

## Abstract

Copyright © 2017 LANCEVILLE TECHNOLOGY GROUP CO., LIMITED. All rights reserved.

This process is licensed under the Libre Silicon public license; you can redistribute it and/or modify it under the terms of the Libre Silicon public license as published by the Libre Silicon alliance, either version 1 of the License, or (at your option) any later version.

This design is distributed in the hope that it will be useful, but WITHOUT ANY WARRANTY; without even the implied warranty of MERCHANTABILITY or FITNESS FOR A PARTICULAR PURPOSE. See the Libre Silicon Public License for more details.

This document is part of the specification of the free silicon manufacturing standard for manufacturing the LibreSilicon standard logic cells<sup>1</sup> and related free technology nodes from the LibreSilicon project.

For this initial revision 0.1 a gate-first approach has been chosen which led to the choice of polysilicon as the gate electrode material because of the simplicity of the gate alignment. For better isolation properties of the transistors and gates in overall a box-isolation approach has been chosen. All of these choices have been made with the future scale down from the recent  $1\mu m$  to smaller structure sizes. **This process is for manufacturing  $1\mu m$  only!** But further releases which will have been tested with smaller structure sizes can be expected.

---

<sup>1</sup><https://github.com/chipforge/StdCellLib>

# Contents

<b>1</b>	<b>Shallow trench isolation</b>	<b>5</b>
1.1	Initial cleaning	6
1.1.1	Sulfuric Cleaning	6
1.1.2	HF dip	6
1.1.3	Drying	6
1.2	Hard mask: Oxide growth	7
1.3	Hard mask: Patterning	7
1.4	Hard mask: Etching	8
1.5	Hard mask: Resist removal	8
1.6	Silicon etching	9
1.7	Hard mask: Removal	9
<b>2</b>	<b>P-well</b>	<b>10</b>
2.1	Mask dioxide layer	11
2.2	Patterning	11
2.3	Etching	12
2.4	Cleaning	12
2.5	Implantation/Doping	13
2.6	Oxide mask removal	13
<b>3</b>	<b>N-well</b>	<b>14</b>
3.1	Mask dioxide layer	15
3.2	Patterning	15
3.3	Etching	16
3.4	Cleaning	16
3.5	Implantation/Doping	17
3.6	Oxide for drive-in	18
3.7	Drive-in	18
3.8	Oxide mask removal	18
<b>4</b>	<b>Field oxide</b>	<b>19</b>
4.1	Oxide growth	20
4.2	Patterning	20
4.3	Etching	21
4.4	Resist removal	21
<b>5</b>	<b>Gate</b>	<b>22</b>
5.1	Gate oxide deposition	23
5.2	Polysilicon deposition	23
5.3	Patterning	24
5.4	Etching	25
5.5	Resist removal	25
<b>6</b>	<b>n+ Implant</b>	<b>26</b>
6.1	Mask dioxide layer	27
6.2	Patterning	27
6.3	Etching	28
6.4	Cleaning	28
6.5	Implantation/Doping	29
6.6	Oxide removal	29
<b>7</b>	<b>p+ Implant</b>	<b>30</b>
7.1	Mask dioxide layer	31
7.2	Patterning	31
7.3	Etching	32
7.4	Cleaning	32
7.5	Implantation/Doping	33
7.6	Oxide removal	33
<b>8</b>	<b>Silicification</b>	<b>34</b>

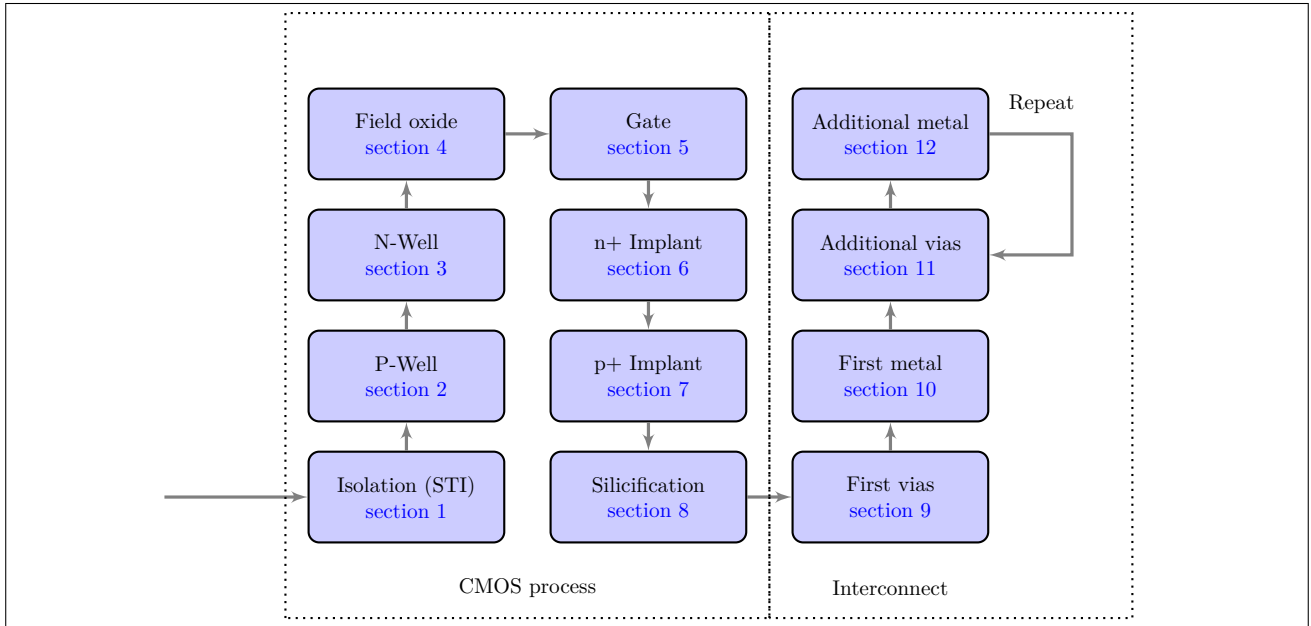
8.1	Oxide deposition . . . . .	35
8.2	Silicide block patterning (optional) . . . . .	35
8.3	Sputter etching(Spacers) . . . . .	36
8.4	Titanium deposition . . . . .	36
8.5	First reaction step . . . . .	36
8.6	Metal removal . . . . .	37
8.7	Second reaction step . . . . .	37
<b>9</b>	<b>First vias</b>	<b>38</b>
<b>10</b>	<b>First metal layer</b>	<b>39</b>
<b>11</b>	<b>Additional vias</b>	<b>40</b>
<b>12</b>	<b>Additional metal layer</b>	<b>41</b>

# Libre Silicon process steps

David Lanzendörfer

May 29, 2018

The general flow chart of the overall process flow can be seen in [Figure 1](#). These process steps will be discussed within the following sections.



**Figure 1:** Frontend and backend process flow

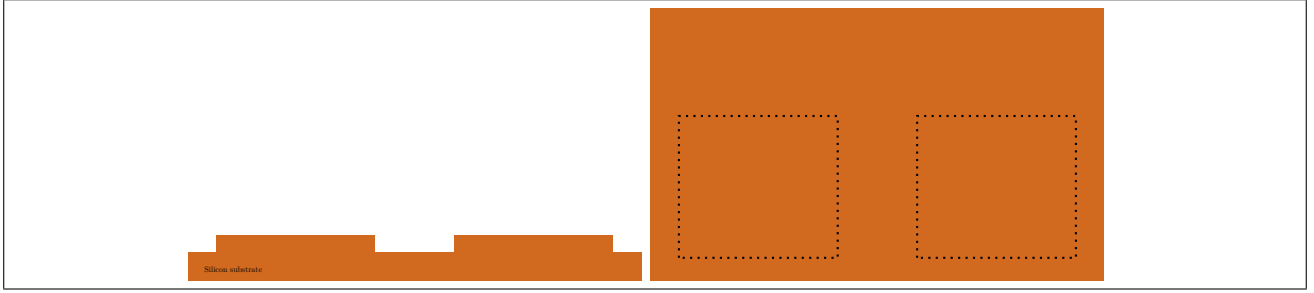
The six overall process steps are part of an active part of the technology, while the final metal (respectively contact) layers will be used for making a contact between the logic gates and macro cells and making them available to the exterior world.

For this process p-substrate is the required basic substrate, but forks and modifications will be very well possible based on a Graphene substrate or alike, still under the LSPL. The starting material is a p-type,  $\langle 100 \rangle$  oriented silicon with a doping concentration of  $\approx 9 \times 10^{14} \text{ cm}^{-3}$ .

If you don't have a plasma etcher you will need  $\langle 111 \rangle$  substrate!

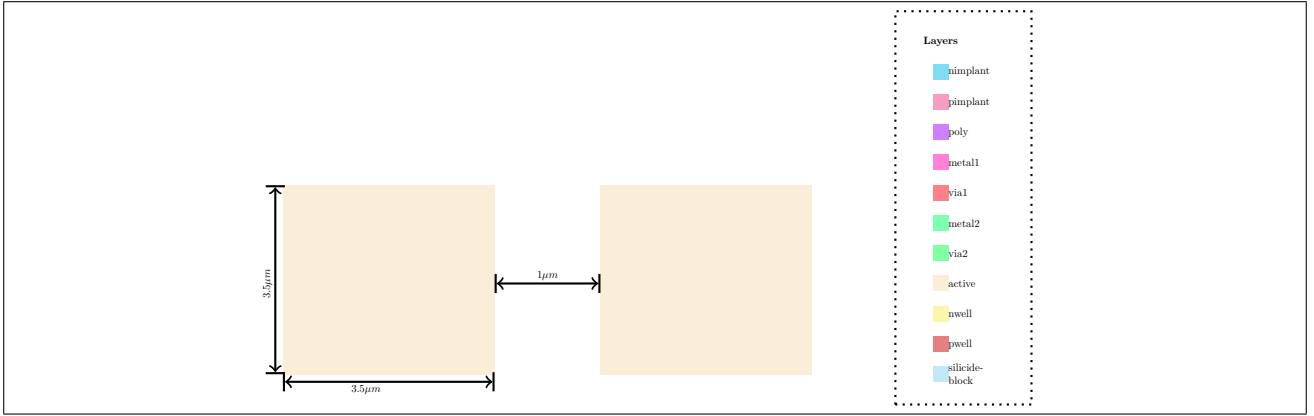
# 1 Shallow trench isolation

The geometry of a substrate with STI implemented can be seen in [Figure 2](#).



**Figure 2:** Shallow trench isolation target geometry

As can be seen in [Figure 19](#), the n-well and the STI trench are supposed to have approximately the same depth but the n-well and p-well go down a little bit further. Because the n-well will be  $\approx 4\mu m$  in depth we have to match this with our trench depth. In order to allow a sufficiently low resistance of the ESD diode but at the same time a sufficient isolation of between the standard cells a trade-off has been done. The targeted depth of the box isolation is  $\approx 2\mu m$ .

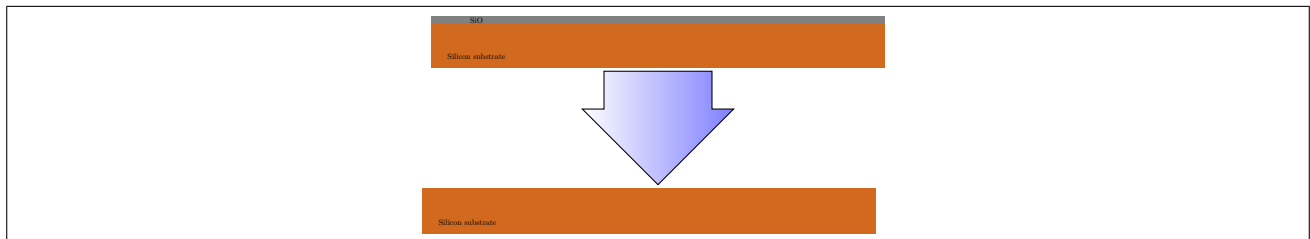


**Figure 3:** Shallow trench isolation layout

In [Figure 3](#) we can see the layout for the STI area. The STI area will be everywhere, where no active areas are. The field oxide needs to be grown out of trenches which can't be etched out of the silicon by using resist as a mask. For that reason we will have to resort to a protective mask made from a silicon dioxide layer which has to be etched before hand. So the mask will be exposed onto positive resist on top of the hard mask oxide layer in order to form a protective mask covering the active areas from having etched trenches into them. After that we can either use a dry etching method or wet etching for cutting into the silicon substrate and making the active area become islands with trenches in between. After these steps we have to remove the hard mask. Our minimum width and height as well as the space between the active areas comes from the line space constrain of the silicon etcher and of course the optical limitations of the stepper which are as well  $0.5\mu m$ .

## 1.1 Initial cleaning

In order to remove the initial naturally grown silicon dioxide from the wafer, acid is being applied to the wafer which leads to a pure silicon substrate wafer as in the process illustration shown in [Figure 4](#).



**Figure 4:** Initial cleaning

This needs to be done because the naturally grown initially existing silicon oxide is not pure and may contain contamination which may render the final product unusable.

### 1.1.1 Sulfuric Cleaning

The sulfuric acid mixture,  $H_2SO_4 + H_2O_2$  is being applied to the wafer for 10 minutes at a temperature of 120 °C.

### 1.1.2 HF dip

After the sulfuric cleaning a HF ( $HF:H_2O, 1:50$ ) dip is being performed for one minute.

Hydrofluoric acid (HF) is used to remove native silicon dioxide from wafers. Since it acts quickly, one needs to only expose the wafer for a short time ("dip").

### 1.1.3 Drying

After that the wafer needs to be dried and quickly processed further before new uncontrolled natural oxide can build up on the wafer through the contact with air.

## 1.2 Hard mask: Oxide growth

We need a thick layer of oxide as protective hard mask to etch the trenches into the silicon.



**Figure 5:** Hard mask growth

In subsection 1.6 we want to etch  $2\mu\text{m}$  deep into the silicon which, depending on the approach, will take a different amount of oxide for an effective hard mask.

For now we only have the plasma etcher variant being verified because chemically etching the silicon wafer with KOH isn't allowed at the HKUST labs for contamination control reasons. In case you can verify this in your lab with a chemical etching method, please update this chapter and make a pull request!

**Possible approaches:**

- **"DRIE Etcher #1" from HKUST**

We can use anisotropic plasma etching for sharper borders.

With this method it takes pretty much a minute to etch  $2\mu\text{m}$  deep.

In one minute the oxide will have been etched away by at most 25nm.

This means we can grow a 100nm thick layer and are well off.

Growing 100nm of dioxide takes around 5 minutes 30 seconds at  $1050^\circ\text{C}$  in wet ambient<sup>1</sup>.

- **Chemical solution**

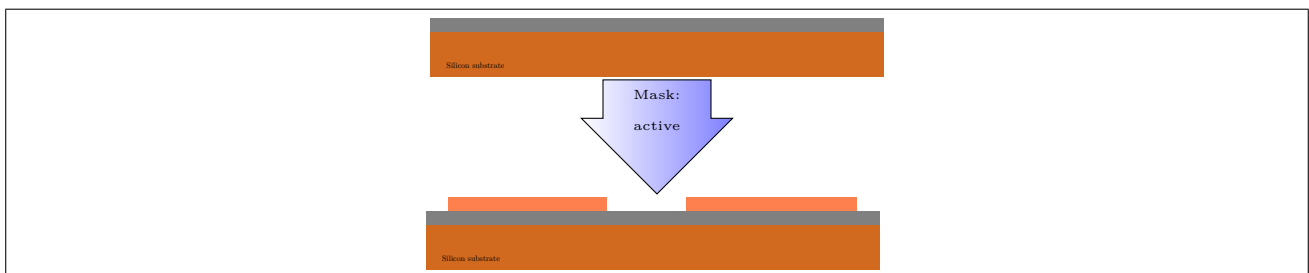
When using KOH acid at  $60^\circ\text{C}$ , it takes 4 minutes and 30 seconds in order to etch  $2\mu\text{m}$  deep.

This means the oxide layer needs to be at least 226 nm thick, so we choose a nice round number of 300nm.

The layer of silicon dioxide of around 300nm thickness is grown in wet ambient for 25 minutes at  $1050^\circ\text{C}$ <sup>2</sup> in the diffusion furnace.

## 1.3 Hard mask: Patterning

The resist is being deposited using spin coating and then soft baked depending on the baking time for the specific resist. The requirement is a **positive** tone resist.



**Figure 6:** Patterning with positive resist

The layout for being exposed onto the resist is being extracted from the "active" layer within the GDS2 file onto a **bright field** mask because we need to use the same mask again in section 4, so alignment needs to be possible.

<sup>1</sup><http://cleanroom.byu.edu/OxideTimeCalc>

<sup>2</sup><http://cleanroom.byu.edu/OxideTimeCalc>

## 1.4 Hard mask: Etching

We open the access to the silicon, outside of the active areas, in order to etch the trenches.



**Figure 7:** Nitride mask etching

There are dry etching and wet etching methods available for etching the oxide hard mask. The downside of wet etching is that it also etches horizontally, however the chemical BHF is readily available and allows for easy implementation of the process.

**Possible approaches:**

- **"DRIE Etcher #1" from HKUST**

We can use anisotropic plasma etching for sharper borders.

- **Chemical solution**

We can use buffered hydrofluoric acid (BOE (1:6)) at room temperature for a little bit over 3 minutes in order to get through the 300nm of oxide or around 1 minute for 100nm. in case RIE is being used for silicon etching

Etching too long might cause under-etch however!

## 1.5 Hard mask: Resist removal

Now we need to remove the contaminants for further processing.



**Figure 8:** Resist removal

We strip the resist, rinse and perform sulfuric cleaning.



## 1.6 Silicon etching

Silicon can only be etched by a very aggressive chemical cocktail of KOH and TMAH (20%) or by plasma etching.

