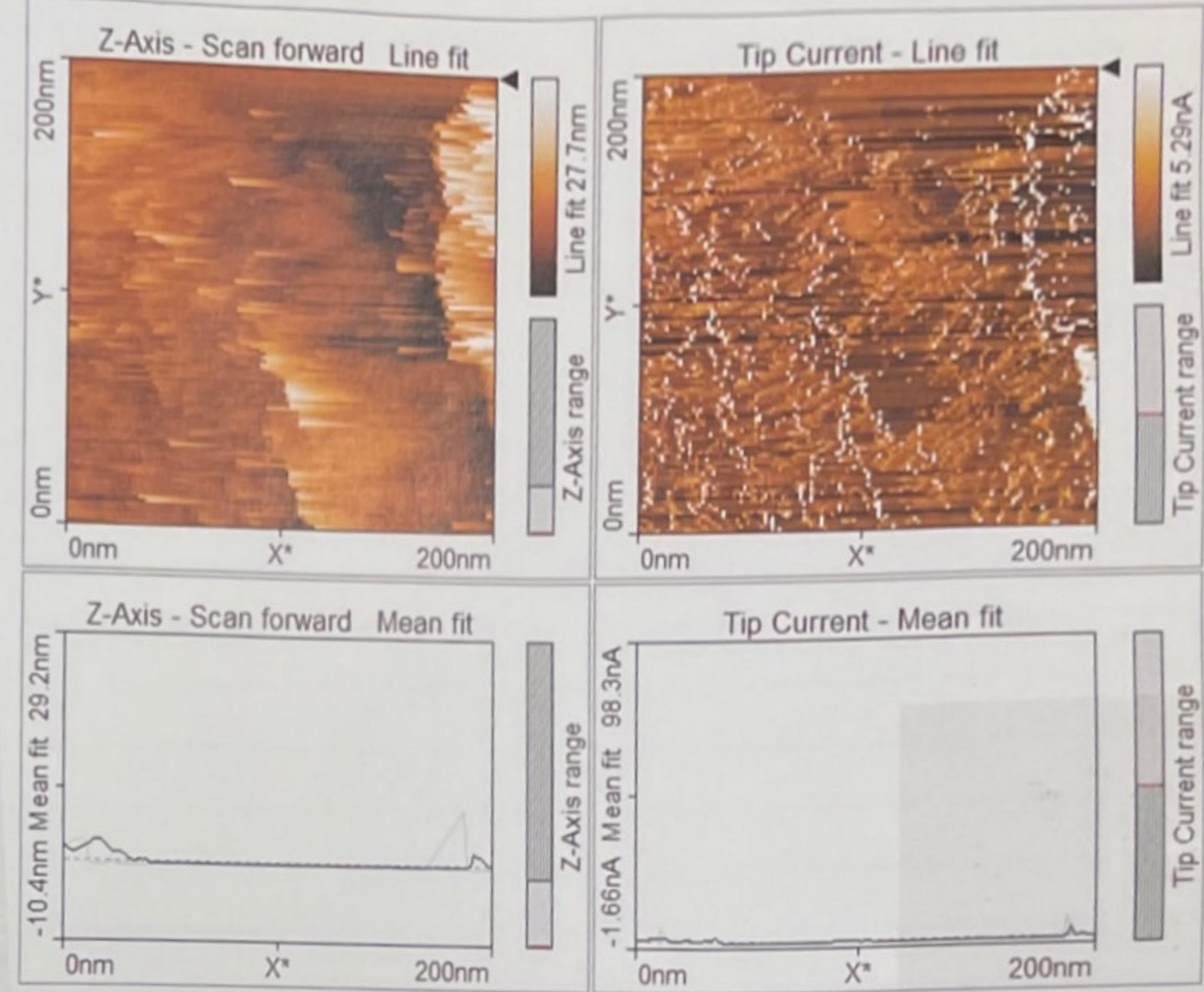
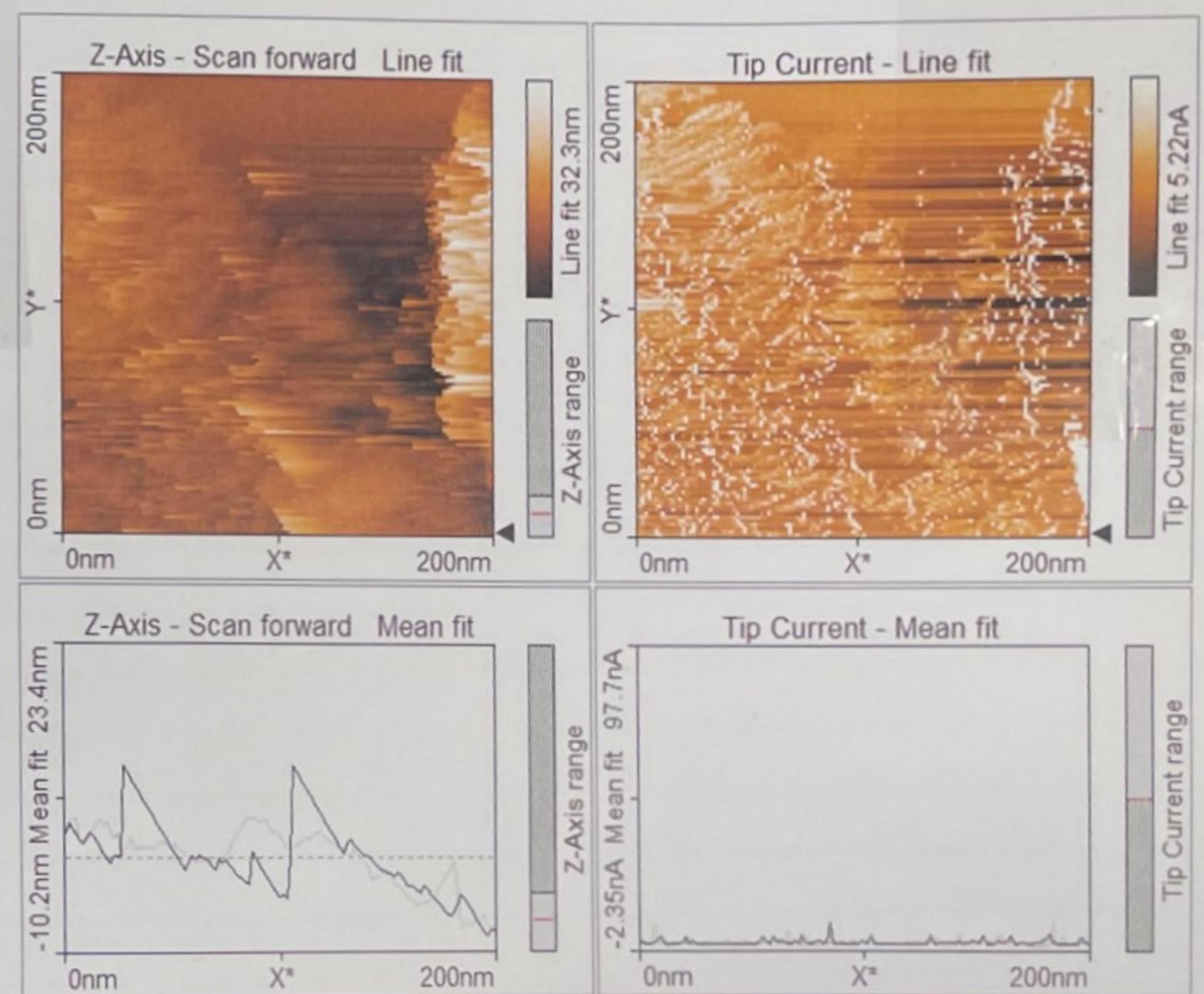
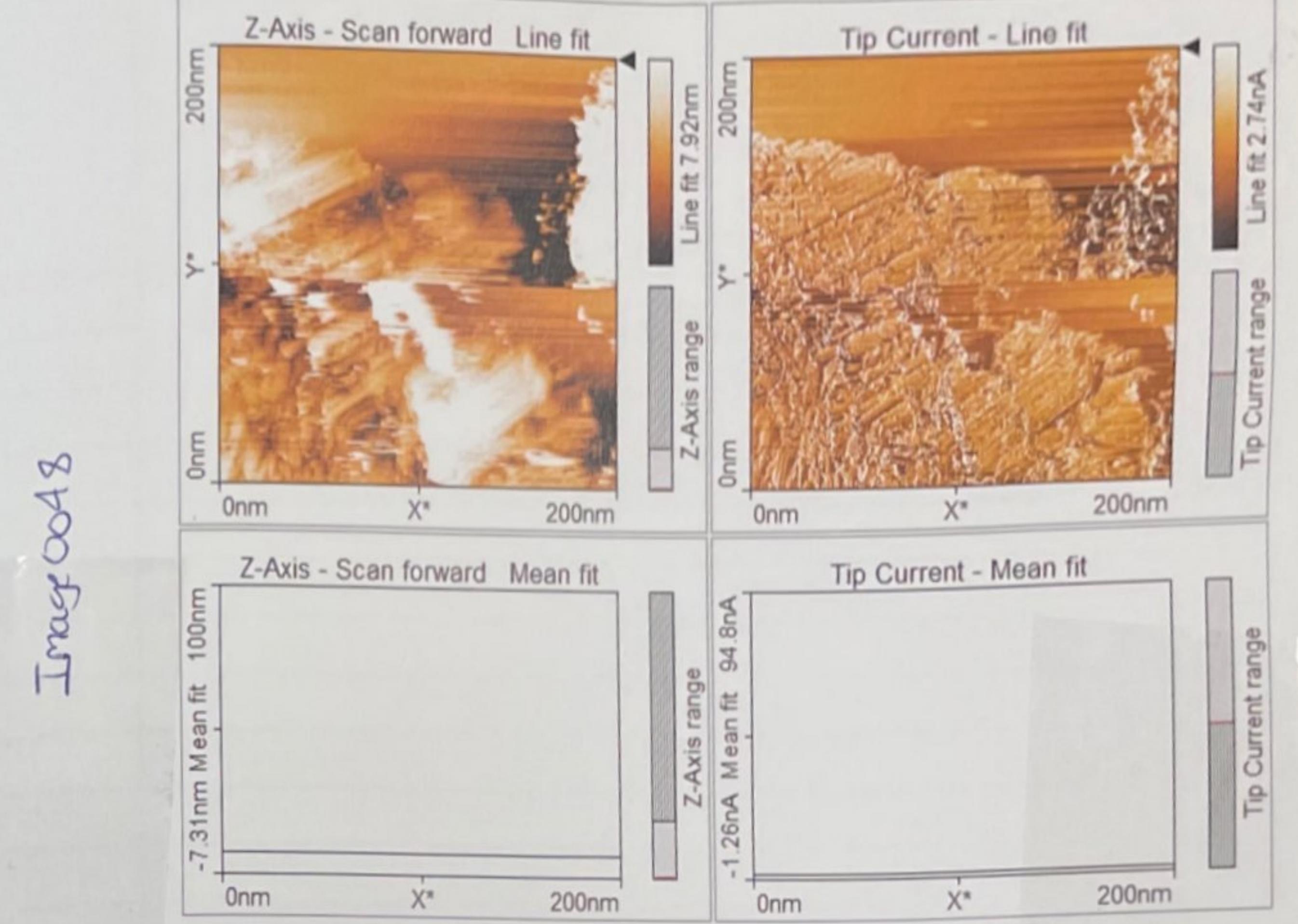


Image 0047



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Image 0050

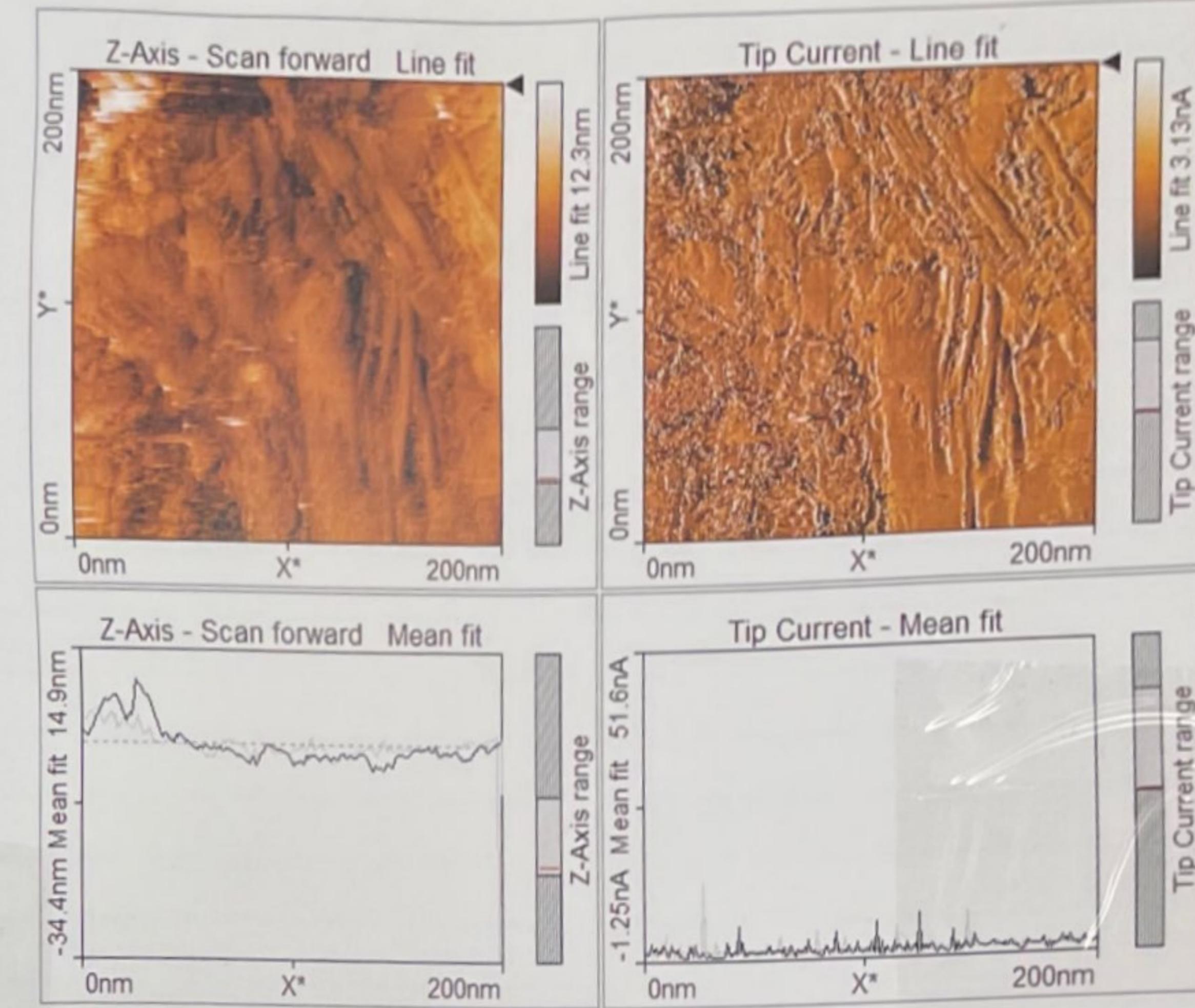
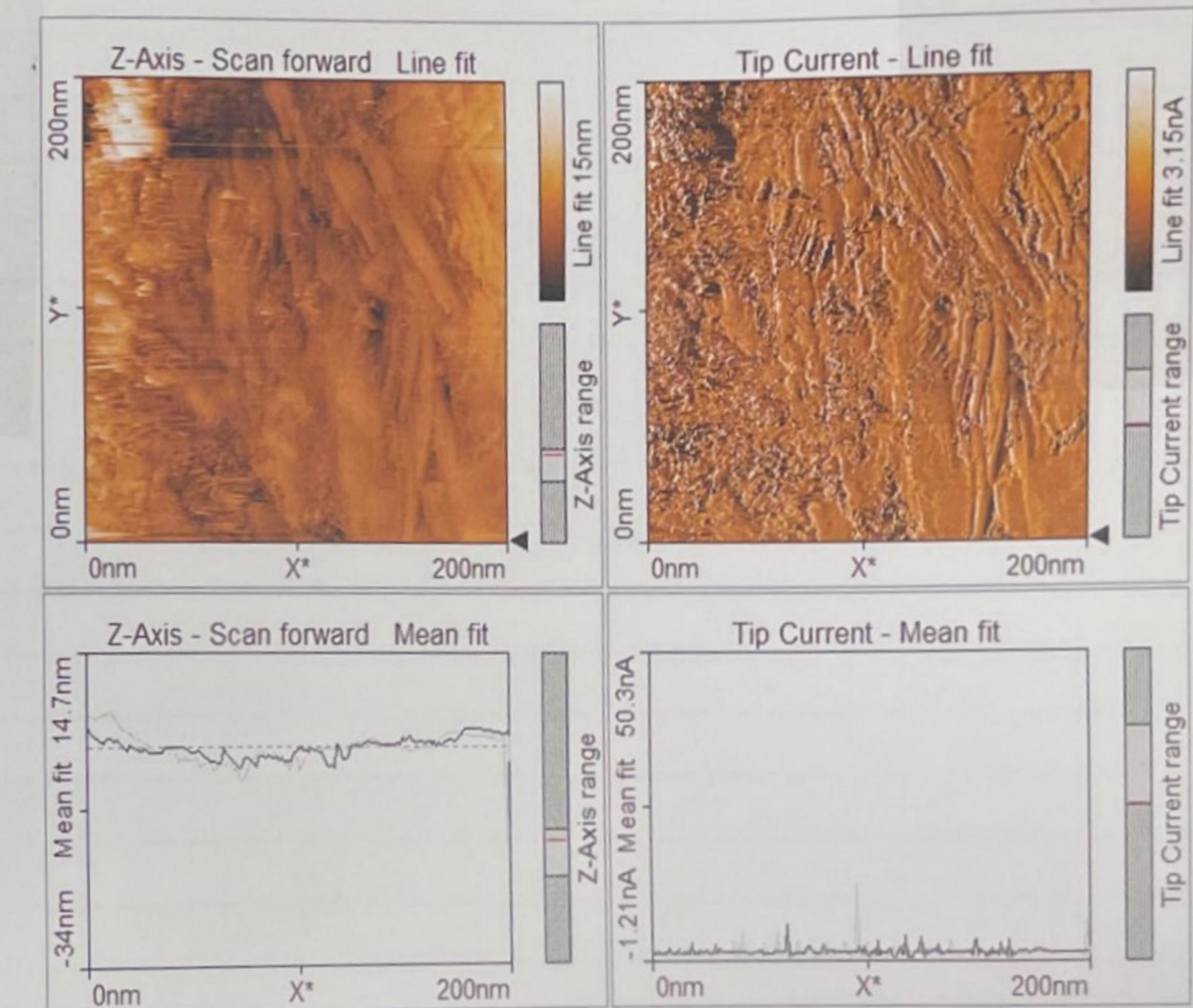
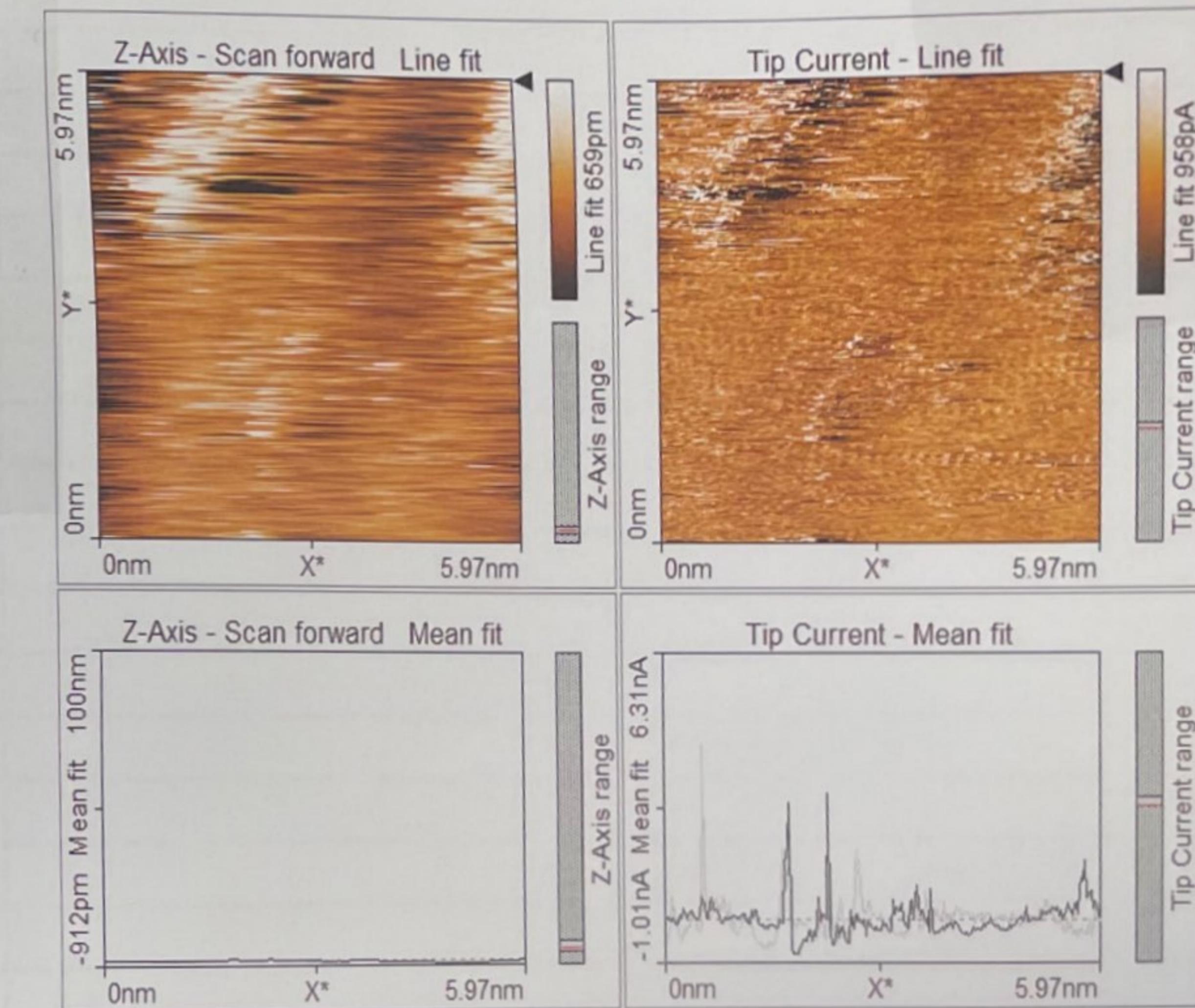
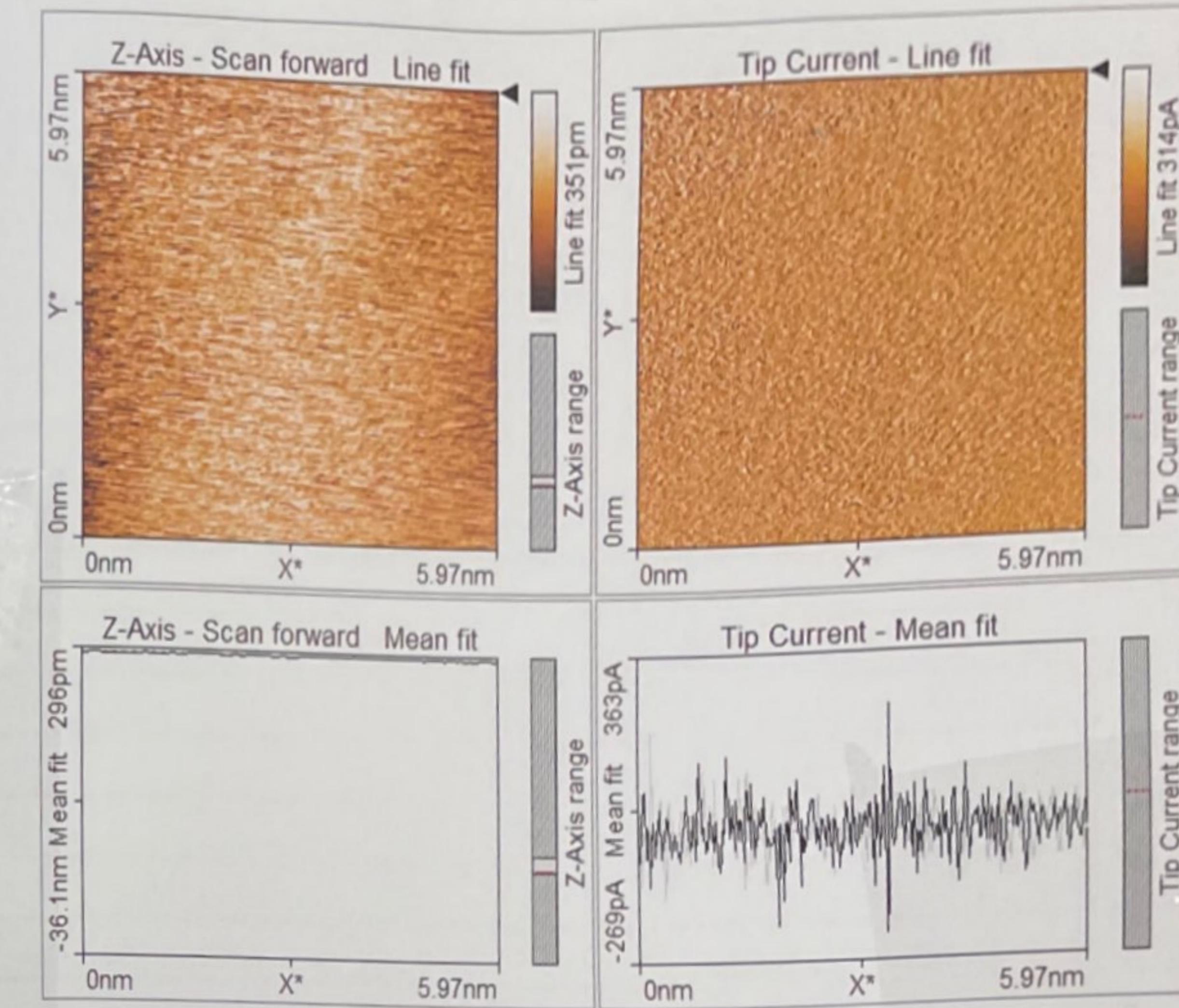


