

# Physical properties of Seawater

Lecture 3



# What is salinity and how is it measured?

- Grams of dissolved material in 1 kg of sea water. Dimensionless. But, gases and chlorides lost in drying.
- Most accurately measured using laboratory titration of chloride,  $S=1.8Cl$ . Ratio of ions in seawater (largely) constant: 50% chloride, 31% sodium, 8% sulphate...
- Most conveniently measured in situ using conductivity. Error (0.003) caused by variations in non-conducting salts like  $SiO_2$ . Units "psu". TEOS10 Absolute Salinity uses geographic information to correct to 0.0003.



# What is temperature and how is it measured?

- Temperature relates the activity level of molecules. More active = higher  $T$  ( $^{\circ}\text{C}$ )
- Measured in situ using a thermistor. Loosely wound, strain-free, pure platinum wire with resistance a function of temperature.



# What is pressure and how is it measured?

- Depth in ocean measured in situ as pressure (decibar), typically with a quartz transducer
- $p = \int \rho g dz \cong \rho g z$ , assuming incompressibility
- $p(1 \text{ m}) = 1027 \text{ kg m}^{-3} \times 9.81 \text{ ms}^{-2} \cong 1 \text{ dbar}$
- $1 \text{ Atm} = 10 \text{ dbar}$



# What is density and how is it measured?

- Density ( $\text{kg m}^{-3}$ ) is derived using the equation of state of seawater, an empirical high-order polynomial. 48 terms. TEOS-10.org. (McDougall, Millero et al., 2011).
- Density,  $\rho$ , is a function of T, S, p. Contours called isopycnals.
- For accuracy, oceanographers use  $\sigma = \rho - 1000$



# Means and ranges in the ocean

- Mean

- $S=34.7$ ;
- $T=3.5\text{ }^{\circ}\text{C}$ ;
- $p=4000\text{ dbar}$ ;
- $\rho=1027\text{ kg m}^{-3}$

- Range

- $5 < S < 40$ ;
- $-2\text{ }^{\circ}\text{C} < T < 30\text{ }^{\circ}\text{C}$ ;
- $0\text{ dbar} < p < 11,000\text{ dbar}$
- $1020\text{ kg m}^{-3} < \rho < 1070\text{ kg m}^{-3}$



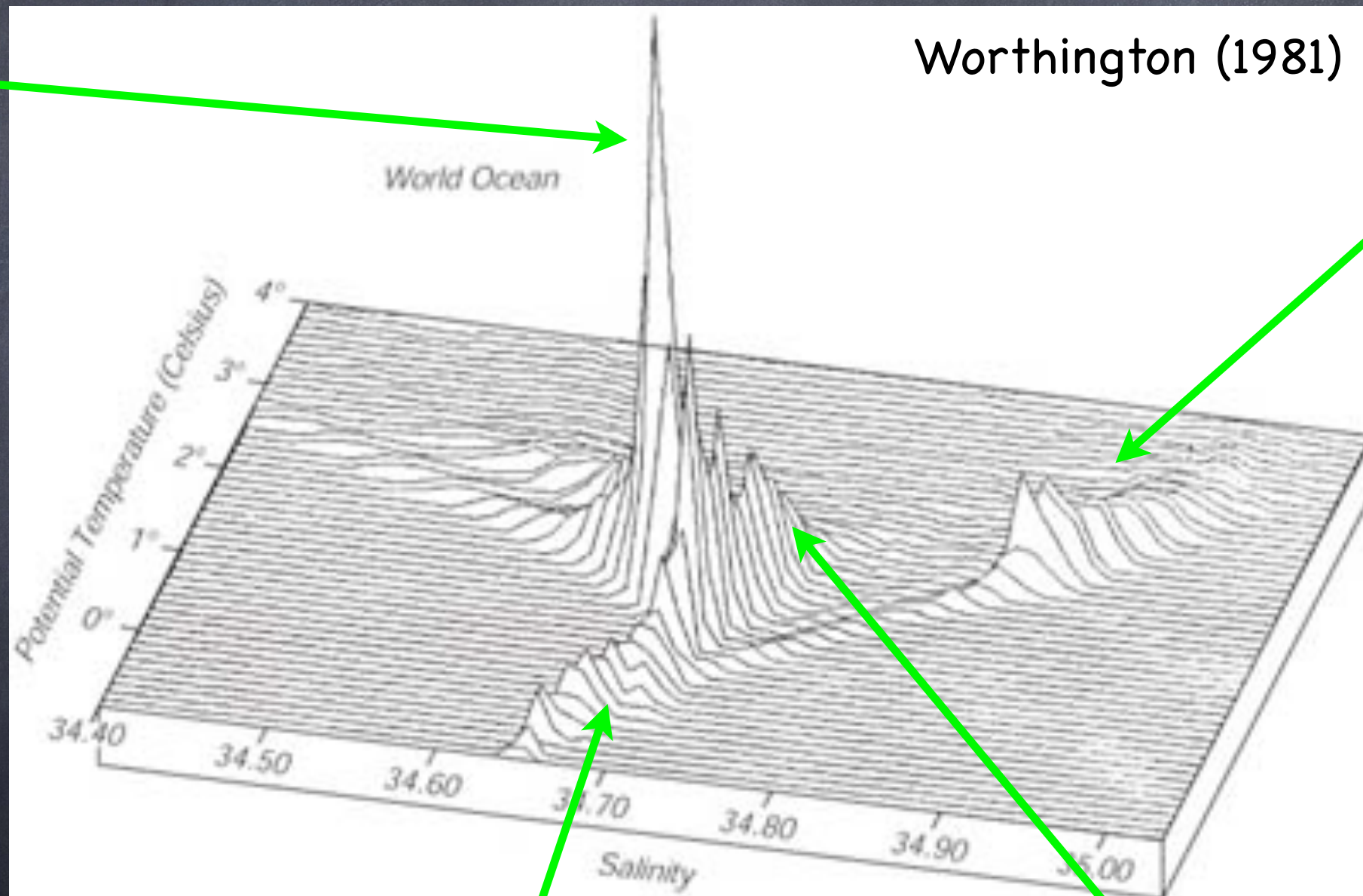
50% water in ocean is deep water!

$1.3^{\circ}\text{C} < T < 3.8^{\circ}\text{C}; 34.6 < S < 34.8$

Pacific

Worthington (1981)

Atlantic

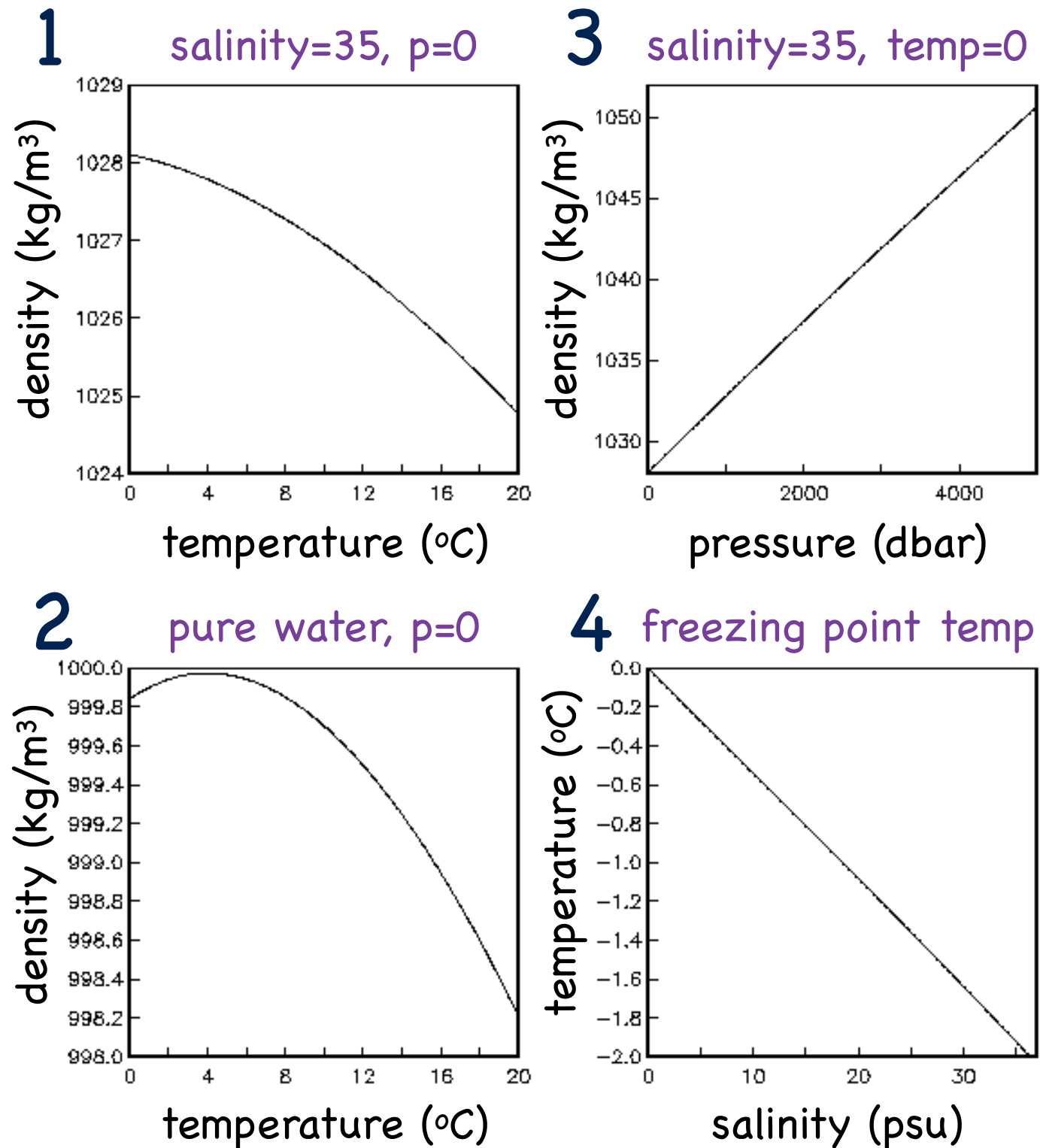


Antarctic

Indian



# Density, T, and S



1. seawater densest at freezing point
2. fresh water densest at 4°C
3. under a lot of pressure seawater compresses and gets denser.
4. freezing point of seawater is -2°C. when ice crystals form salt is rejected. Ice crystals rise to the top leaving behind super-cooled briny water



# Why are temperature and salinity key to understanding the ocean?

- Changes in T and S lead to changes in DENSITY and density gradients drive ocean currents, overturns, and mixing
- Once away from the ocean surface, water masses can be traced around the globe using their distinctive T/S relationship, revealing the global thermohaline circulation.
- Water masses like to preserve their density, moving along surfaces of constant density, or isopycnals, in the ocean interior.