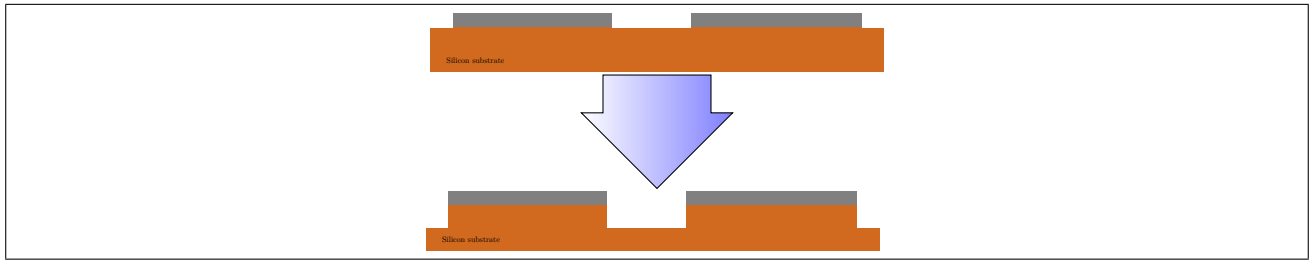


Figure 9: Trench etching

For now we only have the plasma etcher variant being verified because chemically etching the silicon wafer with KOH isn't allowed at the HKUST labs for contamination control reasons. In case you can verify this in your lab with a chemical etching method, please update this chapter and make a pull request!

Possible approaches:

- **"DRIE Etcher #1" from HKUST**

This machine has a normal etching rate of up to  $2 \frac{\mu m}{min}$  for etching silicon.

This means we etch for 1 minute in order to reach the desired depth.

The selectivity to oxide is  $>80:1$  which means the etch speed for the hard mask will be at most  $\frac{1}{80} 2 \frac{\mu m}{min} = \frac{1}{80} 2000 \frac{nm}{min} = 25 \frac{nm}{min}$ .

- **Chemical solution**

Using a KOH solution of 20% at  $60^\circ C$  gives us an etch rate of roughly  $26.57 \mu m$  per hour<sup>3</sup>.

- The  $\langle 100 \rangle$  etch rate is:  $26.57 \text{ micron/hr} = 0.44 \text{ micron/min}$
- The  $\langle 110 \rangle$  etch rate is:  $40.5 \text{ micron/hr}$
- The  $\langle 111 \rangle$  etch rate is:  $0.4932 \text{ micron/hr}$
- The  $\text{SiO}_2$  etch rate is:  $49.92 \text{ nanometers/hr}$

With a desired depth of  $2 \mu m$  we will have to etch around 4 minutes and 30 seconds in order to reach the desired depth. The disadvantage of this approach is the imprecision and under-etch of the mask.

## 1.7 Hard mask: Removal

Now we have to remove the oxide hard mask for further processing in order to proceed with well formation without contamination during oxide growing.

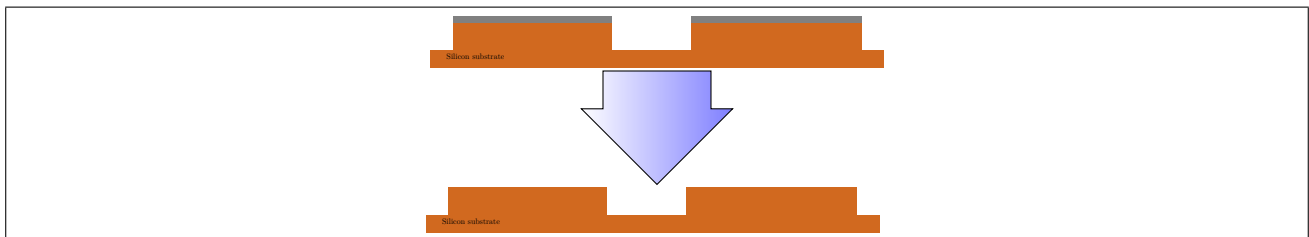


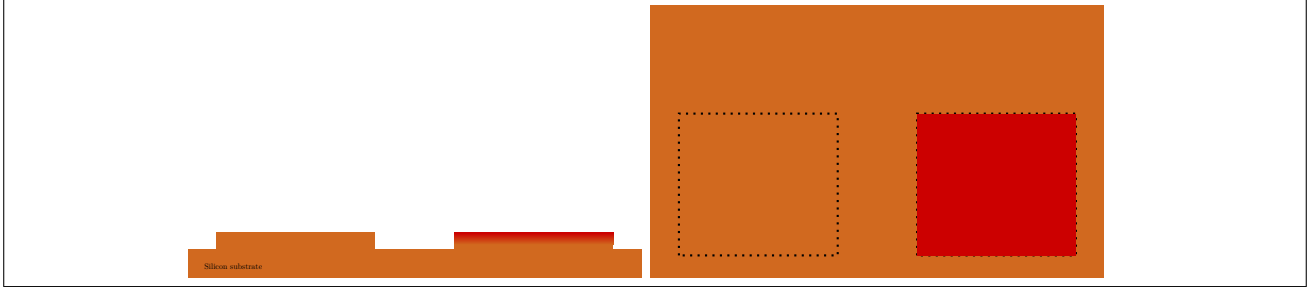
Figure 10: Trench etching

We use buffered hydrofluoric acid (BOE (1:6)) at room temperature for a little bit over 3 minutes in order to remove all of the 300nm thick oxide layer and roughly 1 minute for 100nm.

<sup>3</sup><http://www.lelandstanfordjunior.com/KOH.html>

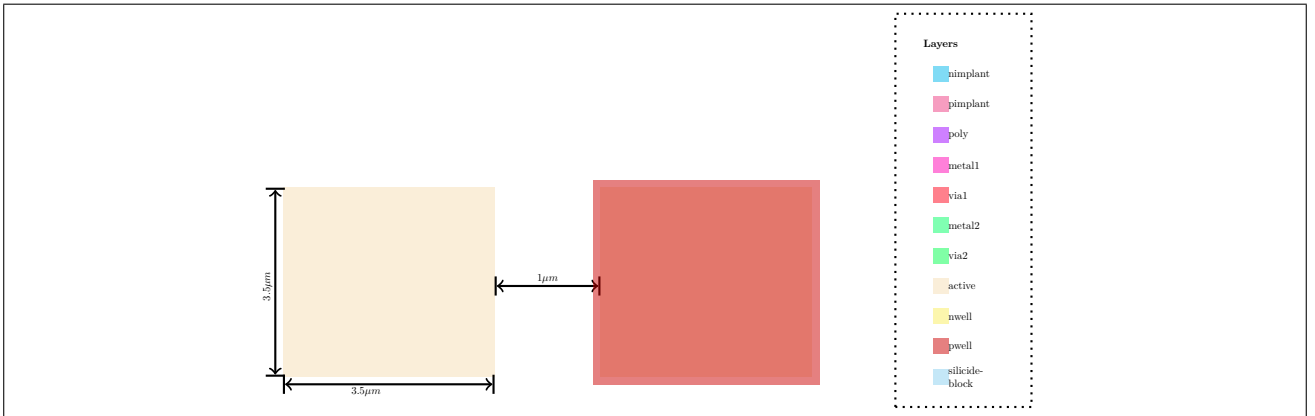
## 2 P-well

In order to build CMOS on the same substrate, a P-well is required for building the complementary N-channel transistor for a n-p-channel logic circuitry. The cross section as well as the top view of the targeted geometry are shown in [Figure 19](#)



**Figure 11:** P-well target geometry

The P-well will serve us as an island of higher p-doped substrate within the slightly p-doped basis substrate. The dopant dose will be  $2.5 \times 10^{12} \text{cm}^{-2}$  as calculated in the documentation of the process design leading to these steps<sup>4</sup>.



**Figure 12:** P-Well layout

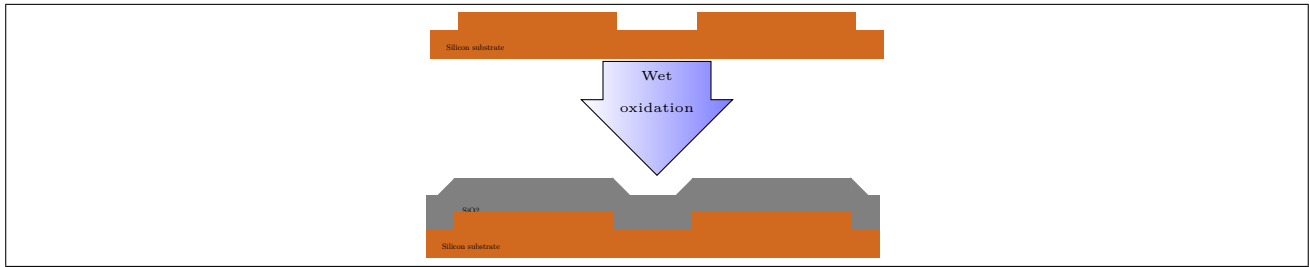
In [Figure 12](#) the layout of the P-well region on top of the active area region can be seen. The p-well is being fit into the active area.

It should even be a little bit bigger than the active area, because of possible alignment offsets

<sup>4</sup>[https://github.com/leviathanch/libresiliconprocess/raw/master/process\\_design/process\\_design.pdf](https://github.com/leviathanch/libresiliconprocess/raw/master/process_design/process_design.pdf)

## 2.1 Mask dioxide layer

In order to selectively inject charge carrying atoms into the crystalline structure a protective dioxide ( $SiO_2$ ) layer needs to be grown on top of a p-type substrate.



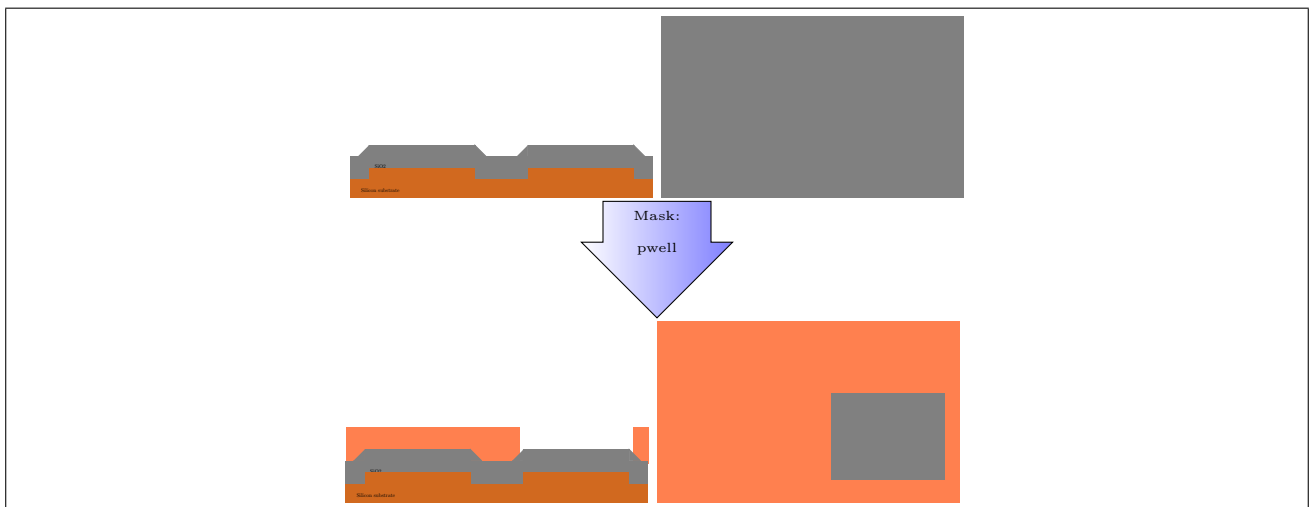
**Figure 13:** Dioxide layer growth

With an energy of 100keV for the implantation performed in [subsection 2.5](#), the projected range of the dopants within the oxide will be 310nm (380nm tops) <sup>5</sup>. This means being on the safe side and having 500nm as the thickness is a good approach.

In order to grow the 500nm thick oxide layer, the wafer is being oxidized for around 56 minutes at 1050°C using wet oxidation which results in a dioxide layer of around 500nm in thickness<sup>6</sup>.

## 2.2 Patterning

The resist is being deposited spray or spin coating (spray coating is better because of the uneven surface!) and then soft baked depending on the baking time for the specific resist. The layout for being exposed onto the resist is being extracted from the "pwell" layer within the GDS2 file onto a **bright field** mask. The requirement is a **negative** tone resist.



**Figure 14:** Cross/top view of P-well layout on resist

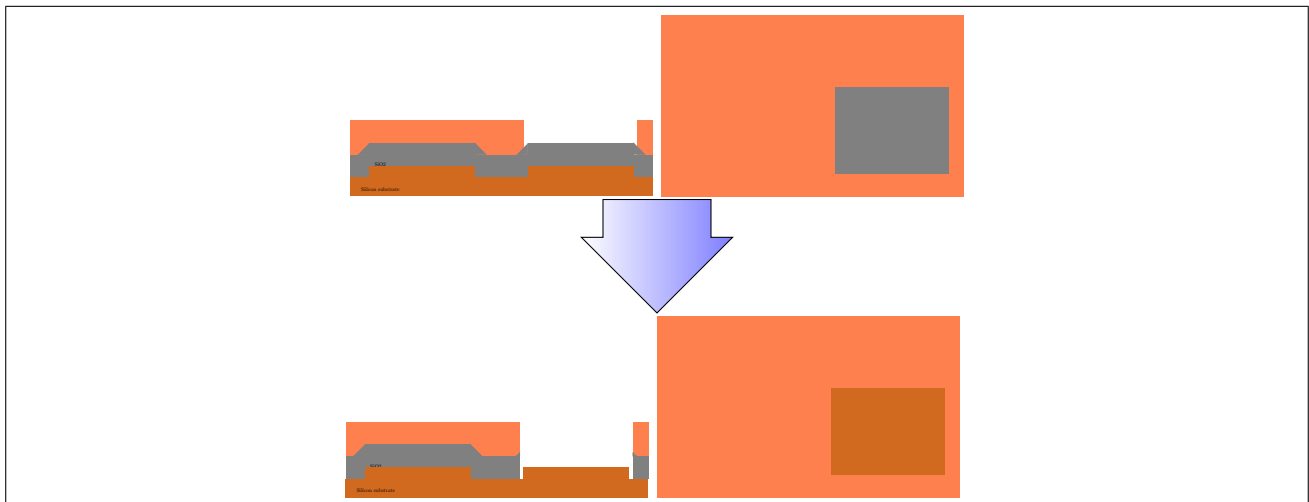
The thickness of the resist layer and the baking duration will variate depending on the specific equipment for which this process will be implemented with. Also after the exposure and development, the hard baking shouldn't be forgotten!

<sup>5</sup><http://cleanroom.byu.edu/rangestraggle>

<sup>6</sup><http://cleanroom.byu.edu/OxideTimeCalc>

## 2.3 Etching

We now need to open a window in the dioxide layer, through which we will inject carrier atoms into the silicon crystal structure.



**Figure 15:** Cross/top view of P-well oxide window

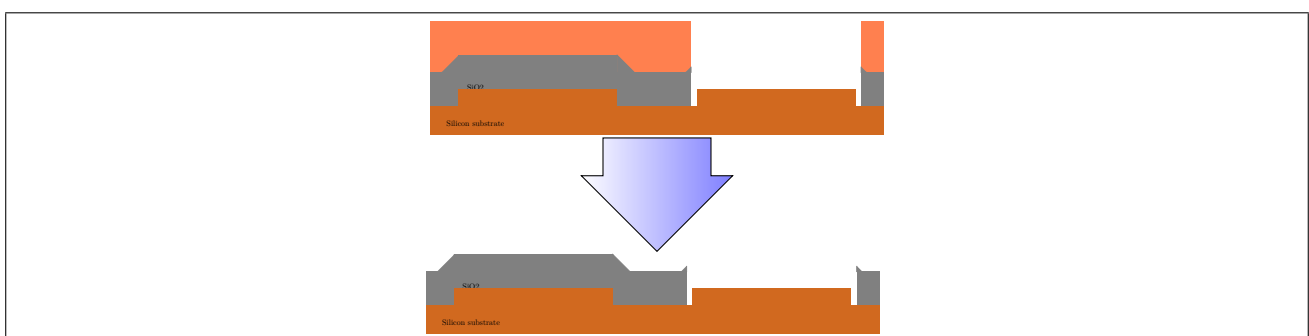
There are multiple possible approaches to etch through these 500nm of oxide.

**Possible approaches:**

- **"AOE Etcher (DRY-AOE)" from HKUST**  
We can use anisotropic plasma etching for sharper borders.
- **Chemical solution**  
We can use buffered hydrofluoric acid (BOE (1:6)) at room temperature for 5 minutes in order to get through the 500nm of oxide.  
Too long over 4 minutes might cause under-etch however!

## 2.4 Cleaning

In order to avoid contamination of the machines we need to make sure all the resist has been stripped off from the wafer.



**Figure 16:** Resist removal

Please just use the solvent for the specific resist.

## 2.5 Implantation/Doping

We now need to inject the carriers into the upper level of the n-channel area so that we can later on drive them into the crystal during the drive-in step.

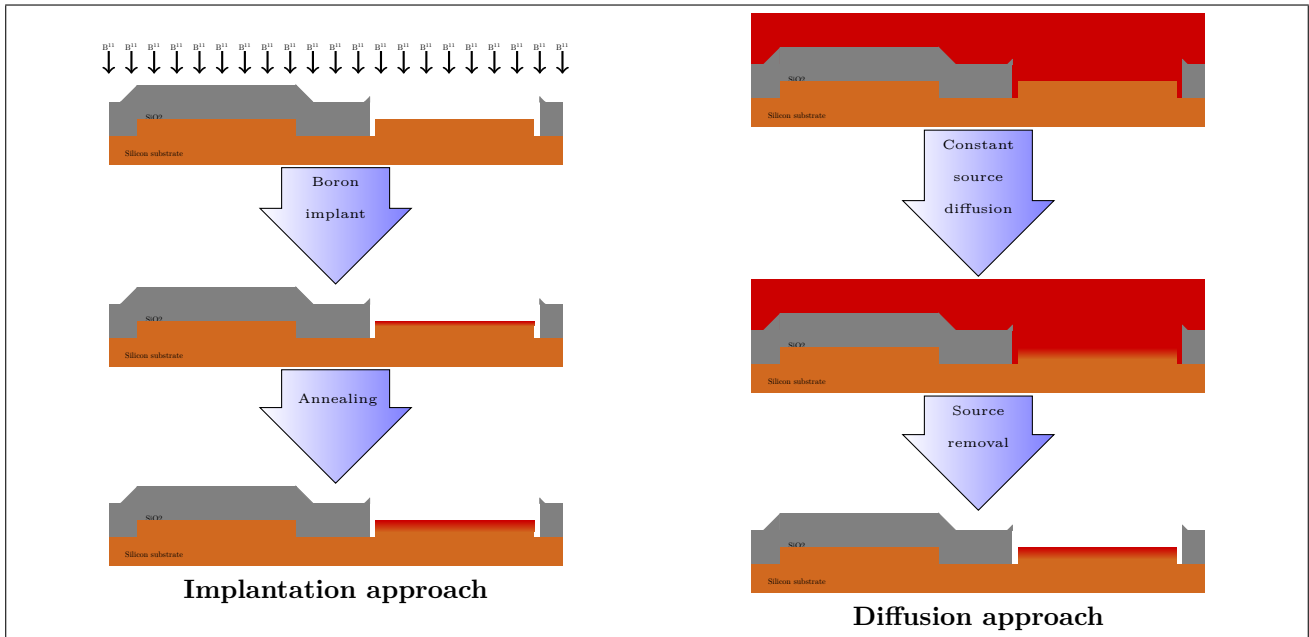


Figure 17: Doping process

Possible approaches:

- **"CF-3000 Implanter (IMP-3000)" from HKUST**

At HKUST we have an implanter which gives us better control over the initial surface concentration. These steps are needed to arrive with the desired geometry:

1. The P-well is implanted with a Boron ( $B^{11}$ ) dose of  $2.5 \times 10^{12} cm^{-2}$  at an energy of 100 keV
2. The P-well is annealed for 30 minutes at  $1050^{\circ}C$  in  $N_2$  environment (DIF-A1)  
After that the P-well will be around  $1\mu m$  deep and will become deeper during [subsection 3.5](#)

- **Constant source diffusion**

We can add a layer of Boron solution and diffusing in order to have an initial concentration in order to reach the desired concentration later by main diffusion.