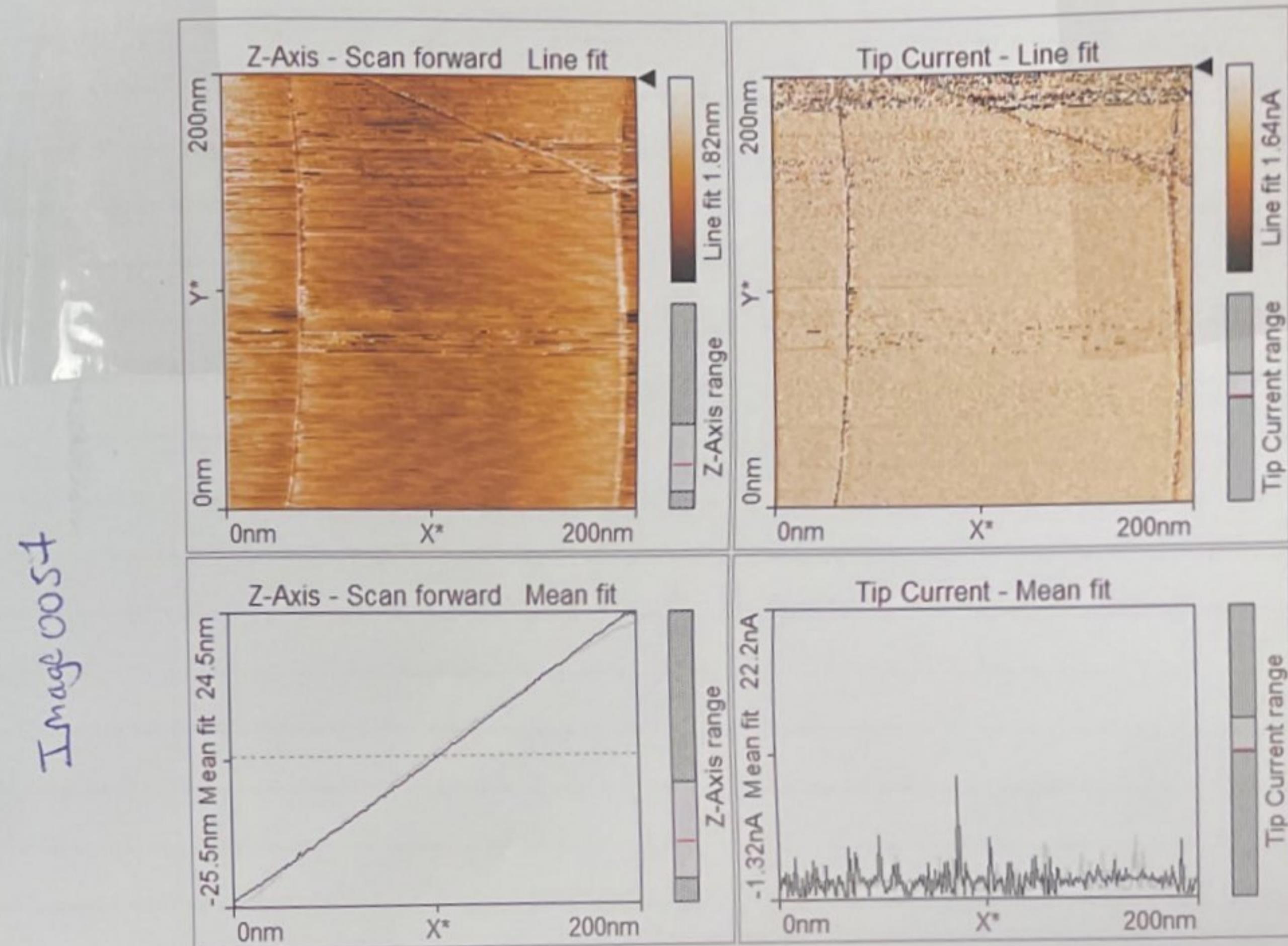
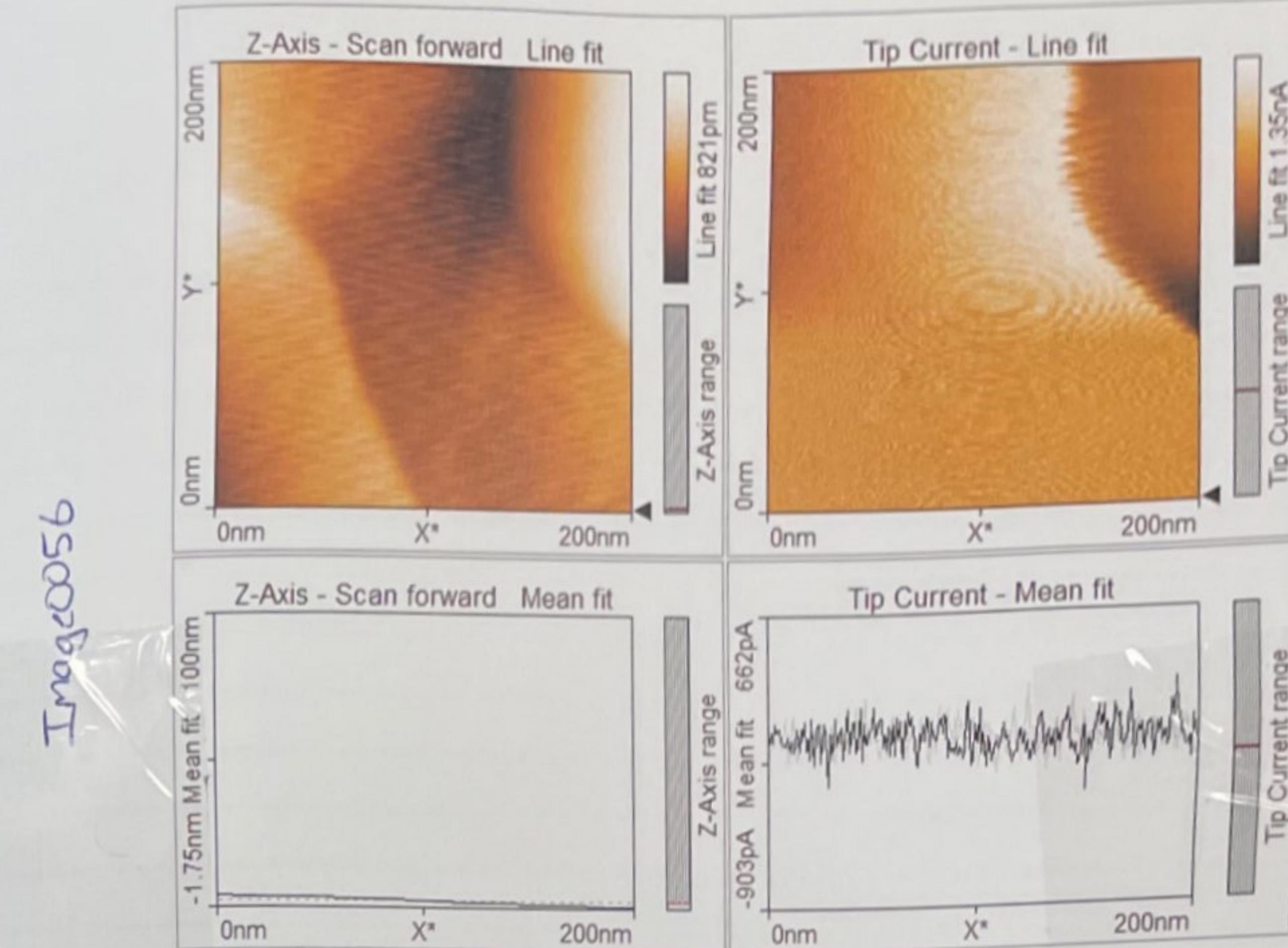
Image 0051



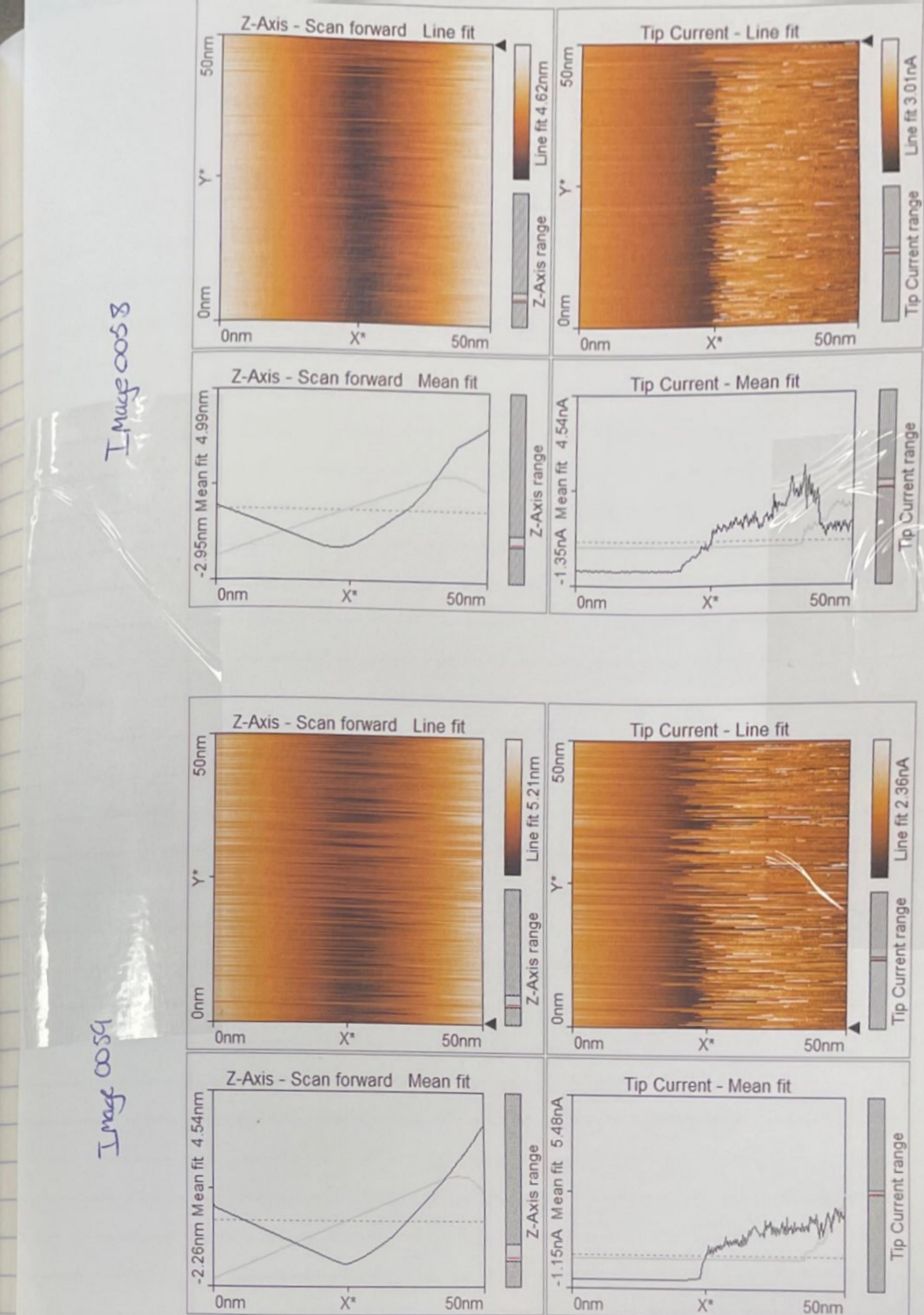




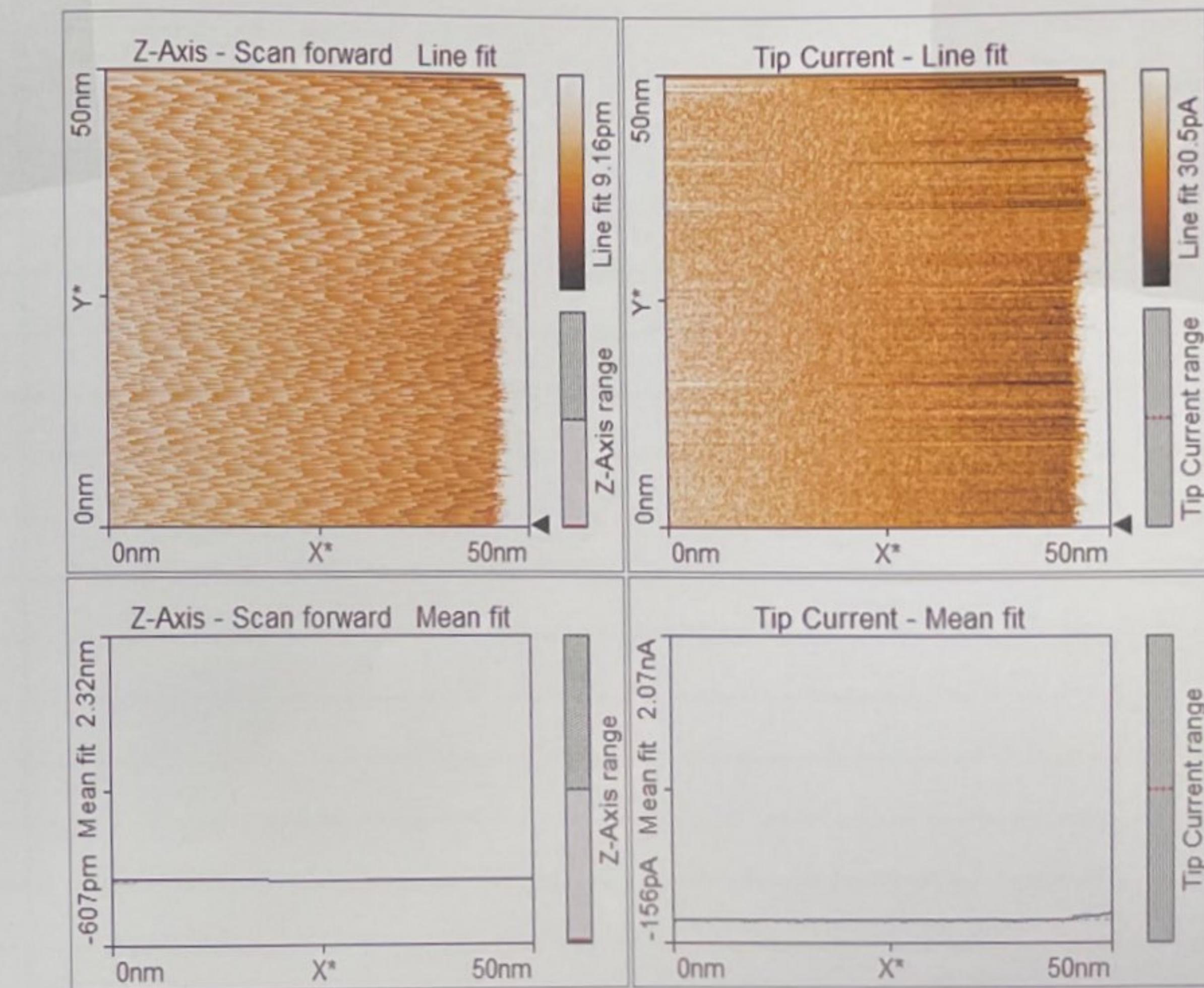
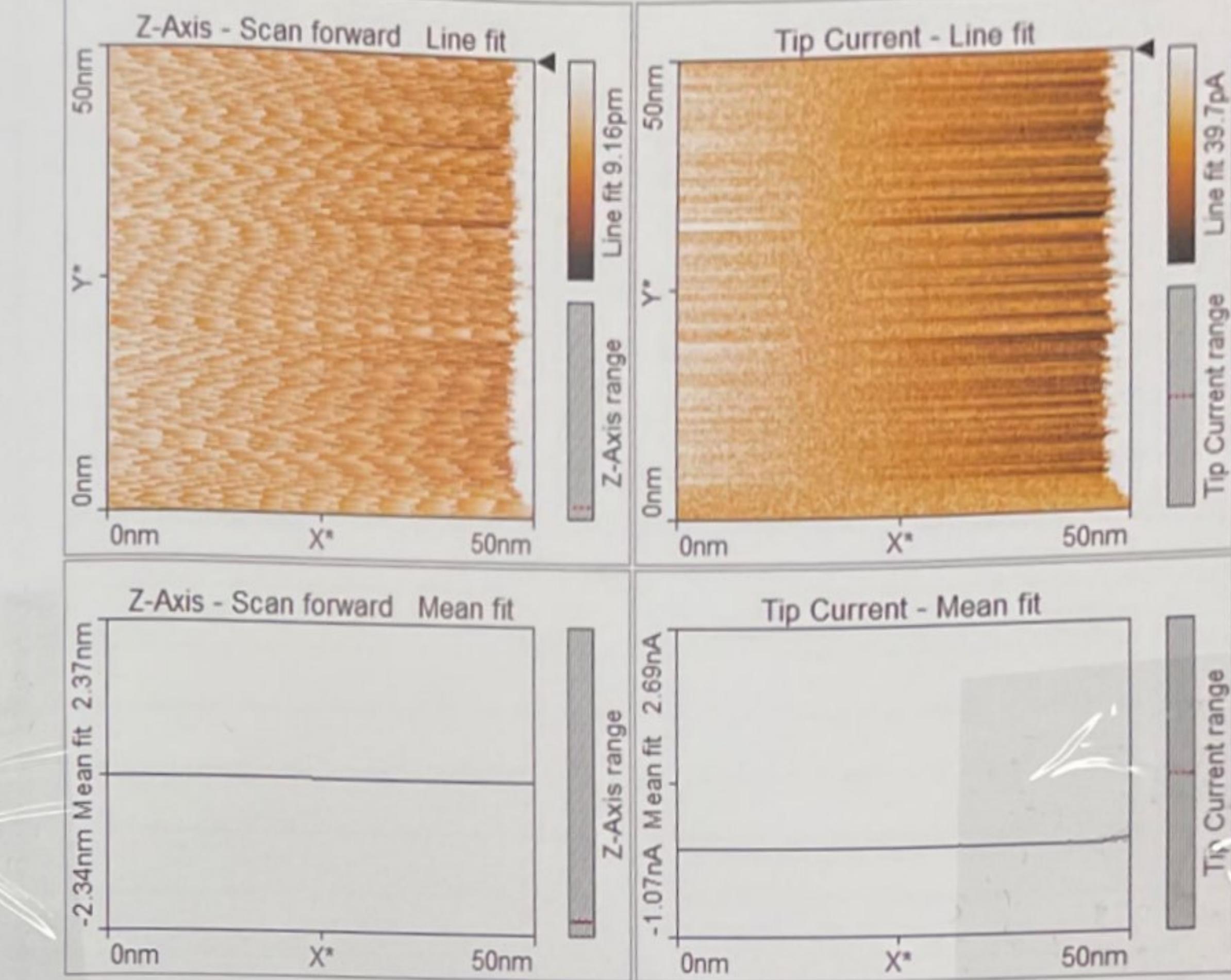
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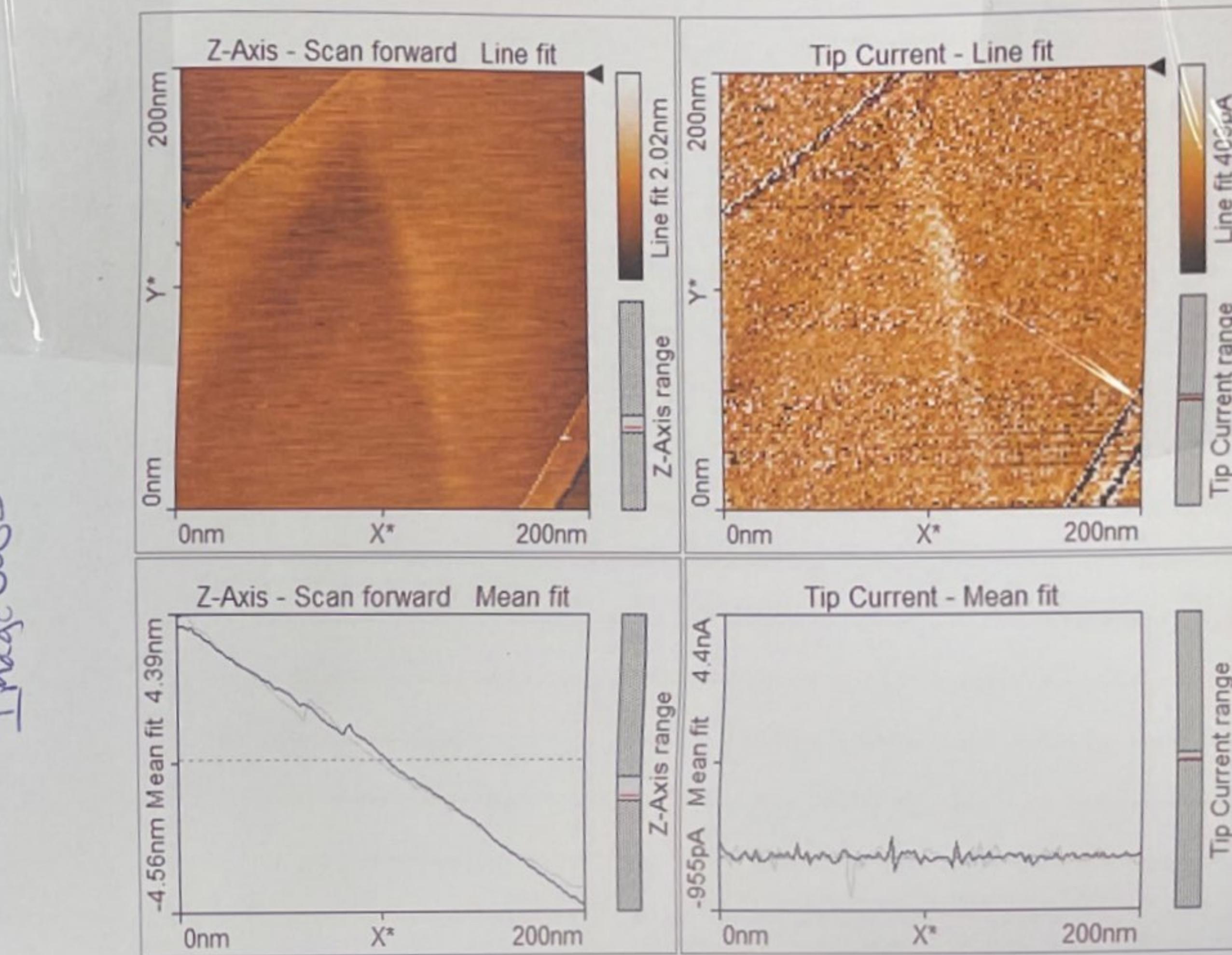
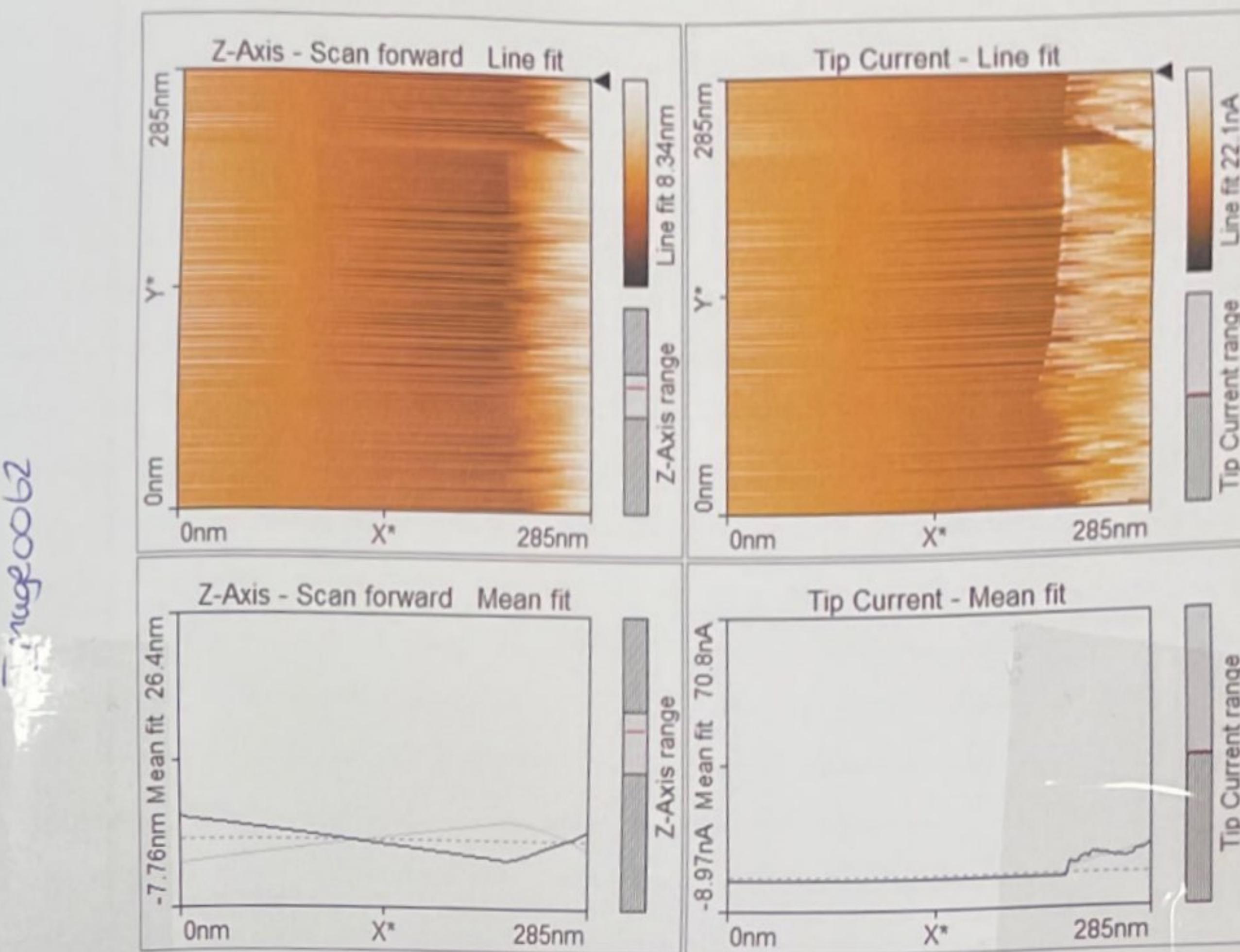


