

2013  
Acker

THESIS ON GYPSUM

RESPECTFULLY SUBMITTED TO

TO

Dr. James Lewis Howe  
Professor of Chemistry

Washington and Lee University

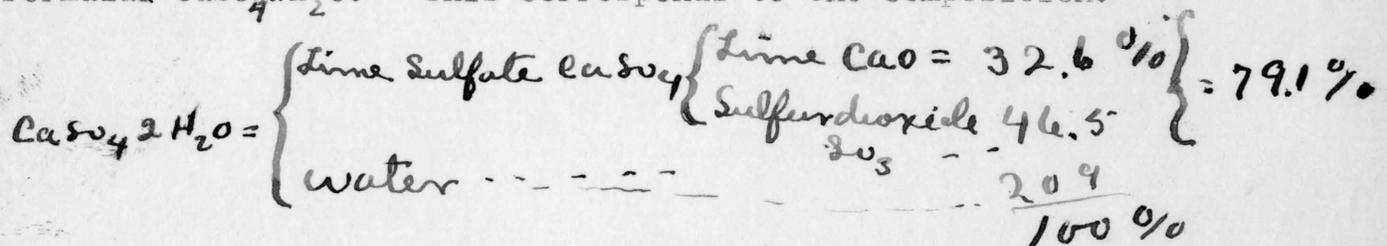
By

J. E. Acker.

December 10, 1913.

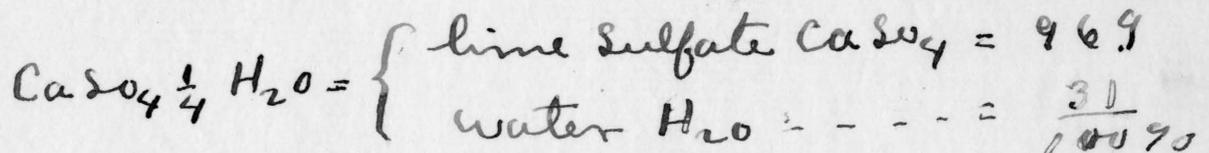
INTRODUCTION.

Calcium Sulfate is frequently found in nature, occurring in lime stone water or in company with common salt, here it is found in anhydrate form. This has the formulax of  $\text{CaSO}_4$ . It is found in both <sup>h</sup>rhombic and semi-crystalline form. However it is found in the hydrate form known as gypsum, ~~thith~~ with the formulax  $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ . This corresponds to the composition.

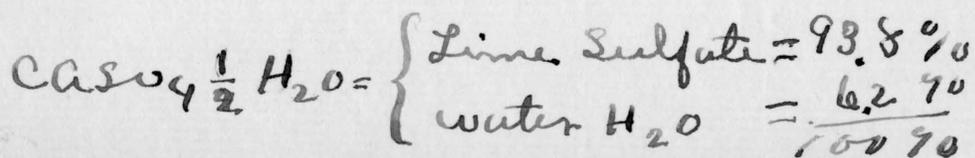


And is usually in a well crystalized form with a hardness of two. Sp. gr. 2.3 it has a silky lusture and iss colorless or a white gray. It has a fracture in three ways.

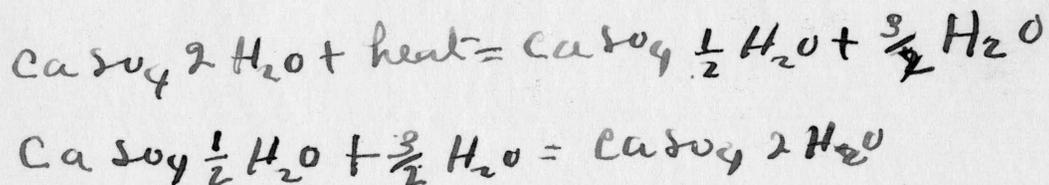
When it is heated above  $120^\circ$  it looses part of its water, probably having the formulax  $\text{CaSO}_4 \cdot \frac{1}{2}\text{H}_2\text{O}$  which corresponds to



This can be proven by heating it in a closed tube, water forming on the side of the tube. But most of the chemists claim it has a formular of  $\text{CaSO}_4 \cdot \frac{1}{2}\text{H}_2\text{O}$  which would correspond to



If it is heated to 120° instead of 110° it gives better results although it is generally conceded that three fourths of the water will go off at 110°. After it has been heated at this temperature it is a soft grayish mass. ~~It~~ It forms first in the orthorhombic system and finally in the monoclinic system. After it has been heated to this temperature and mixed with water it soon sets to a hard mass. This setting is due to the plaster dissolving little by little in the water in which it is slightly soluble, and then immediately crystallizing out in a form of minute interlocking crystals or needles which bind the mass together. It seems to have the hardness of about three. It solidifies very quickly under ordinary circumstances with the evolution of heat and expanding in so doing as to fill any mold in which it is a cast. These qualities make it very valuable, and is used very extensively in forming models of artistic figures as well as ornamentations and used in surgery. This is known as plaster paris. The reaction is



However if it is heated over 200° it loses all of its water as well as its property of hardening with the additional water. This is called dead burnt and the loss of this property is generally conceded to be the lack of nuclei of the more stable form

which has an essential influence on it when present. In this case it lacks the nuclei for the formation of new crystals.