1. A constant source is added (gas or liquid)
2. The source dopant is driven in for 10 minutes at  $1050^{\circ}C$
3. The dopant source is removed by stopping the gas flow or cleaning the surface

## 2.6 Oxide mask removal

Now we want to remove the silicon mask from the wafer and clean it for another clean oxide mask layer in order to perform the implantation of the N-well in the next step.

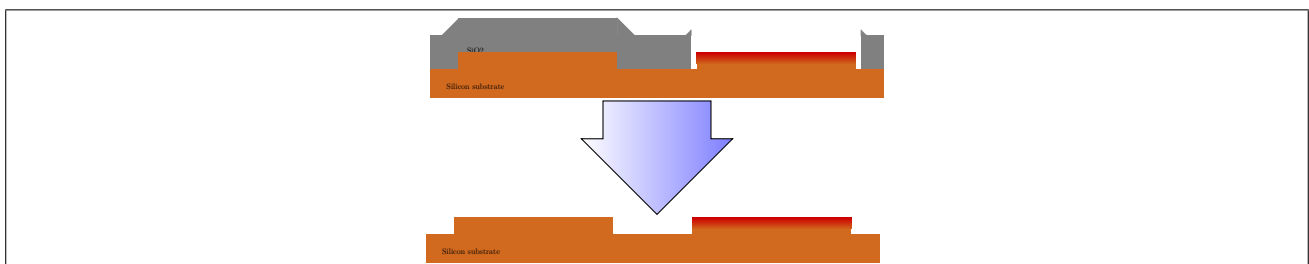
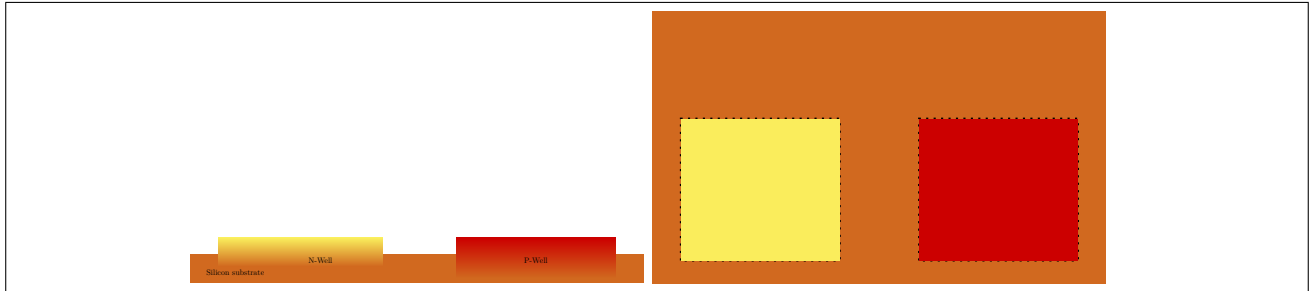


Figure 18: Oxide removal

We use buffered hydrofluoric acid (BOE (1:6)) at room temperature for 5 minutes in order to remove the 500nm of oxide layer.

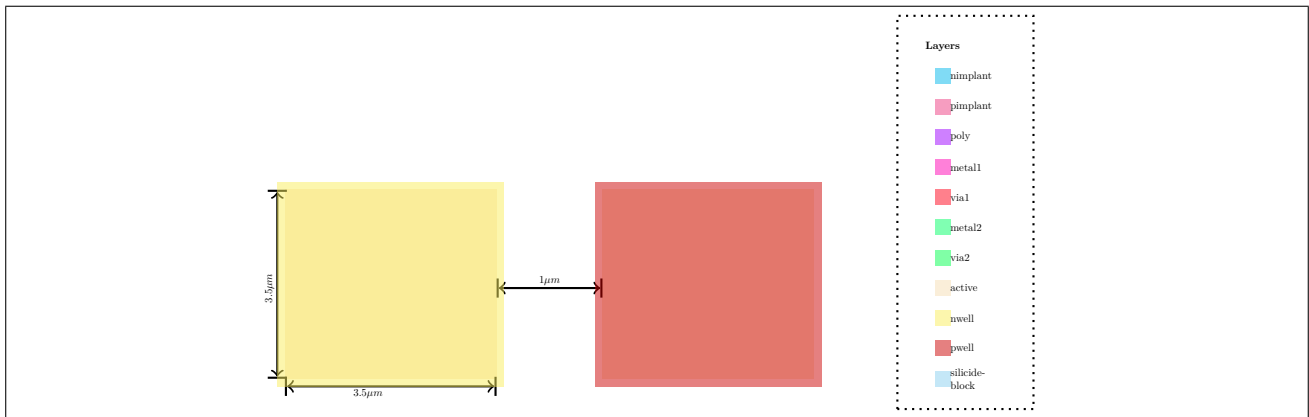
### 3 N-well

In order to build CMOS on the same substrate, an N-well is required for building the complementary P-channel transistor for a n-p-channel logic circuitry as shown above in the example section. The cross section as well as the top view of the targeted geometry are shown in [Figure 19](#)



**Figure 19:** N-well target geometry

The N-well will serve us as an island of N-doped substrate within the P-doped basis substrate. The dopant dose will be  $2.5 \times 10^{12} cm^{-2}$  as calculated in the documentation of the process design leading to these steps<sup>7</sup>.



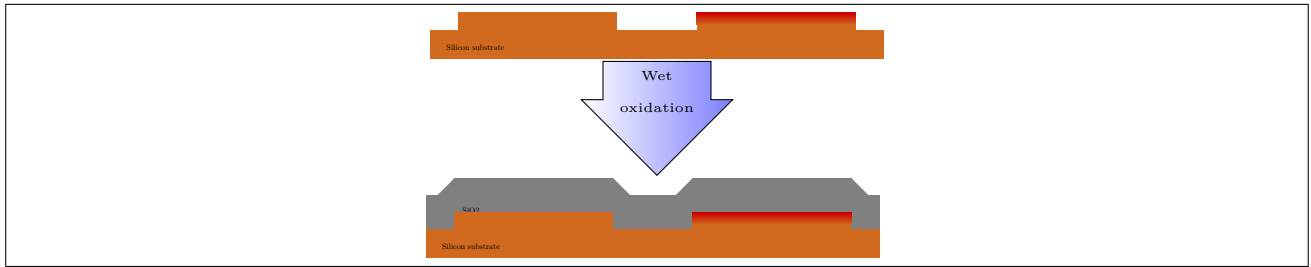
**Figure 20:** N-Well layout

In [Figure 20](#) the layout of the n-well region on top of the active area region can be seen. The n-well is being fit into the active area. It should even be a little bit bigger than the active area, because of possible alignment offsets

<sup>7</sup>[https://github.com/leviathanch/libresiliconprocess/raw/master/process\\_design/process\\_design.pdf](https://github.com/leviathanch/libresiliconprocess/raw/master/process_design/process_design.pdf)

### 3.1 Mask dioxide layer

In order to selectively inject charge carrying atoms into the crystalline structure a protective dioxide ( $\text{SiO}_2$ ) layer needs to be grown on top of a p-type substrate.



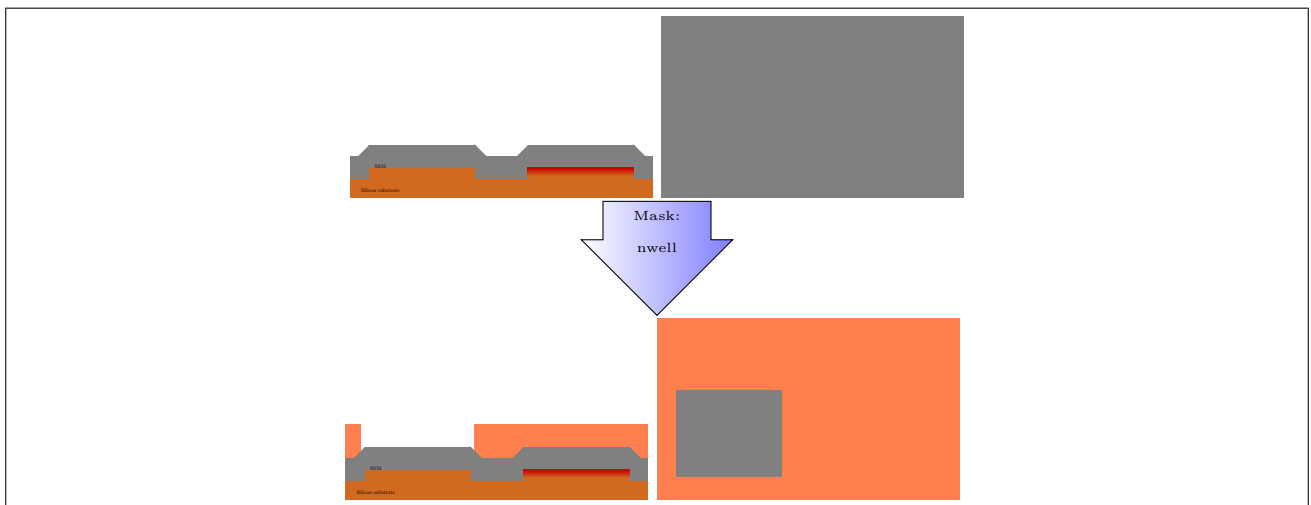
**Figure 21:** Dioxide layer growth

With an energy of 100keV for the implantation performed in subsection 3.5, the projected range of the dopants within the oxide will be 100nm (130nm tops)<sup>8</sup>. This means being on the safe side and having 300nm as the thickness is a good approach.

In order to grow the 300nm thick oxide layer, the wafer is being oxidized for around 25 minutes at 1050°C using wet oxidation which results in a dioxide layer of around 300nm in thickness<sup>9</sup>.

### 3.2 Patterning

The resist is being deposited using spray coating because the uneven nature of the oxide layer. After that the wafer is being soft baked depending on the baking time and temperature for the specific resist. The layout for being exposed onto the resist is being extracted from the "nwell" layer within the GDS2 file onto a **bright field** mask. The requirement is a **negative** tone resist.



**Figure 22:** Cross/top view of n-well layout on resist

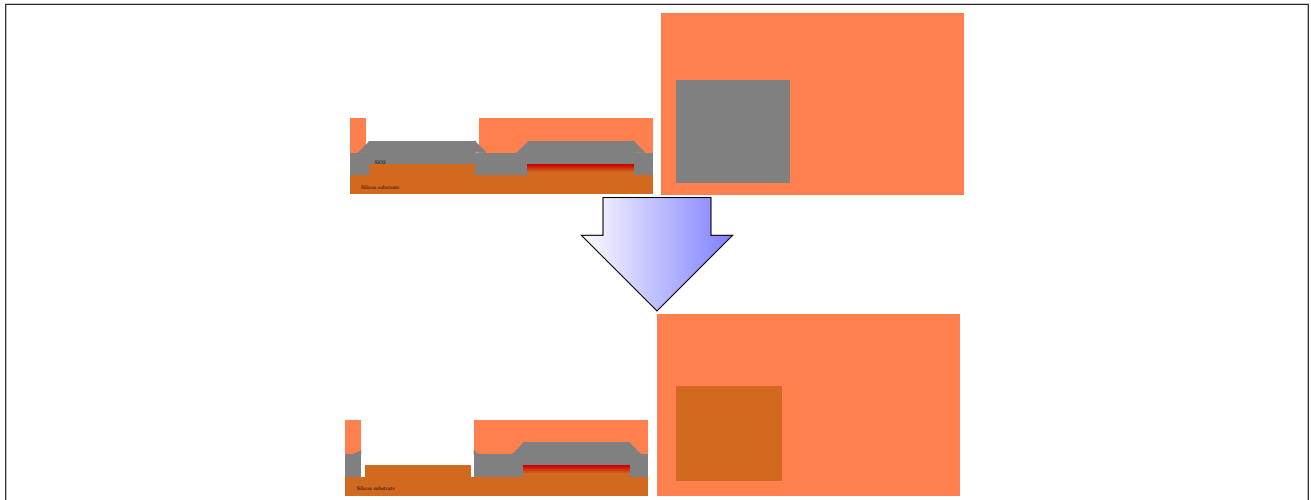
The thickness of the resist layer and the baking duration will variate depending on the specific equipment for which this process will be implemented with. Also after the exposure and development, the hard baking shouldn't be forgotten!

<sup>8</sup><http://cleanroom.byu.edu/rangestraggle>

<sup>9</sup><http://cleanroom.byu.edu/OxideTimeCalc>

### 3.3 Etching

We now need to open a window in the dioxide layer, through which we will inject carrier atoms into the silicon crystal structure.



**Figure 23:** Cross/top view of n-well oxide window

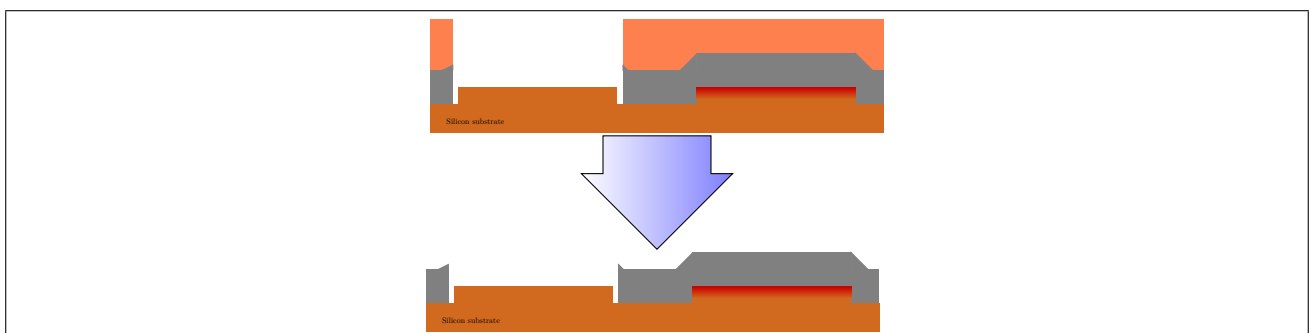
Since the silicon dioxide layer is 300nm thick and we wanna reach the silicon below we can use wet etching as described in the chemistry chapter.

Possible approaches:

- **"AOE Etcher (DRY-AOE)" from HKUST**  
We can use anisotropic plasma etching for sharper borders.
- **Chemical solution**  
We can use buffered hydrofluoric acid (BOE (1:6)) at room temperature for a little bit over 3 minutes in order to get through the 300nm of oxide.  
Too long over 3 minutes might cause under-etch however!

### 3.4 Cleaning

In order to avoid contamination of the machines we need to make sure all the resist has been stripped off from the wafer.



**Figure 24:** Resist removal

Please just use the solvent for the specific resist.



### 3.5 Implantation/Doping

We now need to inject the carriers into the upper level of the n-channel area so that we can later on drive them into the crystal during the drive-in step.