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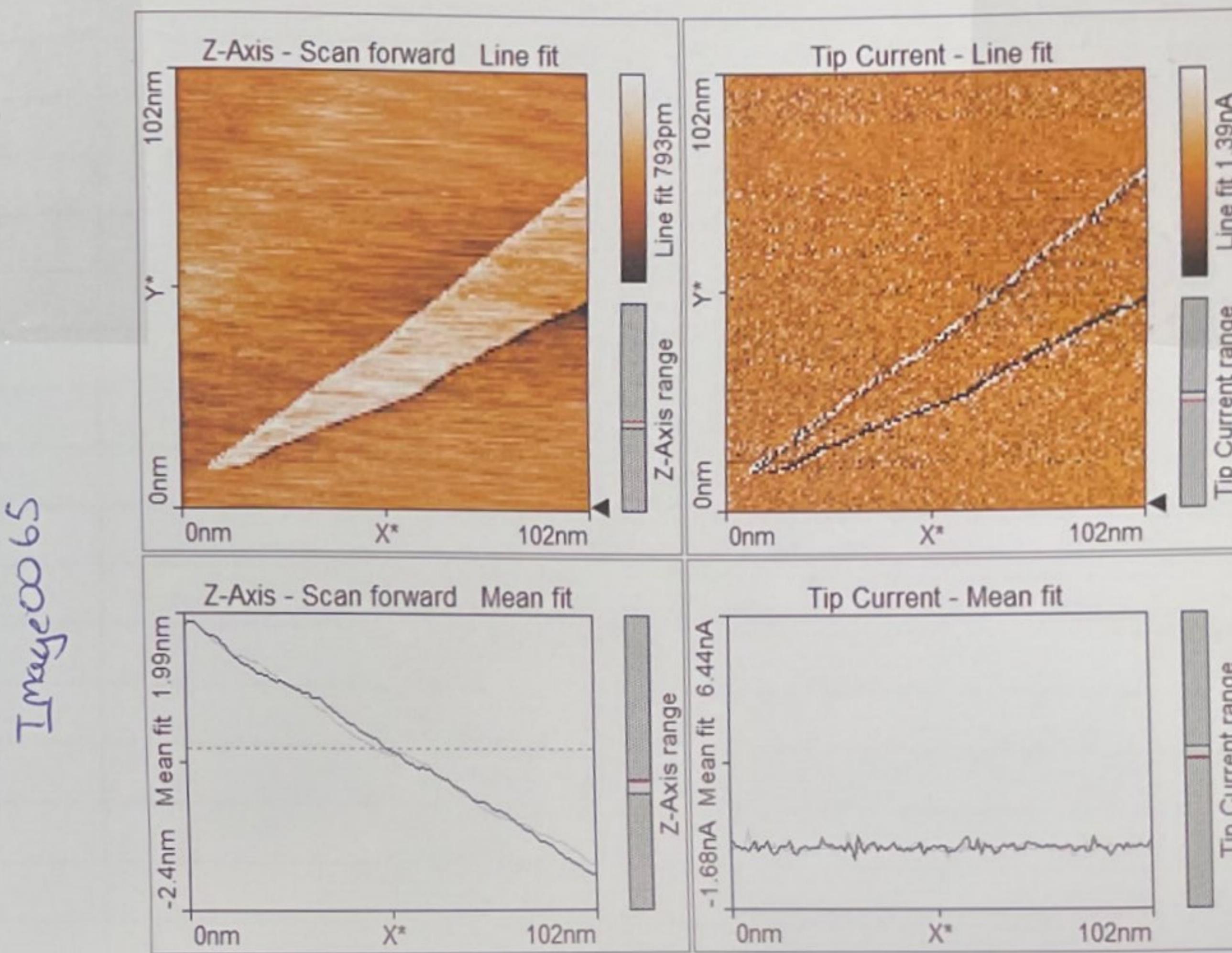
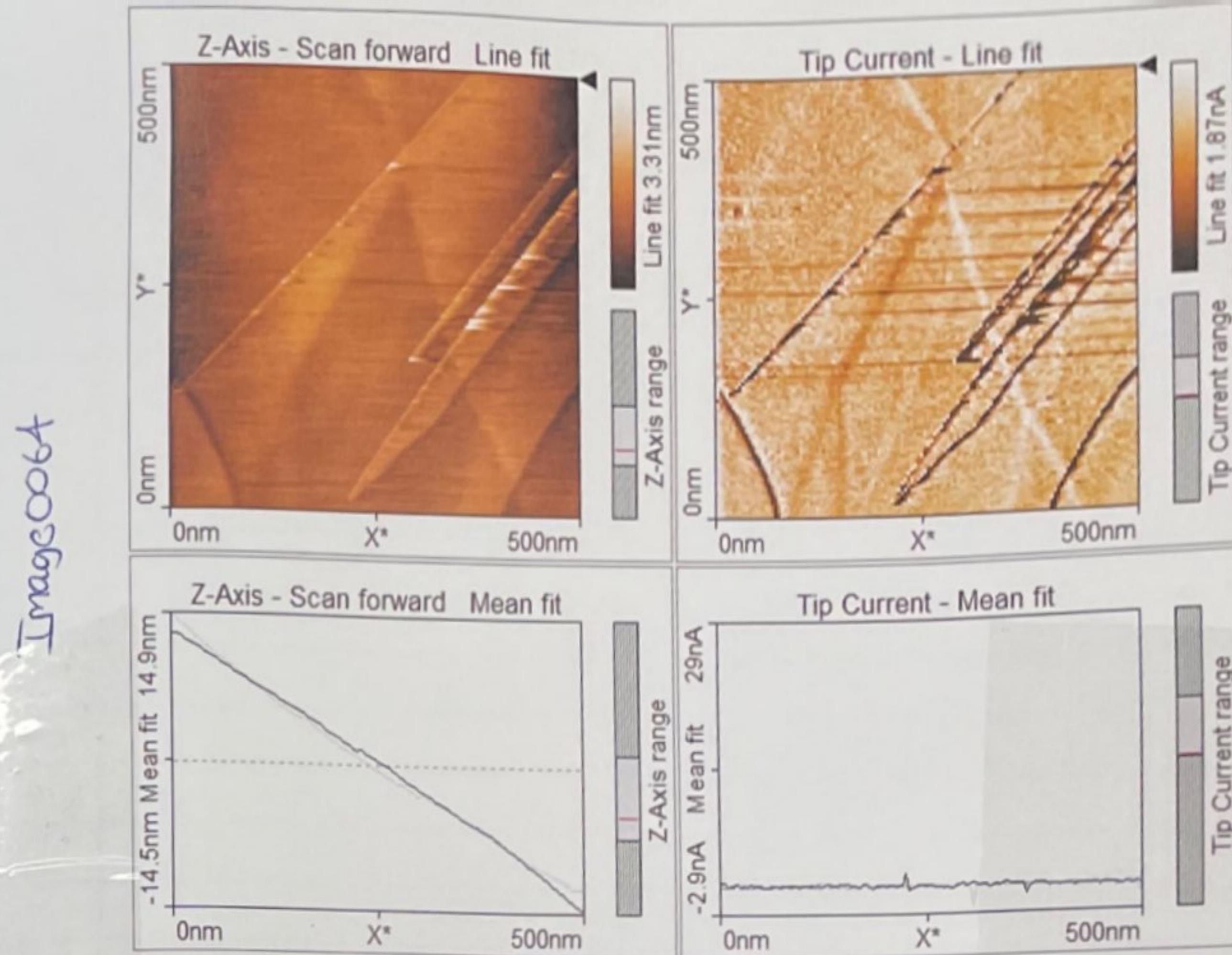


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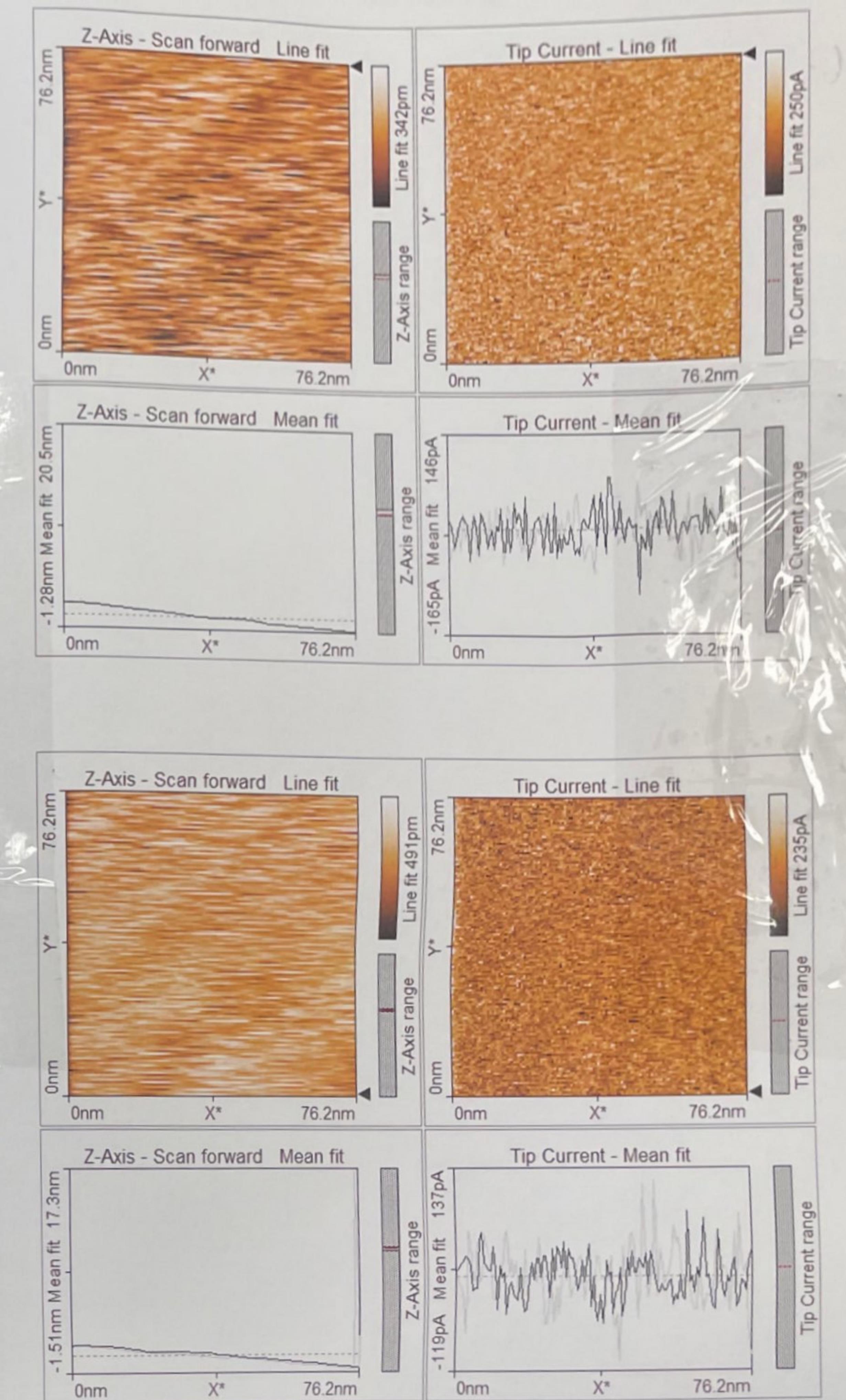
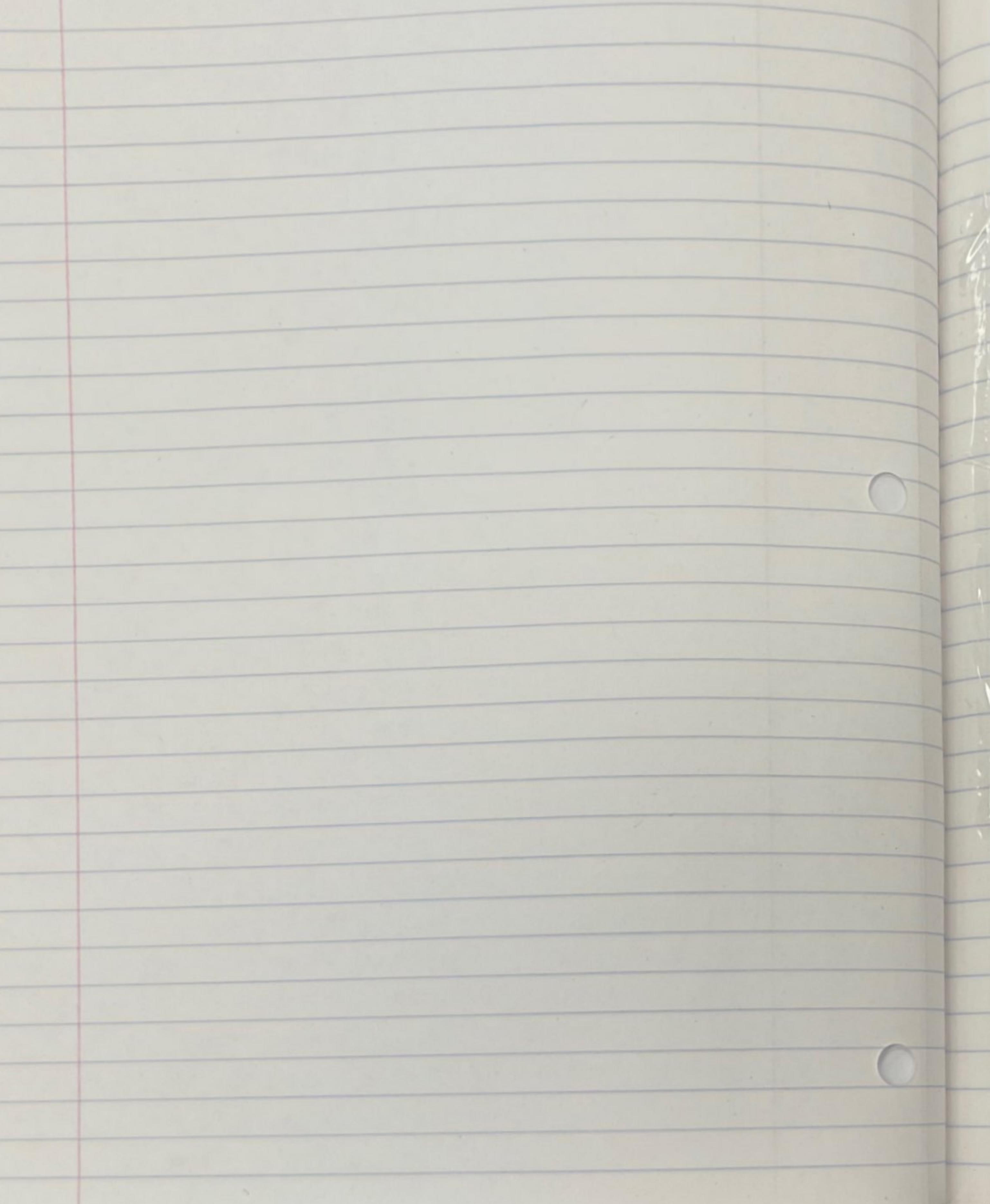
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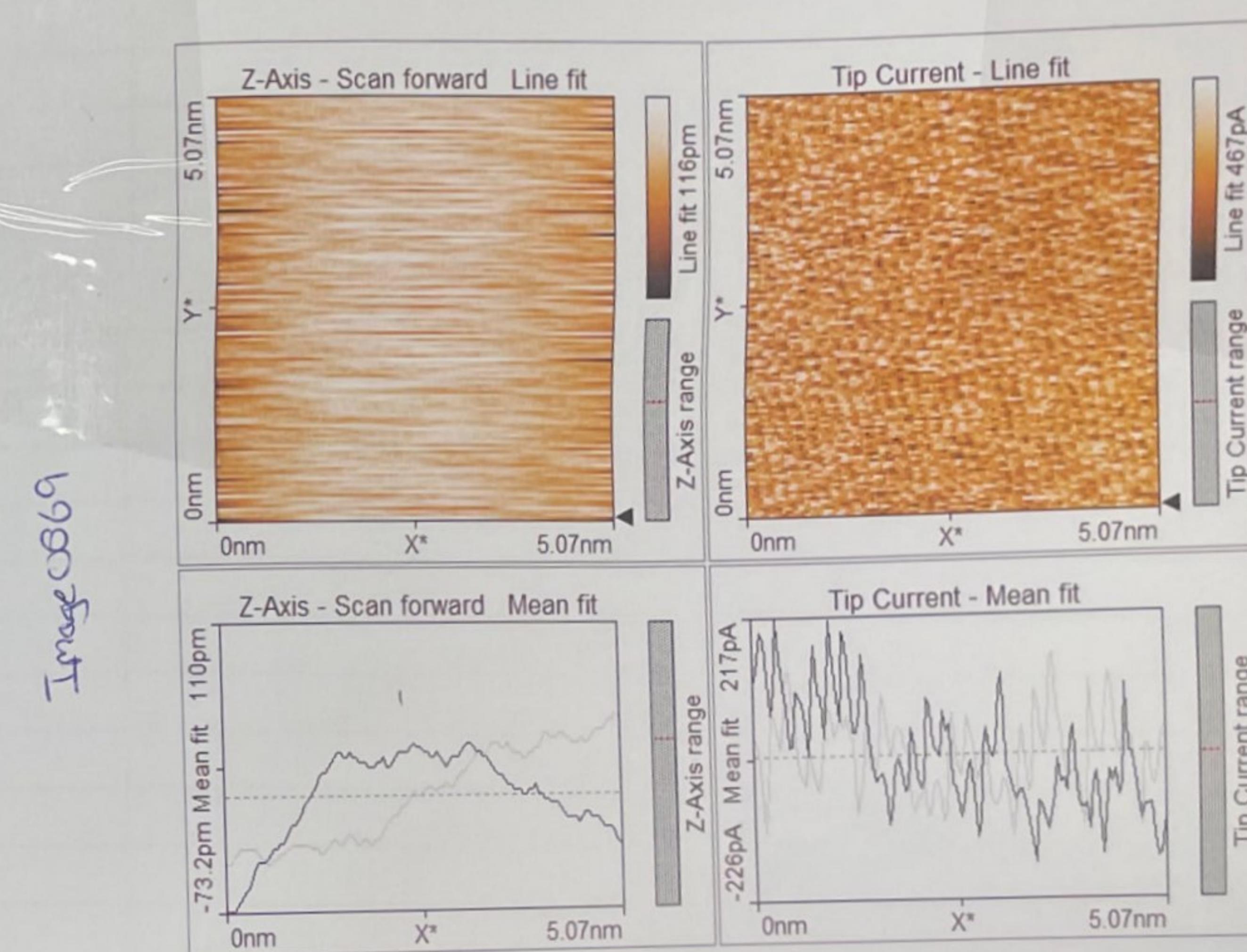
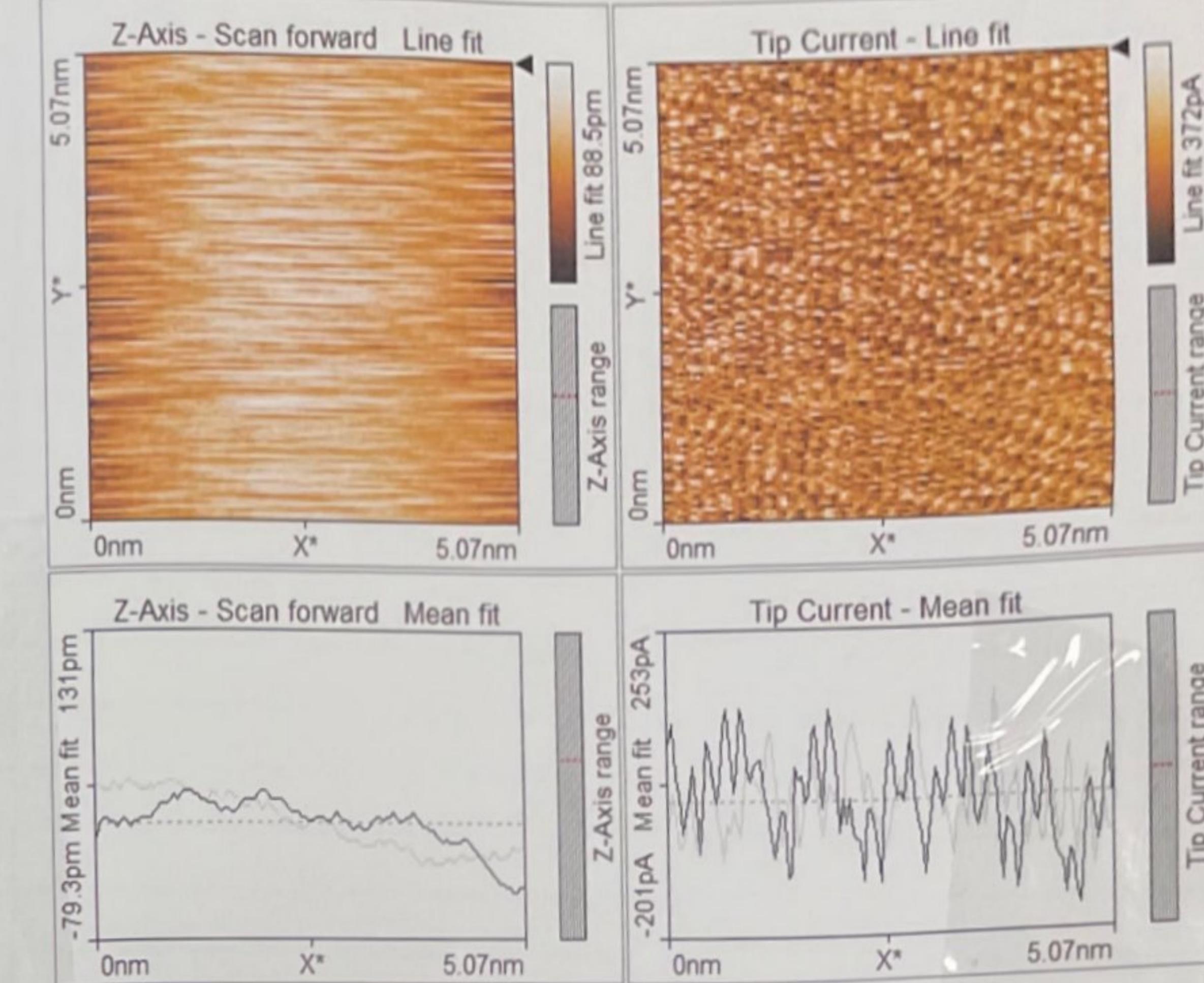
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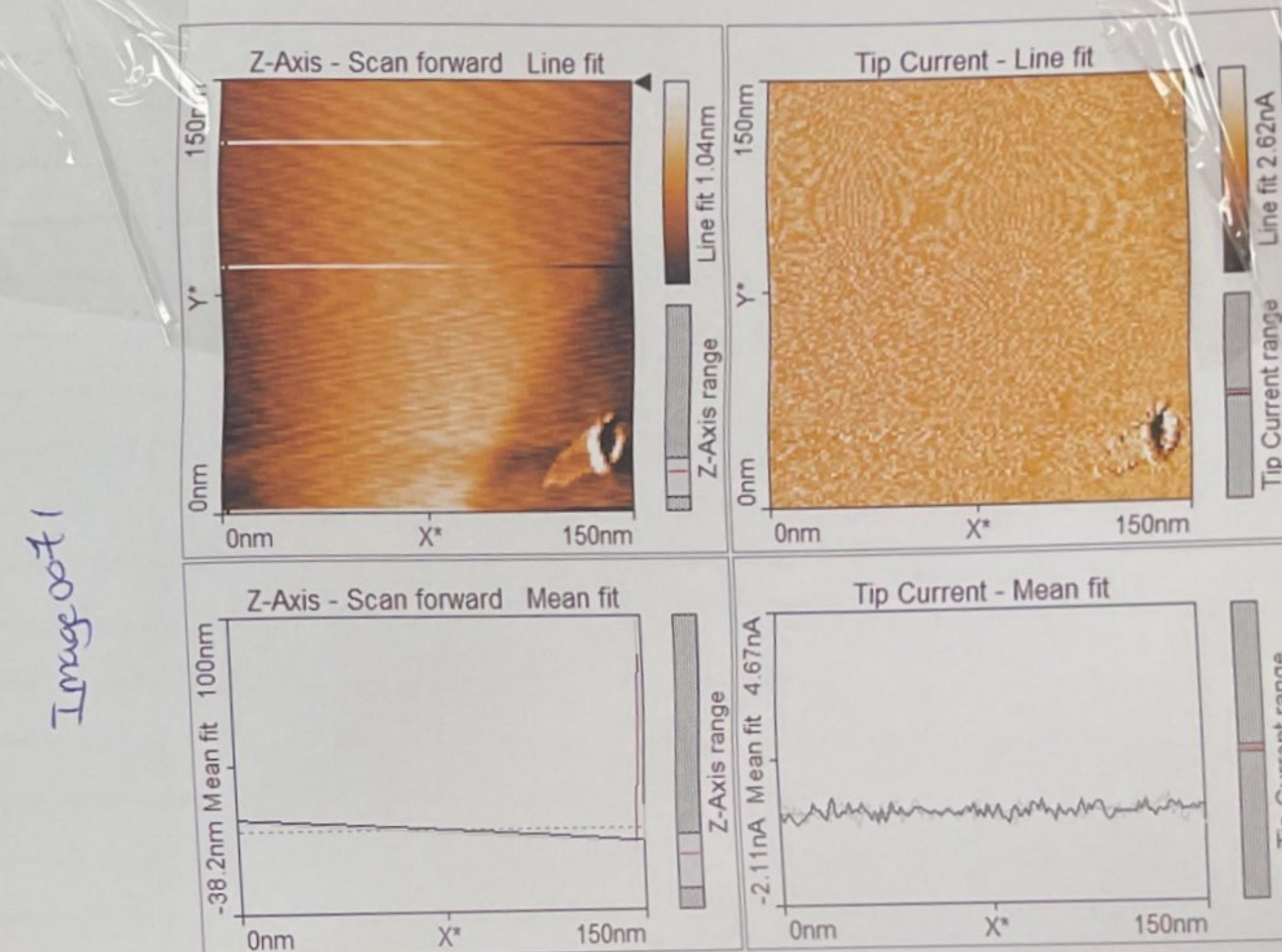
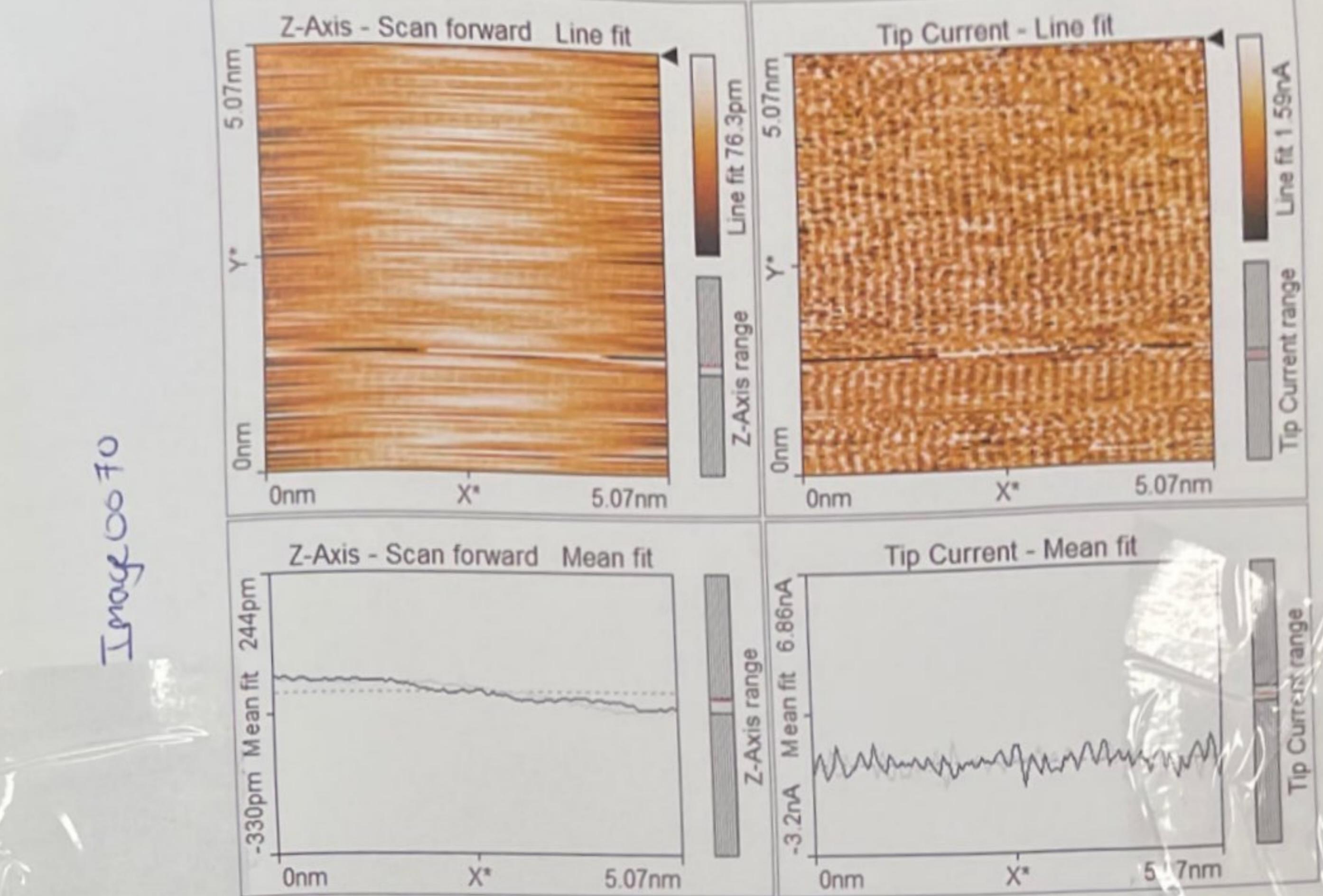
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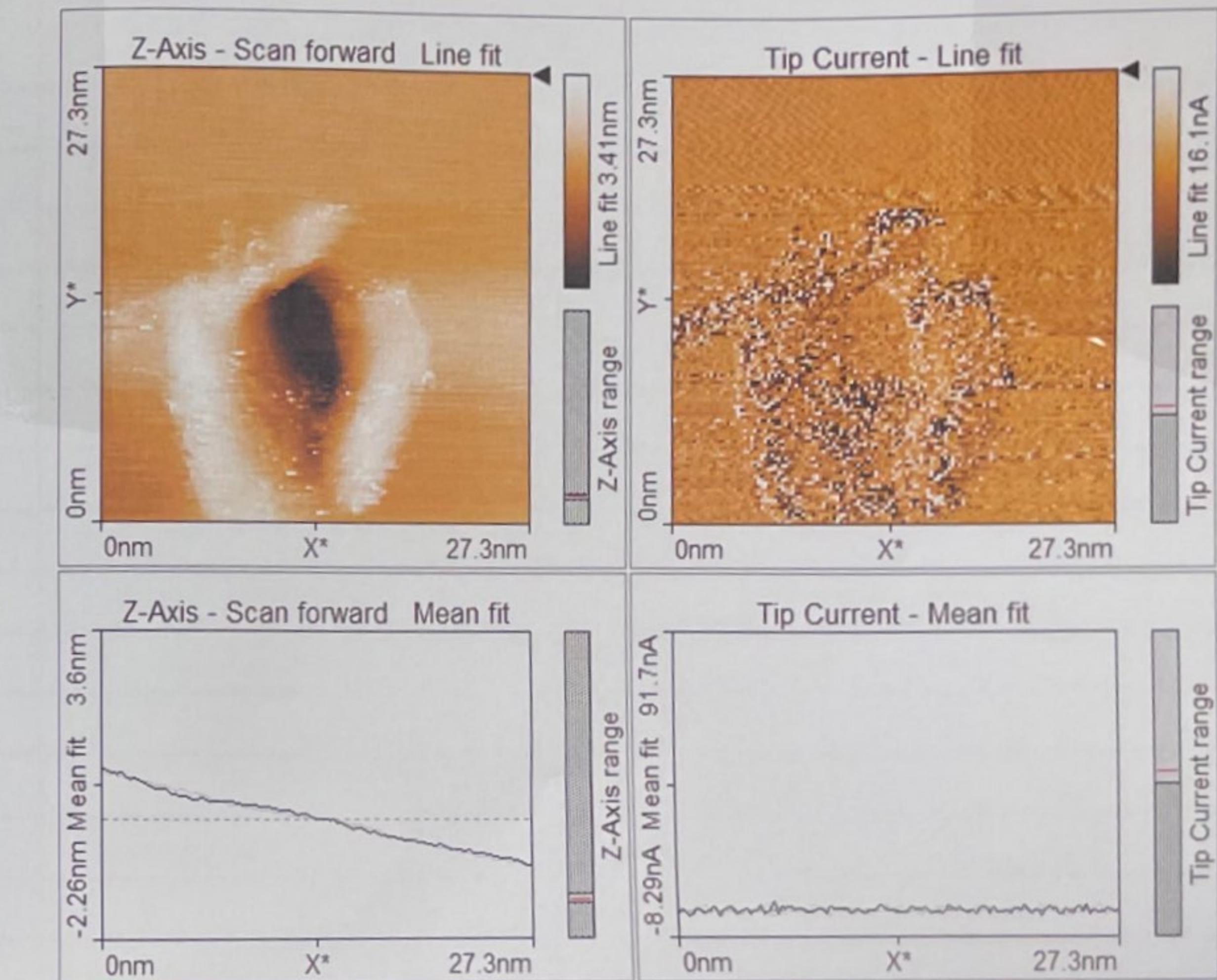
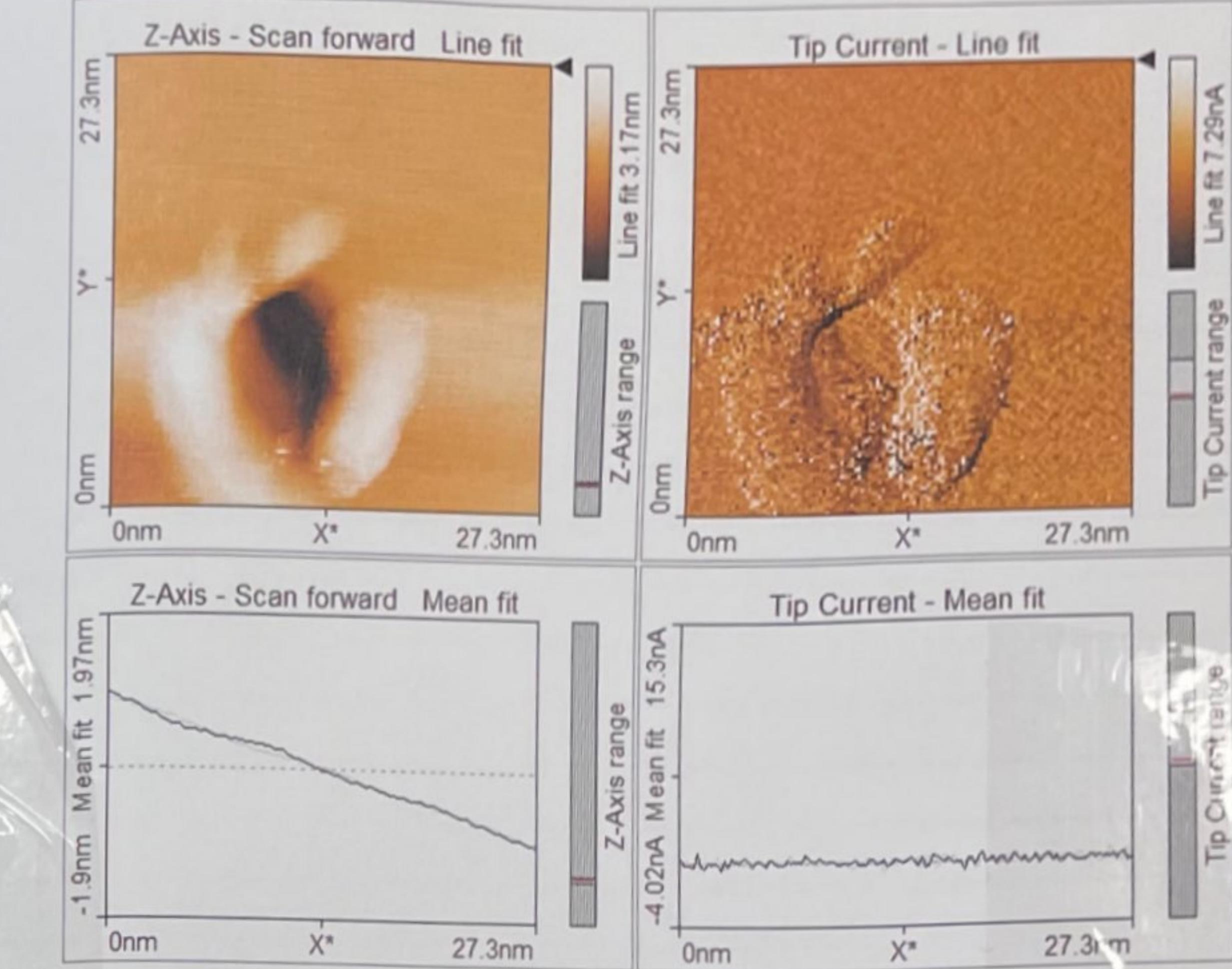
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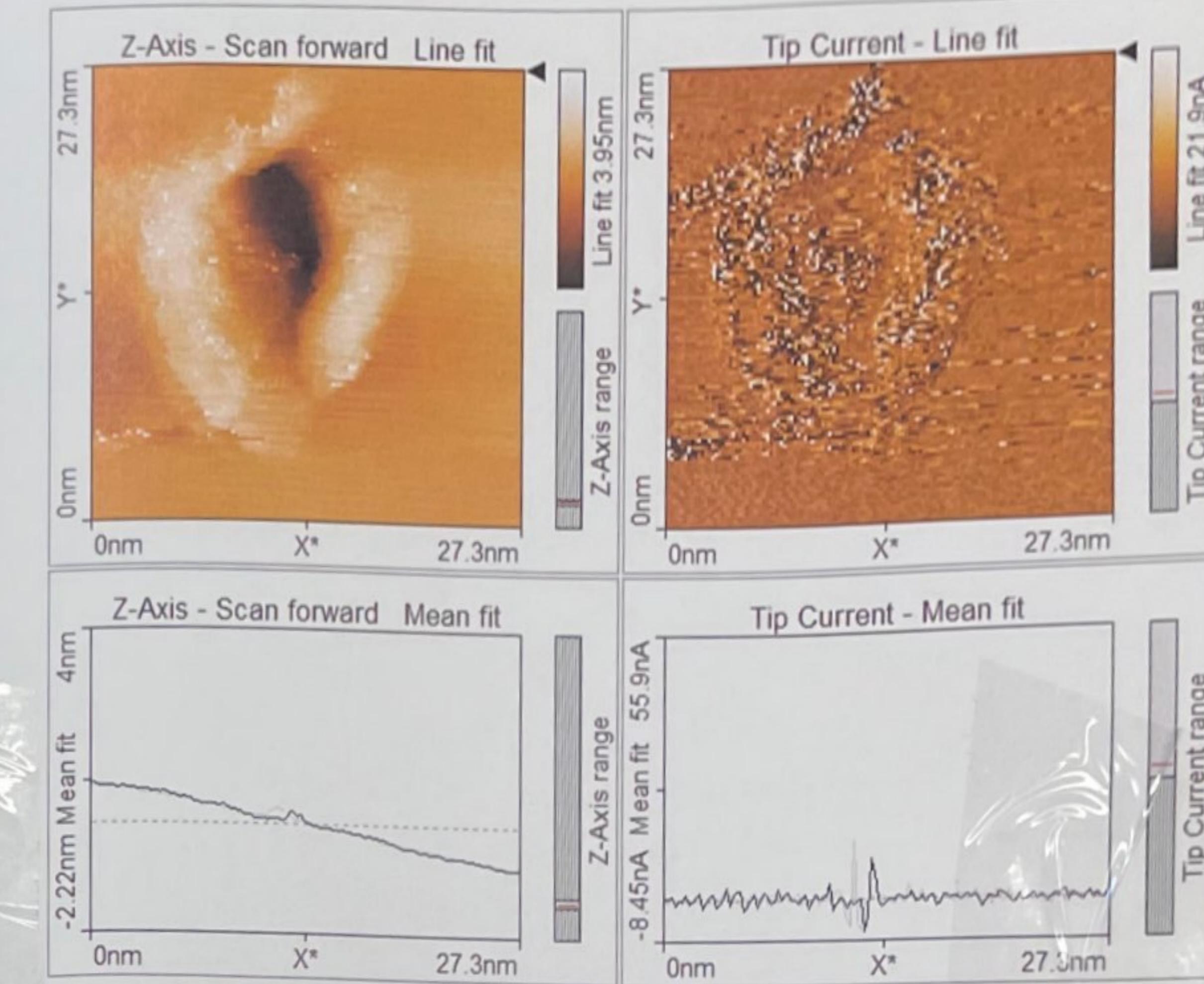