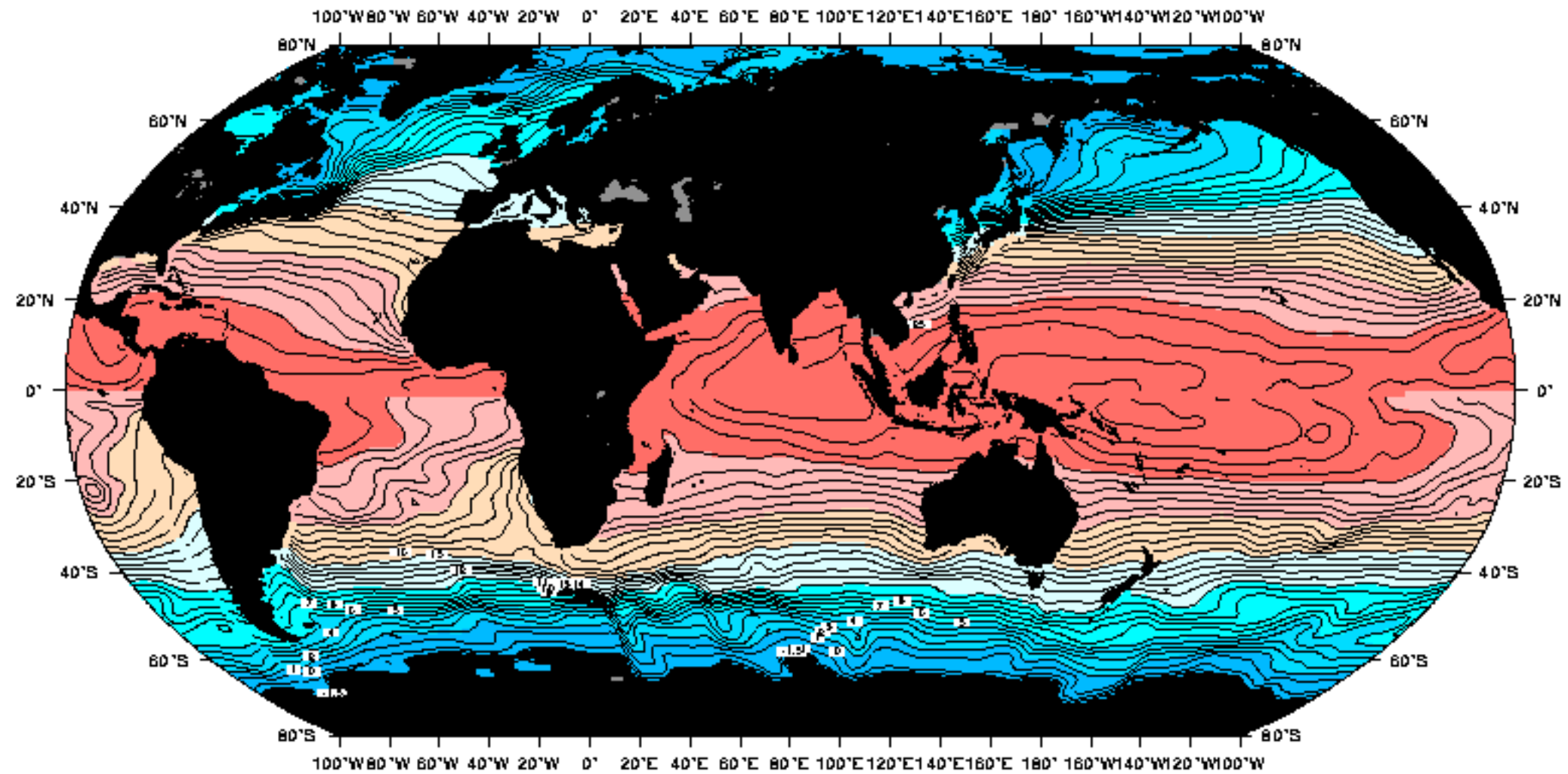


# What can affect the T & S of sea water at the surface of the ocean?

- surface heat fluxes: Incoming solar radiation, outgoing longwave radiation, sensible and latent heat.
- evaporation and precipitation and runoff
- freezing/melting of sea ice
- buoyancy-driven convection
- wind-driven mixing

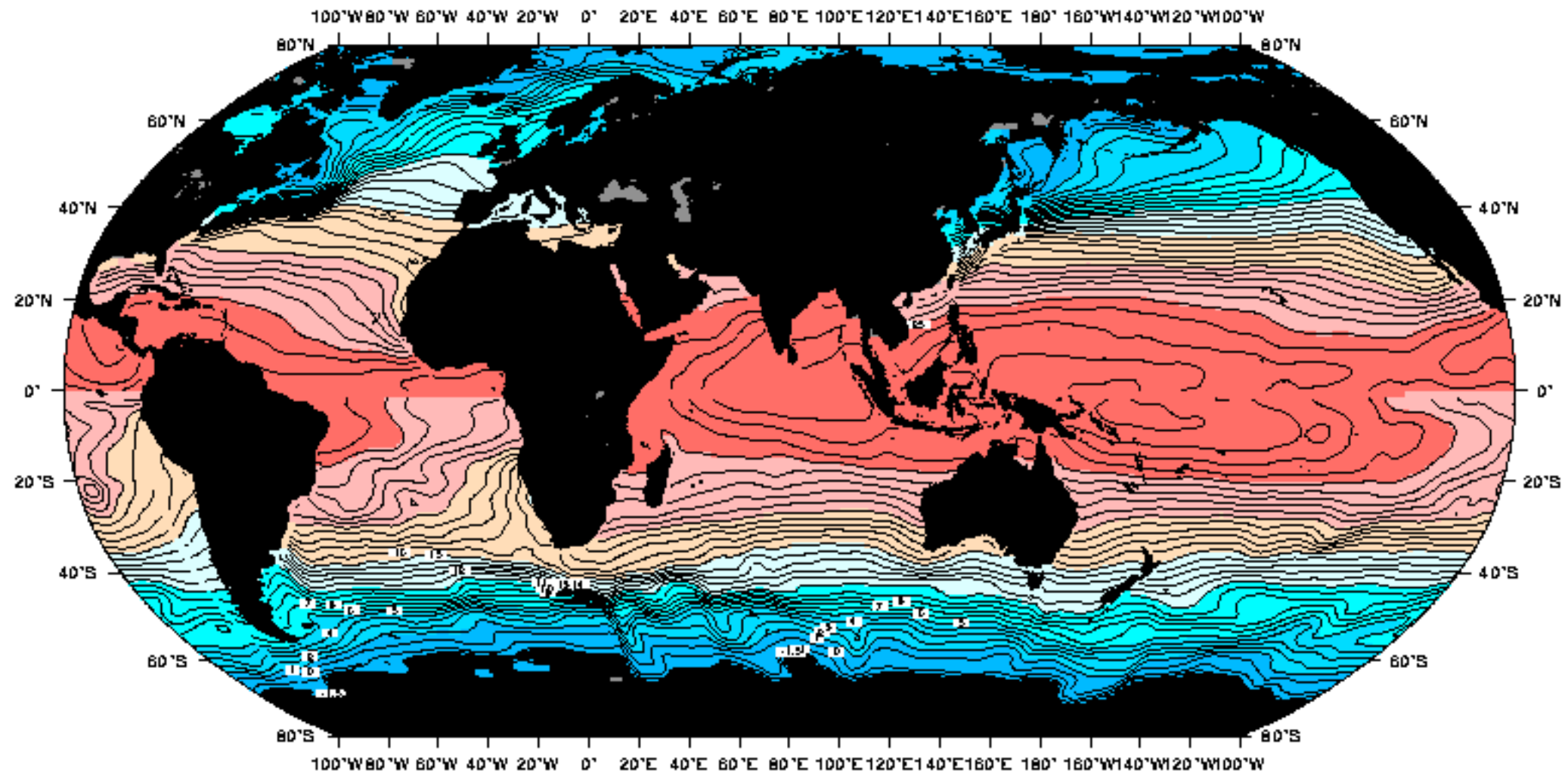


Temperature (JFM - north; JAS - south) 10m (Levitus)