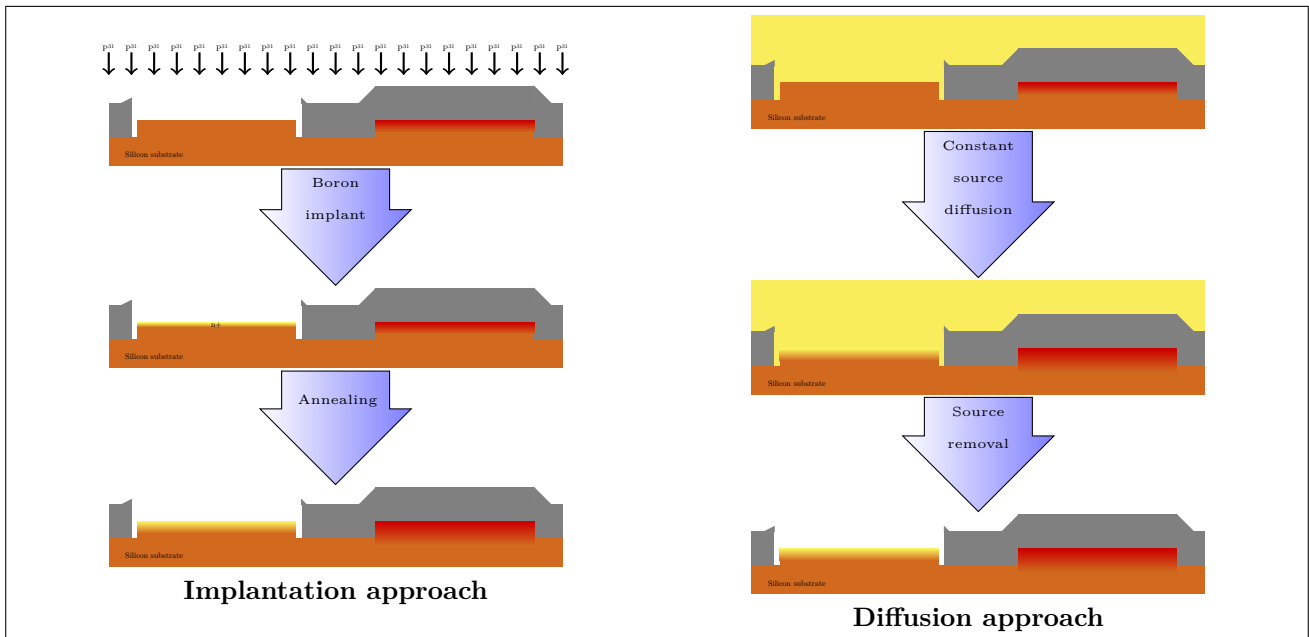


Figure 25: Doping process

Possible approaches:

- **"CF-3000 Implanter (IMP-3000)" from HKUST**

At HKUST we have an implanter which gives us better control over the initial surface concentration. These steps are needed to arrive with the desired geometry:

1. The N-well is implanted with a Phosphorus ( $P^{31}$ ) dose of  $2.5 \times 10^{12} cm^{-2}$  at an energy of 100 keV.
2. The N-well is annealed for 30 minutes at  $1050^{\circ}C$  in  $N_2$  environment (DIF-A1)  
After that the P-well will be around  $2\mu m$  deep and the N-Well around  $1\mu m$  deep

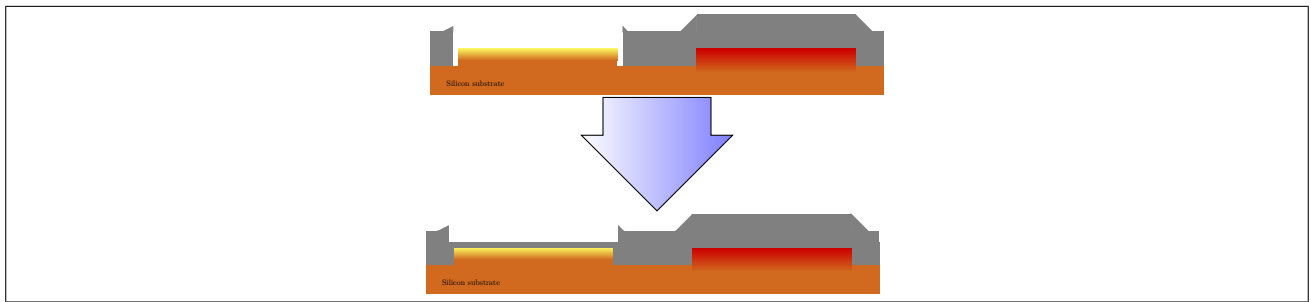
- **Constant source diffusion**

We can add a layer of Phosphorus solution and diffusing in order to have an initial concentration in order to reach the desired concentration later by main diffusion.

1. A constant source is added (gas or liquid)
2. The source dopant is driven in for 10 minutes at  $1050^{\circ}C$
3. The dopant source is removed by stopping the gas flow or cleaning the surface

### 3.6 Oxide for drive-in

Now we need to cover the now doped and annealed area with an oxide layer in order to seal it off and prevent the dopants from diffusing out of the crystal during the drive-in phase.

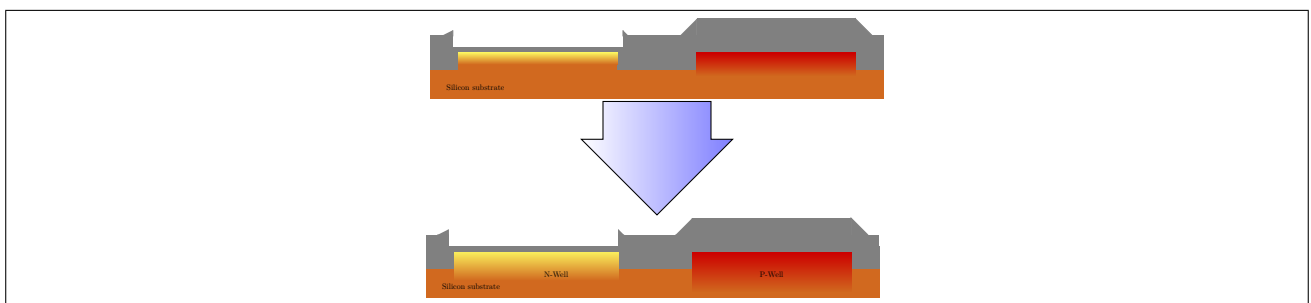


**Figure 26:** Oxide growth

The wafer is being oxidized for around 5 minutes at 1050°C in a wet environment in order to achieve a cover silicon layer of around 90nm thickness.

### 3.7 Drive-in

In order to drive the carrier atoms deeper into the crystalline structure the wafer needs to be driven in after predeposition.

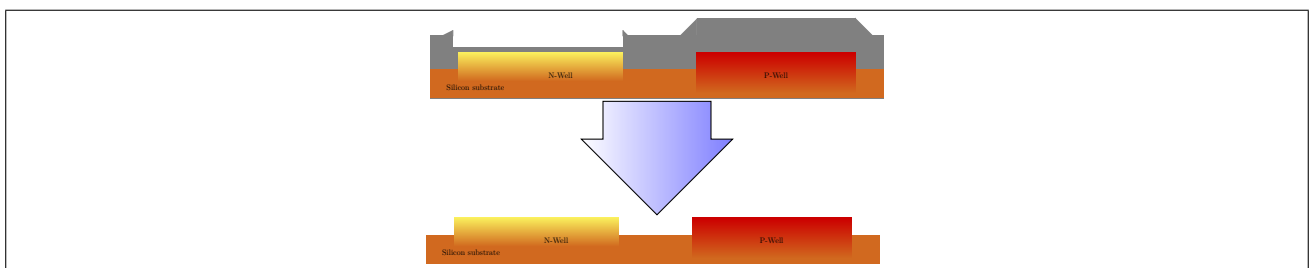


**Figure 27:** Drive-in process

In this step the wafer is driven-in for 90 minutes at 1050°C in an inert ambient<sup>10</sup>

### 3.8 Oxide mask removal

Now we want to remove the silicon mask from the wafer and clean it for another clean oxide mask layer.



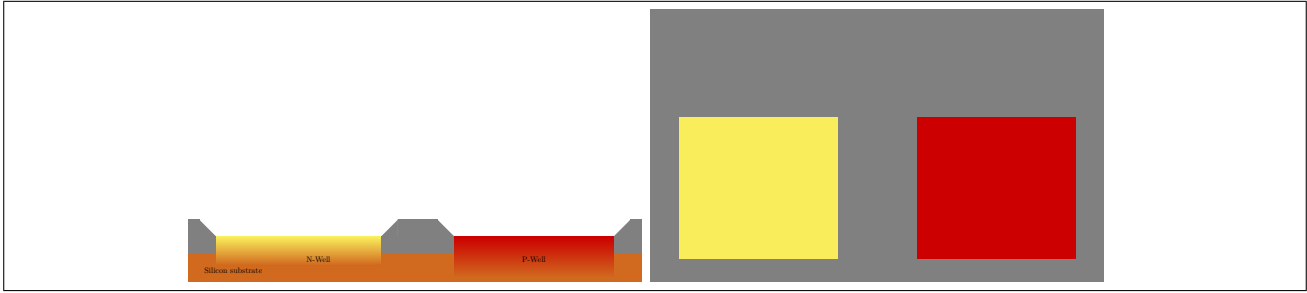
**Figure 28:** Oxide removal

We use buffered hydrofluoric acid (BOE (1:6)) at room temperature for 3 minutes in order to remove the 300nm of oxide layer.

<sup>10</sup><http://cleanroom.byu.edu/DopConCalc>

## 4 Field oxide

The geometry of a substrate with the field oxide filling the shallow trench from [section 1](#) now needs to be made.

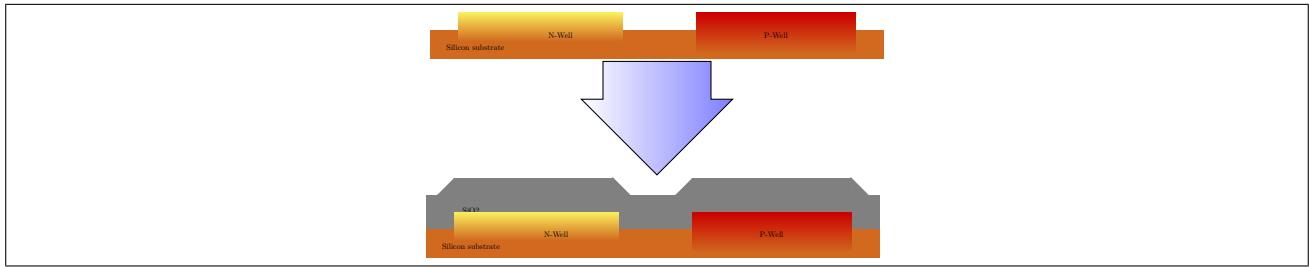


**Figure 29:** Shallow trench isolation target geometry

As can be seen in [Figure 29](#), the STI trenches need to be filled with silicon oxide and windows need to be etched into them so that the gate can be constructed later on. The windows are needed so that the poly silicon is far enough away from the non-active areas so that the threshold voltage of the parasitic FETs is so high that they will never switch. Only within the active areas we want to allow the poly layer to touch down closer to the silicon.

## 4.1 Oxide growth

Now we need to fill the trenches with silicon dioxide which will provide a spacer between the non active area and the polysilicon gate layer. within the non-active areas.

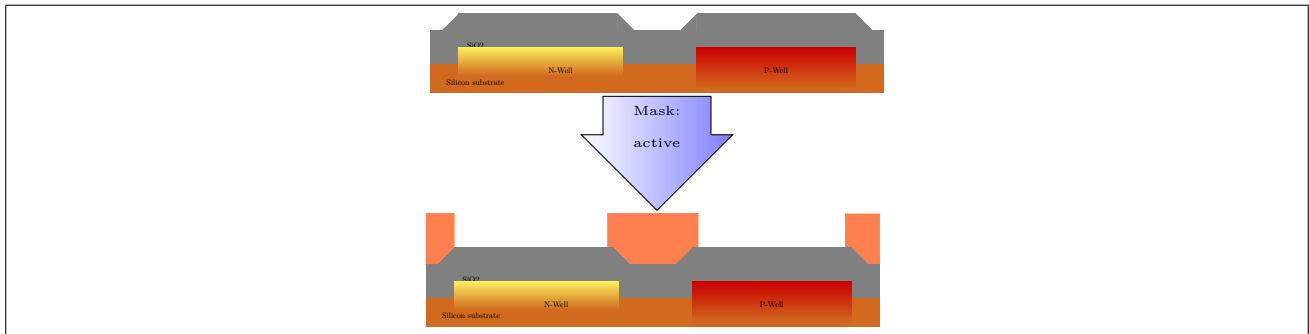


**Figure 30:** Hard mask growth

We grow a  $1\mu m$  thick layer of silicon dioxide by putting the wafer into the furnace at  $1050^{\circ}C$  for 3 hours in a wet environment.

## 4.2 Patterning

We reuse the mask from [section 1](#), because it's exactly the same layout, only inverted. The requirement is a **negative** tone resist.

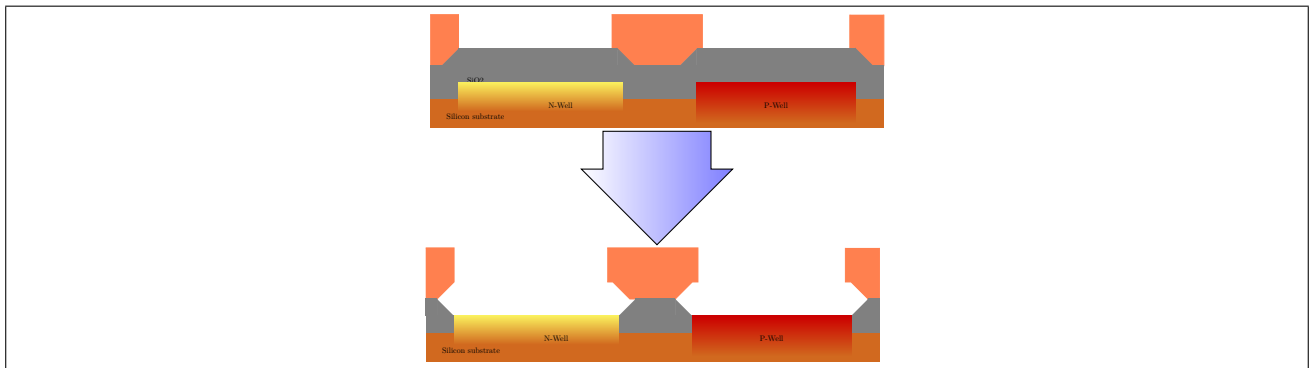


**Figure 31:** Patterning with positive resist

The thickness of the resist layer and the baking duration will variate depending on the specific equipment for which this process will be implemented with. Also after the exposure and development, the hard baking shouldn't be forgotten!

### 4.3 Etching

We open the access to the silicon inside of the active areas in order to touch down with the polysilicon further on.



**Figure 32:** Nitride mask etching

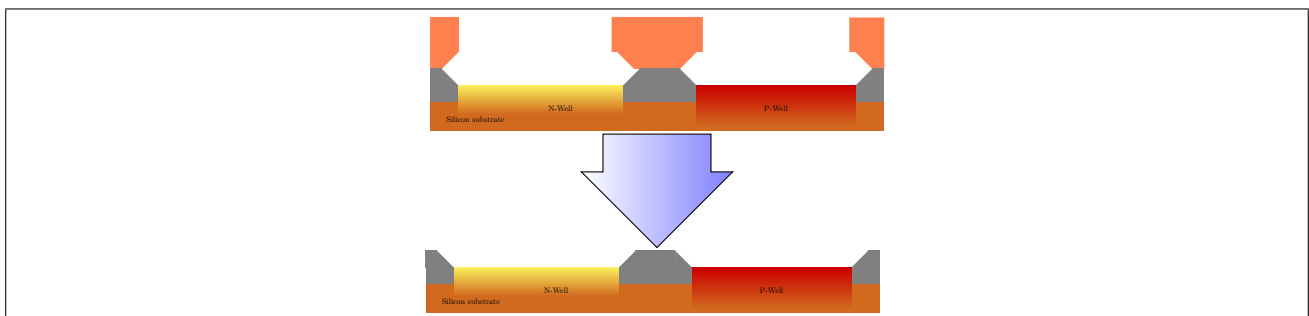
There are dry etching and wet etching methods available for etching the thick field oxide. The downside of wet etching is that it also etches horizontally, however the chemical BHF is readily available and allows for easy implementation of the process.

**Possible approaches:**

- **"AOE Etcher (DRY-AOE)" from HKUST**  
We can use anisotropic plasma etching for sharper borders.
- **Chemical solution**  
We can use buffered hydrofluoric acid (BOE (1:6)) at room temperature for a little bit over 3 minutes in order to get through the 300nm of oxide.  
Too long over 3 minutes might cause under-etch however!

### 4.4 Resist removal

Now we need to remove the contaminants for further processing.

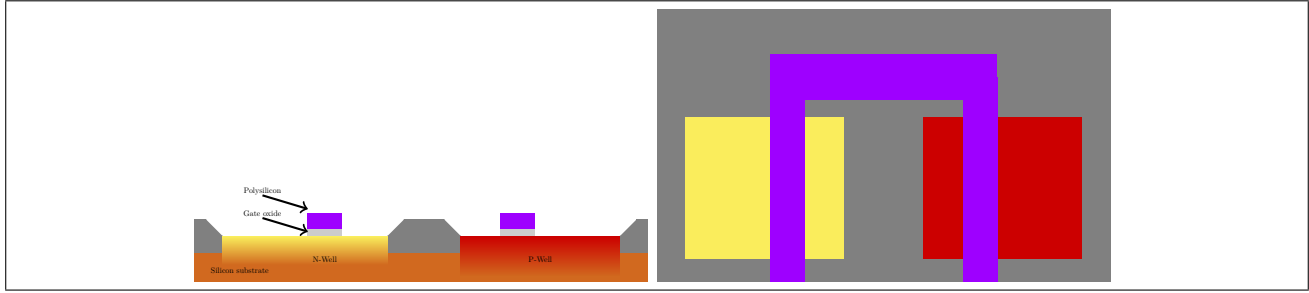


**Figure 33:** Resist removal

We strip the resist, rinse and perform sulfuric cleaning.

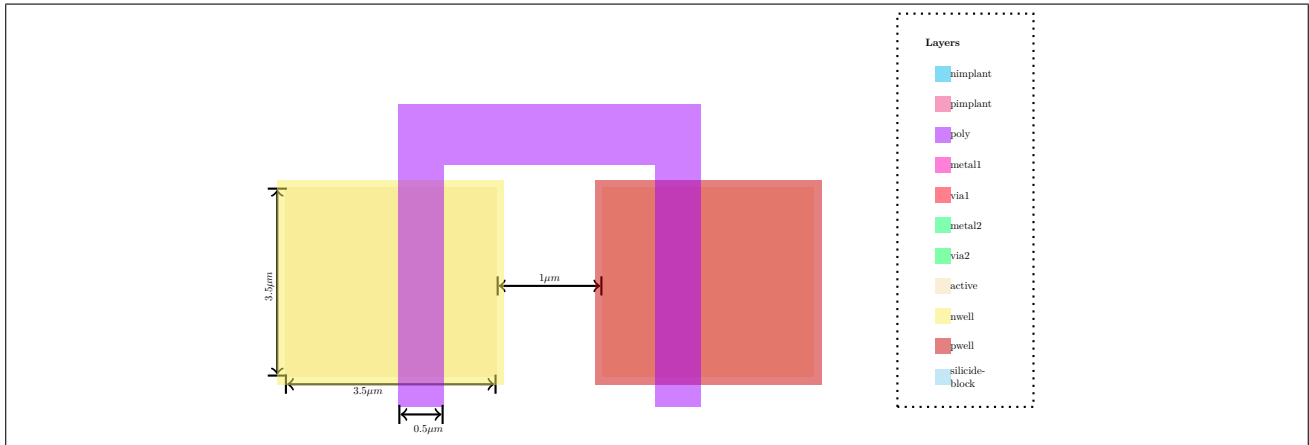
## 5 Gate

Now we have to build the initial gate structure which contains of the 40nm thick dielectric (in our case just silicon dioxide) and the polysilicon electrode.