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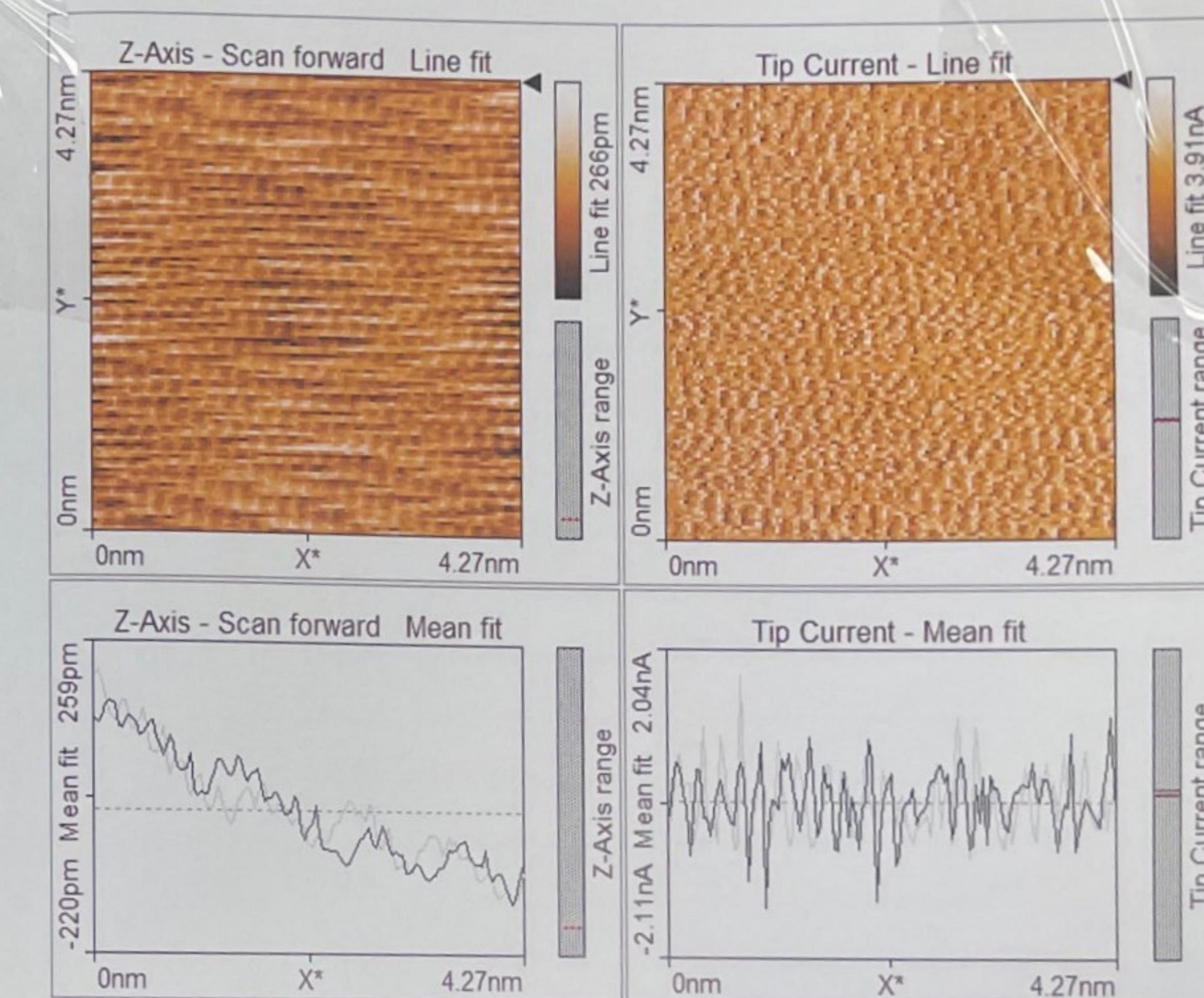
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Sept 10 1981



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Image 0076

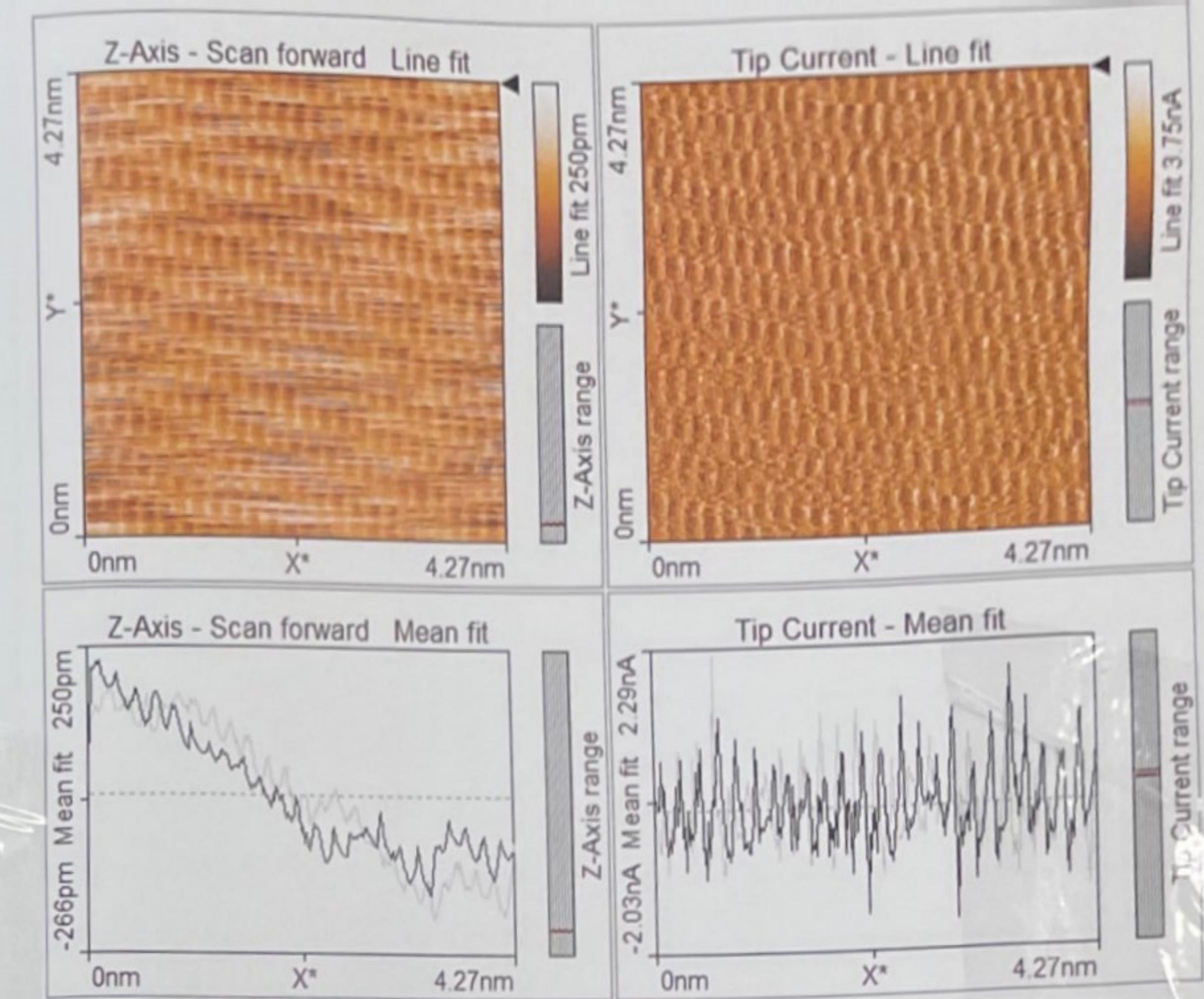
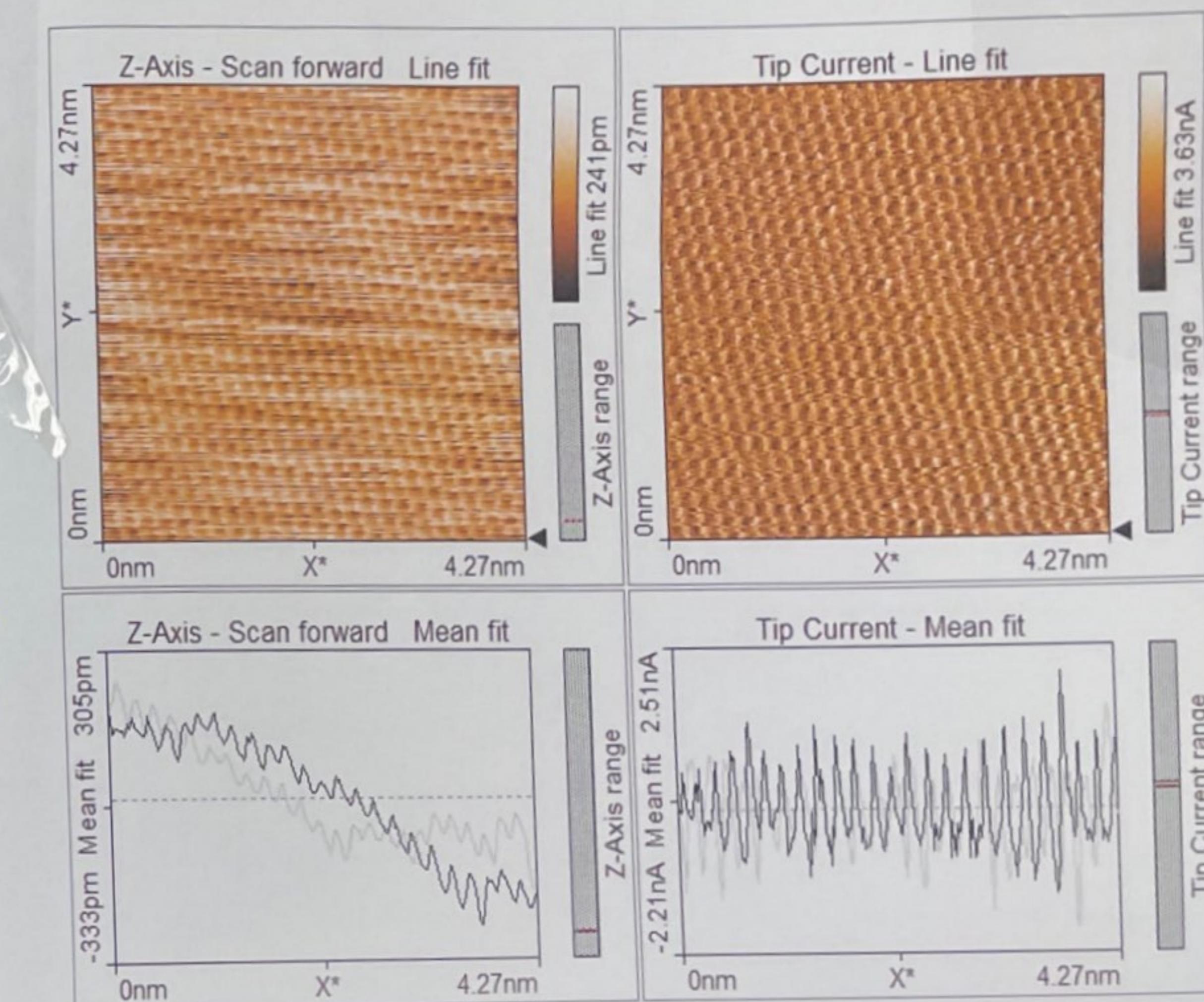


