

Math Diversion Problem 752

P. Reany

August 8, 2025

Math is just hard.
— Alex Wei
(an OpenAI IMO
trainer)

The material here is found at:

Source: The Ether of Great Mathematical Ideas
Title: A lemma on the Carnot cycle of an ideal gas
Presenter: Copilot

1 The Setup

After I had Copilot double check my computations in establishing that for an ideal monoatomic gas in the Carnot cycle, analyzed in the PV plane, that

$$\frac{V_2 V_4}{V_1 V_3} = 1, \quad (1)$$

Copilot suggested that I do the same proof in the TS plane for a comparison. I agreed and went to my resources for a reference on it, but I found nothing. So, I asked Copilot to help me do it. What follows is mostly the work of Copilot, though I edited it a bit.

2 The Problem

Let's reconstruct the volume identity

$$\frac{V_2 V_4}{V_1 V_3} = 1 \quad (2)$$

within the Carnot cycle using thermodynamic principles, focusing on reversible processes and the ideal gas law in both PV and TS planes.

Starting Point: Carnot Cycle Process Overview In the PV diagram, the cycle includes:

- 1 → 2: Isothermal expansion at T_H
- 2 → 3: Adiabatic expansion
- 3 → 4: Isothermal compression at T_C
- 4 → 1: Adiabatic compression

Each leg affects volume and entropy differently depending on temperature and reversibility.

3 The Solution in the PV plane

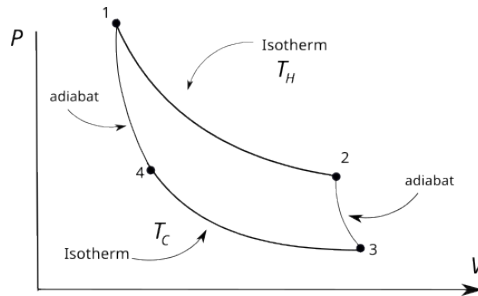


Figure 1. For the path 1 to 2, the ideal gas undergoes reversible isothermal expansion. For 2 to 3, the gas undergoes adiabatic expansion. From 3 to 4, the gas undergoes reversible isothermal compression. And from 4 to 1, the gas undergoes an adiabatic compression.

The ideal gas law

$$PV = nRT \quad (3)$$

applied to an isotherm implies that $PV = \text{constant}$.

During adiabatic processes, $TV^{\gamma-1} = \text{constant}$, for a monoatomic gas where

$$\gamma = \frac{C_P}{C_V} \quad (4)$$

Now let's tie the volume ratios to the adiabatic processes:

From 2 → 3: Adiabatic expansion from high temperature T_H to T_C

$$T_H V_2^{\gamma-1} = T_C V_3^{\gamma-1} \quad (5)$$

⇒

$$\left(\frac{V_2}{V_3}\right)^{\gamma-1} = \frac{T_C}{T_H} \quad (6)$$

From 4 → 1: Adiabatic compression from T_C to T_H

$$T_C V_4^{\gamma-1} = T_H V_1^{\gamma-1} \implies \left(\frac{V_4}{V_1}\right)^{\gamma-1} = \frac{T_H}{T_C} \quad (7)$$

Multiplying the two ratios:

$$\left(\frac{V_2}{V_3}\right)^{\gamma-1} \cdot \left(\frac{V_4}{V_1}\right)^{\gamma-1} = \frac{T_C}{T_H} \cdot \frac{T_H}{T_C} = 1 \quad (8)$$

Which implies:

$$\left(\frac{V_2 V_1}{V_4 V_3}\right)^{\gamma-1} = 1 \longrightarrow \frac{V_2 V_1}{V_4 V_3} = 1 \quad (9)$$

4 The Solution in the TS plane

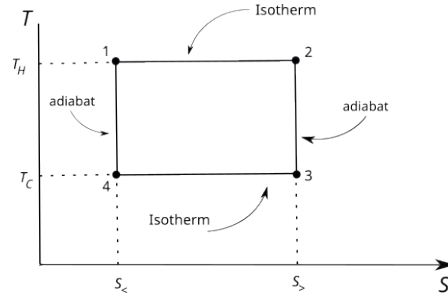


Figure 1. For the path 1 to 2, the ideal gas undergoes reversible isothermal expansion. For 2 to 3, the gas undergoes adiabatic expansion. From 3 to 4, the gas undergoes reversible isothermal compression. And from 4 to 1, the gas undergoes an adiabatic compression.

Although TS diagrams aren't volume-explicit, entropy S changes during isothermal processes are:

For 1 \rightarrow 2:

$$\Delta S = nR \ln \left(\frac{V_2}{V_1} \right) \quad (10)$$

For 3 \rightarrow 4:

$$\Delta S = nR \ln \left(\frac{V_4}{V_3} \right) \quad (11)$$

Equal and opposite entropy exchanges across the isothermal legs confirm that:

$$\ln \left(\frac{V_2}{V_1} \right) + \ln \left(\frac{V_4}{V_3} \right) = 0 \longrightarrow \frac{V_2}{V_1} \cdot \frac{V_4}{V_3} = 1 \longrightarrow \frac{V_2 V_4}{V_1 V_3} = 1. \quad (12)$$