**Figure 34:** Poly silicon gate contacts with gate oxide

The line spacing of the polysilicon electrode shape has to be at least  $0.5\mu m$  because of the resolution of the stepper and also because of the etching process which has  $0.5\mu m$  as the minimum line spacing.

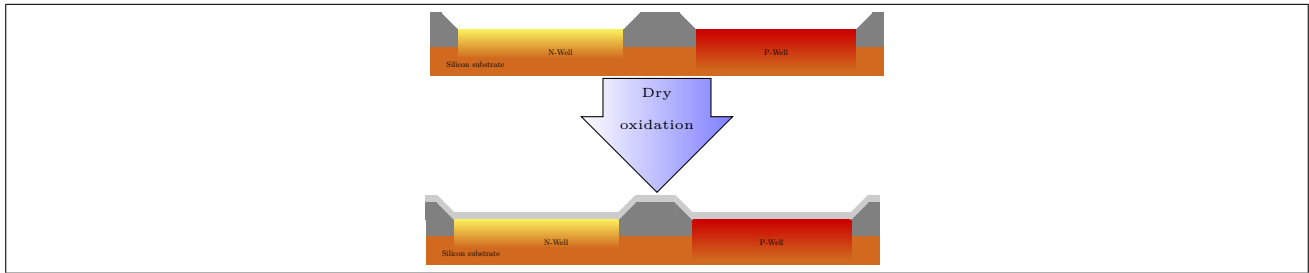


**Figure 35:** Gate layout

In [Figure 35](#) we can see the layout honoring the  $0.5\mu m$  spacing design rule for the gate structure shape and poly-layer interconnect between NMOS and PMOS.

## 5.1 Gate oxide deposition

Now we have to deposit the dielectric isolator between the gate electrode and the channel. As designed in the process design document, the layer will be 40nm thick.

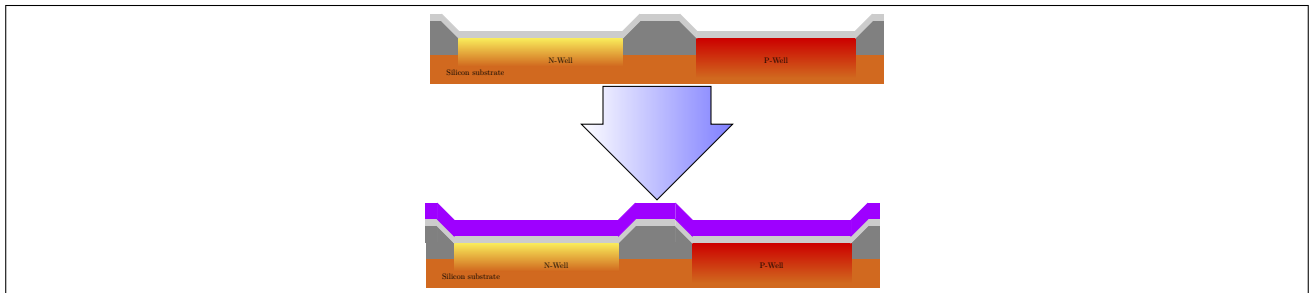


**Figure 36:** Thin oxide

The thickness of this layer decides over many critical key properties of the transistor, hence there should be little to no variation in the thickness of the gate oxide layer. For that reason we put the wafer into the diffusion furnace and perform dry oxidation at  $1050^{\circ}\text{C}$  for 33 minutes and 14 seconds.<sup>11</sup>

## 5.2 Polysilicon deposition

Now we need to add the polysilicon layer for forming the gate structure after etching.



**Figure 37:** Polysilicon

We use the LPCVD machine and deposit a layer of around 600nm polysilicon<sup>12</sup>.

We set the temperature to  $650^{\circ}\text{C}$ , the gas will be Silane ( $\text{SiH}_4$  ( $\text{Si} + 2\text{H}_2$ )), the pressure will be set to 300 mTorr with a flow of 90sccm.

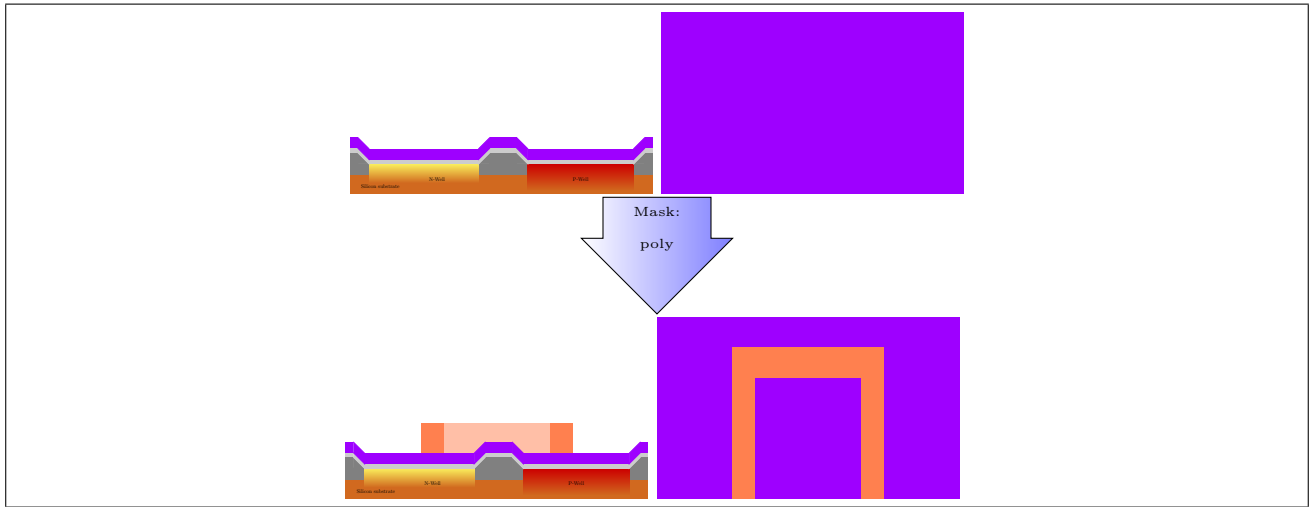
This will give us a growth rate of roughly 23.5 nm per minute, so for 600nm we let it grow half an hour.

<sup>11</sup><http://cleanroom.byu.edu/OxideTimeCalc>

<sup>12</sup>[https://people.rit.edu/lffeee/LPCVD\\_Recipes.pdf](https://people.rit.edu/lffeee/LPCVD_Recipes.pdf)

### 5.3 Patterning

The resist is being deposited using spin coating and then baked depending on the baking time for the specific resist. The layout for being exposed onto the resist is being extracted from the "poly" layer within the GDS2 file onto a **bright field** mask. The requirement is a **positive** tone resist.



**Figure 38:** Resist

The thickness of the resist layer and the baking duration will variate depending on the specific equipment for which this process will be implemented with. Also after the exposure and development, the hard baking shouldn't be forgotten!



## 5.4 Etching

Now we've got to etch the gate structures.

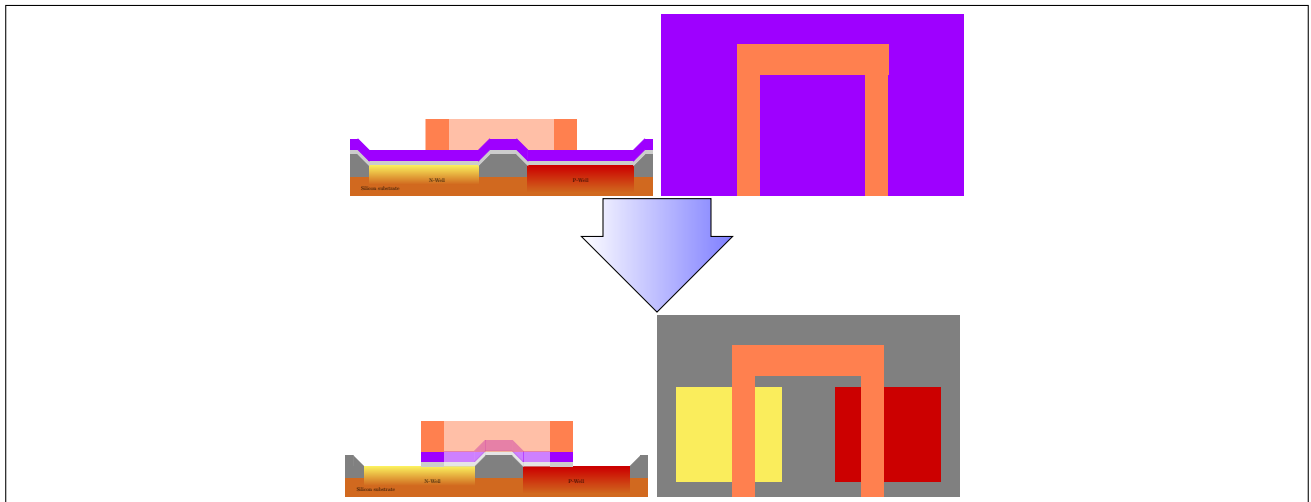


Figure 39: Resist

For now we only have the plasma etcher variant being verified because chemically etching polysilicon isn't allowed at the HKUST labs for contamination control reasons. In case you can verify this in your lab with a chemical etching method, please update this chapter and make a pull request!

Possible approaches:

- **"Poly Etcher (DRY-Poly)" from HKUST**

An anisotropic plasma etcher, in order to etch the polysilicon and gate oxide layer. In [subsection 5.2](#) we've grown 600nm of polysilicon which takes 200 seconds (= 3 minutes 20 seconds) (at 180nm/min) to etch. The selectivity to oxide is 13:1 which leads to an oxide etching speed of around 14nm/min, in [subsection 5.1](#) we've grown a 40nm thick oxide layer which leads to the oxide adding another 2 minutes 51 seconds to the etching time. All together, we will have to etch for around **6 minutes and 10 seconds**.

- **Chemical method**

Please add a verified method here!

## 5.5 Resist removal

Now we need to remove the contaminants for further processing.

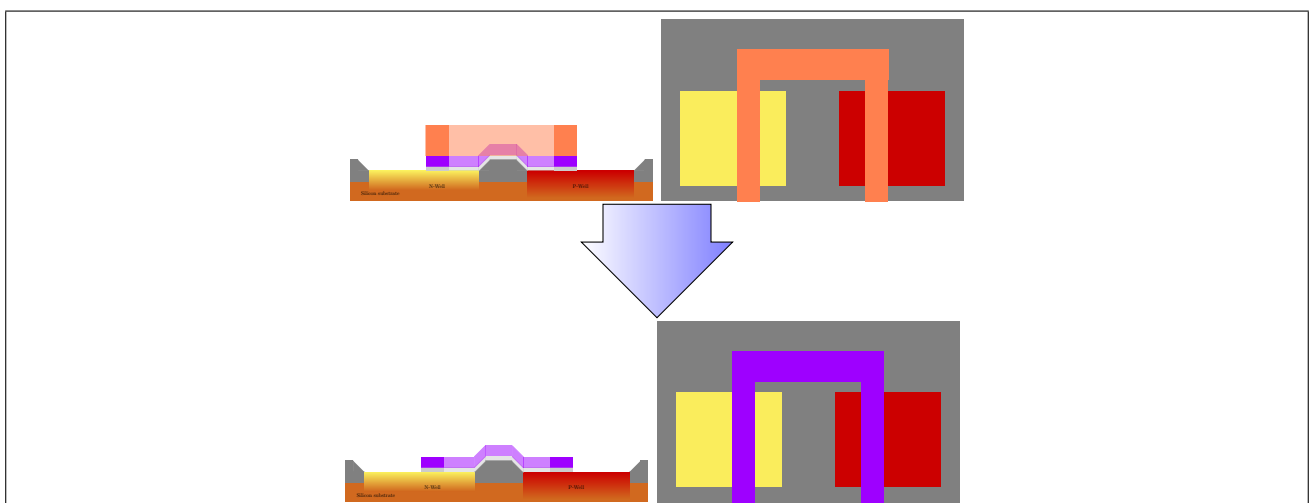
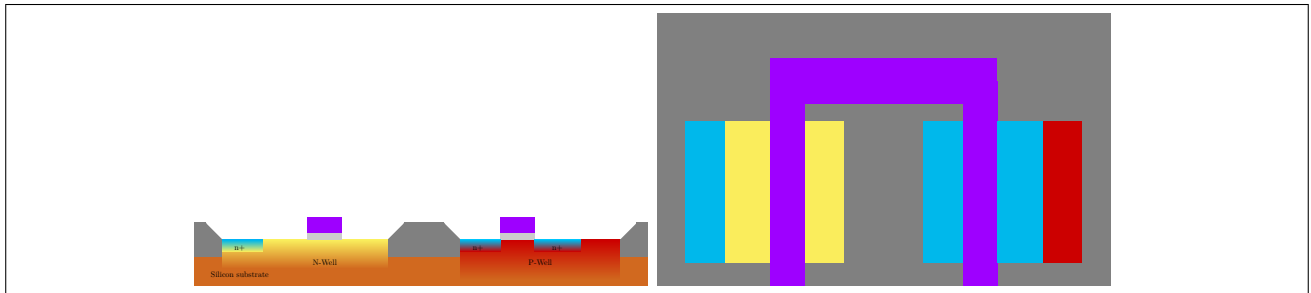


Figure 40: Resist

We strip the resist, rinse and perform sulfuric cleaning.

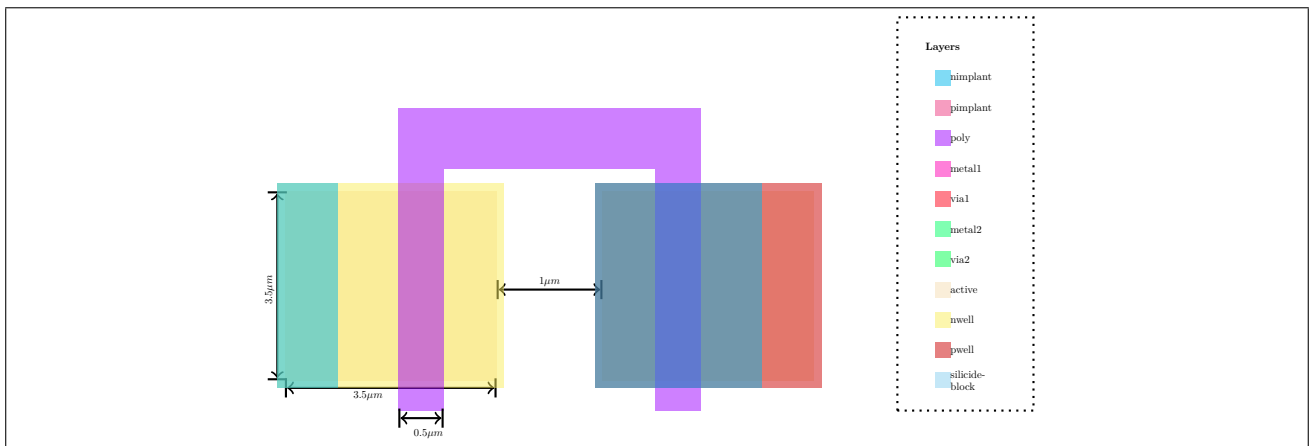
## 6 n+ Implant

For the bulk of the PMOS transistors and for the source and drain of the NMOS transistors highly doped n+ areas are required. In this step we're going to build these.



**Figure 41:** N+ implant geometry target

The tricky thing here is to have a reasonable implant depth but not too deep because the deeper the junction, the higher the junction capacity which in turn limits the switching performance of the CMOS circuitry.



**Figure 42:** N+ layout

An example layout of p-implants can be seen in [Figure 42](#), the mask is being extracted from the layer "n\_plus\_select".

Also important to notice is that this example layout is just for demonstration purposes only, please have a look at the standard cell documentation for the actual layouts.

## 6.1 Mask dioxide layer

In order to selectively inject charge carrying atoms into the crystalline structure a protective dioxide ( $SiO_2$ ) layer needs to be grown on top of a p-type substrate.

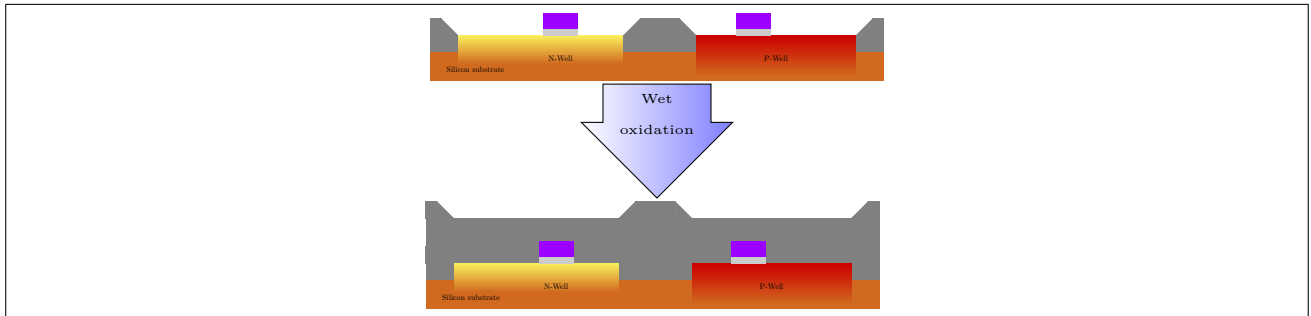


Figure 43: Oxide layer

With an energy of 35keV for the implantation performed in subsection 6.5, the projected range of the dopants within the oxide will be 34nm (47nm tops) <sup>13</sup>. This means being on the safe side and having 100nm as the thickness is a good approach. In order to grow the 100nm thick oxide layer, the wafer is being oxidized for around 5 minutes 30 seconds at 1050°C using wet oxidation which results in a dioxide layer of around 100nm in thickness <sup>14</sup>.

## 6.2 Patterning

The resist is being deposited using spin coating and then baked depending on the baking time for the specific resist. The layout for being exposed onto the resist is being extracted from the "n\_plus\_select" layer within the GDS2 file onto a **bright field** mask. The requirement is a **negative** tone resist.