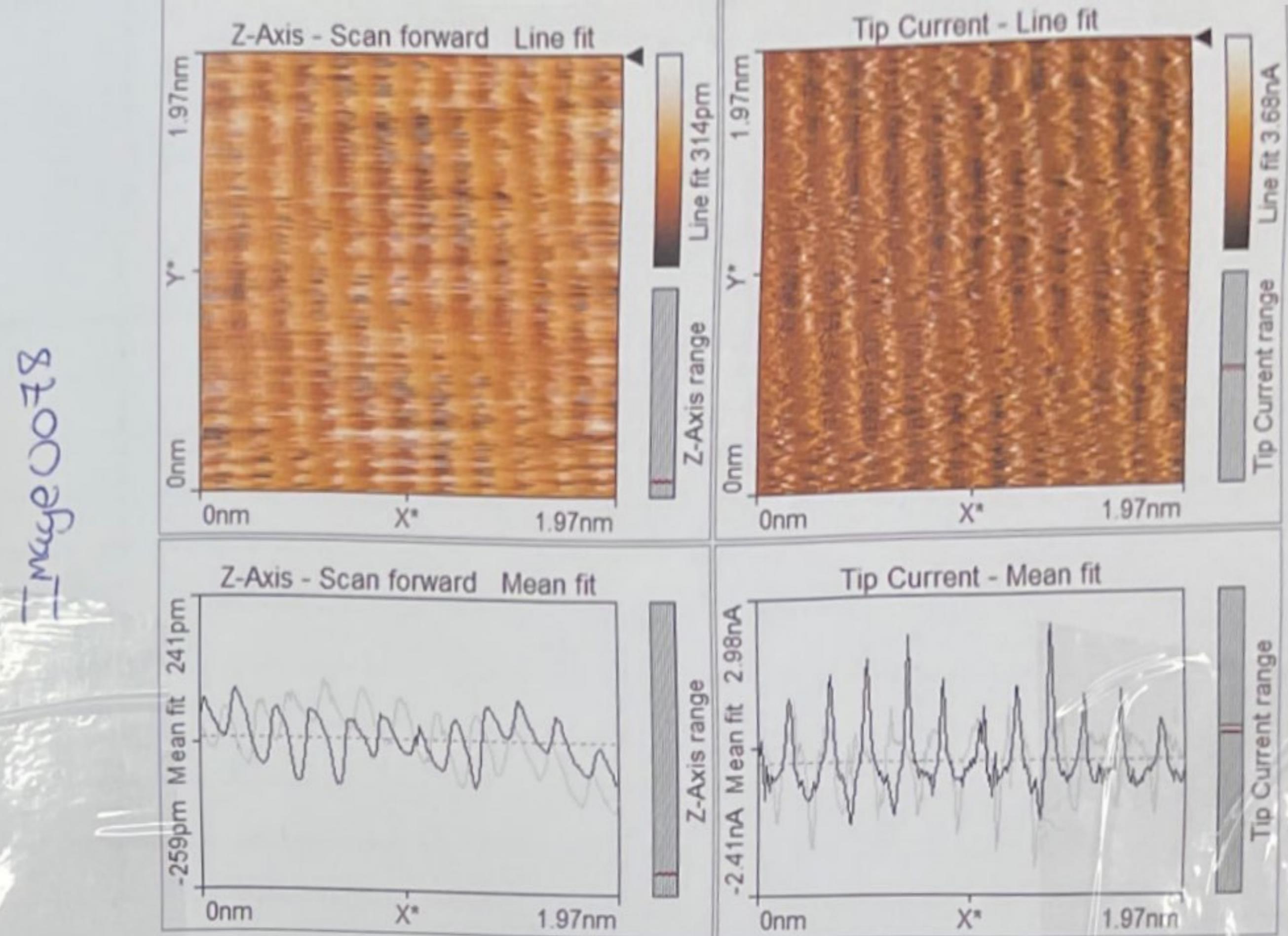
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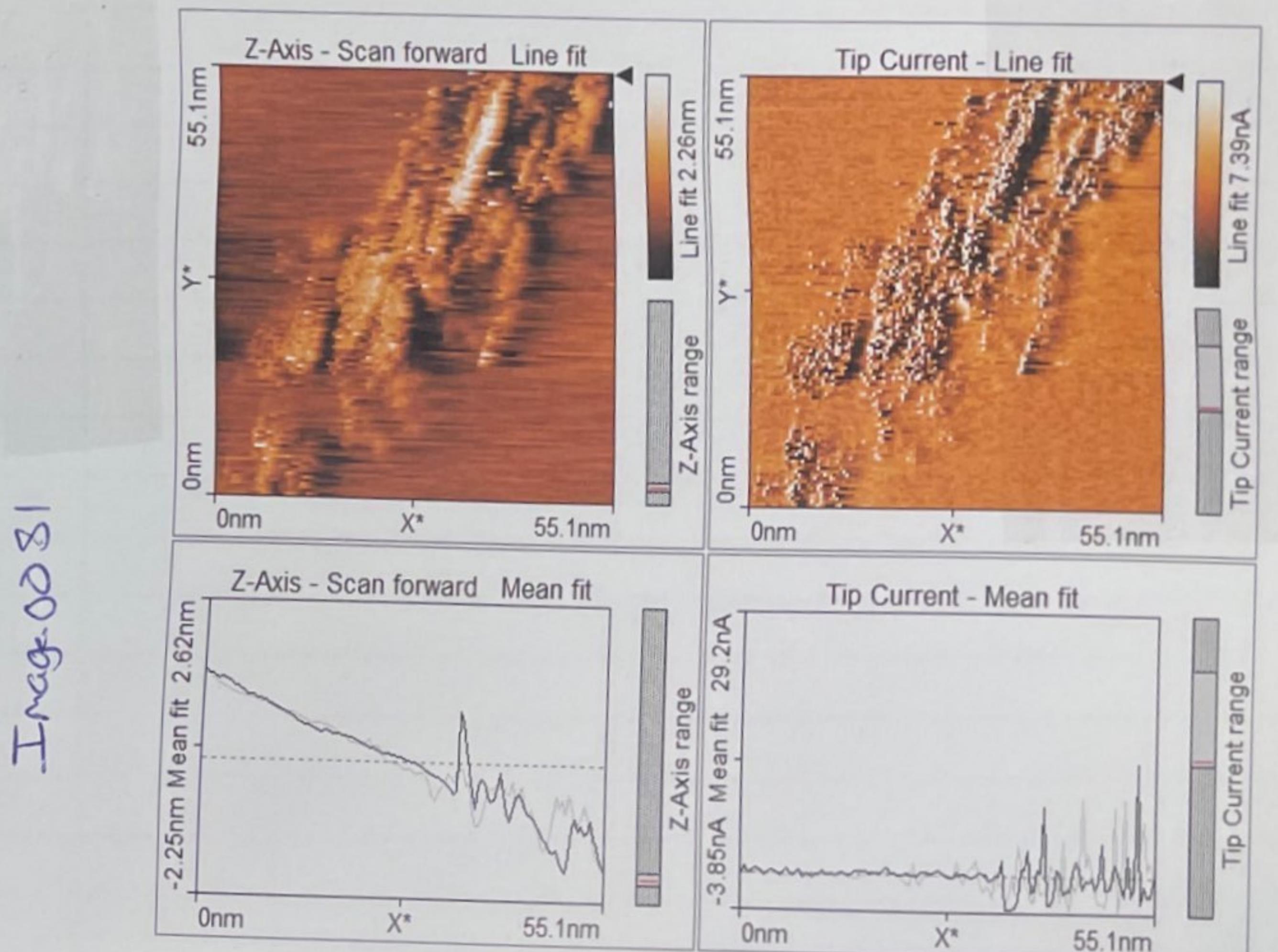
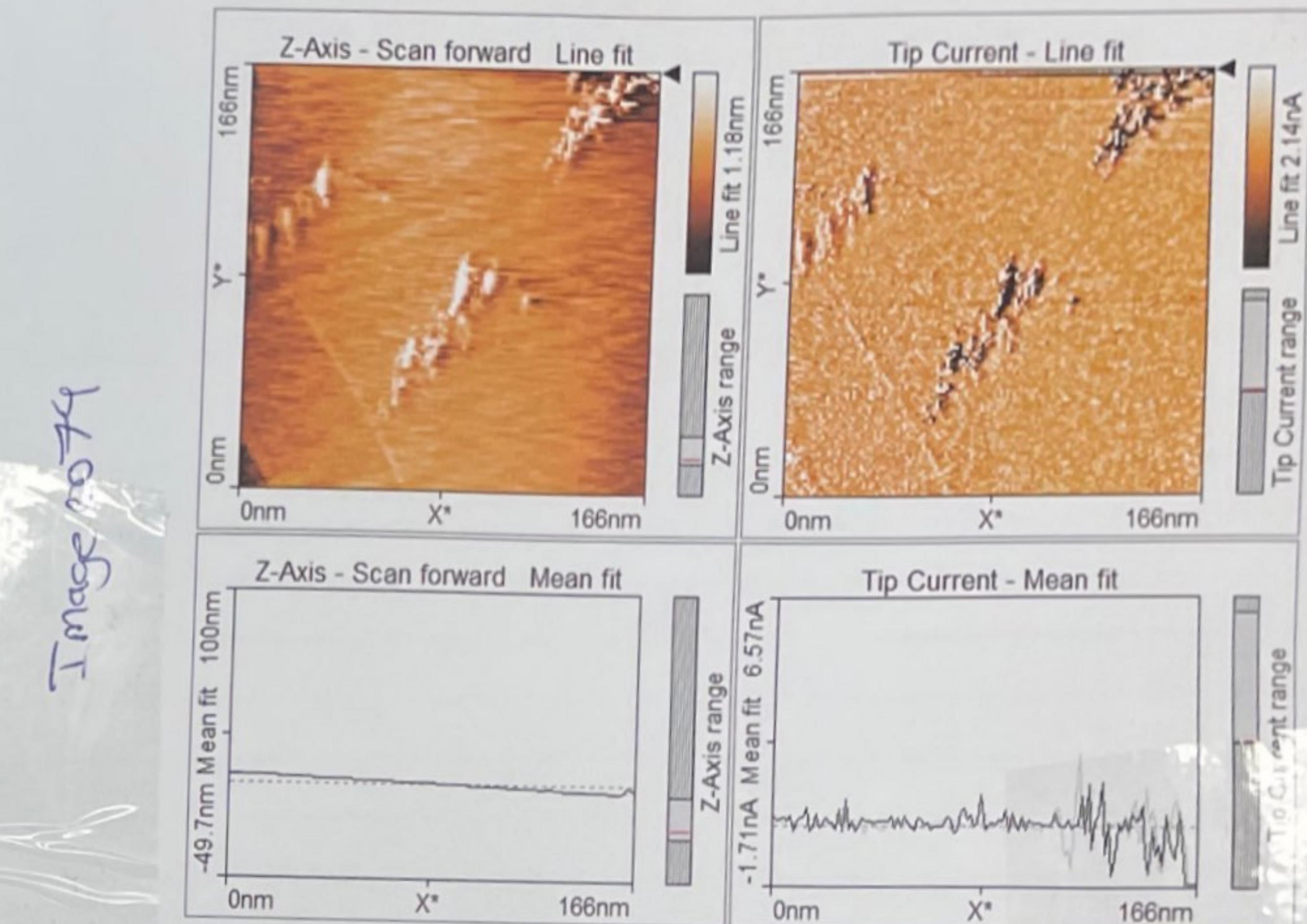
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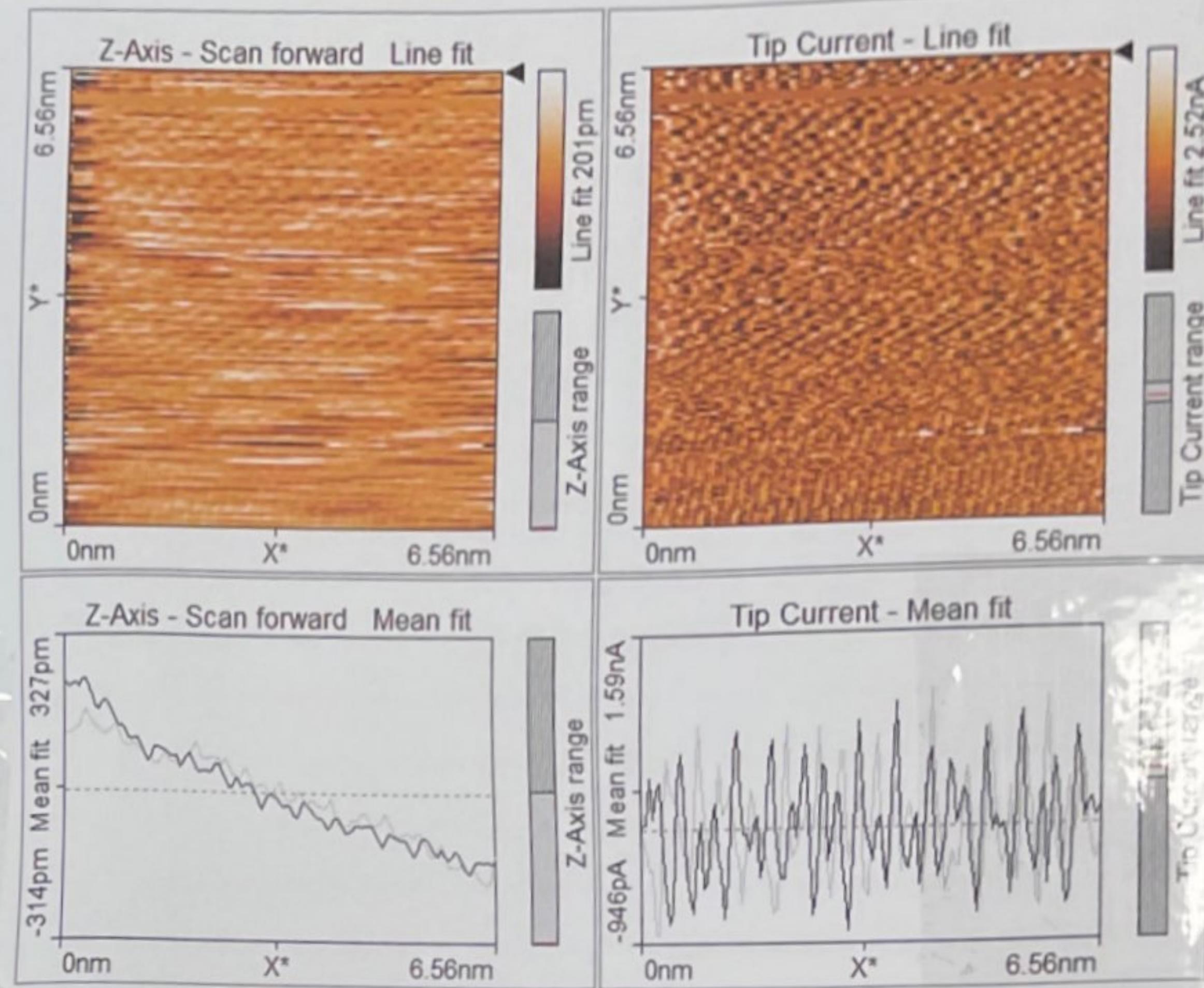
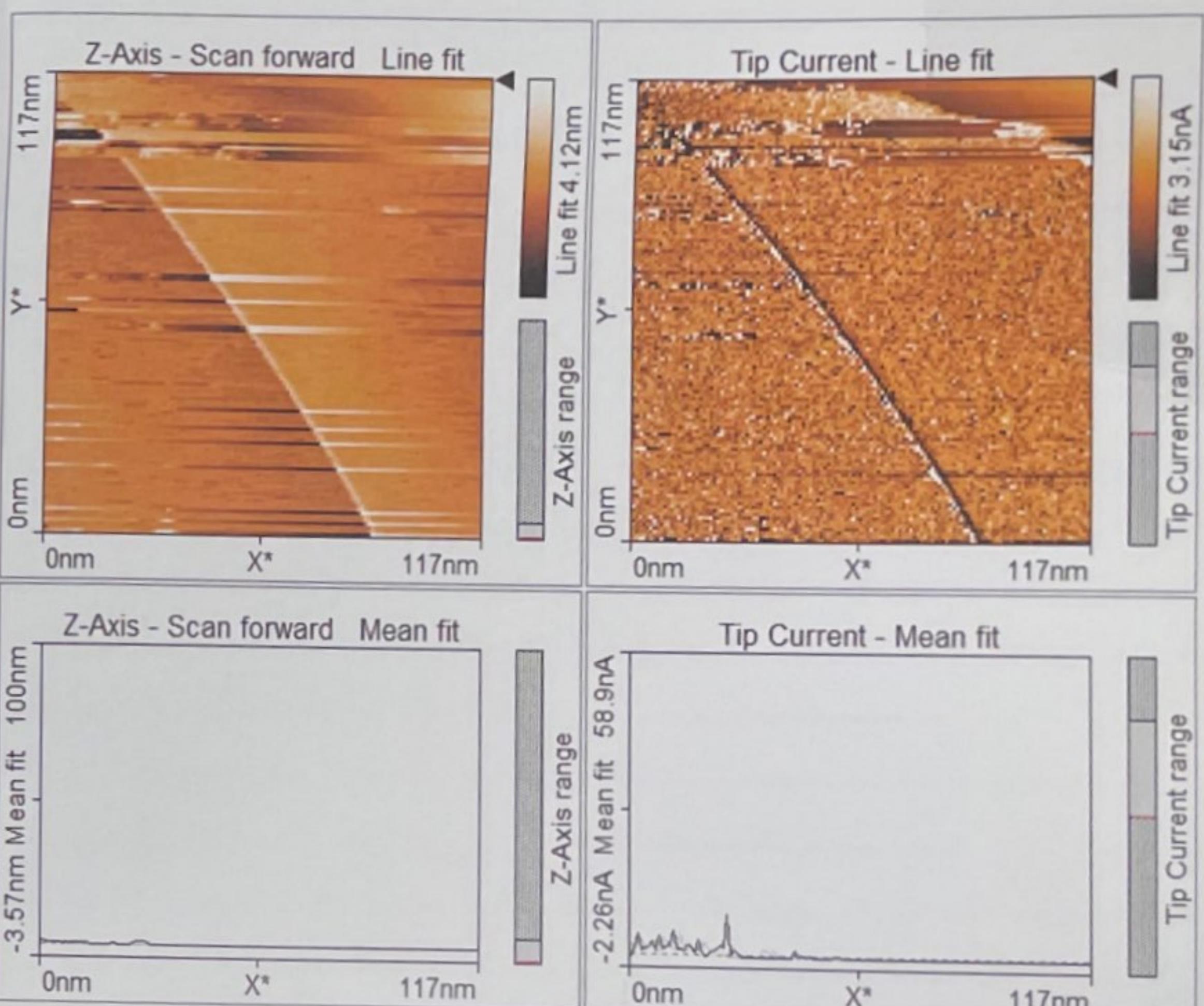


Image 0083



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Image 00 84

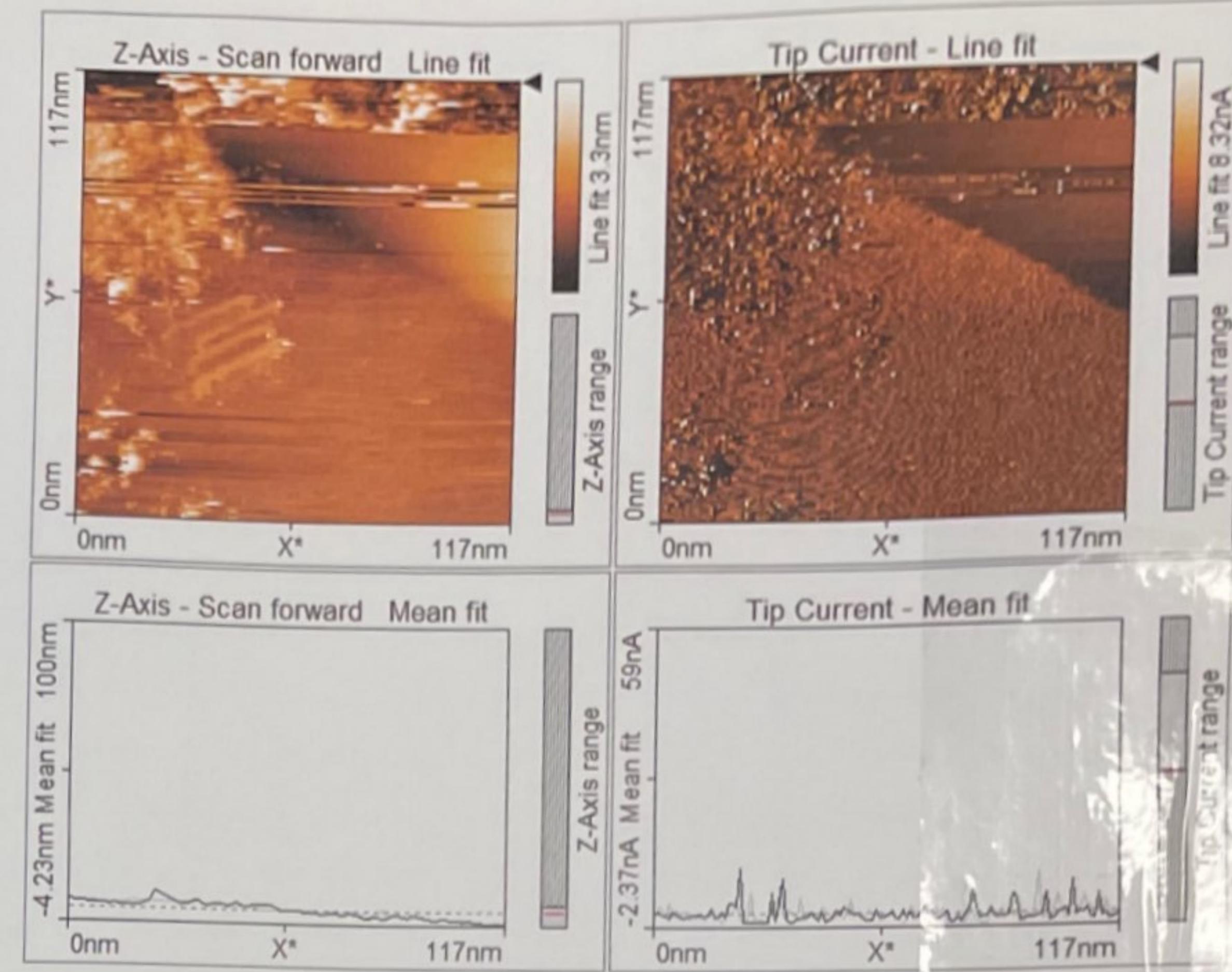
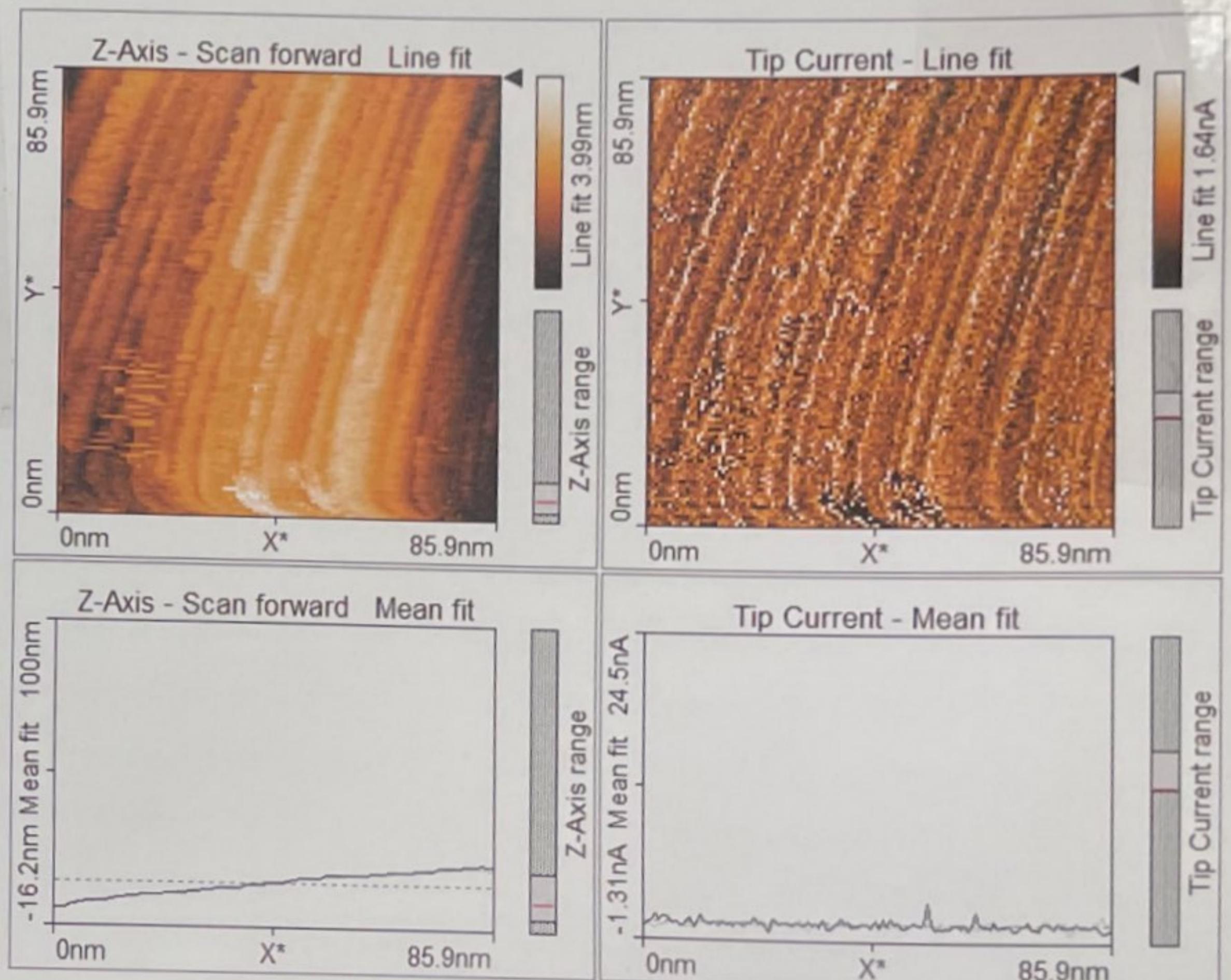