#### REFERENCES.

Principals of Inorganic chemistry. Estwald page 519  
Bloxams Inorganic chemistry page 367.  
Outlines of industrial chemistry. Thorp  
Principals of inorganic chemistry . Jones page 367.  
Inorganic chemistry. Ramay page 421  
General inorganic chemistry . Alex & Smith page 604.  
Inorganic chemistry . Howe page 304  
Dictionary of applied chemistry . Thorp page 611.  
Volume 1.  
General & Industrial chemistry.  
Seletine & Mallinard page 447.  
Cements, limes and plasters . Eckel page 31. and many other inorganic chemistries.

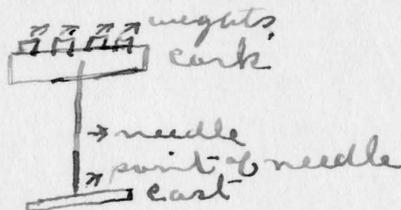
#### SUBJECT.

The subject of this thesis is to try to prove that dead burnt gypsum will set if it has a nuclei to set around and for this nuclei to be furnished with a given % of either gypsum or plaster of Paris. To confirm the theory that the presence of a nuclei of a stable form has an essential influence and this can be accomplished by adding either <sup>dead burnt gypsum</sup> gypsum or plaster Paris to furnish the nuclei for the formation of new crystals.

#### PROCESS.

I took a platinum crucible which was about two thirds

full of plaster of paris and heated it before a blow pipe at white heat for about fifteen minutes it probably being 800° to be sure all of the water had been driven off. Small quantities were used that the heat might penetrate the entire mass. This was allowed to cool and then water added. This was put in a cast with three glass sides three fourths of an inch wide, two inches long and ~~in~~ an inch and a half in height, and allowed to dry out. It dried out very slowly and when dried, tested with *heat* needle as to the resisting power



An ordinary darning needle was used with a sharp point and about three inches long. The blunt end was stuck in a cork and the sharp end placed on the mold and weight added to the cork stopper until the darning needle penetrated the cast. Ten grams were added before it broke. This was *repeated* five or six times in order to prove that it was correct. It was also proven that it was harder just before entirely dry than when very dry. A cast of it was allowed to dry ~~up~~ *out* and wrapped in a damp rag and allowed to stand over night. It was found harder the next morning than the evening before. When entirely dry it only took five *grams* to break it. At high temperature or a very low one it was not as hard as it was at ordinary temperature. It was also proven with all the samples that they

would set quicker and give as good results if heated in a drying closet the temperature between 30° and 40° .

About seven grams of the dead burnt gypsum was used for all the samples in this work and all the casts were practically of the same measurements .

Next I added one per cent of the pure gypsum to the dead burnt . This was put in a mold and allowed to dry.

The weights were:

$$\begin{aligned} \text{Burnt gypsum } & 7.0760 \text{ grams} \\ 7.0760 \div 100 & = 0.07076 = 1\% \end{aligned}$$

This when dry was tested as to hardness and it took 141 grams to brake it making it 14.1 times as hard as the dead burnt. This was also tested several times in order to prove its correctness. All samples were tested under the same conditions.

I next added 1% of plaster paris to the sample of dead burnt and the weights were as follows:

$$\begin{aligned} \text{Dead burnt} & = 7.542 \text{ grams} \\ 7.542 \div 100 & = 0.07542 = 1\% \end{aligned}$$

After this was dry I tested it and found it to be 44.2 times as hard as the dead burnt and little over three times as hard as the sample with 1% of gypsum. And the next ~~was~~<sup>I</sup> added 25% of the gypsum to the dead burnt and found it to be 108 times harder than dead burnt.

I added 25% of plaster of paris to a sample of dead burnt and on testing found it to be 145.2 times harder than the dead burnt and just about 1.3 times as hard as 25% of gypsum.

In this way I had a different percent until I got 50% of each, and there could scarcely be any difference told between the 100% of plaster paris and the 50% of plaster paris .

The results of the whole were

1% of gypsum 14 times harder than the dead burnt.

1% of plaster of paris 45 times harder than 1% of gypsum.

25% of gypsum 108 times harder than dead burnt.

25% of plaster of paris , 145 times harder than a dead burnt, and 11/2 times harder than the 25% of gypsum. It was also found that the gypsum took twice as long to set as it did for the plaster of paris.

Owing to the lack of apparatus it was impossible to get the exact resisting power of the cast, but the relation of one to another could be gotten by means of *wicat* needle. It was proven that the presence of a <sup>nuclei</sup> ~~neutralizing~~ of a more stable form did have an essential influence upon the dead burnt gypsum, ~~that~~ to a certain extent the nuclei furnished or rather caused a hardening of a very considerable degree. This is very likely due to the formation of new crystals of the dead burnt, <sup>furnished</sup> ~~by~~ the nuclei of the more stable form.

Respectfully submitted by

*J. E. Acker*