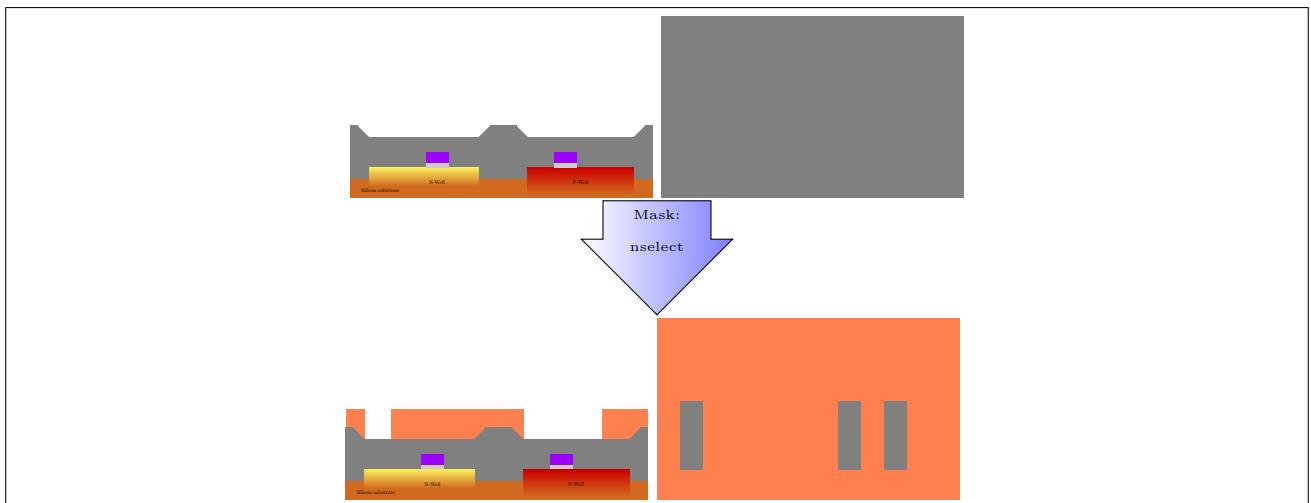


Figure 44: N+ region resist mask

The thickness of the resist layer and the baking duration will variate depending on the specific equipment for which this process will be implemented with. Also after the exposure and development, the hard baking shouldn't be forgotten!

<sup>13</sup><http://cleanroom.byu.edu/rangestraggle>

<sup>14</sup><http://cleanroom.byu.edu/OxideTimeCalc>

## 6.3 Etching

We now need to open a window in the dioxide layer, through which we will inject carrier atoms into the silicon crystal structure.

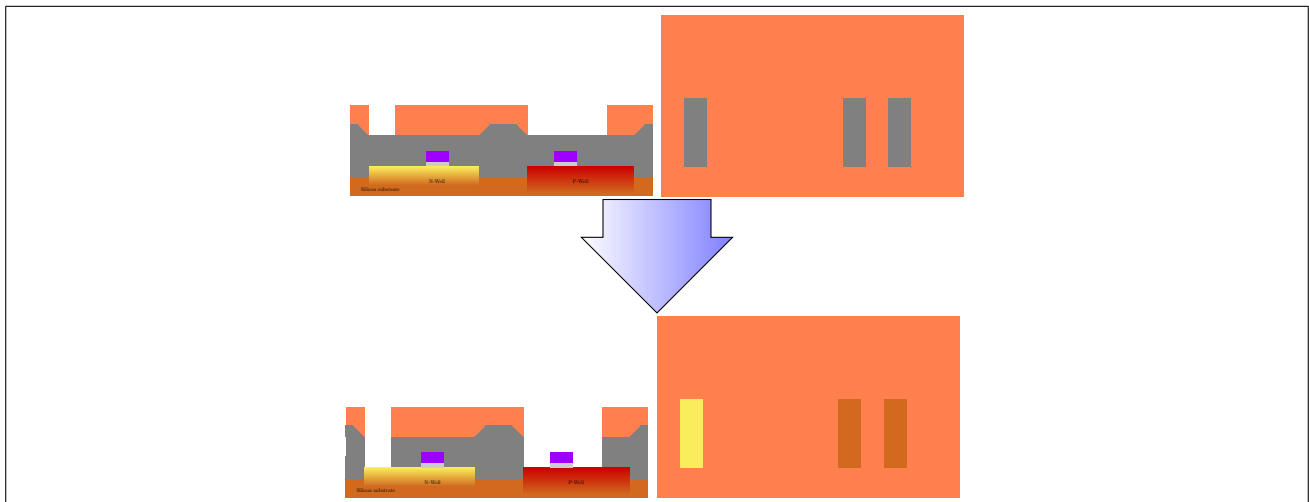


Figure 45: N+ region opened

Since the silicon dioxide layer is 100nm thick and we wanna reach the silicon below we can use wet etching as described in the chemistry chapter.

Possible approaches:

- **"AOE Etcher (DRY-AOE)" from HKUST**  
We can use anisotropic plasma etching for sharper borders.
- **Chemical solution**  
We can use buffered hydrofluoric acid (BOE (1:6)) at room temperature for a little bit over 3 minutes in order to get through the 100nm of oxide.  
Too long over 1 minutes might cause under-etch however!

## 6.4 Cleaning

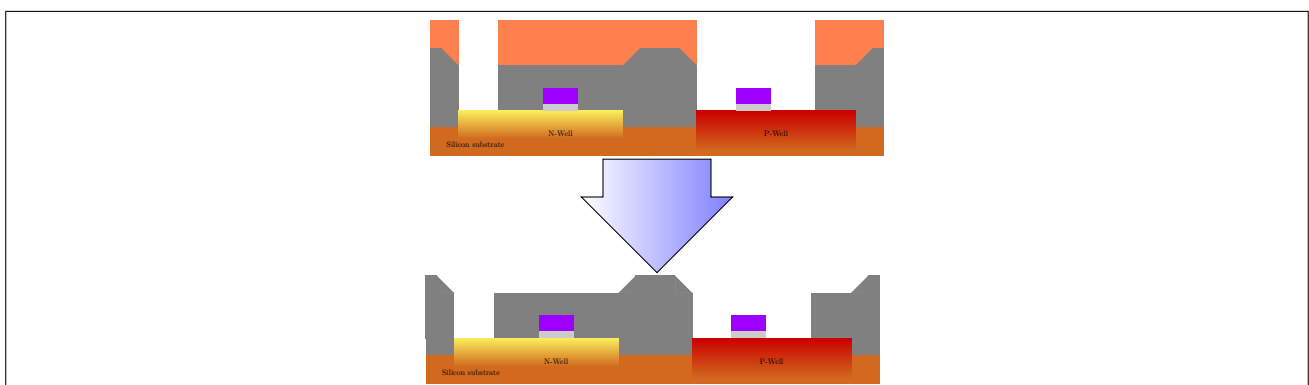


Figure 46: Resist removal

## 6.5 Implantation/Doping

We now need to bring in the carriers in order to build the n-junctions.

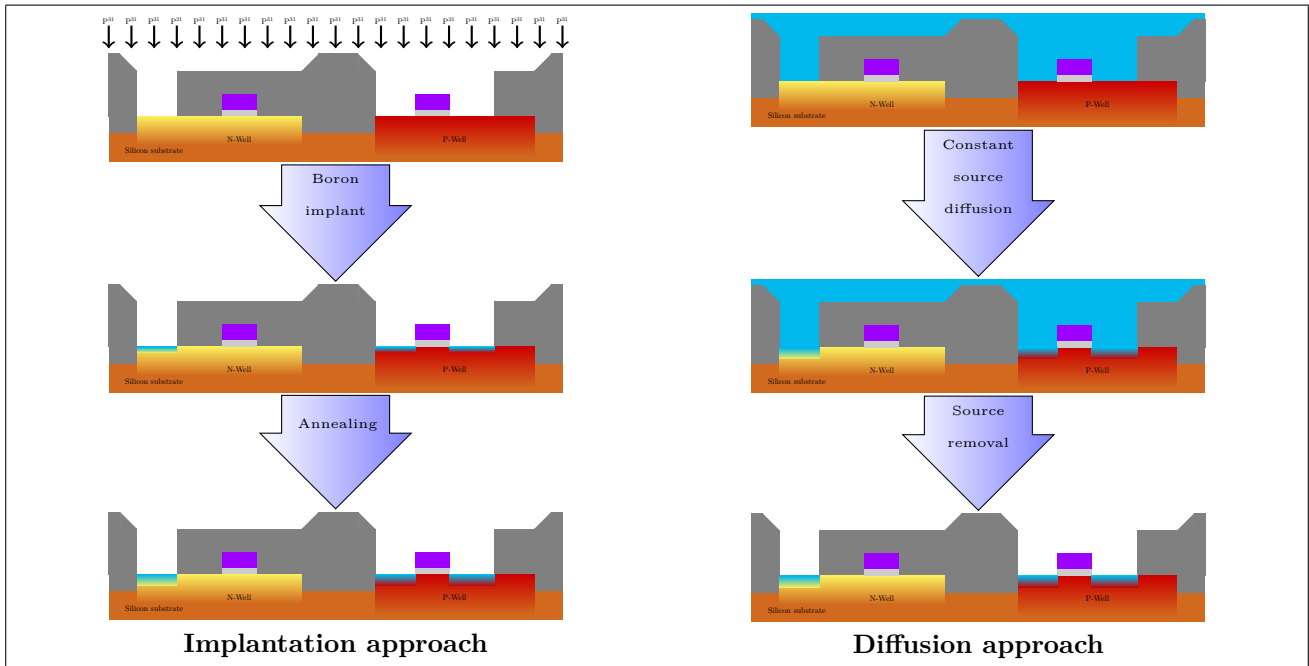


Figure 47: N+ doping process

Possible approaches:

- **"CF-3000 Implanter (IMP-3000)" from HKUST**

At HKUST we have an implanter which gives us better control over the initial surface concentration. These steps are needed to arrive with the desired geometry:

1. The nselect is implanted with a Phosphorus ( $P^{31}$ ) dose of  $2.5 \times 10^{12} cm^{-2}$  at an energy of 35 keV (43nm±18nm deep)
2. The nselect is annealed for 10 minutes at 1050°C in  $N_2$  environment (DIF-A1)

- **Constant source diffusion**

We can add a layer of Phosphorus solution and diffusing in order to have an initial concentration in order to reach the desired concentration later by main diffusion.

1. A constant source is added (gas or liquid)
2. The source dopant is driven in for 5 minutes at 1050°C
3. The dopant source is removed by stopping the gas flow or cleaning the surface

## 6.6 Oxide removal

Now we want to remove the silicon mask from the wafer and clean it for another clean oxide mask layer.

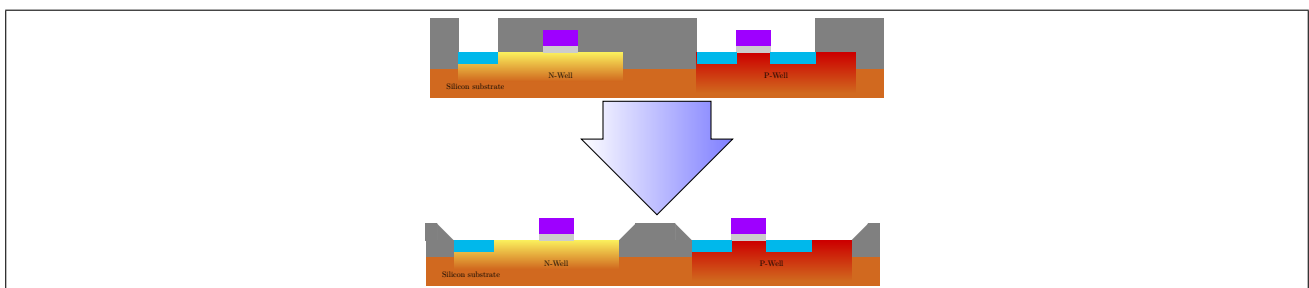
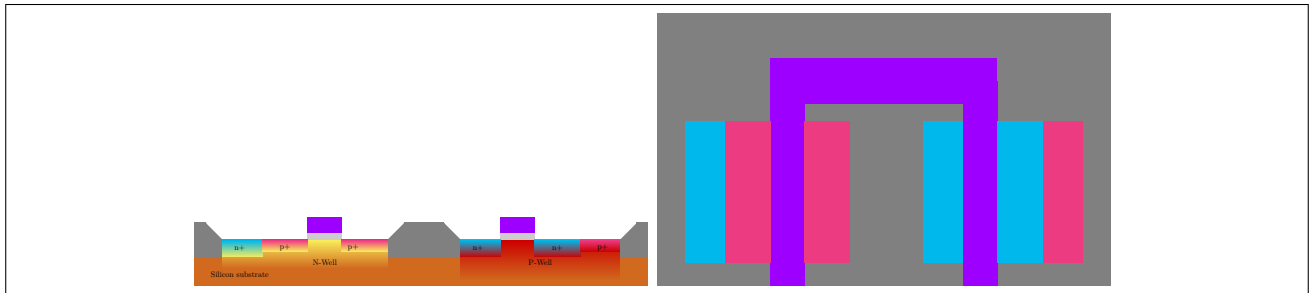


Figure 48: Oxide removal

We use buffered hydrofluoric acid (BOE (1:6)) at room temperature for 1 minute in order to remove the 100nm of oxide layer.

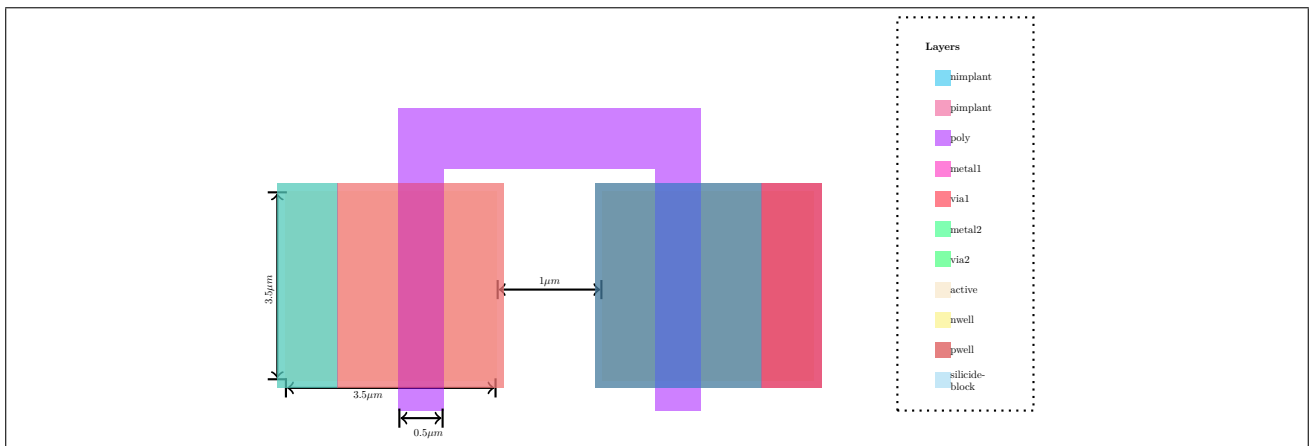
## 7 p+ Implant

For the bulk of the NMOS transistors and for the source and drain of the PMOS transistors highly doped p+ areas are required. In this step we're going to build these.



**Figure 49:** P+ implant geometry target

The tricky thing here is to have a reasonable implant depth but not too deep because the deeper the junction, the higher the junction capacity which in turn limits the switching performance of the CMOS circuitry.



**Figure 50:** P+ layout

An example layout of p-implants can be seen in [Figure 50](#), the mask is being extracted from the layer "p\_plus\_select"

Also important to notice is that this example layout is just for demonstration purposes only, please have a look at the standard cell documentation for the actual layouts.

## 7.1 Mask dioxide layer

In order to selectively inject charge carrying atoms into the crystalline structure a protective dioxide ( $SiO_2$ ) layer needs to be grown on top of a p-type substrate.

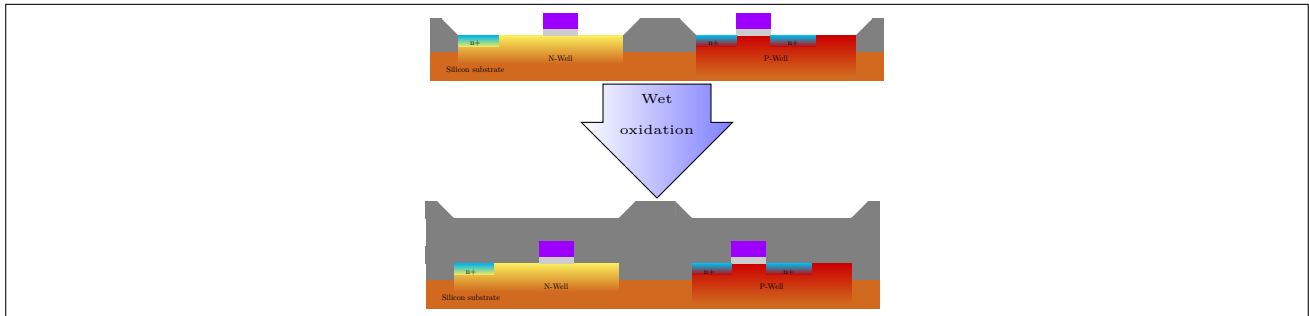


Figure 51: Oxide layer

With an energy of 13keV for the implantation performed in subsection 6.5, the projected range of the dopants within the oxide will be 40nm (58nm tops) <sup>15</sup>. This means being on the safe side and having 100nm as the thickness is a good approach. In order to grow the 100nm thick oxide layer, the wafer is being oxidized for around 5 minutes 30 seconds at 1050°C using wet oxidation which results in a dioxide layer of around 100nm in thickness <sup>16</sup>.

## 7.2 Patterning

The resist is being deposited using spin coating and then baked depending on the baking time for the specific resist. The layout for being exposed onto the resist is being extracted from the "p\_plus\_select" layer within the GDS2 file onto a **bright field** mask. The requirement is a **negative** tone resist.