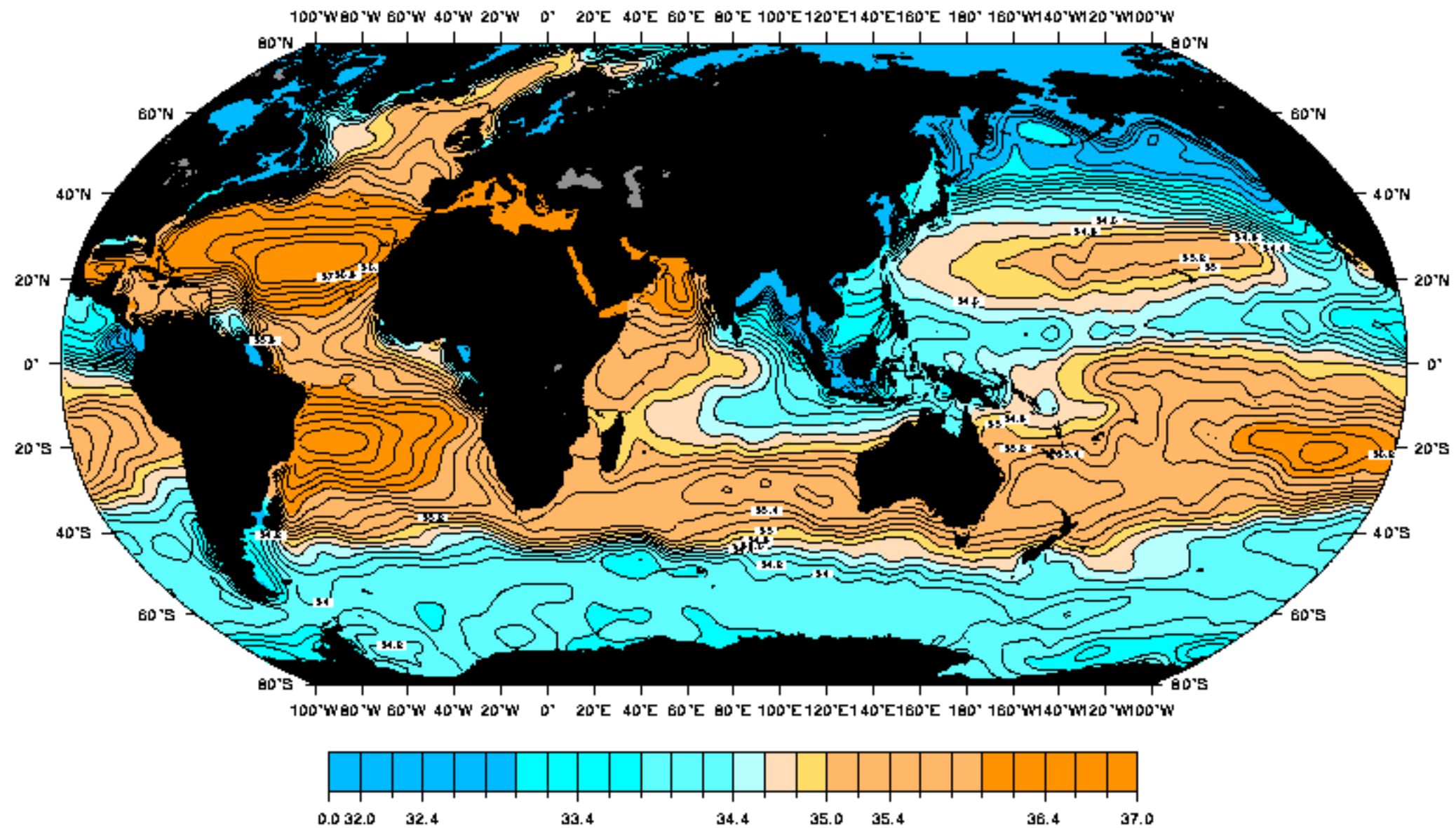


Temperature (JFM - north; JAS - south) 10m (Levitus)

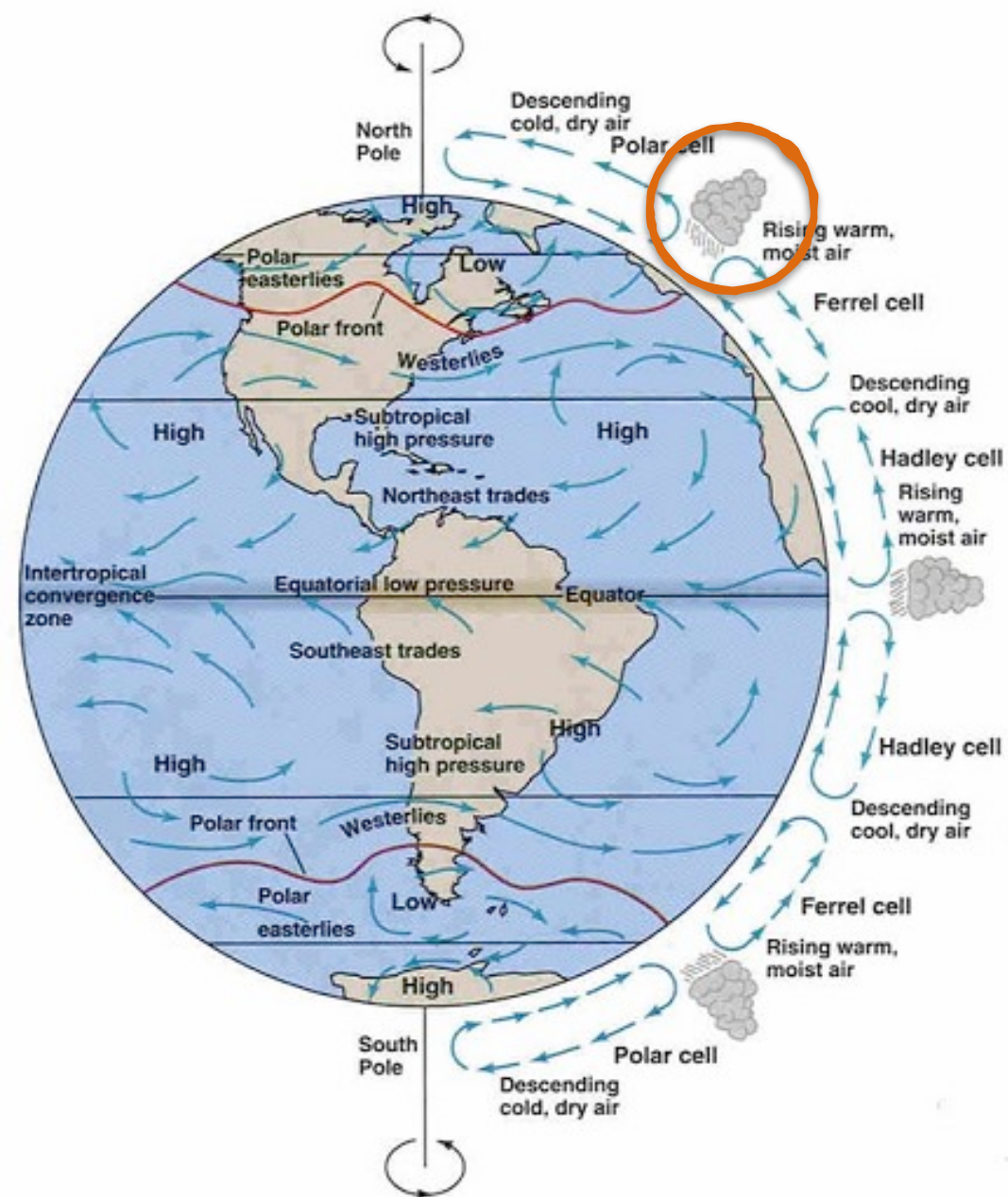


Surface temperature is dominated by net heating in the tropics and cooling at higher latitudes. The total range of temperature is from the seawater freezing point ( $-2^{\circ}\text{C}$ ) up to about  $30^{\circ}\text{C}$ .