Image 00 85



Scanned with CamScanner

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Image 086

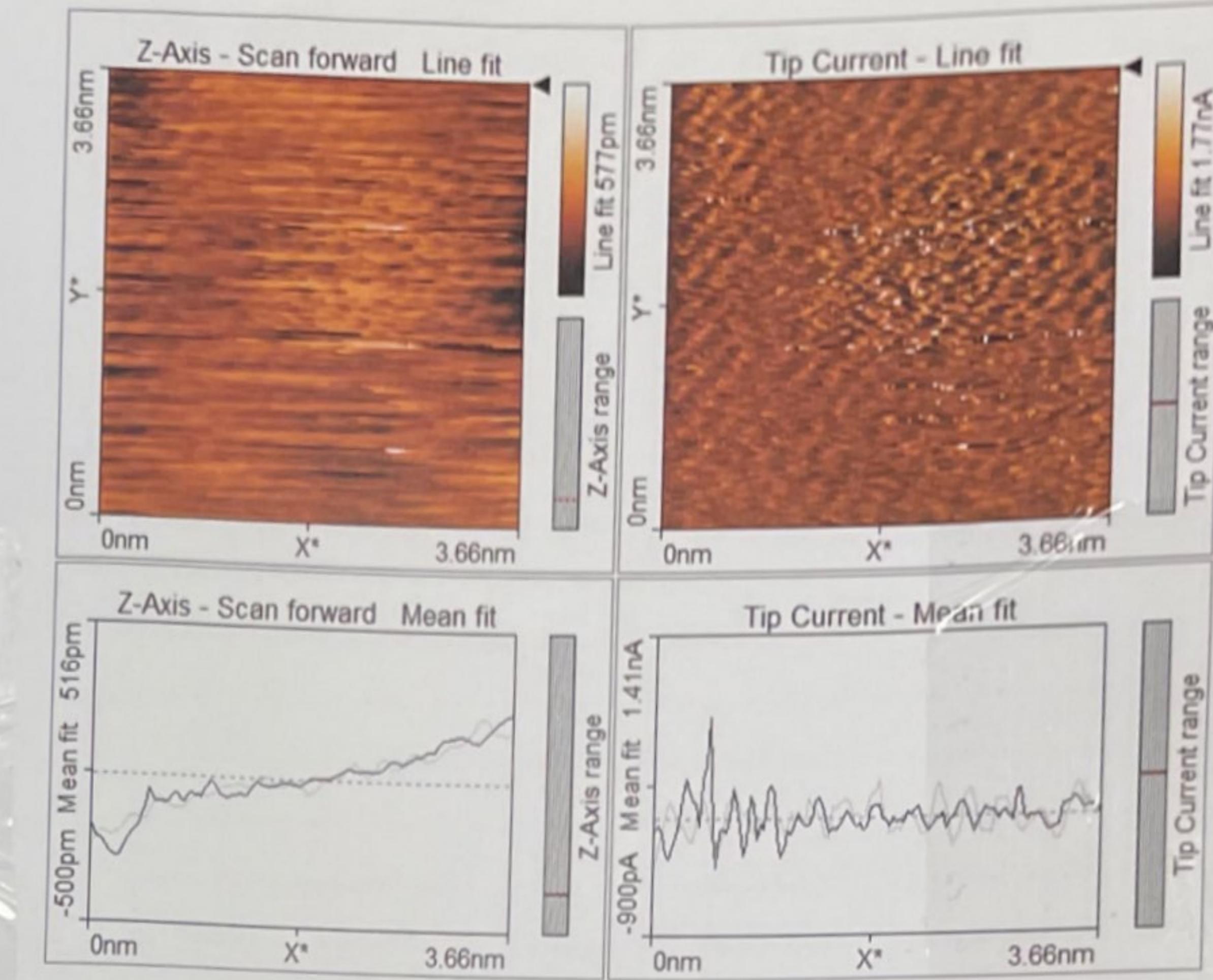
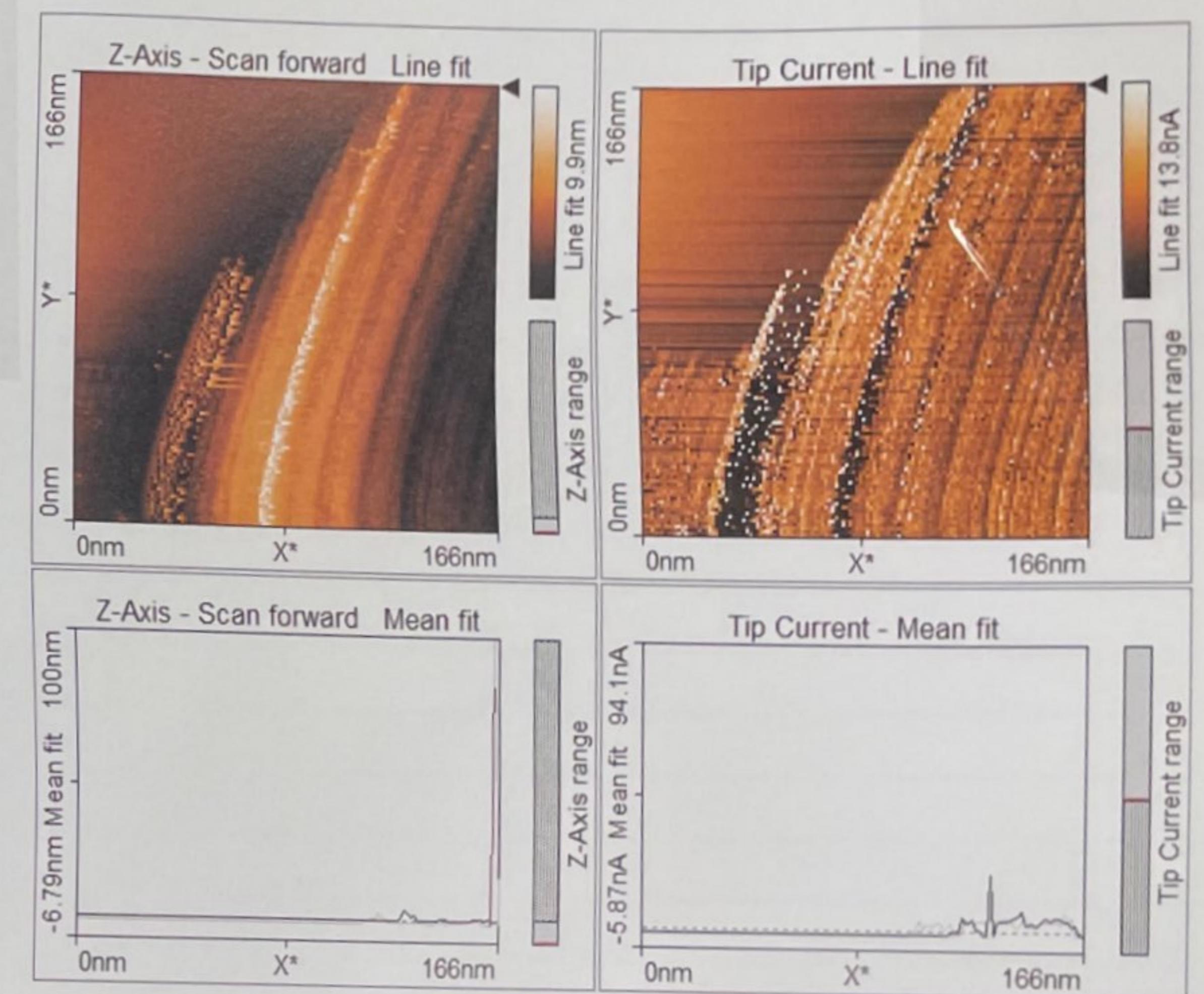
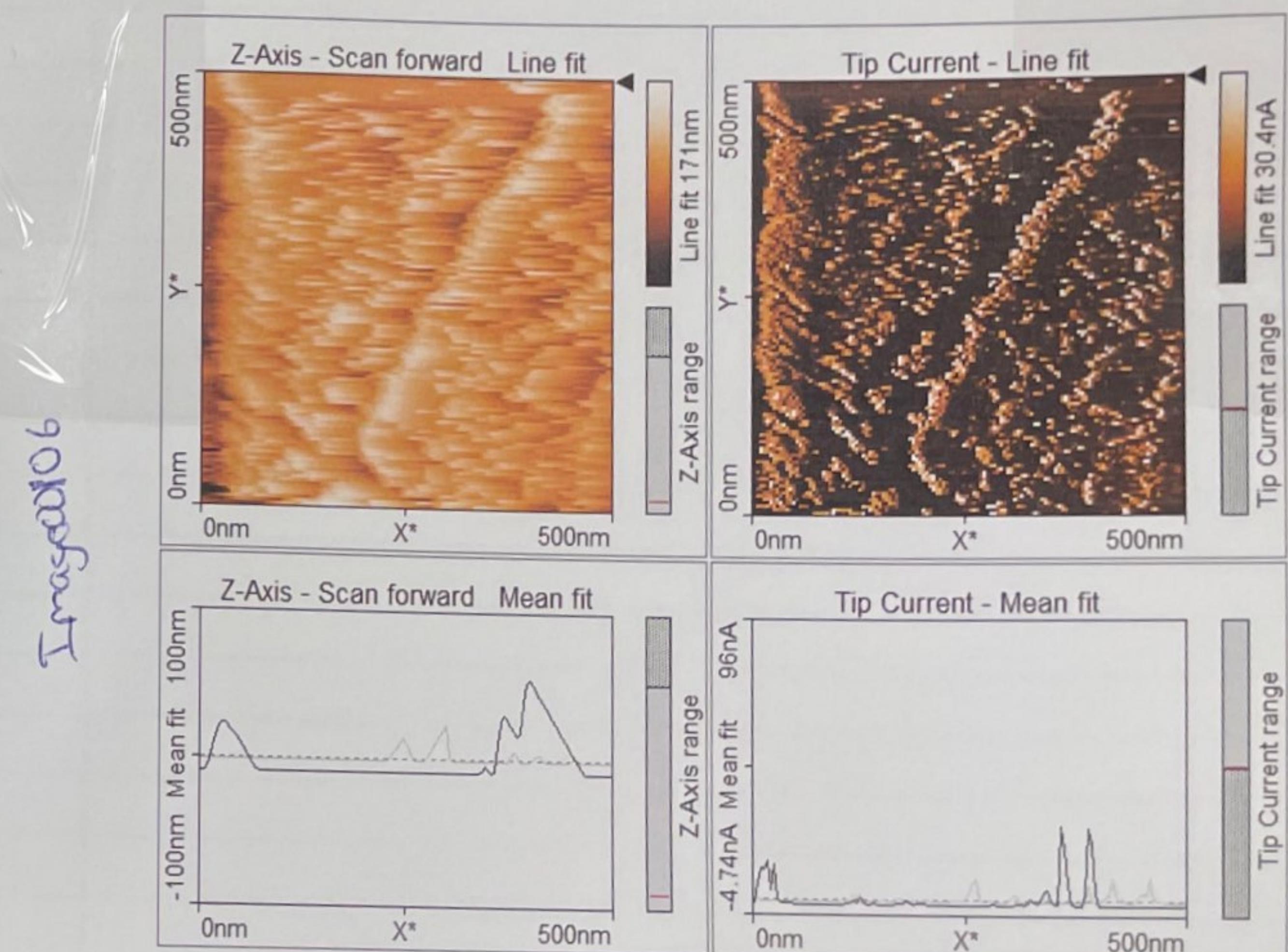
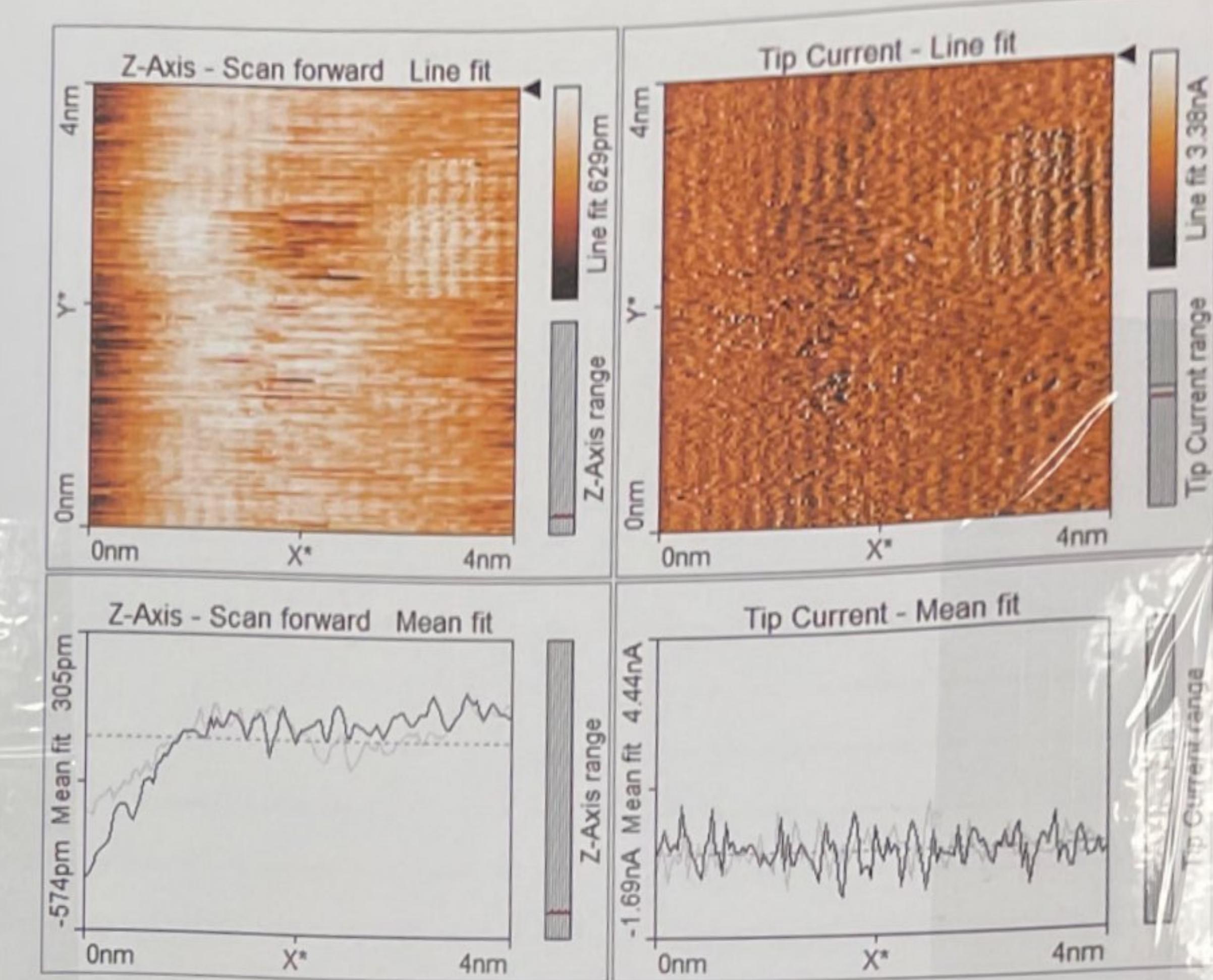