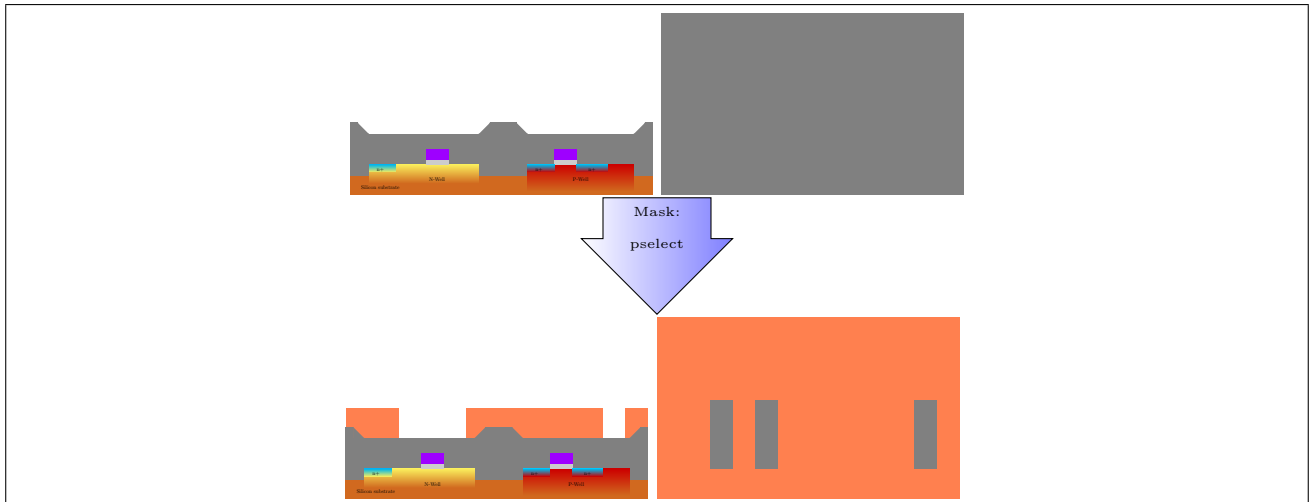


Figure 52: P+ region resist mask

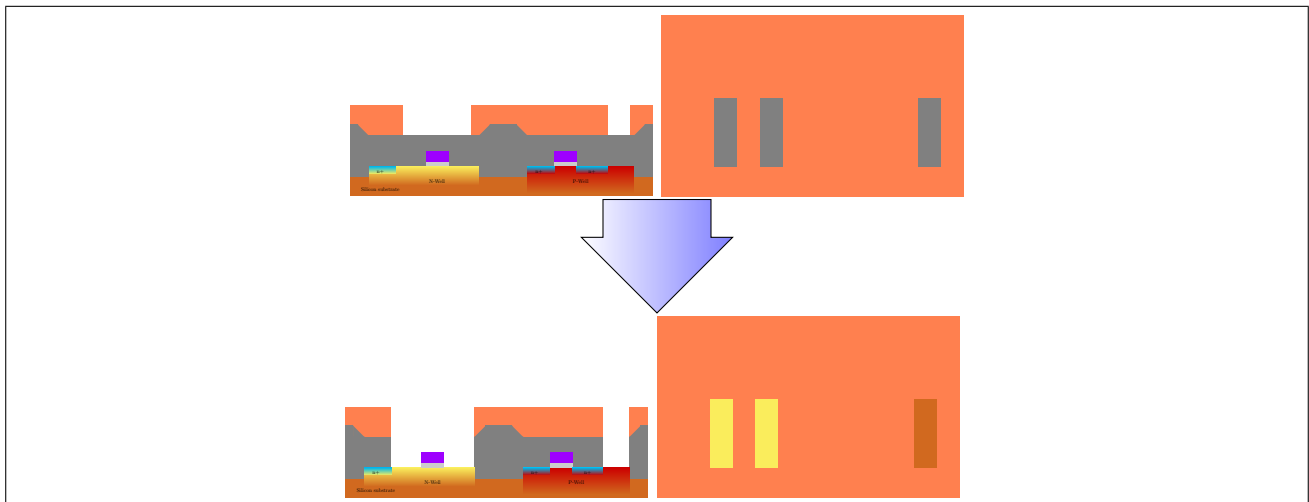
The thickness of the resist layer and the baking duration will variate depending on the specific equipment for which this process will be implemented with. Also after the exposure and development, the hard baking shouldn't be forgotten!

<sup>15</sup><http://cleanroom.byu.edu/rangestruggle>

<sup>16</sup><http://cleanroom.byu.edu/OxideTimeCalc>

### 7.3 Etching

We now need to open a window in the dioxide layer, through which we will inject carrier atoms into the silicon crystal structure.



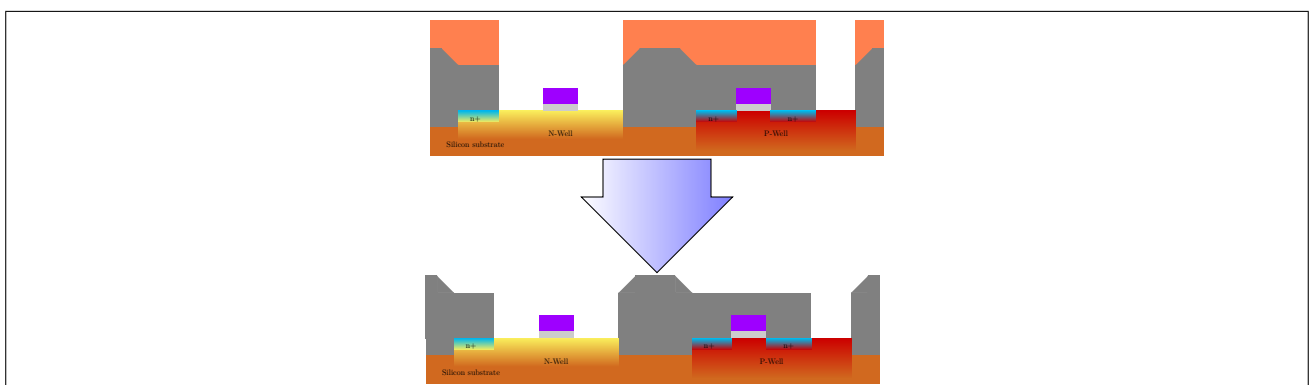
**Figure 53:** P+ region opened

Since the silicon dioxide layer is 100nm thick and we wanna reach the silicon below we can use wet etching as described in the chemistry chapter.

**Possible approaches:**

- **"AOE Etcher (DRY-AOE)" from HKUST**  
We can use anisotropic plasma etching for sharper borders.
- **Chemical solution**  
We can use buffered hydrofluoric acid (BOE (1:6)) at room temperature for a little bit over 3 minutes in order to get through the 100nm of oxide.  
Too long over 1 minutes might cause under-etch however!

### 7.4 Cleaning



**Figure 54:** Resist removal



## 7.5 Implantation/Doping

We now need to bring in the carriers in order to build the p-junctions.

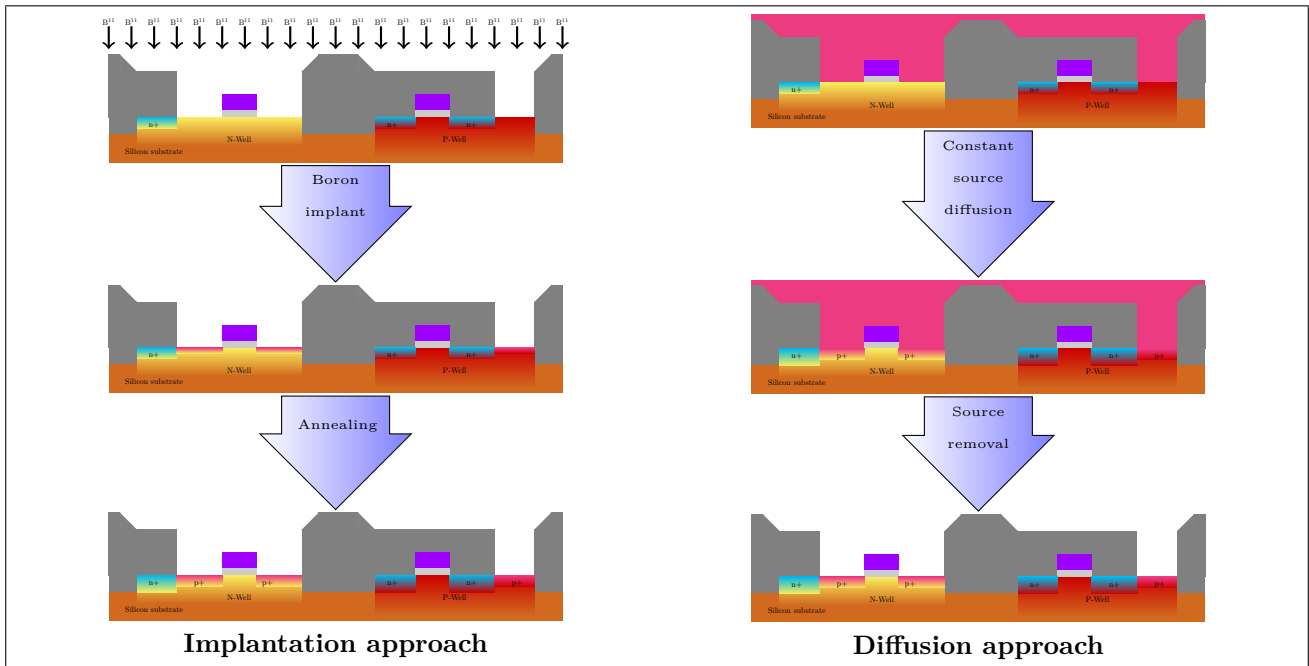


Figure 55: P+ doping process

Possible approaches:

- **"CF-3000 Implanter (IMP-3000)" from HKUST**

At HKUST we have an implanter which gives us better control over the initial surface concentration. These steps are needed to arrive with the desired geometry:

1. The pselect is implanted with a Boron ( $B^{11}$ ) dose of  $2.5 \times 10^{12} cm^{-2}$  at an energy of 13 keV ( $43nm \pm 18nm$  deep)
2. The pselect is annealed for 10 minutes at  $1050^{\circ}C$  in  $N_2$  environment (DIF-A1)

- **Constant source diffusion**

We can add a layer of Phosphorus solution and diffusing in order to have an initial concentration in order to reach the desired concentration later by main diffusion.

1. A constant source is added (gas or liquid)
2. The source dopant is driven in for 5 minutes at  $1050^{\circ}C$
3. The dopant source is removed by stopping the gas flow or cleaning the surface

## 7.6 Oxide removal

Now we want to remove the silicon mask from the wafer and clean it for another clean oxide mask layer.

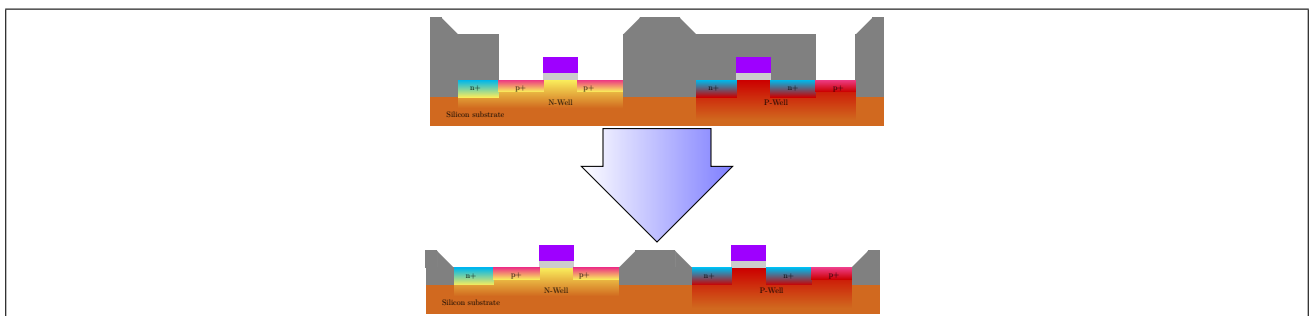
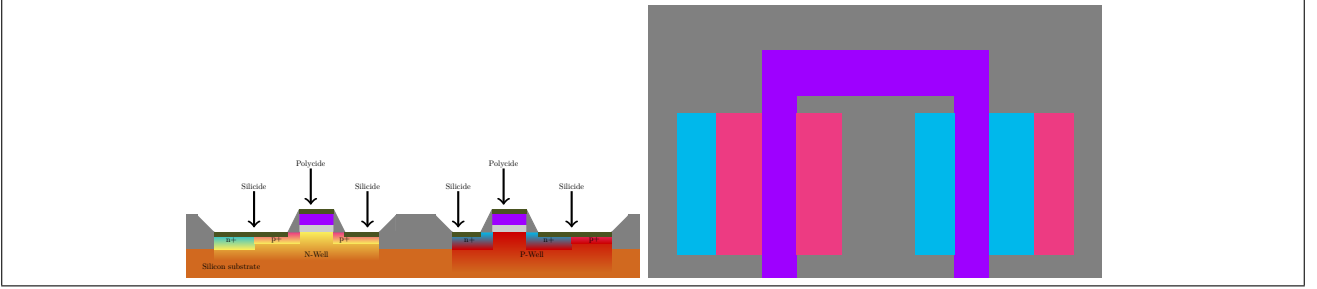


Figure 56: Oxide removal

We use buffered hydrofluoric acid (BOE (1:6)) at room temperature for 1 minute in order to remove the 100nm of oxide layer.

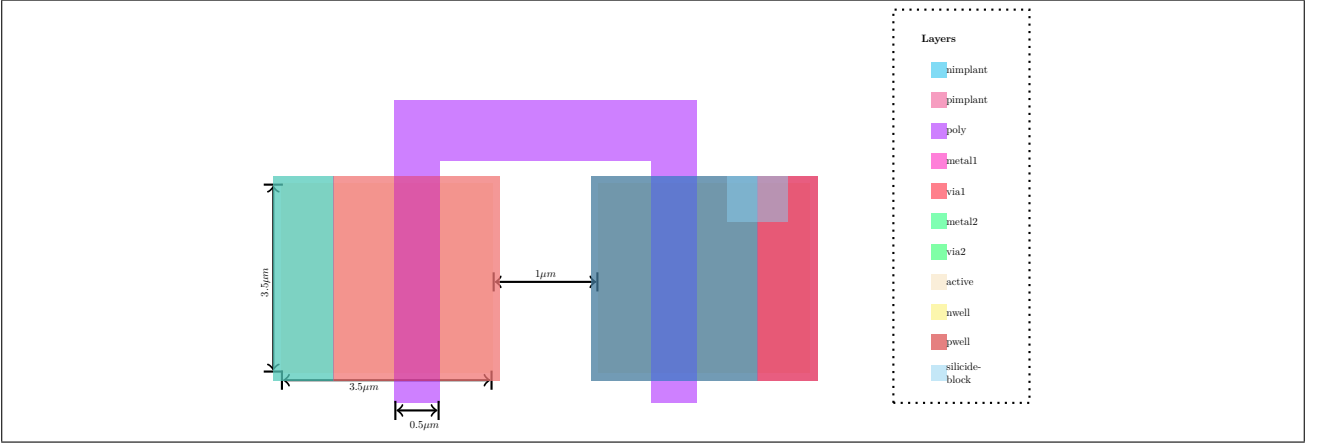
## 8 Silicification

Titanium silicide is one of the first SALICIDE material introduced in ULSI devices owing to its low resistivity, high thermal stability, ease in deposition and compatibility with silicon processes. Titanium has been one of the familiar materials in ULSI productions, which is also an important advantage in practical use of titanium SALICIDE.<sup>17</sup>



**Figure 57:** Silicide geometry target

In order to reduce the gate contact resistance as well as the source and drain resistance and in order to provide a more effective etch stop when plasma etching the contact windows to drain, source and gate, silicide/polycide is being added to the wafer as shown in Figure 57.



**Figure 58:** Silification layout

When titanium and silicon are brought into contact and heated at temperatures above 500 °C (in the presence of excess silicon) the higher-resistivity  $C49-TiSi_2$  phase forms before the low-resistivity phase.

The  $C49-TiSi_2$  phase has an orthorhombic base-centered structure with 12 atoms per unit cell and a resistivity of  $60 - 90\mu\Omega - cm$ .

The  $C54-TiSi_2$  phase has an orthorhombic face-centered structure having 24 atoms per unit cell and a significantly lower resistivity ( $12 - 20\mu\Omega - cm$ ) than the  $C49-TiSi_2$ .

The basic formation process of titanium SALICIDE is as follows:

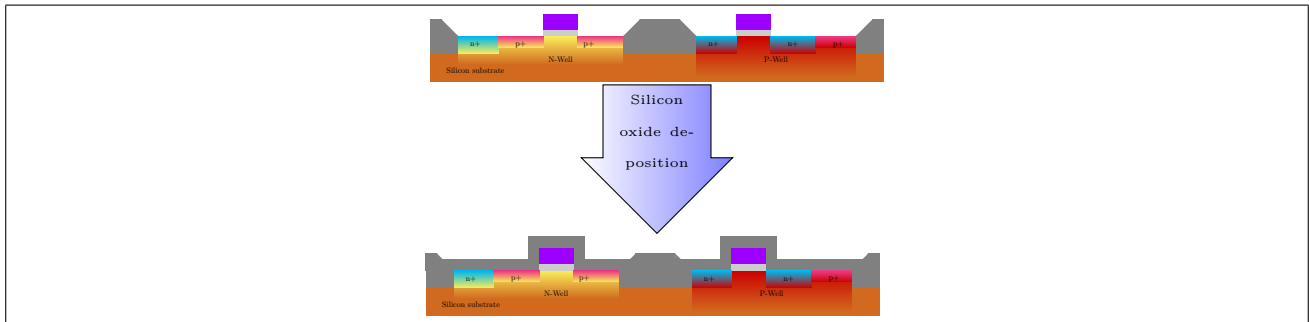
A thin titanium film with 20-60 nm thickness is deposited on an entire wafer with MOSFETs structure. The deposited Ti film reacts with the exposed silicon areas such as the source/drain area and polysilicon gate electrodes by the first anneal at 600-700°C in  $N_2$  ambient. In first anneal,  $C49-TiSi_2$  phase is formed. Then, the unreacted titanium film on the dielectric layer such as  $SiO_2$  or SiN is selectively etched by APM (Ammonia and Hydrogen Peroxide Mixture) solution. The final step is second anneal at 800°C or above to transform high-resistivity  $C49-TiSi_2$  phase to low-resistivity  $C54-TiSi_2$  phase at the gate electrodes and source/drain areas.

<sup>17</sup>A Study on Formation of High Resistivity Phases of Nickel Silicide at Small Area and its Solution for Scaled CMOS Devices, 07D53437, Ryuji Tomita

## 8.1 Oxide deposition

The thickness of this CVD deposited oxide layer will be the width of the spacer after having used highly anisotropic etching in the next few steps, for this reason the thickness of the oxide decides over the distance between the silicide and the gate oxide.