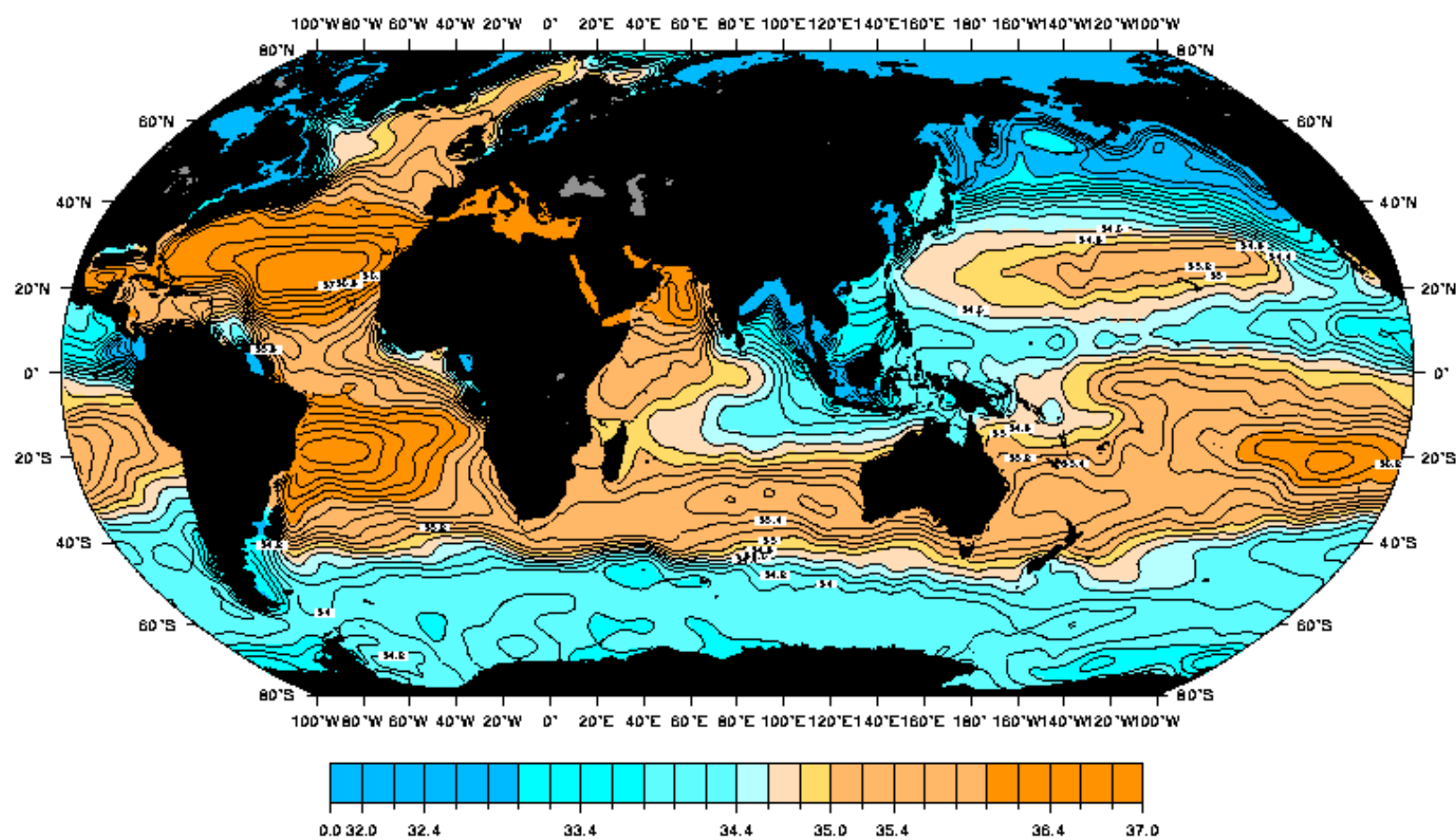


**Salinity at the sea surface (annual mean) (Levitus)**



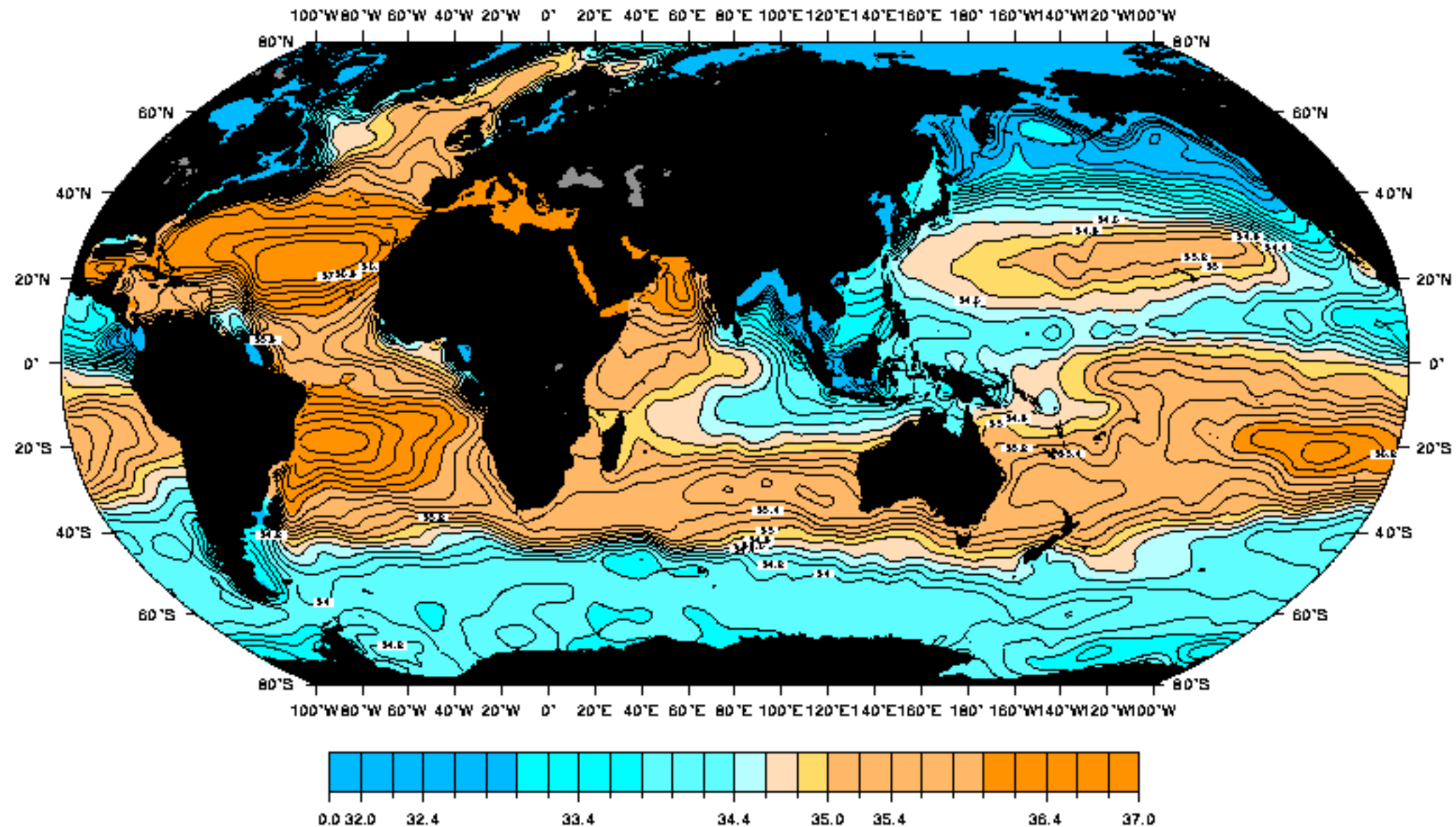


Salinity at the sea surface (annual mean) (Levitus)





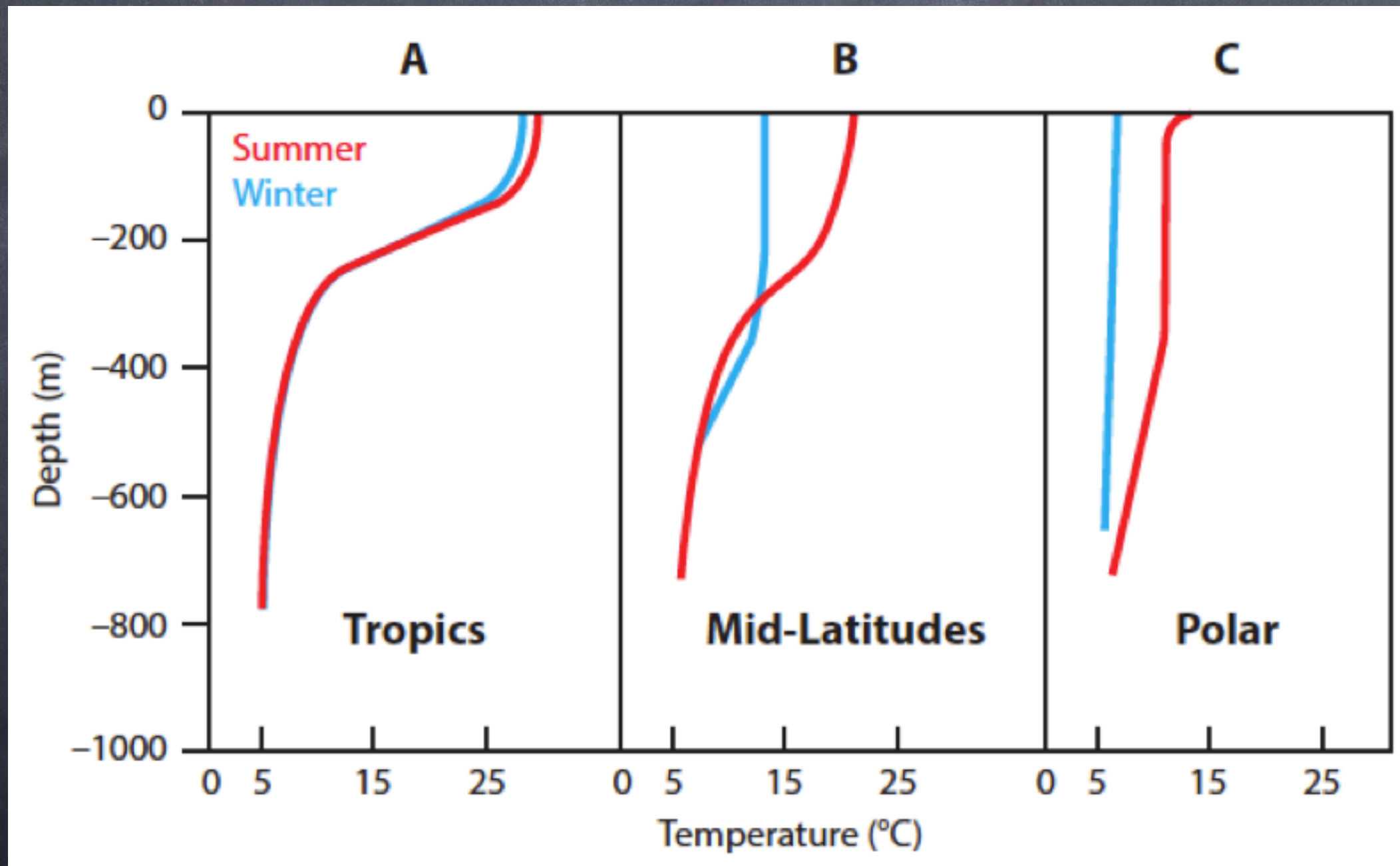
Salinity at the sea surface (annual mean) (Levitus)



Surface salinity is dominated by net evaporation in the subtropical regions, and net precipitation/runoff at higher latitudes and in the tropics. Range in open ocean is 31 to 38.

