that the six sides of a cleavage rhombohedron of calcite always showed the same figures and solubility rates, whereas dolomite showed unlike solubility of the rhombohedral cleavage faces. The crystallographic or entation of the unequal reaction rims, as determined optically on the remaining calcite core of our specimens, shows that the c axis does not pass through the intersection of the three similarly reacted faces. The crystal lographic anisotropy of the reaction implies more than a loss of the symmetry center.

## DOLOMITE AND HYDROGEN

Dolomite and hydrogen were reacted under the following conditions: (1) fragment size 40–60 mesh; (2) 520°C to 835°C; (3) 2000 to 5000 psi initial hydrogen pressure; and (4) experiments lasting between 2 and 12 hours. Solid reactants observed include: CaCO<sub>3</sub>(calcite); Ca(OII)<sub>2</sub>; CaO; non-crystalline Mg(OH)<sub>2</sub>; elemental carbon or graphite, and "soot". Gas analyses show: CH<sub>4</sub>; C<sub>2</sub>H<sub>6</sub>; CO; and CO<sub>2</sub>. Water is the oxygenated product from all of the experiments.

The experimental results are summarized in Table 4 and plotted in Figures 9 and 10. The weight percent CO<sub>2</sub> remaining in the solid and the mole percent CH<sub>4</sub> formed are again plotted against time at 5000 psi for 620°C, and against temperature at 5000 psi for 4-hour runs.

In the thermal dissociation of dolomite under the vapor pressure of the system, the reaction occurs in two stages. The first dissociation, stoichiometrically simplified, is given as follows:

$$CaMg(CO_3)_2 + \Delta = MgO + CaCO_3 + CO_2$$

With increasing temperature, the remaining calcite decomposes to lime and carbon dioxide with the exact dissociation temperature determined by the carbon dioxide pressure. If the  $P_{\rm CO_2}$  never exceeds 100 mm, dolomite decomposes in a single stage dissociation as follows:

$$CaMg(CO_3)_2 + \Delta = MgO + CaO + 2CO_2$$

In the dolomite-hydrogen reaction CO<sub>2</sub> is not a reaction product under most of the experimental conditions and appears only as a minor component of the reaction gases at lower hydrogen pressures (Table 4). It likely is incidental to the main reaction of direct methanation. The basis for this statement is found in the dolomite-hydrogen gas analyses and the reasoning previously used in discussing the calcite-hydrogen system.

The dolomite-hydrogen reaction occurs in two steps. The first stage is represented by the following equation:

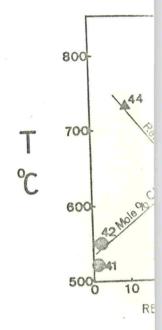


Fig. 9. Plot of mole % CH<sub>4</sub> generative-hydrogen system at 5000 psi (H 620, and 735°C.

$$4H_2 + CaMg(CO_3)_2 =$$

At 5000 psi (H<sub>2</sub>) reaction informed from dolomite persist alone in the calcite-hydrogen thermal dissociation of carbohydrogen. In the dolomite-hydrogen in a manner similar to cite. This possibility will be seen as the cite of the control of the cite.

Mg(OH)<sub>2</sub> or MgO formed Chemical analyses by atomic reaction products.

A very minor amount of action products. It is similar the calcite experiments. "S bomb, and again was part

The reaction gases were to (Table 4). Carbon dioxide