

$$0.02500 \text{ L} = 0.0972 \text{ M}$$

Discussion

This section is critical for demonstrating your understanding.

Discuss:

- Sources of error and how they affected your results
- Why the chosen indicator was appropriate
- Comparison of your results with expected values
- Precautions taken to minimize errors
- Suggestions for improving the experiment

Conclusion

State whether the objective was achieved and provide the final answer with appropriate units and significant figures.

COMMON ERRORS IN ACID BASE TITRATION AND HOW TO AVOID THEM

Even experienced students make mistakes in titration experiments. Being aware of these common errors will help you produce a more accurate **acid base titration lab report matriculation** submission.

Parallax Error

Reading the burette volume at an angle rather than at eye level leads to inaccurate readings. Always read the meniscus at eye level and record the bottom of the meniscus for clear liquids.

Air Bubbles in the Burette

Air trapped in the burette tip can cause volume readings to be inaccurate. Always remove air bubbles before taking the initial reading by running a

small amount of titrant through the tip.

Over-Titration

Adding too much titrant past the endpoint is a common mistake. Slow down your addition as you approach the expected endpoint, and use a wash bottle to rinse down any droplets on the inner walls of the flask.

Improper Swirling

Insufficient mixing of the solution during titration can lead to localized color changes and premature endpoint detection. Swirl continuously and thoroughly throughout

the titration.

these strategies:

Using the Wrong Indicator

Different acid-base combinations require different indicators based on the pH at the equivalence point. For strong acid-strong base, phenolphthalein or methyl orange works. For weak acid-strong base, phenolphthalein is appropriate. For strong acid-weak base, methyl orange is preferred.

TIPS FOR MAXIMIZING YOUR SCORE ON THE LAB REPORT

To excel in your **acid base titration lab report matriculation** assessment, consider

1. **Record data immediately** — never rely on memory; write readings directly into your data table as you take them.
2. **Use appropriate significant figures** — burette readings are typically recorded to two decimal places (e.g., 24.30 mL, not 24.3 mL).
3. **Show all working** — even simple calculations should be written out completely to demonstrate your understanding.
4. **Discuss errors honestly** — acknowledging limitations of

your experiment shows scientific maturity and critical thinking.

5. **Include relevant chemical equations** — balanced equations with state symbols demonstrate thoroughness.
6. **Format your report neatly** — a well-organized report with clear headings and properly labeled tables creates a positive impression.

CHOOSING THE RIGHT INDICATOR

The selection of indicator is a key decision in any acid base titration and should be justified in your **acid base titration lab report**

matriculation. The pH range of the indicator must match the pH at the equivalence point of the titration.

Phenolphthalein

Color change from colorless to pink over pH 8.2–10.0. Suitable for strong acid-strong base and weak acid-strong base titrations. This is the most common indicator used in matriculation titration experiments.

Methyl Orange

Color change from red to yellow over pH 3.1–4.4. Suitable for strong acid-strong base and strong acid-weak base titrations. Not

suitable for weak acid-strong base titrations.

Bromothymol Blue

Color change from yellow to blue over pH 6.0–7.6. Suitable for strong acid-strong base titrations where the equivalence point is near pH 7.

DATA ANALYSIS AND ERROR CALCULATION

A thorough **acid base titration lab report matriculation**

includes error analysis. Calculate the percentage error using

the formula:

$$\text{Percentage Error} = \frac{|(\text{Experimental Value} - \text{Theoretical Value})|}{\text{Theoretical Value}} \times 100\%$$

If no theoretical value is provided, you can still discuss precision by calculating the range or standard deviation of your titre volumes. Concordant results typically fall within 0.1 mL of each other, indicating good precision.

Common sources of systematic error include:

- Impure reagents
- Incorrectly calibrated volumetric glass