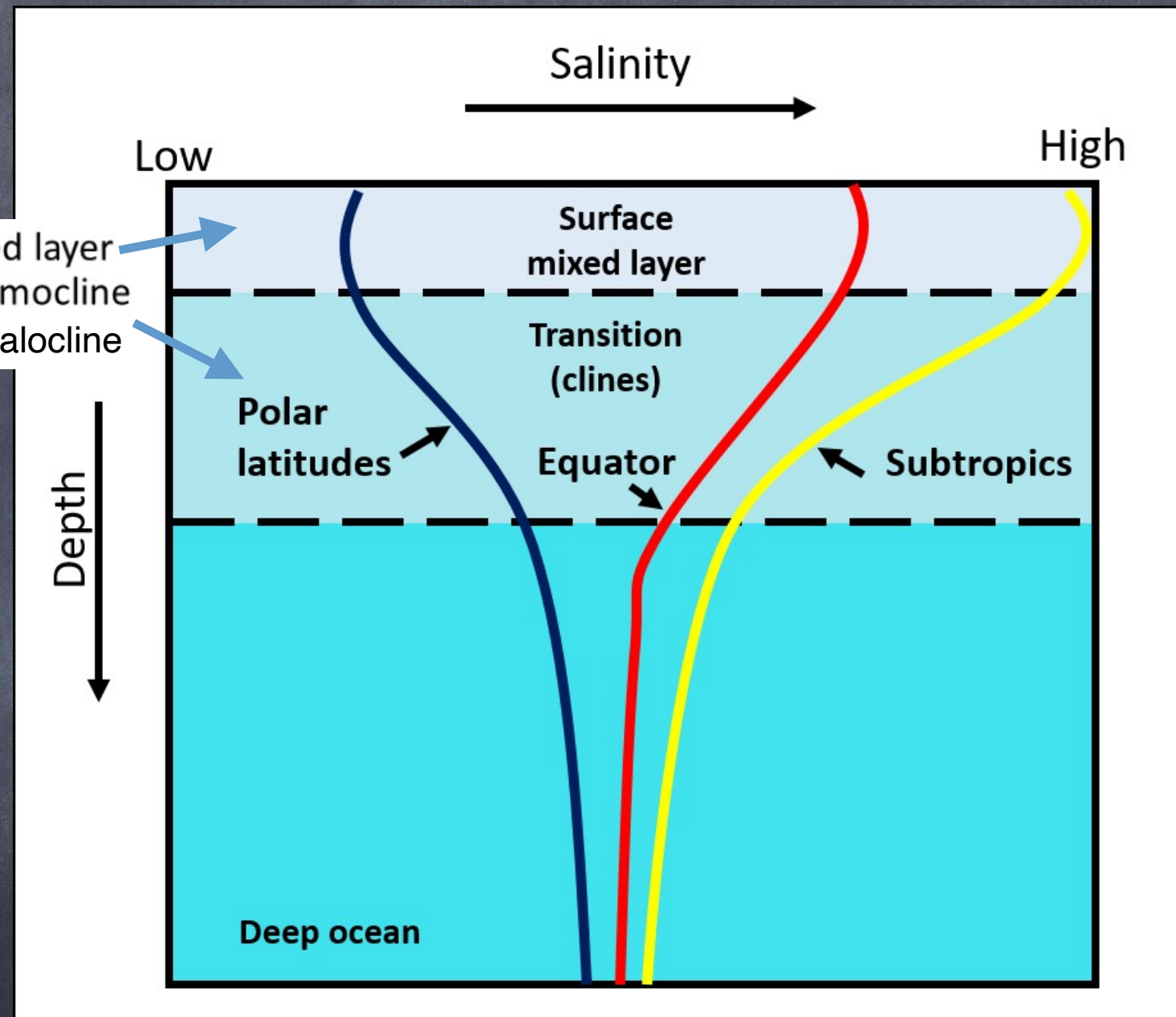
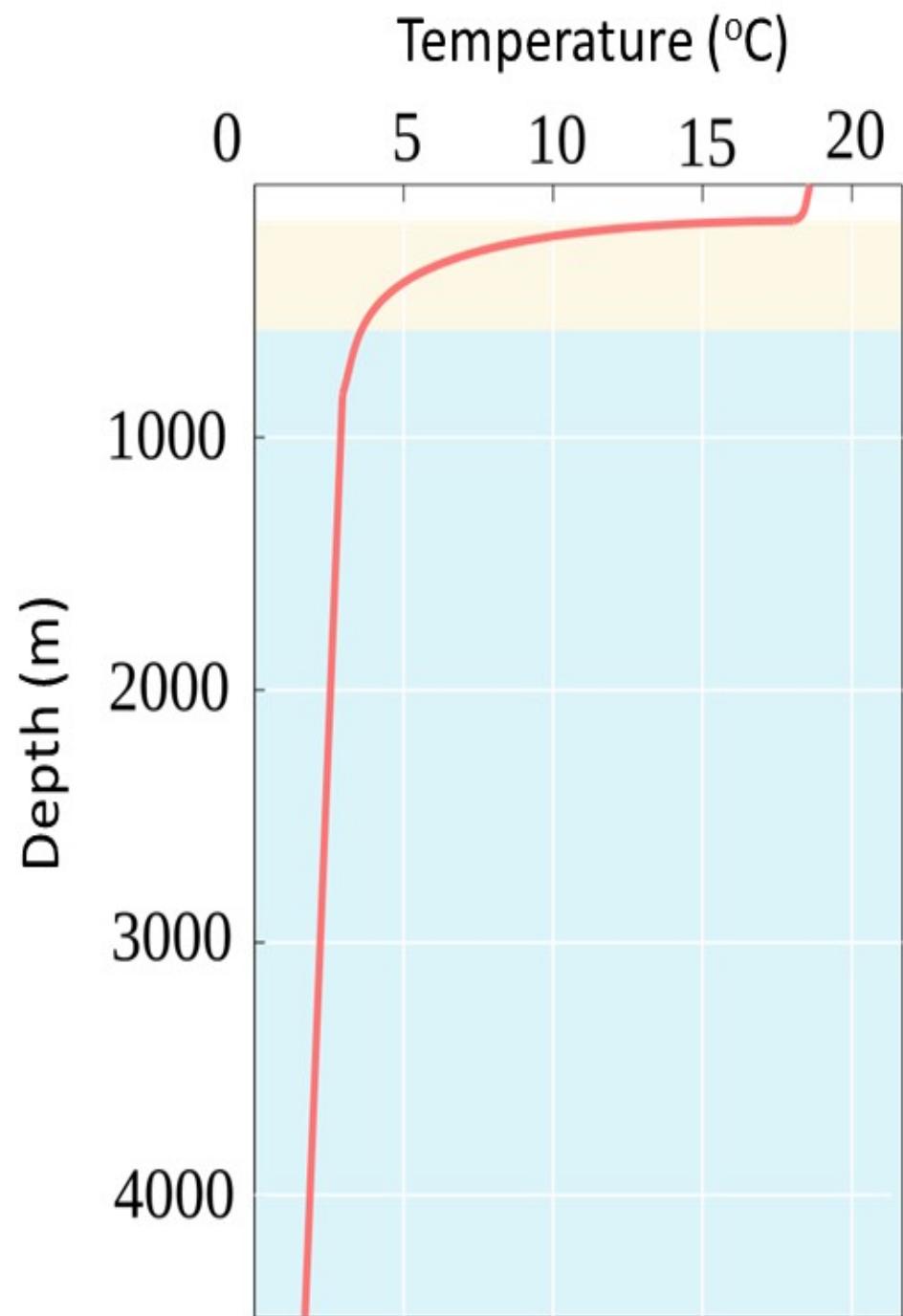


