

Name: \_\_\_\_\_ **Conversions Station Conversion Table** 1 nanoliter (nL)= .001 µL 1 milliliter (mL)= **1000 µL** 1 liter (L) = **1000000 \muL** → 2 mL= 2000 µL → .000000001 mL= µL → .75 mL= \_\_\_\_ µL → .095 mL= µL → .0025 mL= µL → 24 nL= \_\_\_\_µL → .009 nL= µL → .0001 L= \_\_\_\_µL → .0253 L= µL → 28 µL= L → 856 µL=\_\_\_\_L → 20 µL= nL → 0.20 µL=\_\_\_\_\_nL → 634 µL= \_\_\_\_ mL

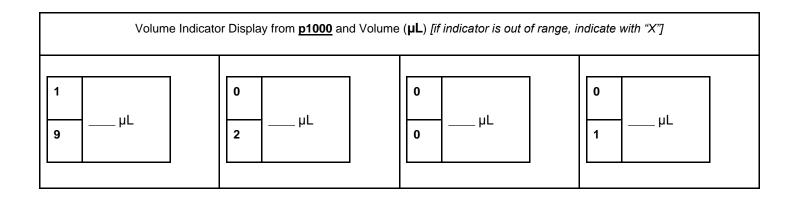
Name: \_\_\_\_\_ **Conversions Station Conversion Table** 1 nanoliter (nL)= .001 µL 1 milliliter (mL)= **1000 μL** 1 liter (L) = **1000000 µL** → 2 mL= 2000 µL → .000000001 mL= µL → .75 mL= \_\_\_\_ µL → .095 mL= µL → .0025 mL= µL → 24 nL= µL → .009 nL= µL → .0001 L= \_\_\_\_µL → .0253 L= µL → 28 µL= \_\_\_\_ L → 856 µL= \_\_\_\_ L → 20 µL= \_\_\_\_\_ nL → 0.20 µL= \_\_\_\_\_ nL → 634 µL= \_\_\_\_ mL

	→ 29000 μL= mL	→ 29000 µL= mL	→ 29000 μL= mL
--	----------------	----------------	----------------

ame:	NEVER c	rank the micropipets	above or below their ran	licropipette Ider	itilication Static
	Range: <b>0.5-10 μl</b>	Range: 2-20 μl	Range: 20-200 μΙ	Range: 200-1000 μl	
		20	200	P.000	
Volume	Required Micropipette	Volume	Required Micropipette	Volume	Required Micropipette
28 µL	p200	856 μL		9.9 µL	
199 µL		10.01 μL		6 µL	
.5 mL		.05 mL		.0089 mL	
		.00001 L		.000006 L	htification Static
.00000089 L			Above or below their ran	/licropipette Ider	ntification Static
				/licropipette Ider	ntification Static
	Range: 0.5-10 µl	rank the micropipets Range: 2-20 μl	above or below their ran Range: 20-200 μl	Aicropipette Ider ges!! 200-1000 μΙ	ntification Statio
ame:	Range: 0.5-10 µl Ploop Required	rank the micropipets Range: 2-20 μl	above or below their ran Range: 20-200 μl P 200 P 200 P 200 Required	Aicropipette Ider ges!! 200-1000 μI	Required
ame: Volume	Range: 0.5-10 µl	rank the micropipets Range: 2-20 μl Polume	above or below their ran Range: 20-200 μl P 200 P 200 P 200 Required	Aicropipette Ider ges!! Range: 200-1000 μΙ Uolume	Required
ame: Volume 28 μL	Range: 0.5-10 µl	rank the micropipets Range: 2-20 μl Po Po Volume 856 μL	above or below their ran Range: 20-200 μl P 200 P 200 P 200 Required	Aicropipette Ider ges!! 200-1000 μΙ	Required

Volume Indicator Display from <b>p20</b> and Volume ( <b>µL</b> ) <i>[if indicator is out of range, indicate with "X"]</i>			
1	0	0	0
9	2	0	2
5	0	1	1
3	0	2	2
5	1	0	0
9	0	0	5

