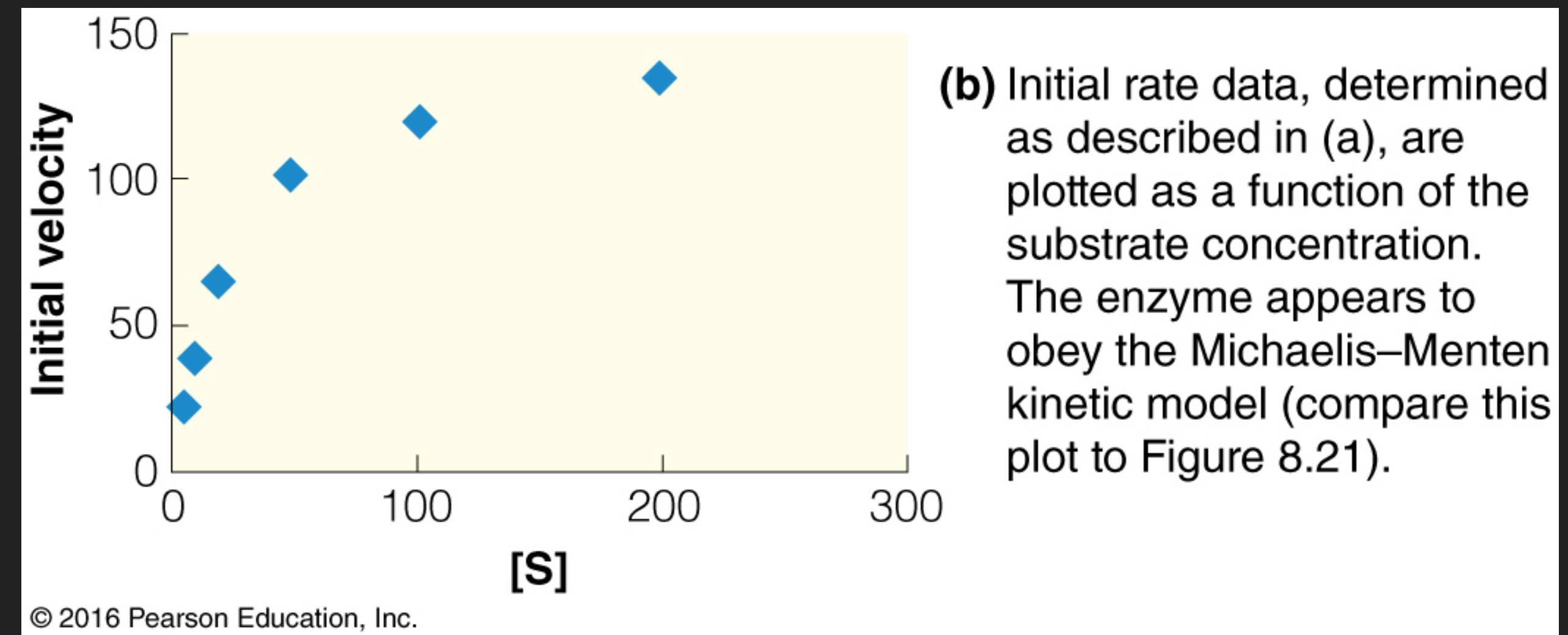


- ▶ Various reactions are performed at known Enzyme and Substrate concentrations.
- ▶ The initial rates of the reactions are plotted against the substrate concentrations
- ▶ If the graph is hyperbolic, then it can be assumed the enzyme obeys the Michaelis-Menten model.
- ▶ We can calculate K_M and V_{max}



- ▶ V_{max} is inherently hard to calculate or measure because the graph of enzyme velocities is hyperbolic.
 - ▶ Velocity asymptotically approaches its maximum
 - ▶ Even at 10-20 times the K_M we are only reaching 90-95% of the maximum
- ▶ To solve this issue, Hans Lineweaver and Dean Burk rearranged the Michailis-Menten equation into a linear function ($y = mx + b$), and then took the reciprocal.

$$\frac{1}{v} = \left(\frac{K_M}{V_{max}} \right) * \frac{1}{[S]} + \frac{1}{V_{max}}$$
$$y = m x + b$$

- ▶ This double reciprocal plot is called the Lineweaver-Burk Plot
 - ▶ It can be used to approximate both K_M and V_{max}