We make the oxide layer 50nm thick.



**Figure 59: Oxide layer**

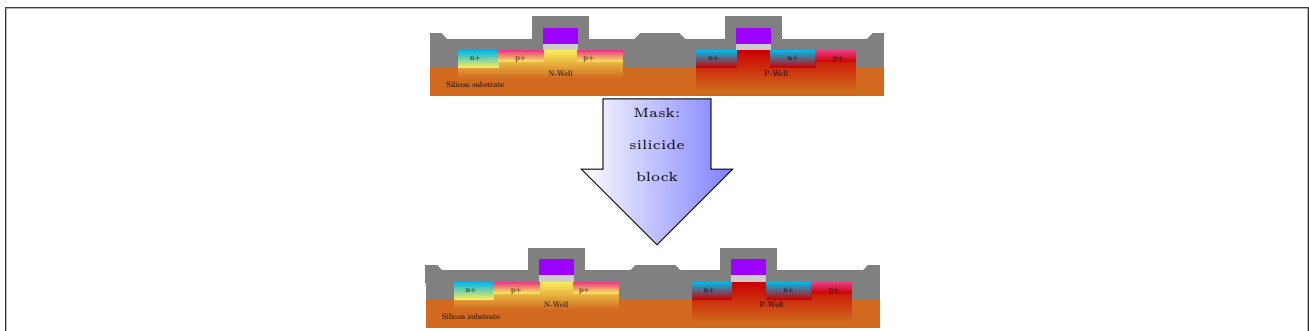
We use the machine LPCVD machine from HKUST and deposit around 50nm of silicon dioxide with the following recipe<sup>18</sup>:

- Temperature: 400 °C ( $SiH_4 + O_2 = SiO_2 + 2H_2$ )
- Pressure = 250 mTorr
- Silane ( $SiH_4$ ) flow = 40sccm
- Oxygen ( $O_2$ ) flow = 48sccm

This will give a rate of 7nm ( $\pm 1nm$ ) per minute, so we deposit for roughly seven minutes (7 min).

## 8.2 Silicide block patterning (optional)

We now have to pattern the mask for the silicide block layer which will produce oxide wherever no silicide is not desired within active areas.



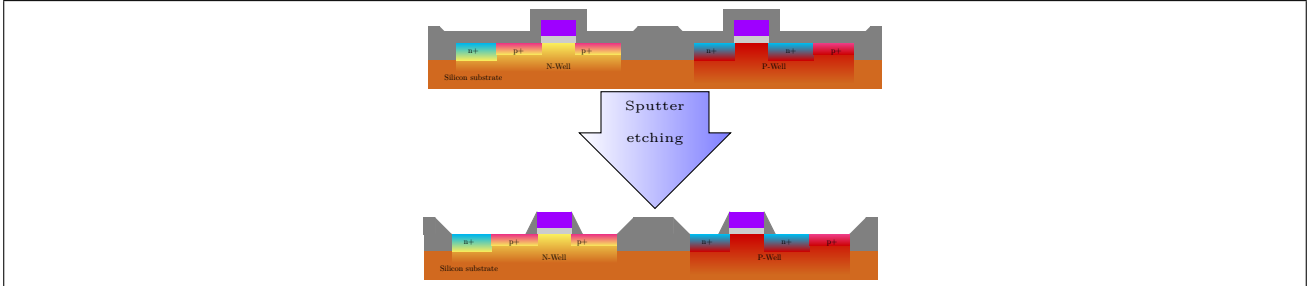
**Figure 60: Patterning (silicide block)**

It is not yet clear whether we will be needing this feature in the future, which means we have not yet finally decided whether we want to keep this step and layer within our process and technology.

<sup>18</sup>[https://people.rit.edu/lfsee/LPCVD\\_Recipes.pdf](https://people.rit.edu/lfsee/LPCVD_Recipes.pdf)

### 8.3 Sputter etching(Spacers)

Now we have to etch our oxide as anisotropic as possible. This means that the etching mostly only comes "from above with a few to nearly none horizontal etching. That means the etching process only "sees" the sidewall as a "thicker layer" and starts etching downward. With an etching speed of 35 nm/min for thermal oxide and an oxide thickness of around 50nm and given that the polysilicon is much higher than 50nm we will have our desired spacer geometry forming as well as any potentially resist covered are (given silicide block is being used) with sharp etches.

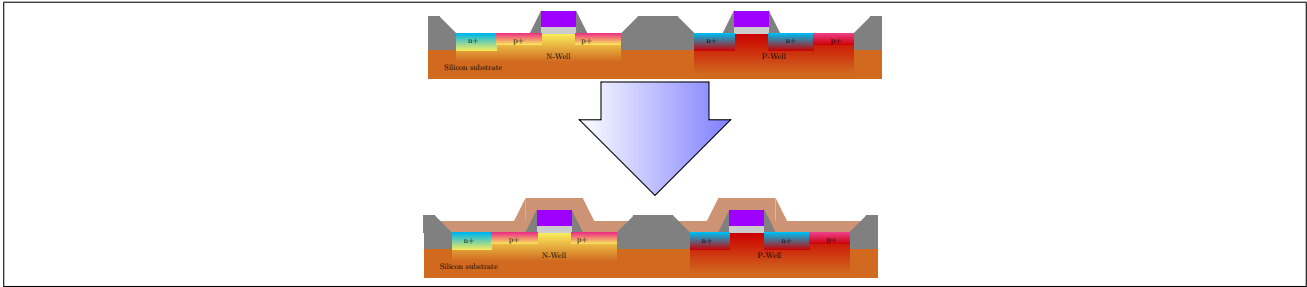


**Figure 61:** Anisotropic etching

The above mentioned machine is the "Trion RIE Etcher" at the NFF HKUST lab. The etching process runs on the oxide for 2 minutes.

### 8.4 Titanium deposition

We deposit a layer of titanium with a thickness of around 20-60nm which will then be reacted into titanium-silicide and titanium-polycide respectively in the further steps.



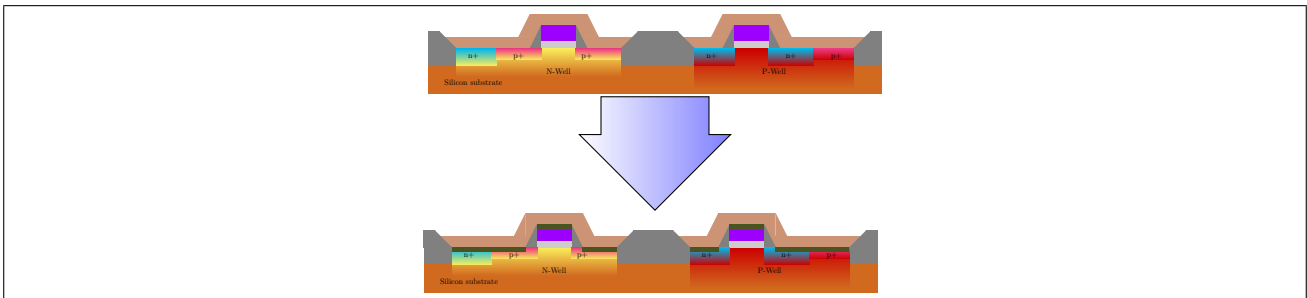
**Figure 62:** Titanium deposition

For this purpose we use the "Denton Sputter (SPT-Denton)" at HKUST NFF lab which has a sputter rate of around 8.8 nm/min for titanium.

This means we run the deposition process for around 5 minutes.

### 8.5 First reaction step

The deposited Ti film reacts with the exposed silicon areas such as the source/drain area and polysilicon gate electrodes by the first anneal at 600-700°C in  $N_2$  ambient. In this first anneal, the  $C49-TiSi_2$  phase is formed.

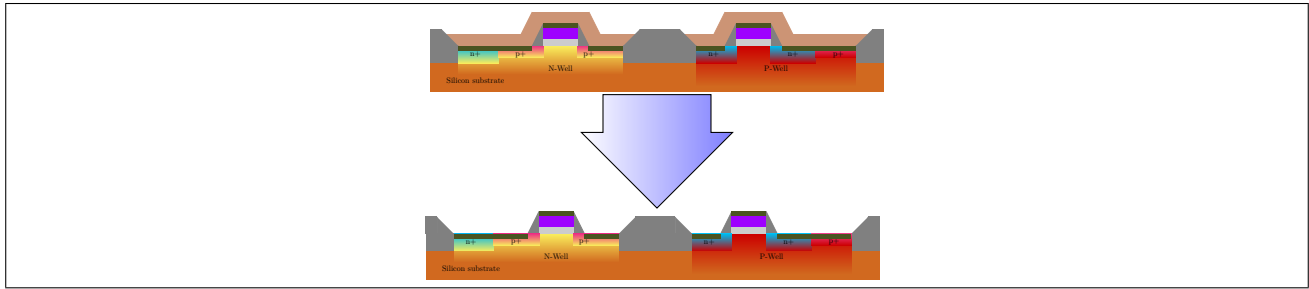


**Figure 63:** Reaction 1

We use the "AG610 RTP (DIF-R2)" from the HKUST at 700°C for 240 seconds.

## 8.6 Metal removal

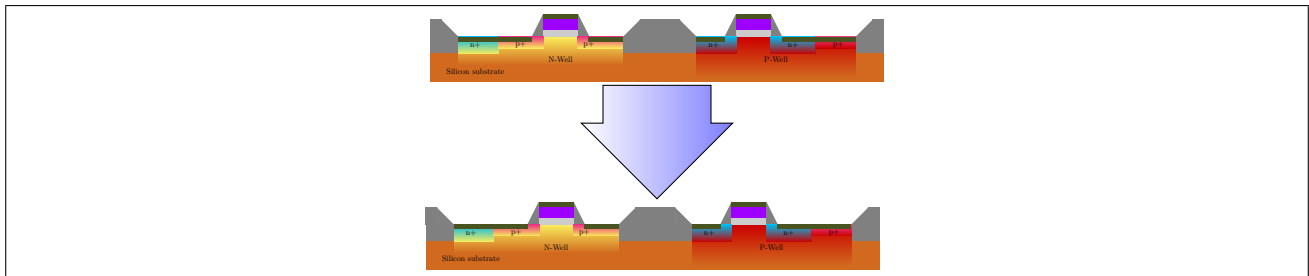
The unreacted titanium film on the dielectric layer such as  $SiO_2$  or  $SiN$  is selectively etched by APM (Ammonia and Hydrogen Peroxide Mixture) solution.



**Figure 64:** Titanium etch

## 8.7 Second reaction step

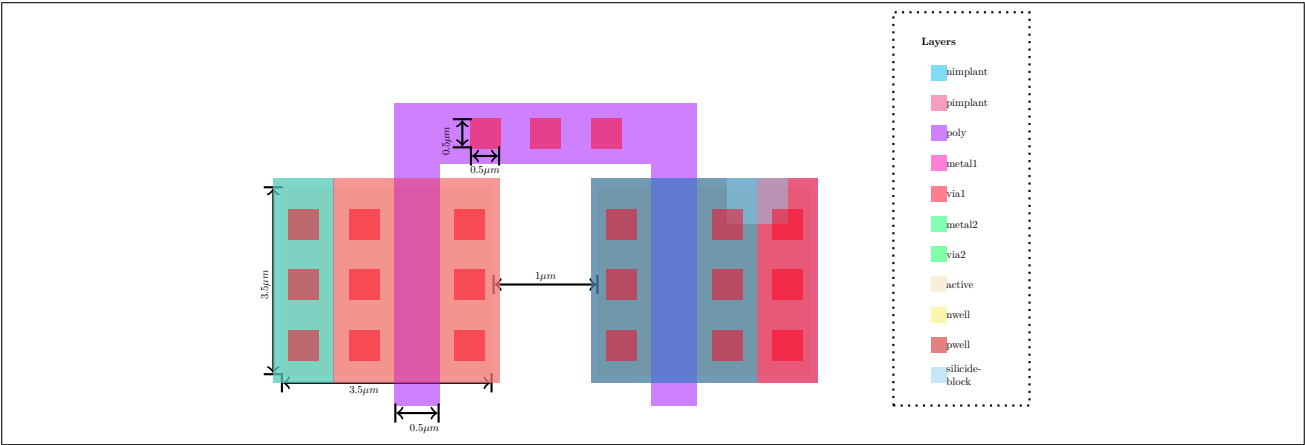
The final step is a second anneal at 800 °C or above to transform the high-resistivity  $C49-TiSi_2$  phase to the low-resistivity  $C54-TiSi_2$  phase at the gate electrodes and source/drain areas.



**Figure 65:** Reaction 2

We use the "AG610 RTP (DIF-R2)" again at 800°C for 240 seconds.

# 9 First vias



10 First metal layer

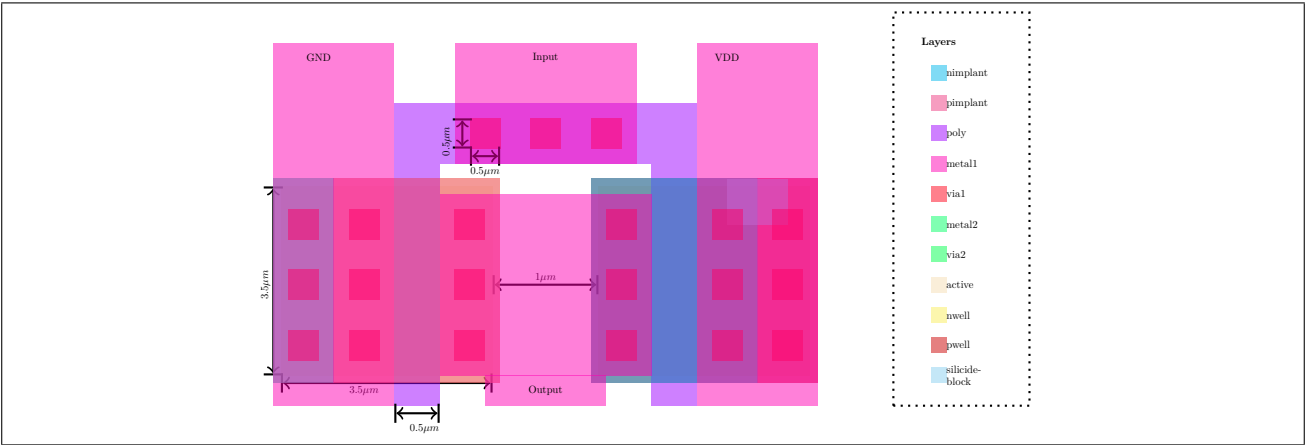


Figure 67: First metal layout

# 11 Additional vias

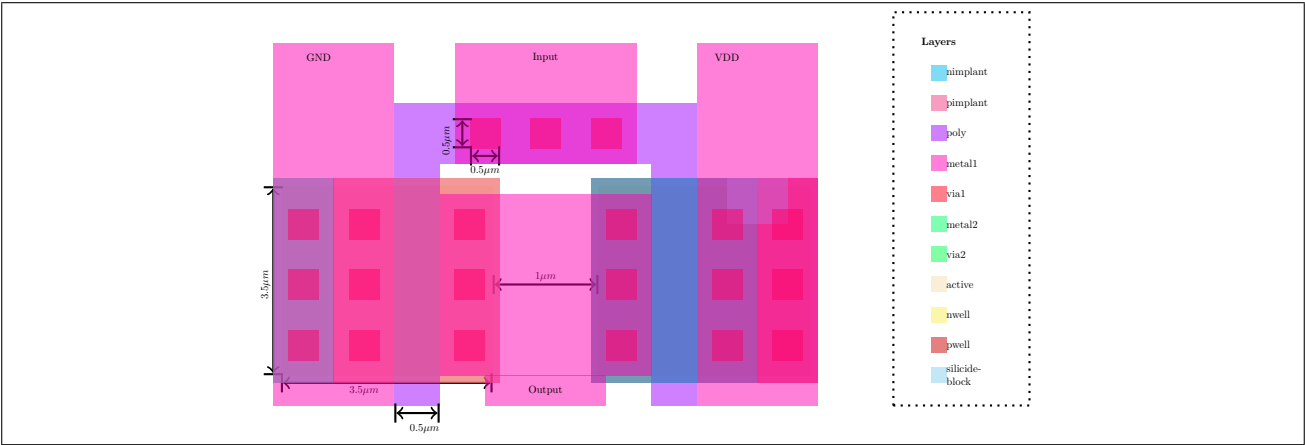


Figure 68: Additional via layout



# 12 Additional metal layer

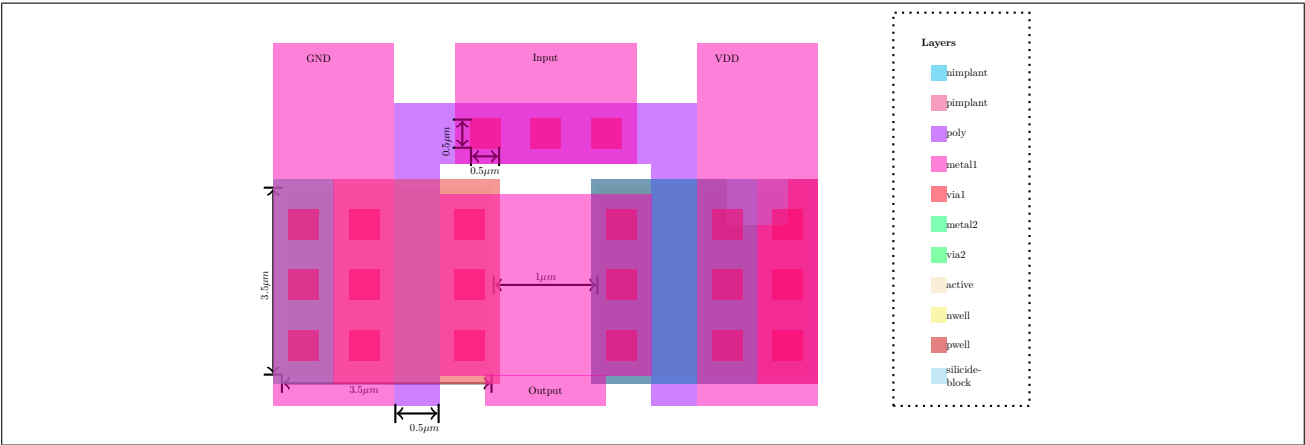


Figure 69: Additional metal layout