1. a) 
$$\frac{\partial S}{\partial \gamma}$$

3. a) 
$$I = (1-e)^2 U = (1-e)eB$$
,  $I = B_2$ 
 $I = I = (1-e)U = B$ ,  $I = B_2$ 
 $I = I = (1-e)U = (1-e)eB$ ,  $I = B_2$ 
 $I = I = (1-e)U = (1-e)eB$ ,  $I = B_2$ 
 $I = I = (1-e)U = (1-e)eB$ ,  $I = B_2$ 
 $I = I = (1-e)U = (1-e)eB$ ,  $I = B_2$ 
 $I = I = (1-e)U = (1-e)eB$ ,  $I = B_2$ 
 $I = I = (1-e)U = (1-e)eB$ ,  $I = B_2$ 

Add first two equations to eliminate  $I = I = I$ 
 $I = I = I = I$ 

Multiply second equation by  $I = I = I$ 
 $I = I = I = I$ 

Multiply second equation by  $I = I = I$ 
 $I = I = I$ 

Add these two new equations to eliminate  $I = I$ 
 $I = I = I$ 

Solve for  $I = I$ 

1) = 2+e I

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c) In class we used  $T = 240 \text{ W/m}^2$ , e = 0.9. This gives  $U = 633 \text{ W/m}^2$ 

Way too hot! What e gives Tg = 14°6?

Solving for e gives

$$e = 2 \left( \frac{\sigma T^* - I}{\sigma T^* + I} \right)$$

Substituting T= 14'C = 287 K gives e = 0.5.

4. as ar = 5.4 1/m2

X co. (t.) = 315 ppm for t. = 1958

Xcol (t) = 45 ppm for t = 2020

△H = 1.5 W/m2

b) Approximate the average extra heating as

AH 2 2 AH = 0.75 W/m2 The total time, Dt, is G3 years \$ 2 × 109 sec.

The upper ocean has depth, b= soom, and surface area A.

The temperature change, DT, is then

$$\Delta T = \frac{\Delta H \Delta t A}{D A g C \rho} = \frac{\Delta H \Delta t}{D g C \rho}$$
macs in kg

5. 
$$Q_{LH} = 175 \text{ W/m}^2$$
,  $\Delta t \approx 3 \times 10^7 \text{ sec}$ 

Latert heat of vaporization:  $L_V = 2.5 \times 10^6 \text{ J/kg}$ 

Rate of evaporation in  $\frac{\text{kg}}{\text{m}^2\text{s}}$  is  $E = \frac{Q_{LH}}{L_V}$ .

Over one year,  $E\Delta t = \frac{Q_{LH}}{L_V} = 2 \times 10^3 \frac{\text{kg}}{\text{m}^2}$ 

Thickness of layer with that mass and area:

$$\frac{2\times10^{3} \text{ kg/m²}}{1\times10^{3} \text{ kg/m²}} = \frac{2m}{5}$$

6. When | Up | increases:

- · T in creases as square of the wind speed.
- · Q<sub>SH</sub> increases,
- · E increases.
- · Q<sub>LH</sub> increases.
  - · B increases with E having a dual effect on heat and freshwater fluxes.

- 7. When T increases;
  - · Q<sub>sh</sub> increases most efficient when sea is warmer than air
  - · E increases exponentially as specific humidity increases exponentially with T.
  - · Qu increases exponentially in proportion to E.
  - · QLW decreases due to increased backradiation from near-surface evaporation despite greater outgoing LW from increased T.
  - · B increases due to increaces in heat flux and evaporation.

8. a)  $\vec{z} = C_d g_a |\vec{V}_{10}| \vec{V}_{10}$   $C_d \approx 1.2 \times 10^{-3}$   $|\vec{z}| = 7 \times 10^{-2} N_{m^2}$ 

b)  $m = \frac{F_0}{g} = \frac{7 \times 10^{-2} \,\text{N}}{10^{\,\text{m/s}\,2}} = 7 \times 10^{-3} \,\text{kg}$ c)  $V = \frac{m}{e} = 7 \times 10^{-6} \,\text{m}^3$ 

d)  $\delta = \frac{\sqrt{10^{-6}}}{10^{-6}}$  m or 7 microns

e) Gravity is strong! The tradewinds apply a force per unit area on the ocean that is equivalent to the downward gravitational pressure of an extremely thin layer of water. It is important to remember, however, that these forces operate in orthogonal directions.

- 9. a) Molecular diffusion is very slow over the vast spatial scales we consider in the ocean. Eddy diffusion approximates mixing at macro scales, which are relevant for ocean observations and models.
  - b) Eddy diffusion occurs via turbulent notions of water parcels or "blobs". The fluid properties (heat, salt, etc.) are conserved within each blob, i.e., each blob retains its properties as it moves. Thus, eddy diffusivity is a property of the flow field, not a characteristic of the property in question. Molecular diffusion depends on the thermal properties of the substance being diffused.
  - c) Stable stratification in the vertical suppresses vertical turbulent eddy motion, leading to a lower vertical eddy difficivity. Molecular difficion occurs due to random thermal motion, which is isotropic.