# T, S, rho of sea water with depth in the ocean





# T, S, rho of sea water with depth in the ocean

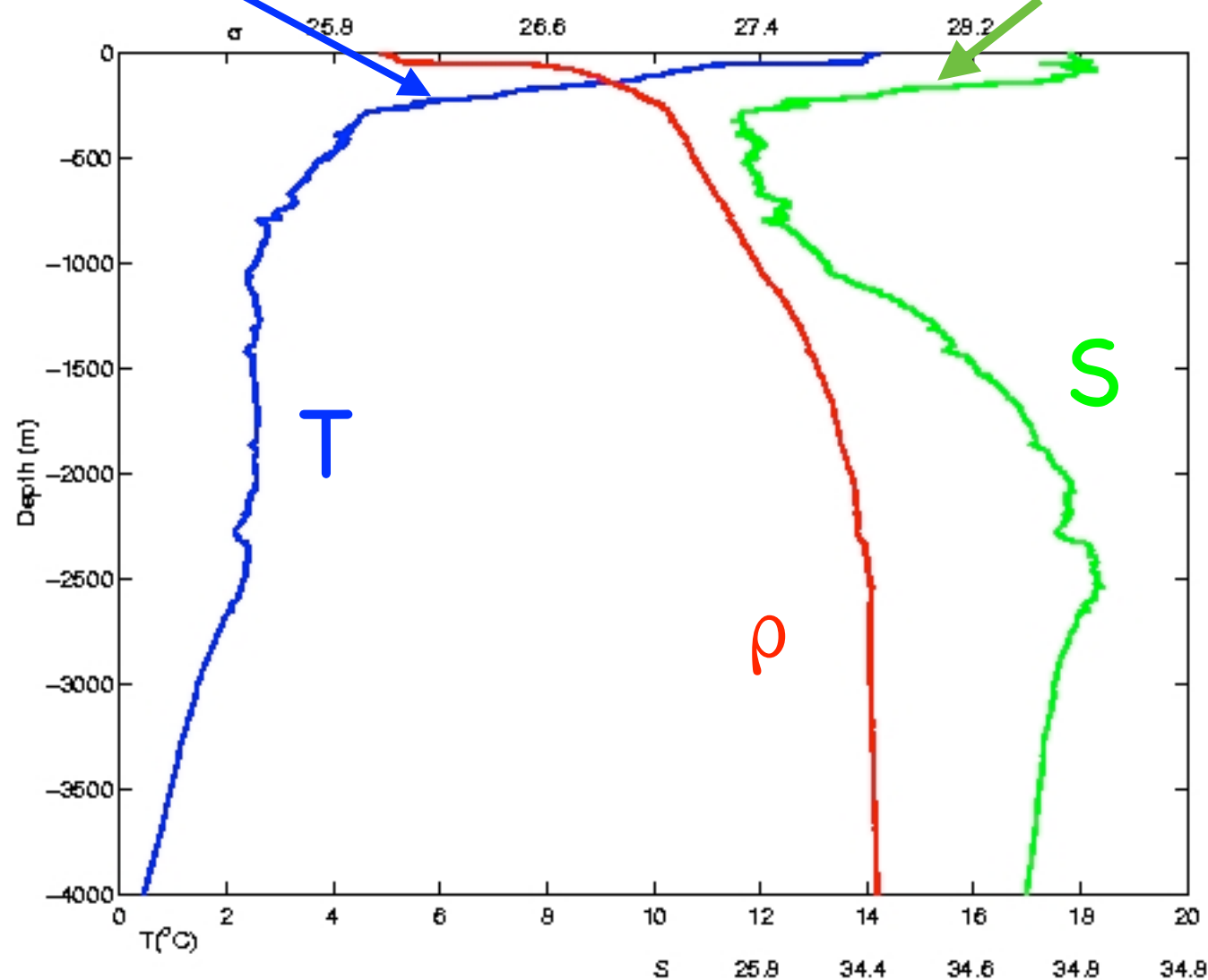




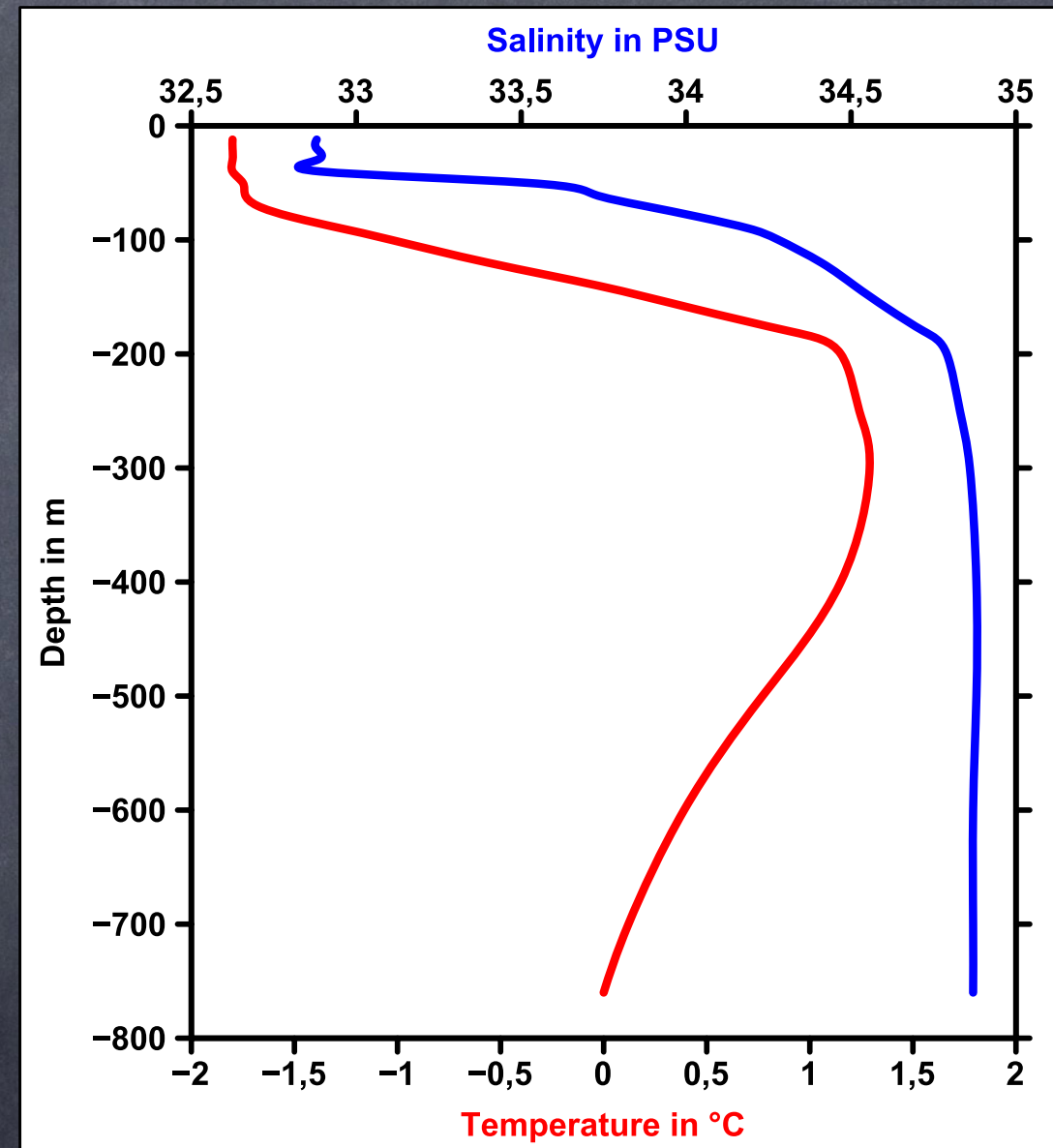
# T, S, rho of sea water with depth in the ocean

thermocline

halocline



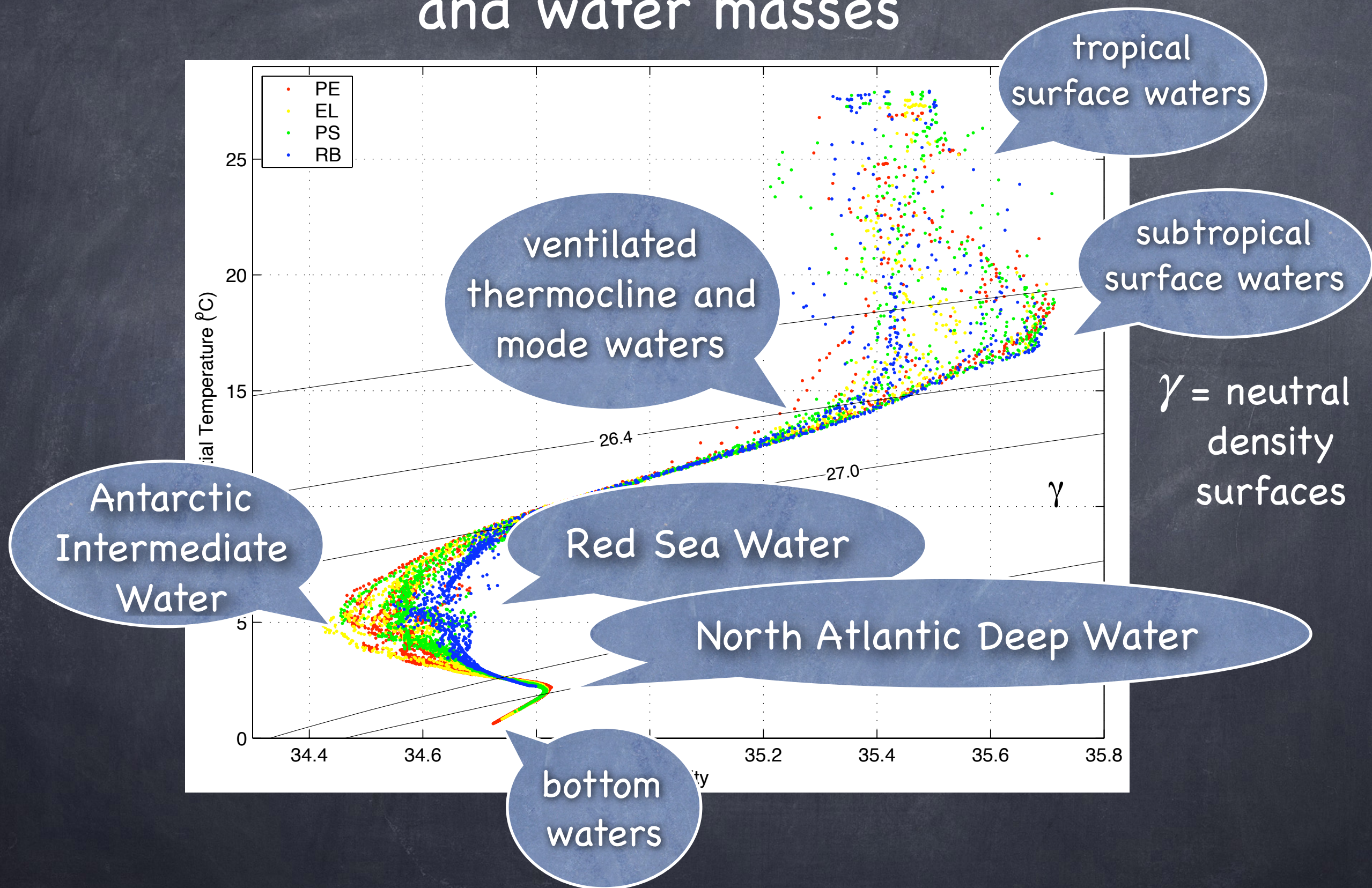
mid-latitude summer



polar winter



# Example of a T/S (or T-S) diagram and water masses





# Potential Temperature, $\Theta$

- A parcel of water moving from one pressure to another will be **compressed** or **expanded**
- When compressed adiabatically (without exchange of heat with surroundings) the parcel's temperature will increase. When expanded, its temperature will decrease.
- But, this T change does not represent a true change in energy, or heat content, of the fluid parcel
- To compare, or trace, parcels at different pressures we need to remove the effect of adiabatic contraction or expansion, called "Lapse Rate"