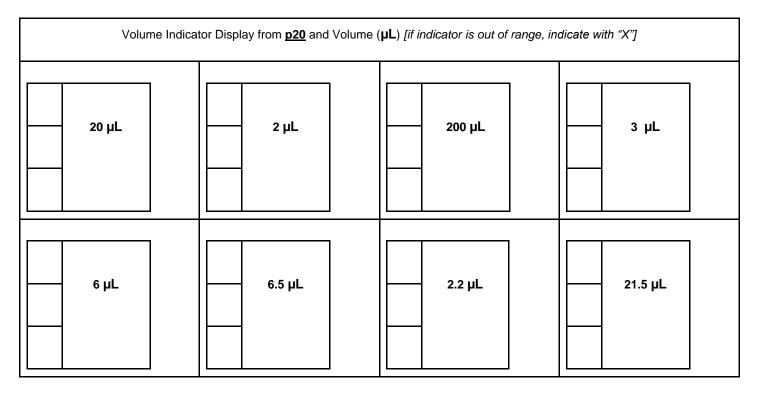
Volume Indicator Display from <u>p200</u> and Volume (µL) [if indicator is out of range, indicate with "X"]			
1	0	0	9
9	2	0	0
5	0	1	0
3	0	3	2
5	1	0	0
9	0	0	5



5	0	9	9
3	0	0	2
5	1	0	0
9	0	1	5

Name:\_\_\_\_\_

## **Volume Indicator B Station**



Volume Indicator Display from <u>p200</u> and Volume (µL) <i>[if indicator is out of range, indicate with "X"]</i>			
200 μL	20 μL	18 μL	

- 199 μL	89.5 µL	199.8 µL	21.5 µL

Volume Indicator Display from <b>p1000</b> and Volume ( <b>µL</b> ) <i>[if indicator is out of range, indicate with "X"]</i>			
1000 µL	200 µL		42 μL
420 μL	999 µL	140 μL	300 µL

Name: \_\_\_\_\_

<ul> <li>Your protocol calls for 1000 uL of distilled water.</li> <li>1) How many uL of diluted water do you need?µL</li> <li>2) Which micropipette will you need to use? p</li> <li>3) What should your volume indicator display look like?</li> </ul>	<ul> <li>Your protocol also calls for .097 mL of dNTPs.</li> <li>1) How many uL of dNTPs do you need?µL</li> <li>2) Which micropipette will you need to use? p</li> <li>3) What should your volume indicator display look like?</li> </ul>
Your protocol also requires .000005 L of taq polymerase.          1) How many uL of taq polymerase do you need?µL         2) Which micropipette will you need to use? p         3) What should your volume indicator display look like?	<ul> <li><u>Review Questions:</u> <ol> <li>Should you keep the same tip between reagents, yes or no? Why?</li> </ol> </li> <li><u>When micropipetting your solution "up" (retrieving), you should</u> <ol> <li>First, depress the plunger (<u>before/after</u>) the tip is in the liquid</li> <li>Next, depress the plunger to the (<u>first/second</u>) stop.</li> </ol> </li> <li>Lastly, Release the plunger (<u>slowly/quickly</u>) while the tip is still in liquid. <ol> <li><u>When transferring solution to the new tube, you should</u></li> <li>First, have the tip (<u>hang freely/touch the bottom</u>) in the tube.</li> <li>Next, press the plunger to the (<u>first/second</u>) stop.</li> </ol> </li> </ul>

Micropipette Prep, Summary Station

Your protocol calls for <b>1000 uL</b> of distilled water.	Your protocol also calls for <b>.097 mL</b> of dNTPs.
1) How many uL of diluted water do you need?µL	1) How many uL of dNTPs do you need?µL
2) Which micropipette will you need to use? p	2) Which micropipette will you need to use? <b>p</b>
3) What should your volume indicator display look like?	3) What should your volume indicator display look like?
Your protocol also requires .000005 L of taq polymerase.          1) How many uL of taq polymerase do you need?µL         2) Which micropipette will you need to use? p         3) What should your volume indicator display look like?	<ul> <li><u>Review Questions:</u> <ol> <li>Should you keep the same tip between reagents, yes or no? Why?</li> </ol> </li> <li><u>When micropipetting your solution "up" (retrieving), you should</u> <ol> <li>First, depress the plunger (<u>before/after</u>) the tip is in the liquid</li> <li>Next, depress the plunger to the (<u>first/second</u>) stop.</li> </ol> </li> <li><u>When transferring solution to the new tube, you should</u> <ol> <li>First, have the tip (<u>hang freely/touch the bottom</u>) in the tube.</li> <li>Next, press the plunger to the (<u>first/second</u>) stop.</li> </ol> </li> </ul>