Image 087



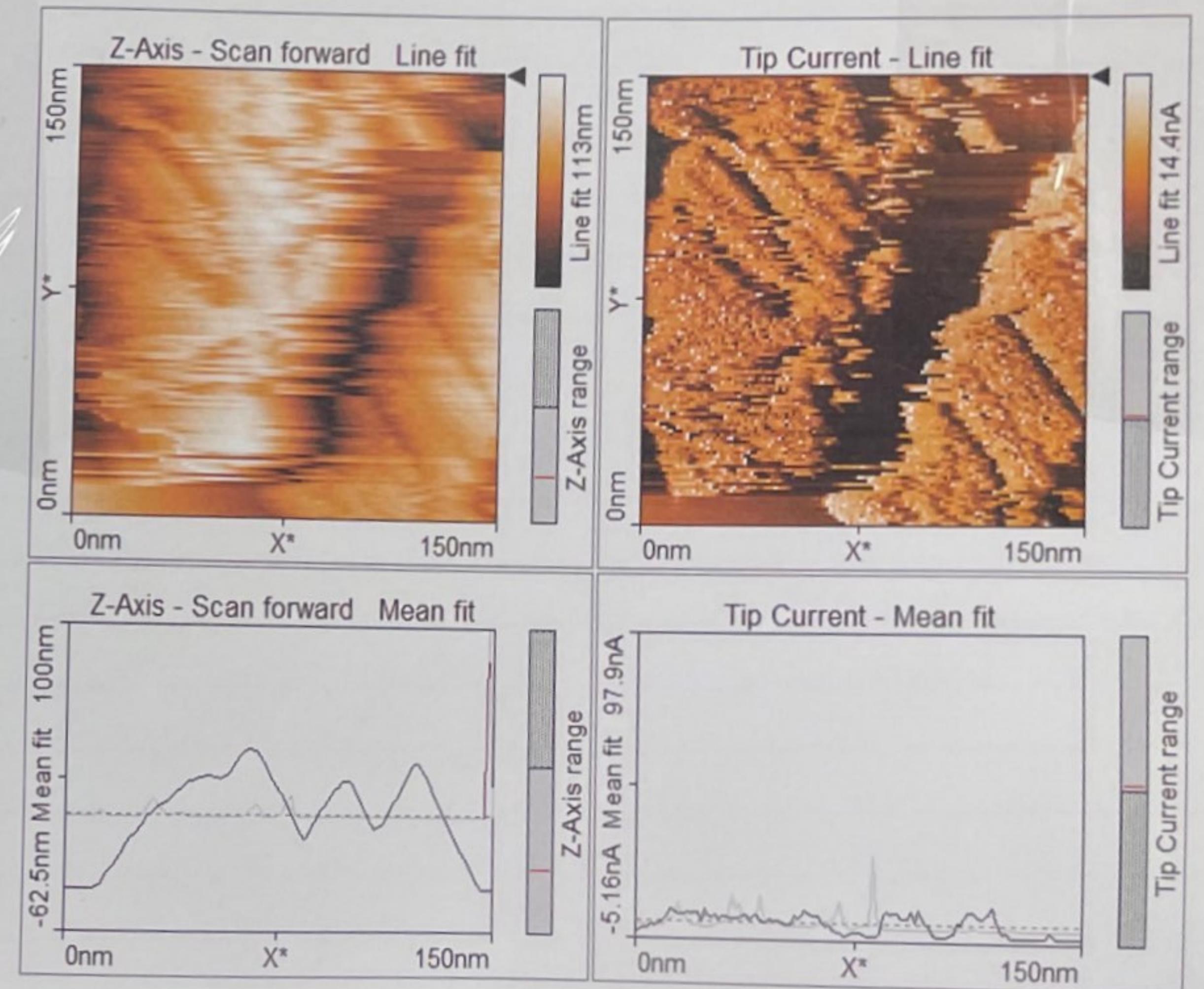
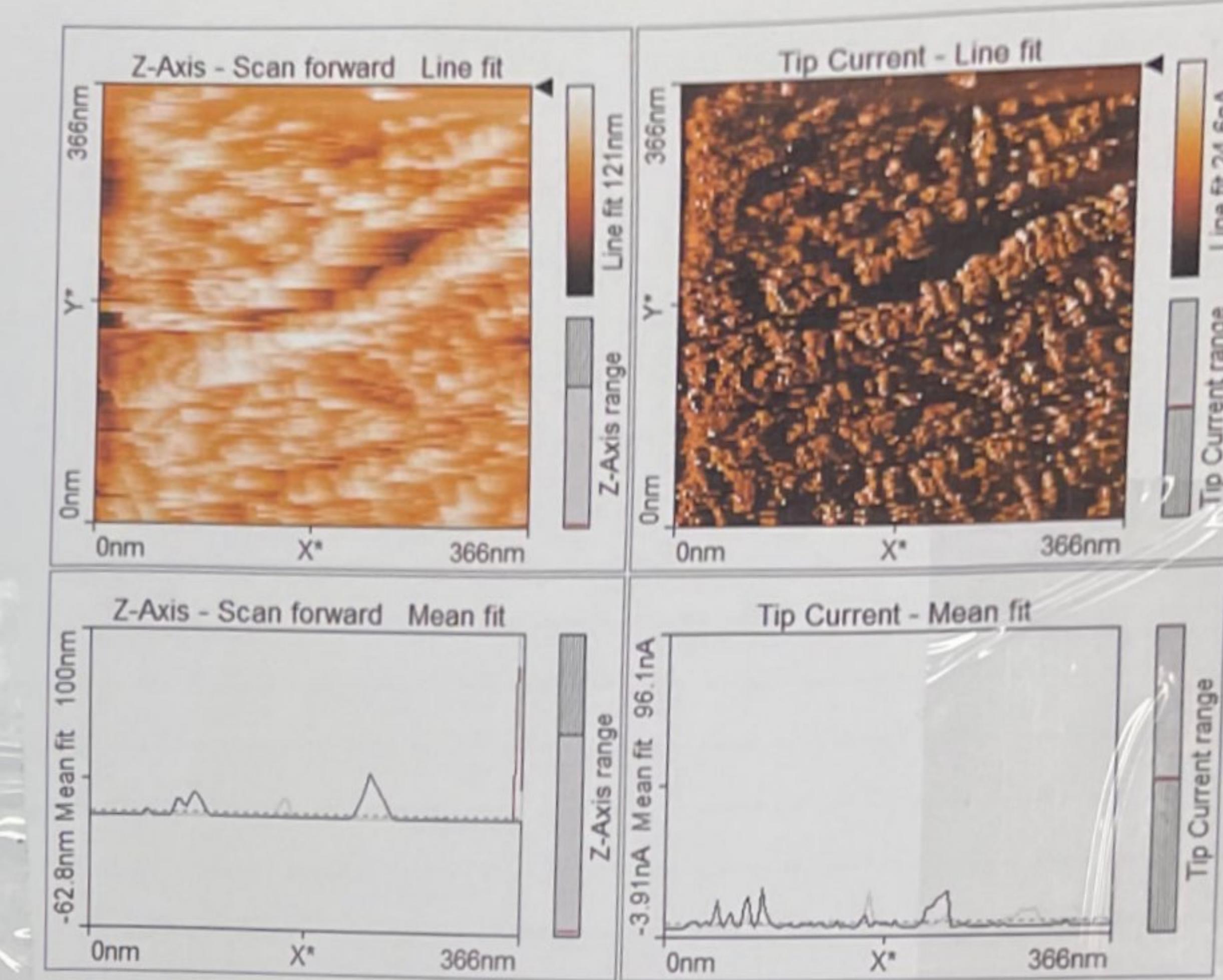
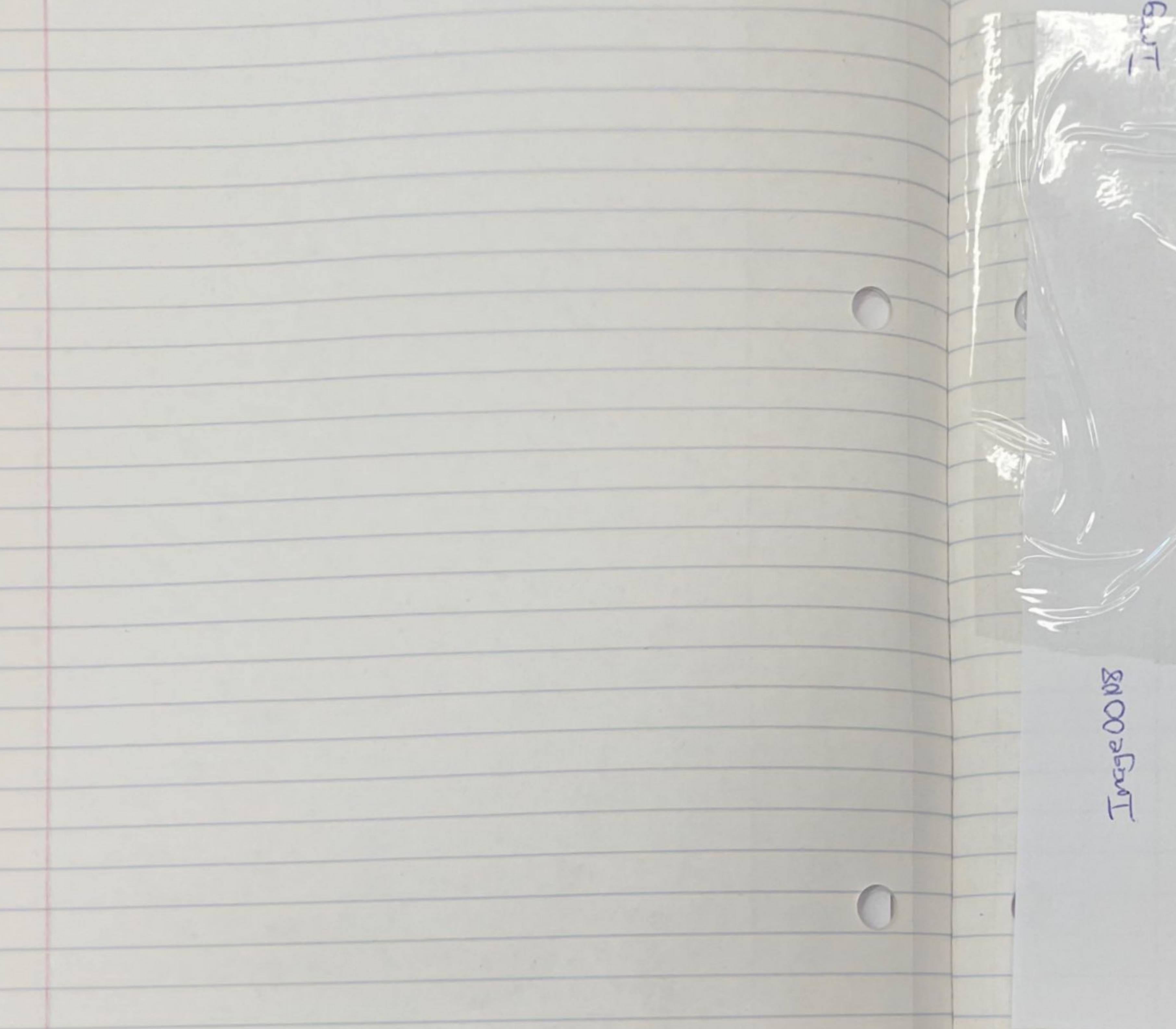
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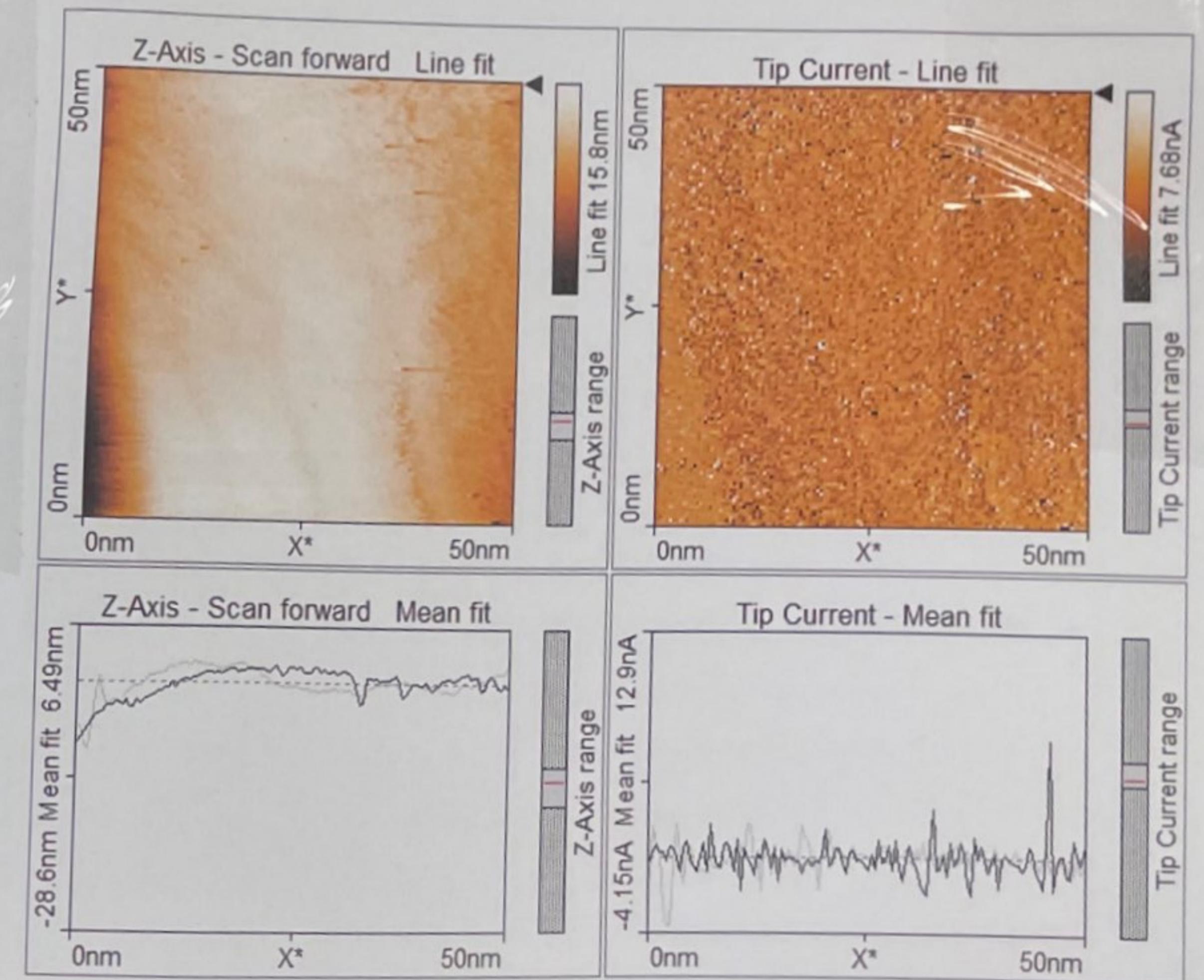
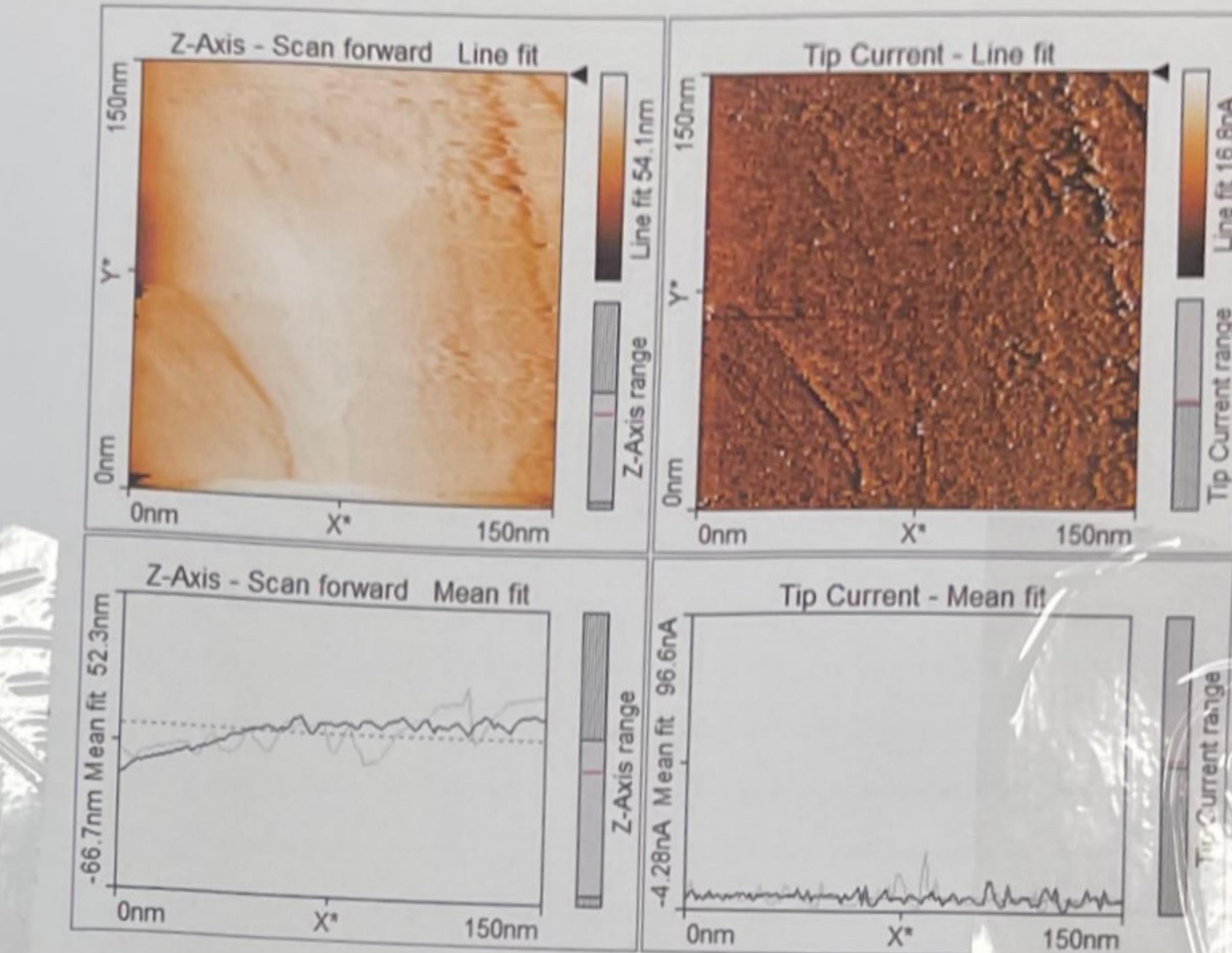
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Image 00111

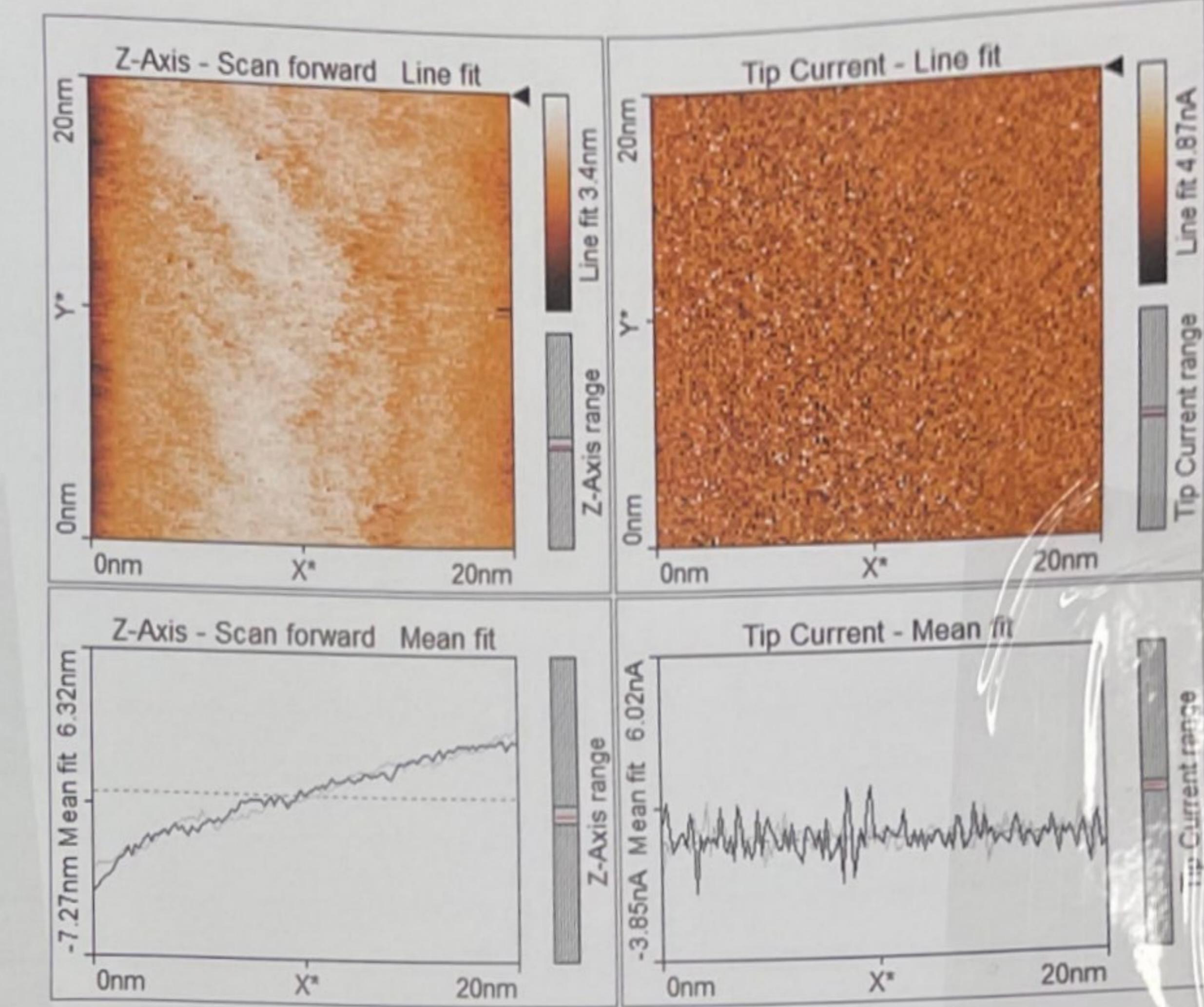
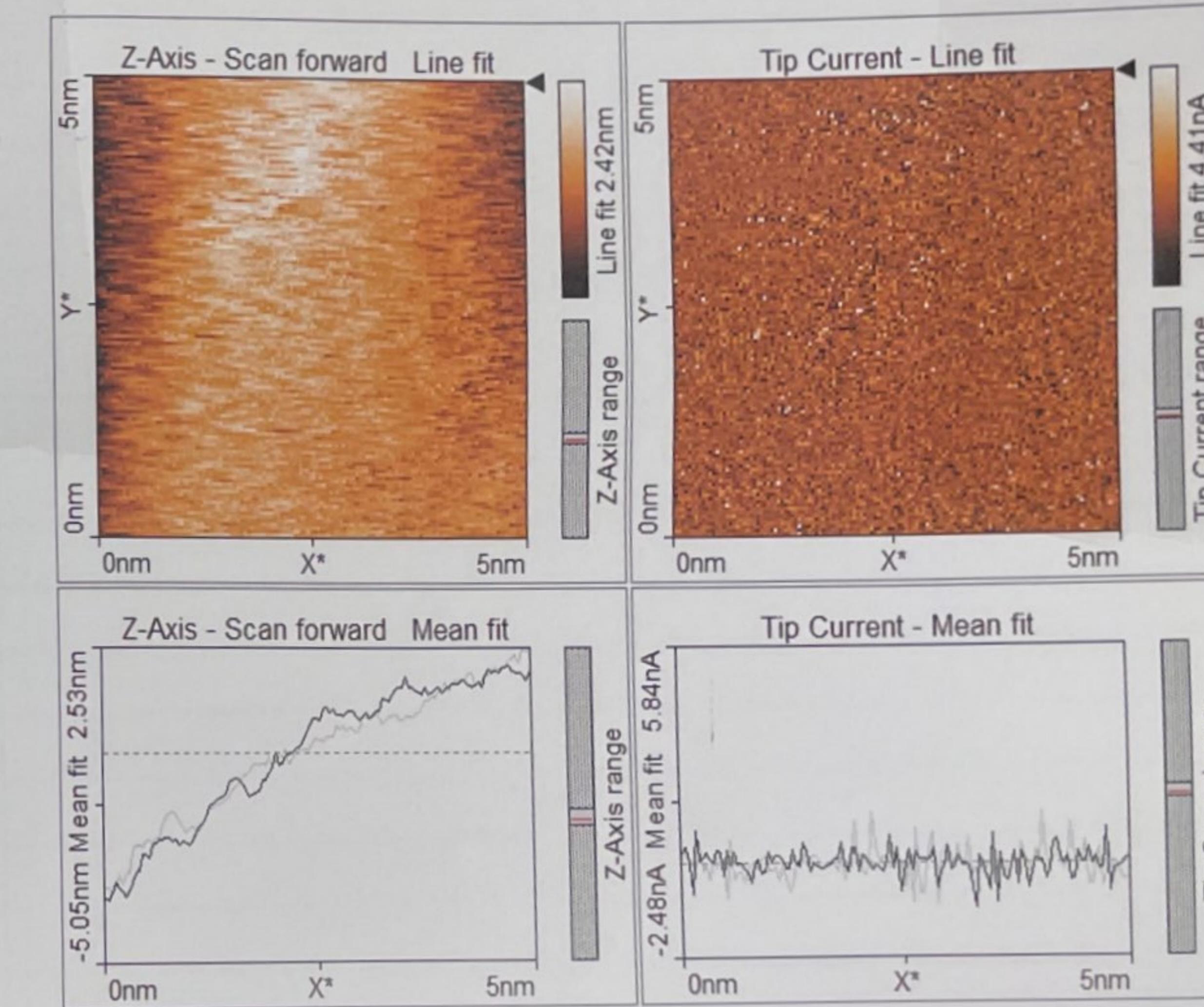
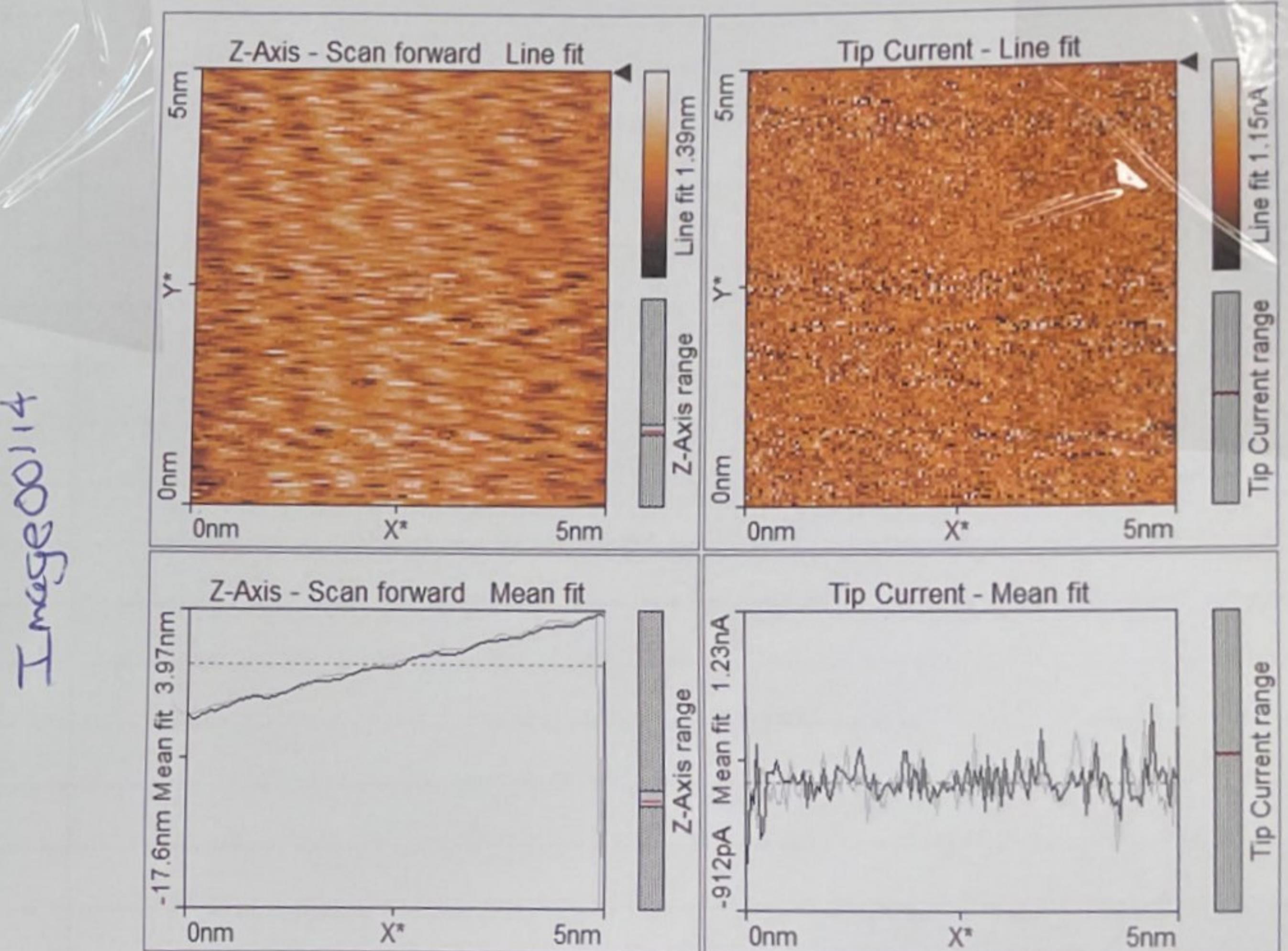
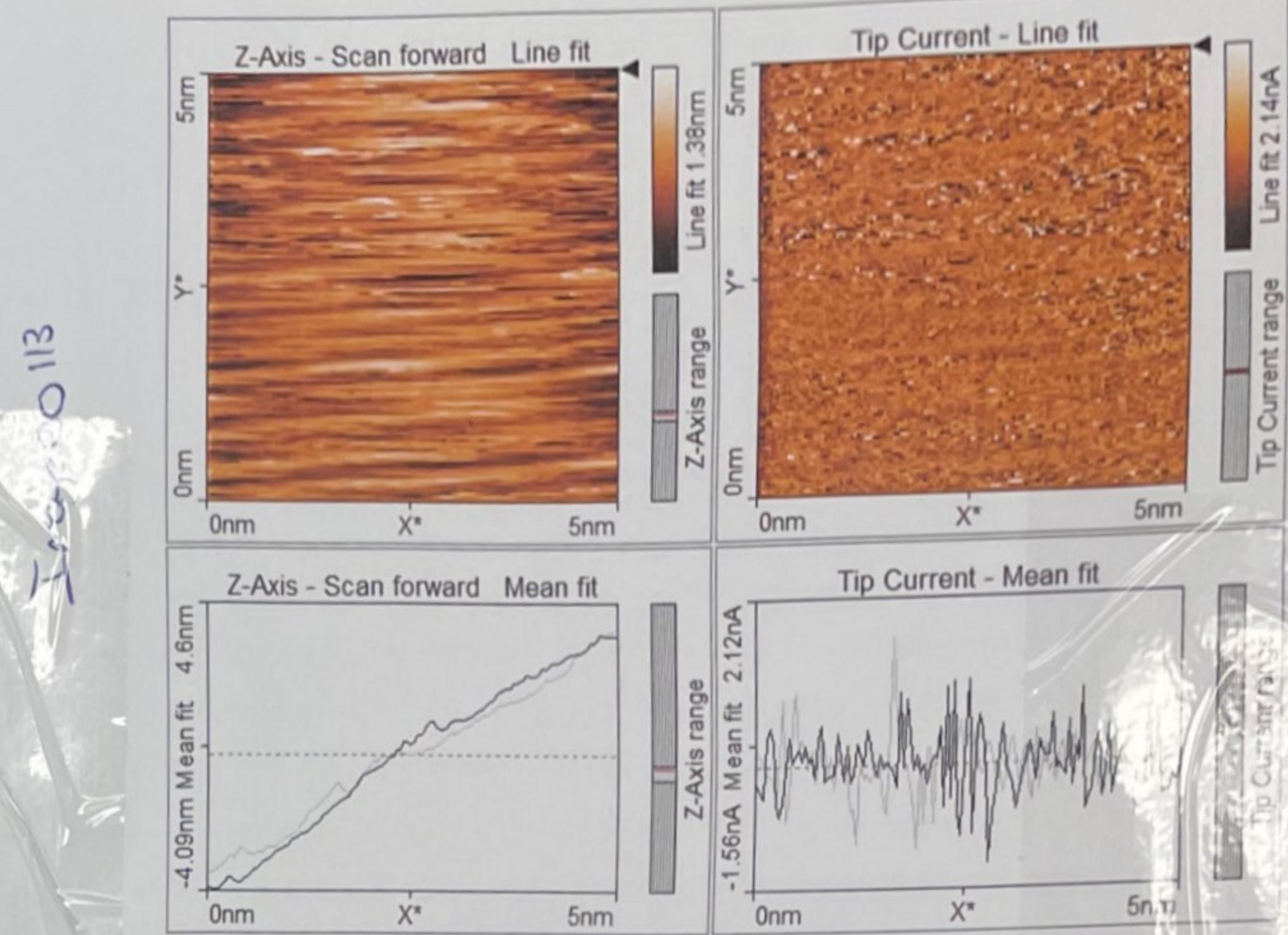