# Potential Density, $\sigma_\theta$

- A parcel of water moving from one pressure to another will be **compressed** or **expanded**, changing its density.
- Potential density,  $\sigma_\theta$  is the density of a fluid parcel that has been moved adiabatically to a reference pressure.
- But, compressibility is dependent on T (and S). e.g. cold water is more compressible than warm.
- To minimize the non-linear effect of thermal expansion and saline contraction on compressibility, we must choose a nearby reference:  $\sigma_1 = 1000 \text{ dbar}$ ,  $\sigma_2$ ,  $\sigma_3, \dots$

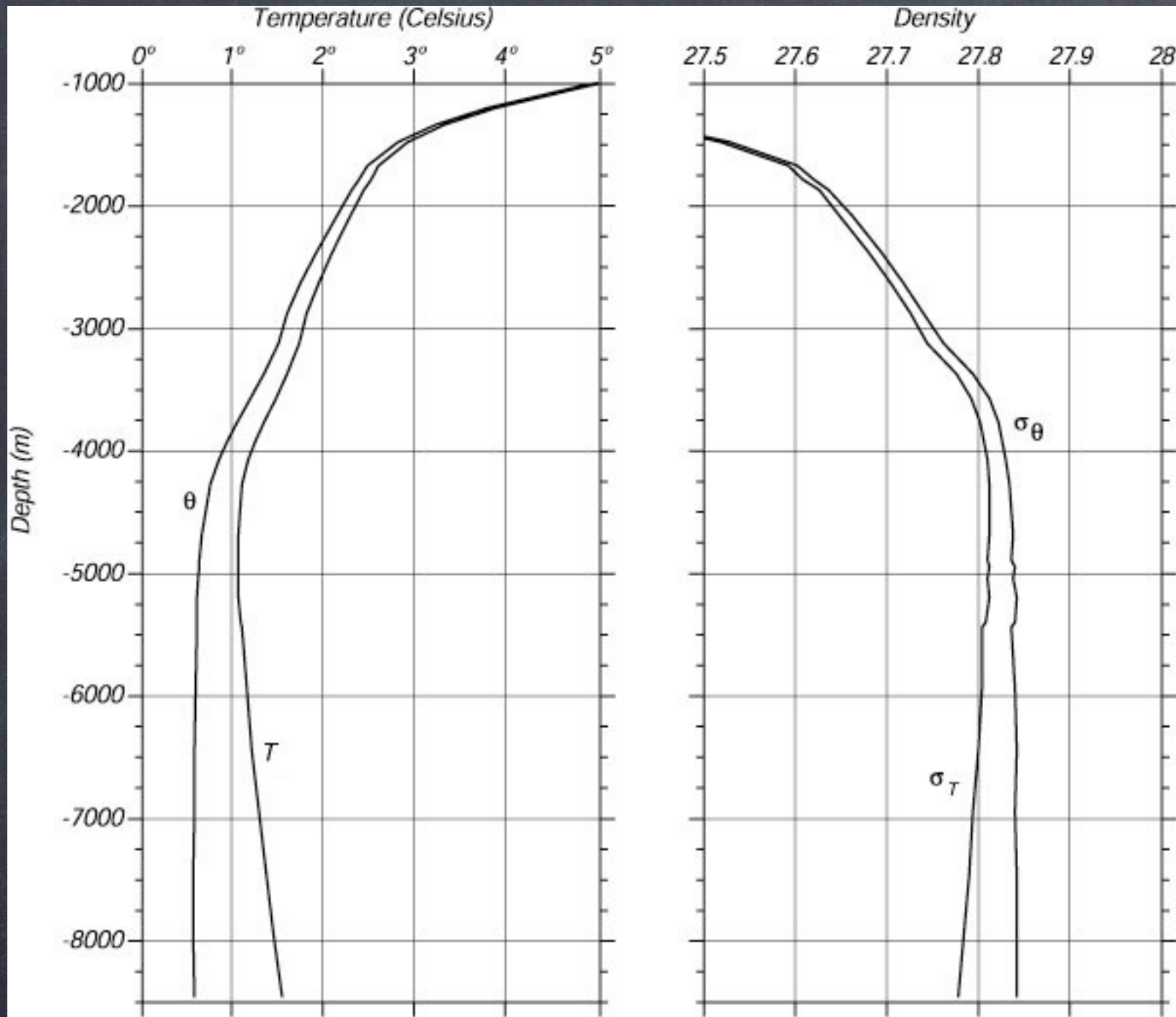


# Neutral Density, $\chi$

- If water moves with no exchange of heat or salt, it must be moving on an isentropic, or neutral, surface.
- A neutral surface can be approximated as a constant potential density referenced to a local pressure.
- Jackett and McDougall (1997) developed these neutral surfaces and they are available as a look-up table

- $$\chi = f(S, T, p, \text{lon}, \text{lat})$$



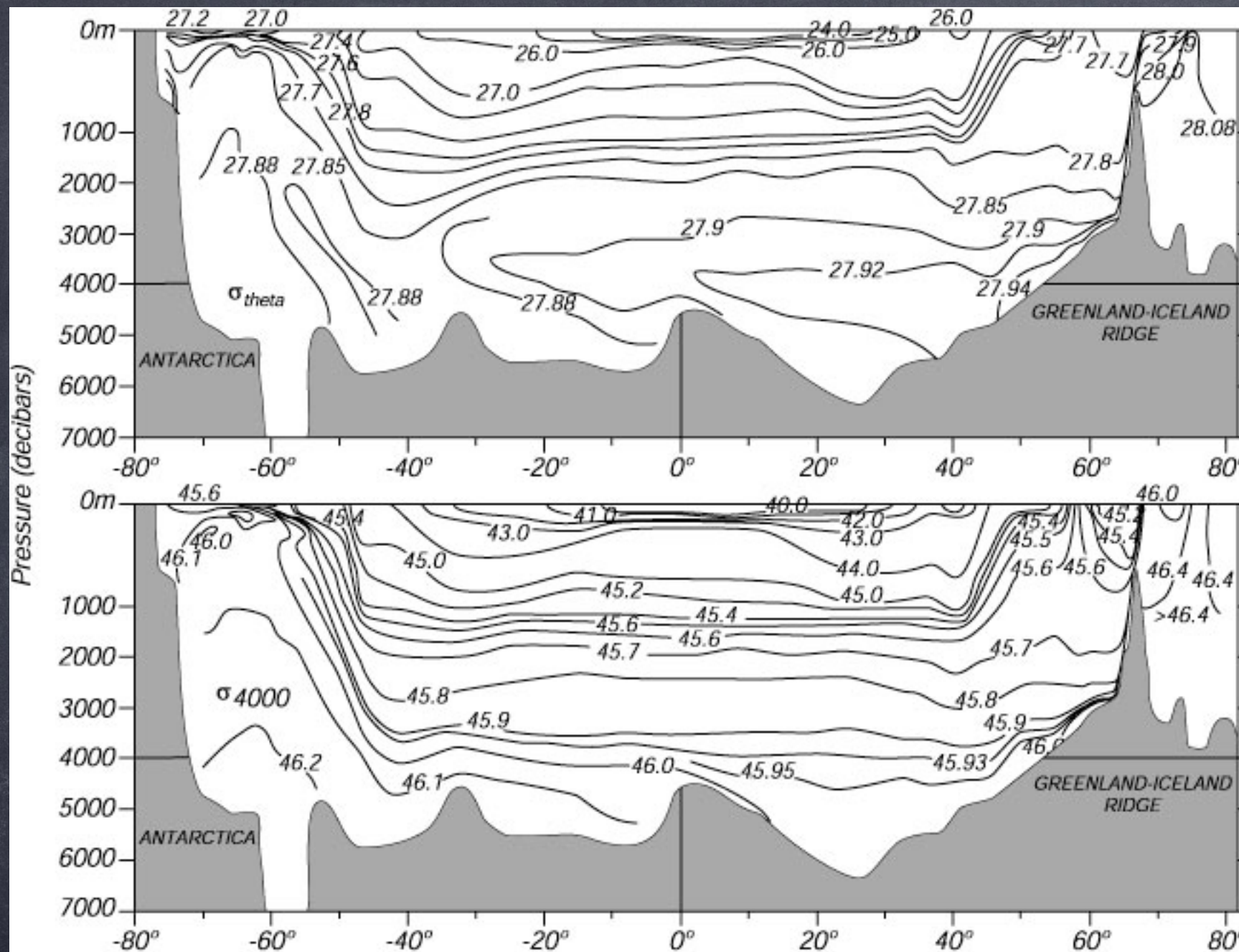


Potential  
temperature and  
potential density  
=  $T$  and  $\rho$  with  
effects of  
pressure  
removed.

Temperature and  
potential  
temperature

Density and  
potential  
density





Potential density must be referenced to a nearby density surface. Best to use neutral density.

