bon monoxide and chlorine are evolved. The production of carbon monoxide by an electrolytic method is unique, as it has heretofore not been obtained by electrolysis.

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WHAT IS AN ACID?

In terms of the theory of electrolytic dissociation, an acid is any substance which yields the hydrogen ion in water solution, and a base any substance capable of yielding the hydroxide ion under similar circumstances. But Franklin has shown that this definition of a base is too narrow, and that a comprehensive definition should include substances capable of yielding the amide ion in ammonia solution. The question may, then, be legitimately asked, if bases may include substances that do not yield the hydroxide ion, may there not exist acids that do not yield the hydrogen ion?

I have recently been forced to the conclusion that the answer to this question is in the affirmative. The evidence, which I shall discuss more fully in another paper, consists in the following facts. Aluminium chloride is very soluble in liquid phosgene, $COCl_2$, and several phosgenates of aluminium chloride have been described in the literature. I have shown that this solution conducts electricity and that the products of electrolysis are carbon monoxide and chlorine. I have also shown that this solution reacts with metals, with metallic chlorides, oxides, sulfides and carbonates to form compounds, containing the metal as chloride, and aluminium chloride, with phosgene of crystallization, and that carbon monoxide, carbon dioxide, carbon oxysulfide are split off.

These observations become intelligible if we assume that the active substance in the solution is a phosgenate of aluminium chloride, such as 2AlCl_a.COCl_a which may be compared with the behavior of $2SO_8.H_2O_7$, or with $SO_8.H_2O$ in water solution; it is customary to write H₂S₂O₇ or H₂SO₄ to represent the condition of the hydrates of sulfur trioxide, and in an analogous manner we are led to write COAl, Cl. to represent the condition of the phosgenate of aluminium chloride in phosgene solution. This compound, carbonyl di-chloraluminate, reacts with metals to produce salts of the type M^{II}Al₂Cl₂.xCOCl₂ (using a bivalent metal as example), with liberation of carbon monoxide, in precisely the same manner as sulfuric acid produces, with metals, salts of the type $M^{II}SO_4$, xH_2O , with liberation of hydrogen. With metallic chlorides, carbonyl di-chloraluminate reacts to produce salts of the same type, splitting off phosgene, in precisely the same manner as sulfuric acid

produces sulfates with the metallic oxides, splitting off water. The analogy may be carried on to the other reactions and to the electrolysis of the two solutions, and the conclusion becomes inevitable that these solutions have properties so precisely similar that we must classify them in the same group—in other words, carbonyl di-chloraluminate is an acid. But this acid contains no hydrogen; in its place, we have carbon monoxide. The metallic chlorides must be regarded as bases, and the chloride ion in these bases corresponds to the hydroxide ion among the aquo-bases, and to the amide ion among the ammonobases.

What, then, is an acid?

First of all, an acid involves a solvent, and as far as we know an ionizing solvent-such as water, ammonia, phosgene, to mention only those we have referred to in this note. In the second place, an acid is a substance dissolved in the solvent, which is chemically combined with a portion of the solvent, as SO₃ is combined with water, as AlCl₃ is combined with phosgene, or as C_aN_4 is combined with ammonia (cyanamide, H₂CN₂, or di-cyanimide, HN(CN)₂). Finally, solutions of these acids in their respective solvents give certain definite reactions-they conduct electricity and the solvent is decomposed in the process; metals displace the ion common to the acid and the related solvent (H, CO, etc.) forming salts; they react with bases derived from the related solvent to form salts, splitting off a molecule of the solvent; etc.

In short, an acid is the substance formed when an ionizing solvent unites chemically with a suitable compound and is capable of forming salts with metals and with bases derived from the same solvent.

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MIOCENE MARINE VERTEBRATES IN KERN COUNTY, CALIFORNIA

PROBABLY the first vertebrate fossils collected in California were the teeth of sharks picked up by W. P. Blake, geologist of the Williamson contingent of the Pacific Railroad Survey, in 1853.

The original location was given as "Depot Camp" on the south side of Poso Creek (called Posé Creek in the original reports; also called Ocoya Creek and spelled Ocoga on the map) "Lat. 305° 30' 27", Long. 118° 53' 02"," Kern County, California. The specimens were described and illustrated by L. Agassiz.¹

¹ Am. Journ. Sci. Arts, 1856, pp. 272-275; Pac. B. R. Reports, Vol. 5, 1857, pp. 313-316, plate 1, figs. 1-44; for a full account of the occurrence of the specimens see pp. 164-173.