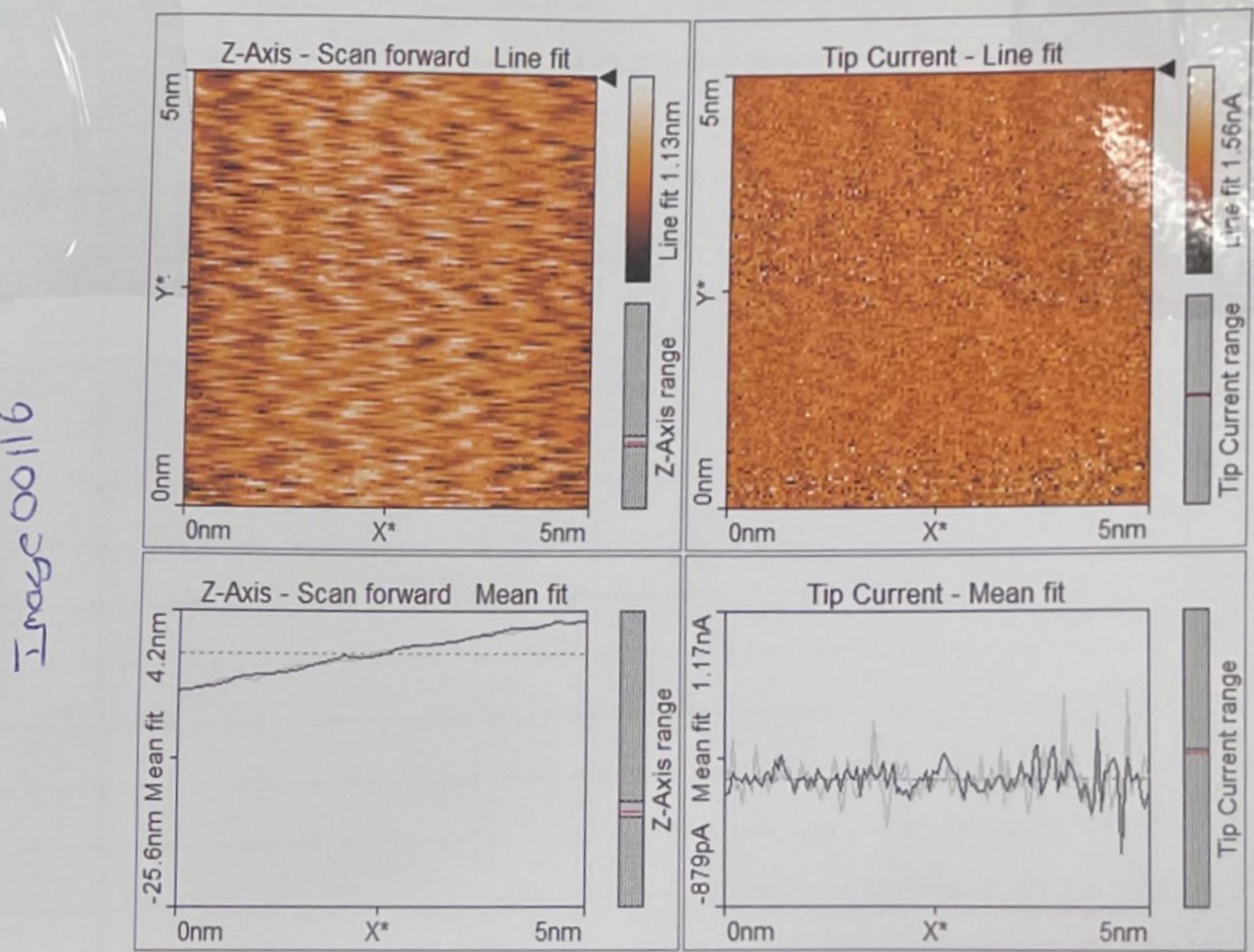
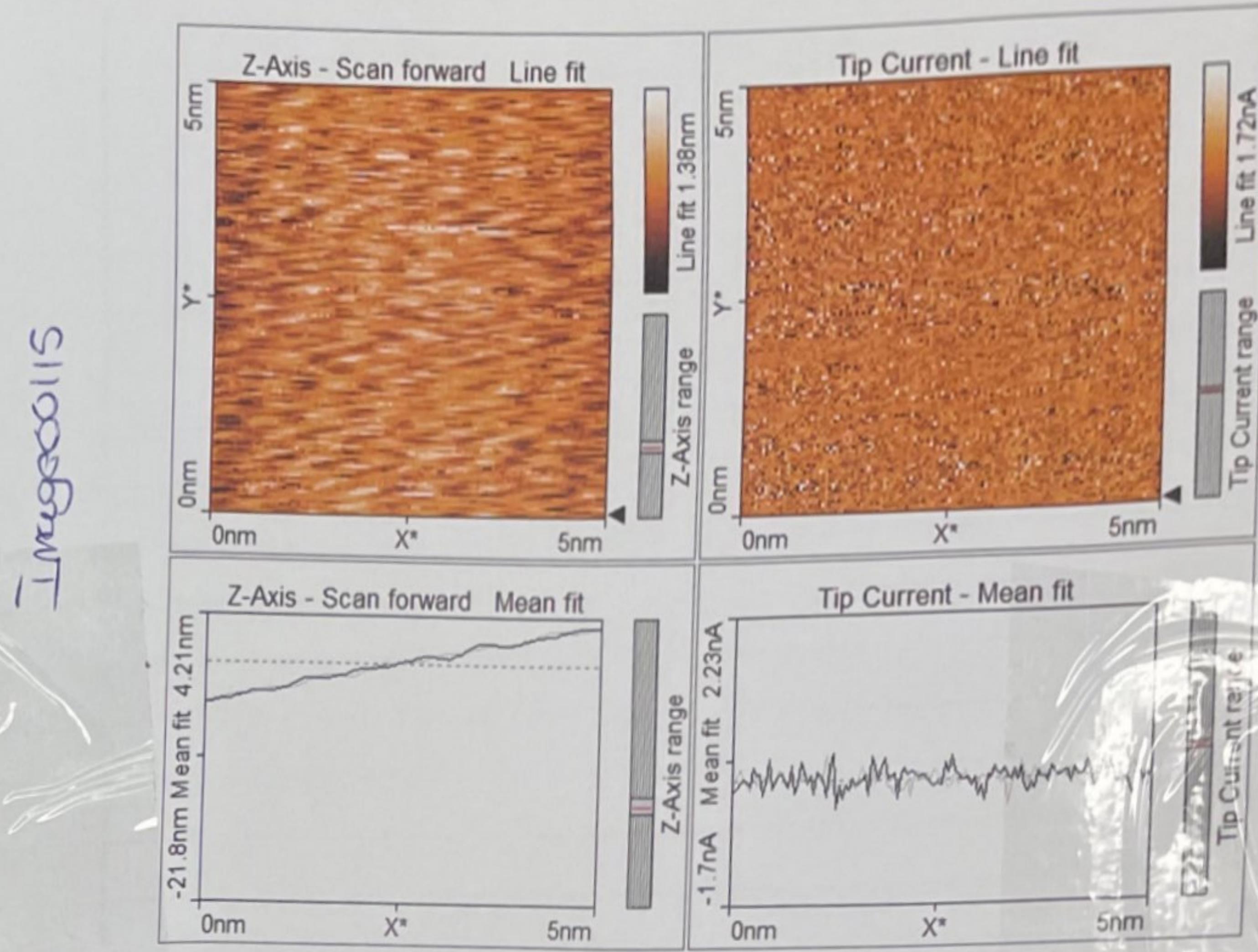
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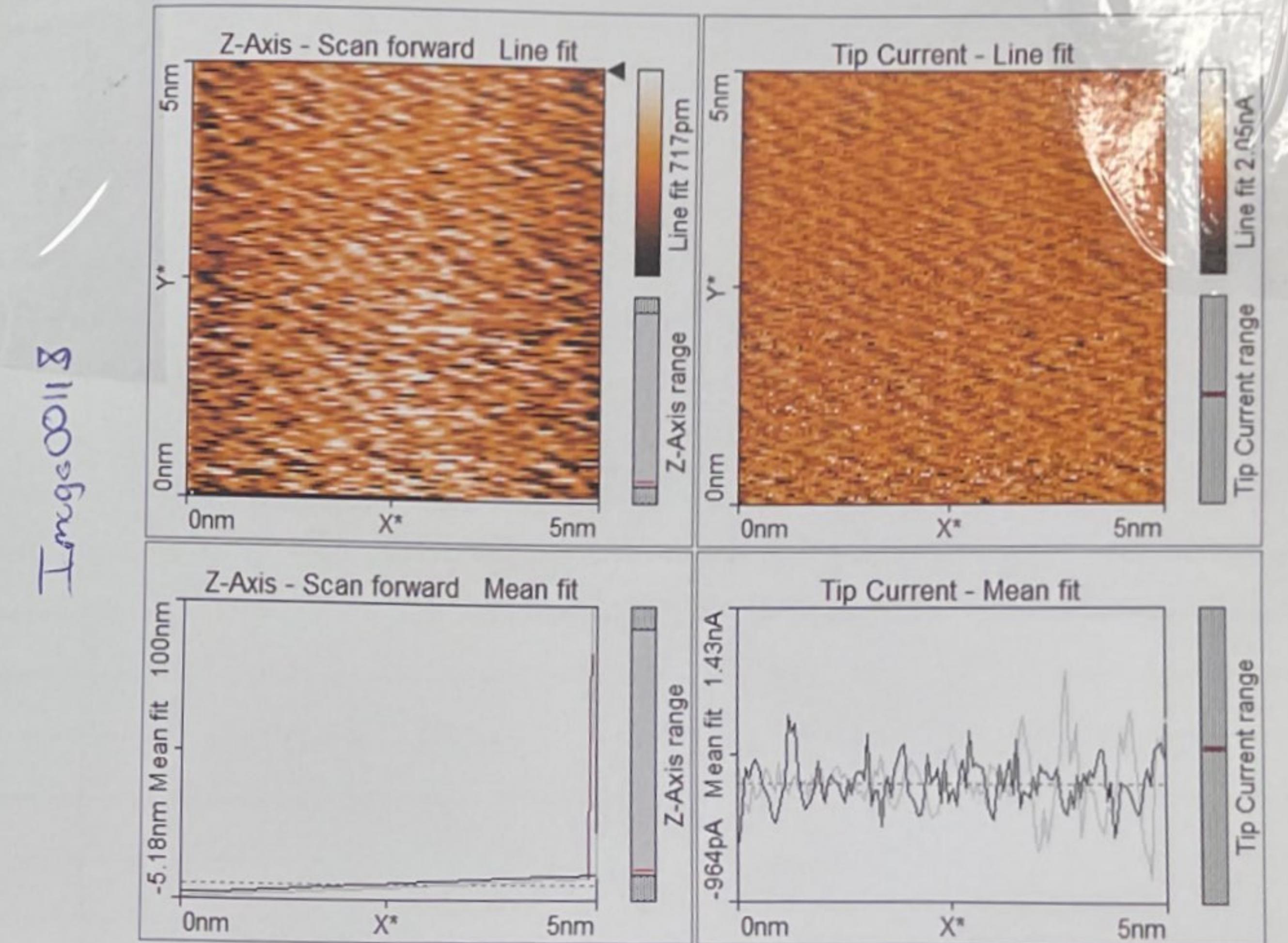
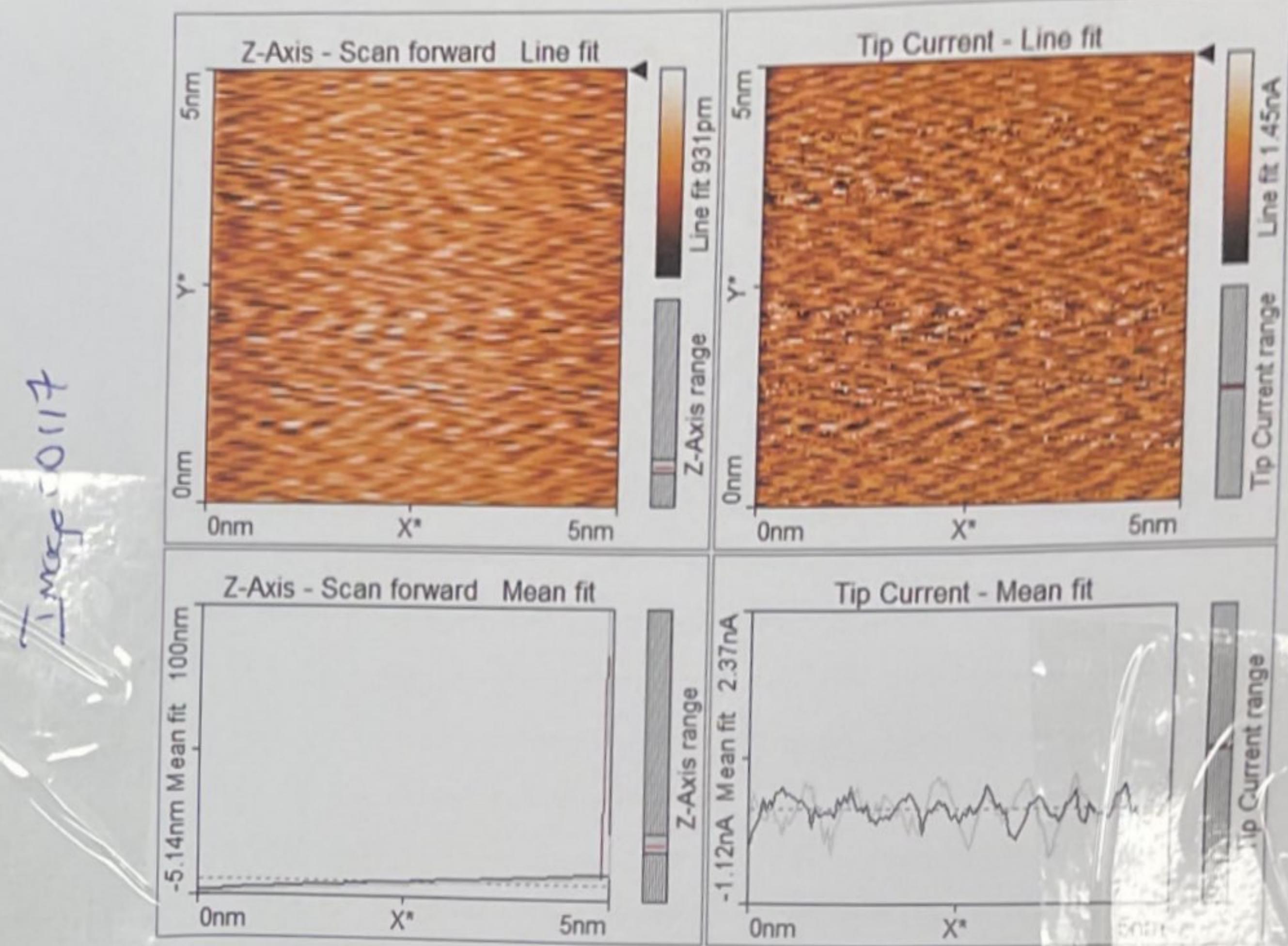
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Image 00119

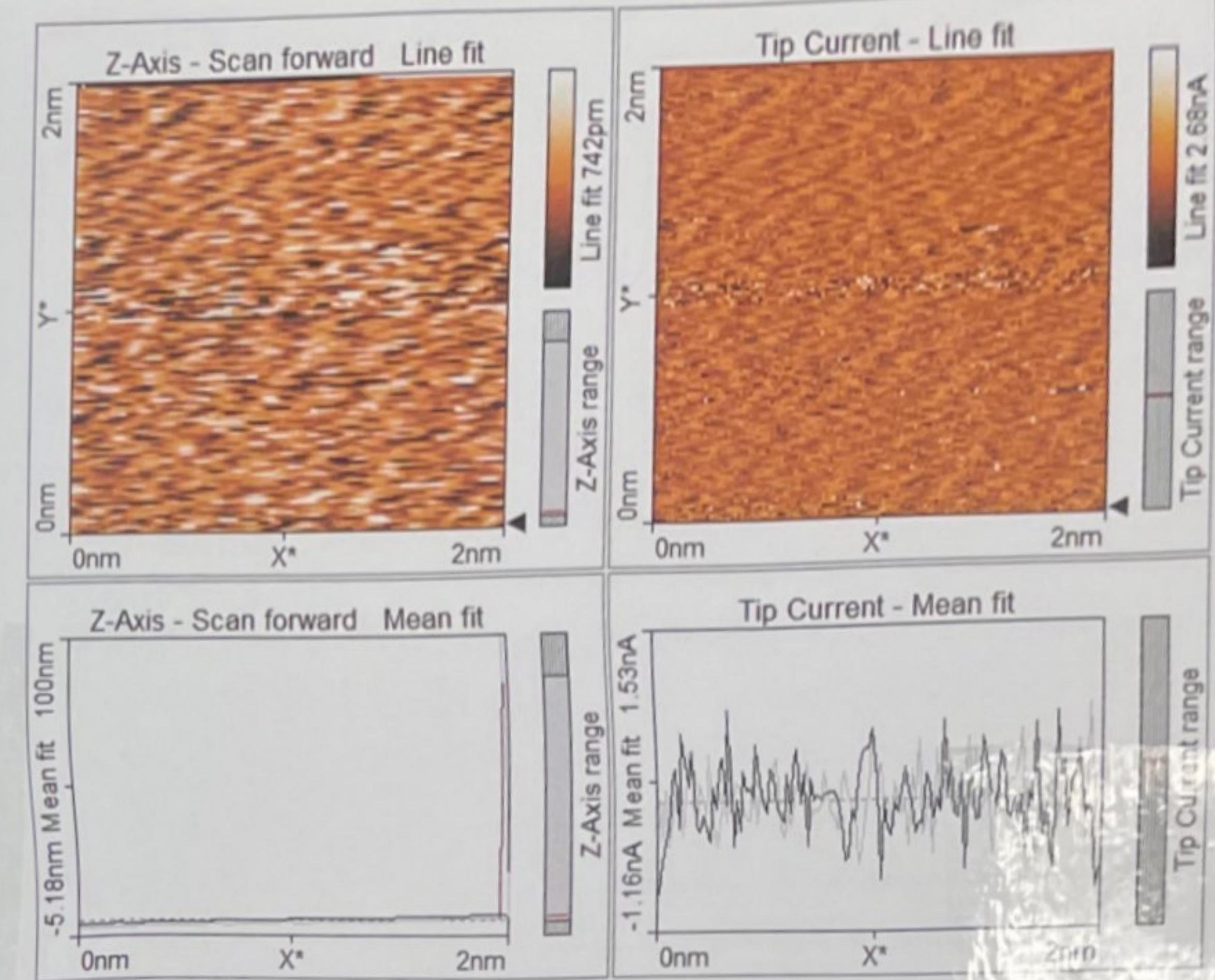
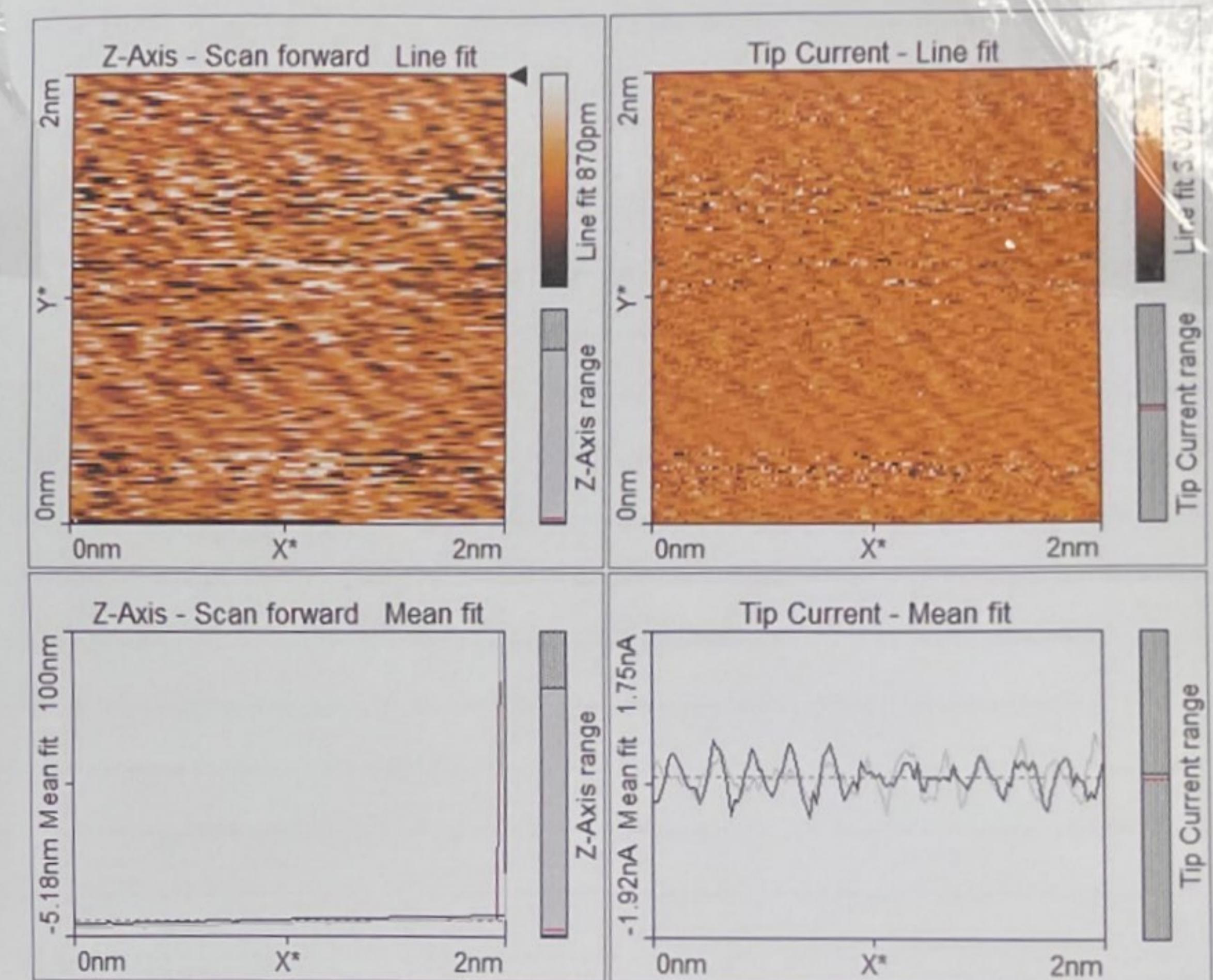


Image 00120



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Image 0125

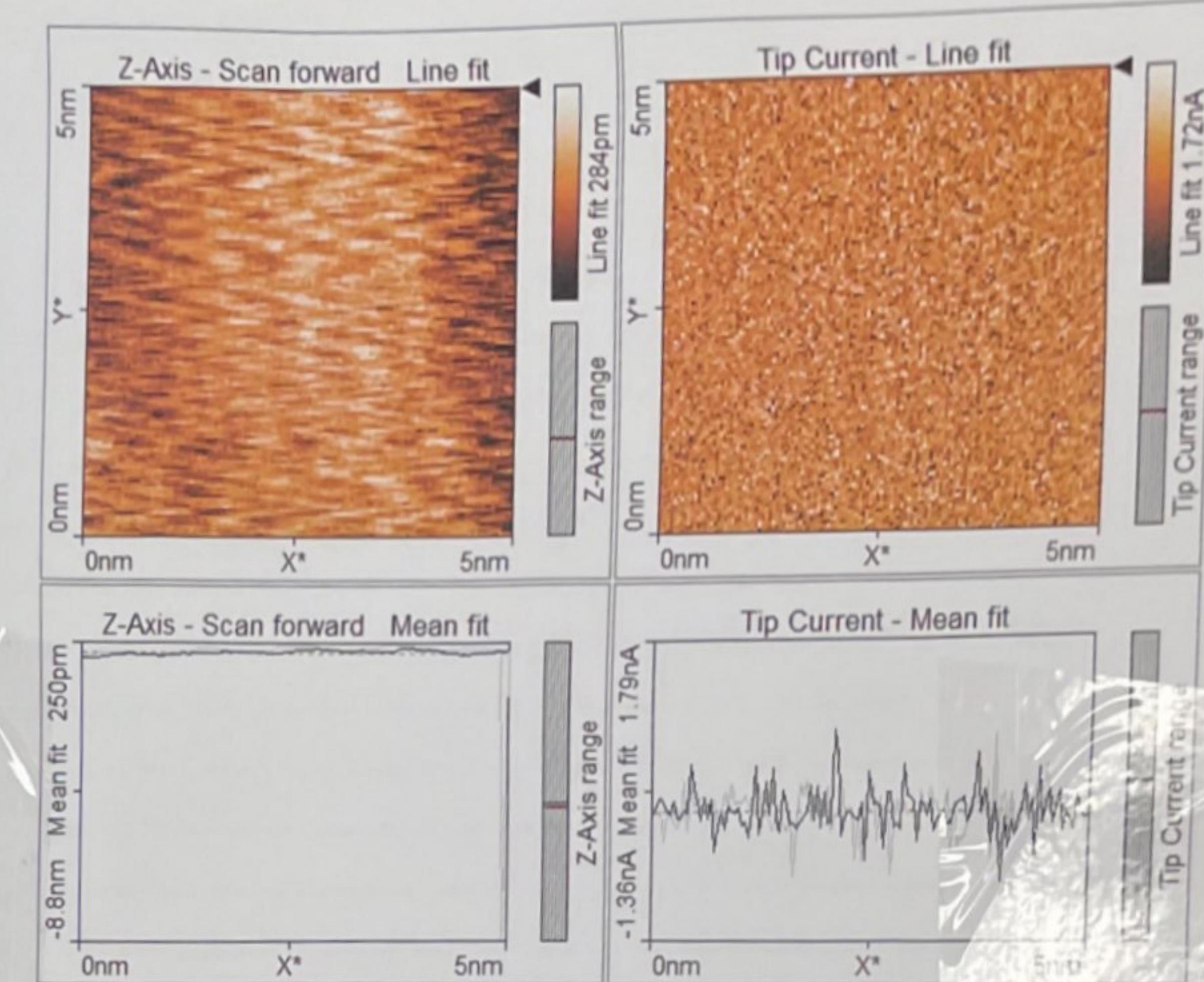


Image 0126