Density versus depth, Western Atlantic. Upper:  $\sigma_\theta$ , showing an apparent density inversion below 3,000 m. Lower:  $\sigma_4$  showing continuous increase in density with depth. Depth scale changes at 1000 m depth. From Lynn and Reid (1968).



# Stability and Buoyancy Frequency

$$\text{Stability, } E = -\frac{1}{\rho} \frac{d\rho}{dz} \quad \text{Buoyancy Freq., } N^2 = -\frac{g}{\rho} \frac{d\rho}{dz}$$

- Buoyancy force is the difference between the density of the water parcel and the surrounding fluid

$$a = \frac{F}{m} = \frac{V \partial \rho g}{V \rho}$$

$$\frac{\partial^2 z}{\partial t^2} = \frac{g}{\rho} \frac{\partial \rho}{\partial z} \partial z \quad \text{take initial } z = 0$$

$$\frac{\partial^2 z}{\partial t^2} + N^2 z = 0, \quad \text{where } N^2 = -\frac{g}{\rho} \frac{\partial \rho}{\partial z} = gE$$

Harmonic with solution  $z(t) = \sin(Nt)$



# Stability and Buoyancy Frequency

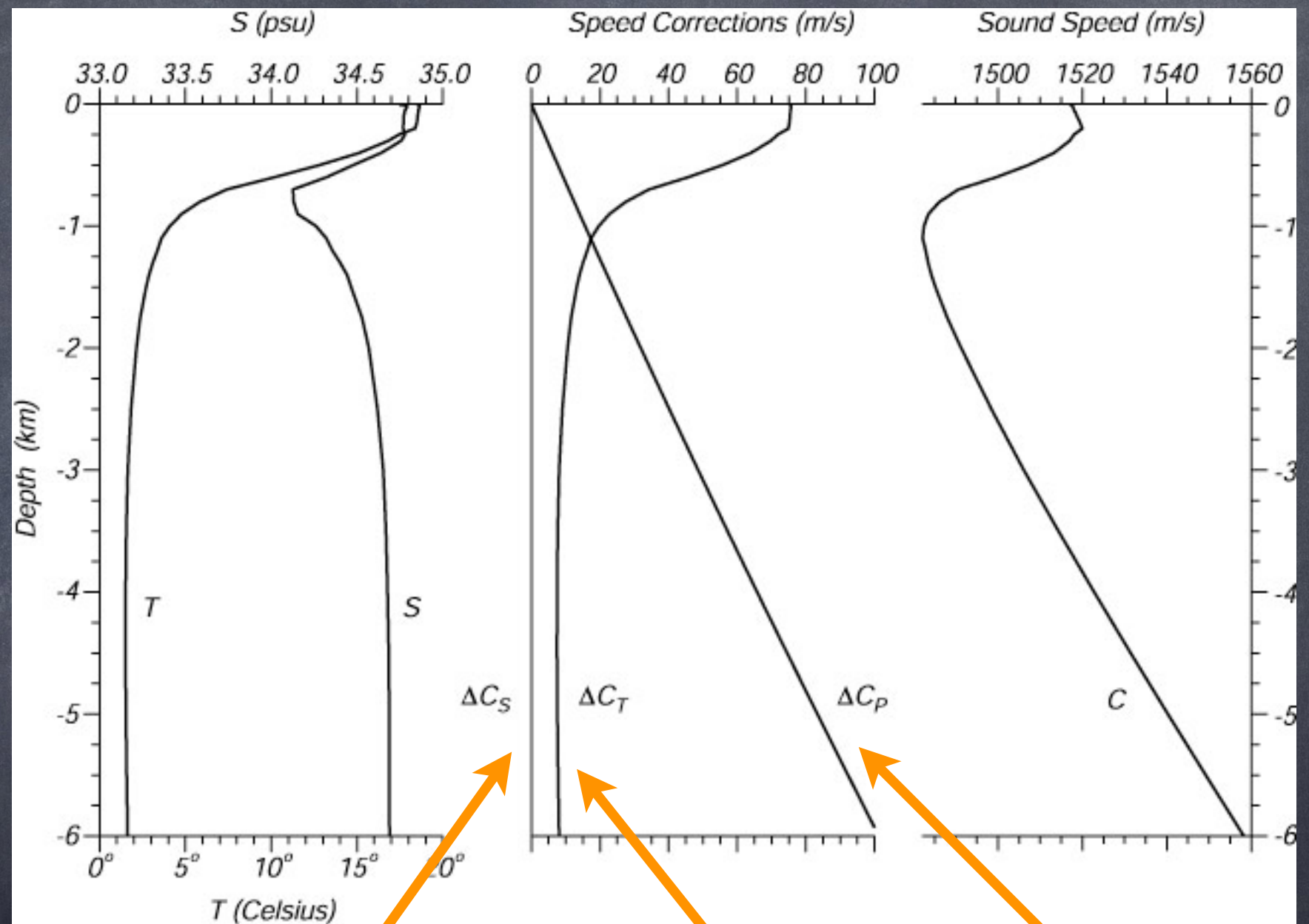
$$\text{Stability, } E = -\frac{1}{\rho} \frac{d\rho}{dz} \quad \text{Buoyancy Freq., } N^2 = -\frac{g}{\rho} \frac{d\rho}{dz}$$

- E and  $N^2$  are positive for stable stratification, negative for unstable
- A water parcel is more stable if the local density gradient is large
- Water parcels oscillate at frequency  $N^2$  in stably stratified fluids. Period is minutes in thermocline and hours in deep ocean



# Effect of p,T, and S on sound speed.

Sound is a compression wave that travels 5X faster through water than air



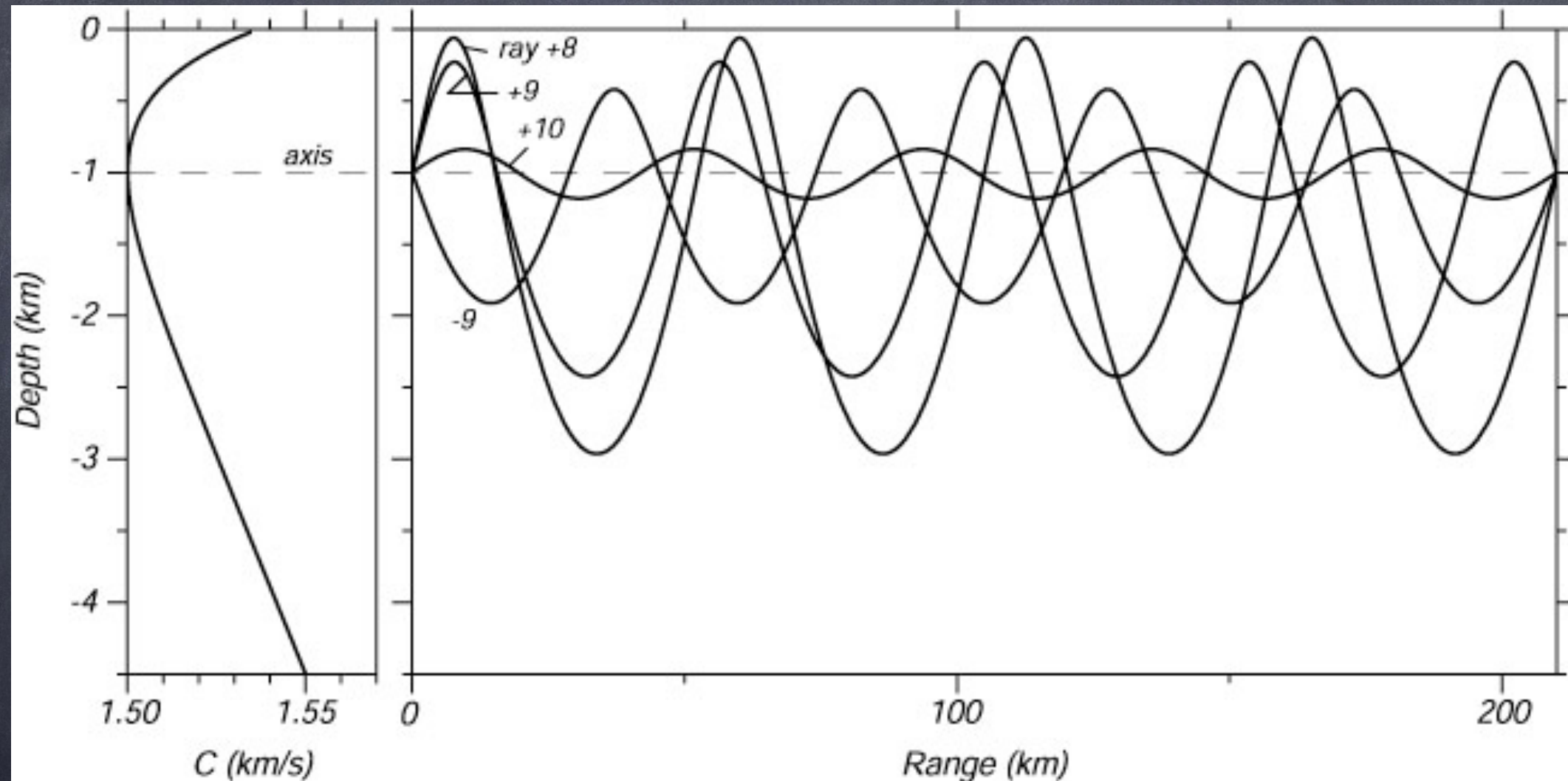
effect of S on  
sound speed

effect of T on  
sound speed

effect of p on  
sound speed



Ray paths of sound in the ocean for a source near the axis of the sound channel. Munk et al. (1995).



Rays are bent (or attracted) towards sound speed minimum = SOFAR channel