#### m22.p10

# Synthesis, Characterization and properties of alkali and earth-alkalisulfonates

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Stochiometric aqueous solutions of benzosulfonic acid and metalcarbonates were evaporated to crystallze different watercontaining salts.

The crystals were characterized by X-ray, thermal, chemical and optical methods.

These layered materials can be used because of their ion-exchange ands intercalation properties, but also for controlling hydration beahviour of cements. A summary of lattice parameters of different phases is given in the table.

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Compound	$a_o[A^\circ]$	<i>b<sub>o</sub></i> [ <i>A</i> °]	$c_o[A^\circ]$
LiC <sub>6</sub> H <sub>5</sub> SO <sub>3</sub>	17,1636	21,8470	14,1591
NaC <sub>6</sub> H <sub>5</sub> SO <sub>3</sub>	29,8833	24,3355	10,2458
KC <sub>6</sub> H <sub>5</sub> SO <sub>3</sub>	15,1770	28,8022	13,2028
$Mg(C_6H_5SO_3)_2$	30,4594	12,6911	14,2171
$Ca(C_6H_5SO_3)_2$	30,0150	20,4244	10,6854
$Sr(C_6H_5SO_3)_2$	31,3182	29,9525	10,1632
$Ba(C_6H_5SO_3)_2$	31,1896	30,1374	10,3097
Ag(C <sub>6</sub> H <sub>5</sub> SO <sub>3</sub> ) <sub>2</sub>	5,1596	5,1979	15,2962
Mn(C <sub>6</sub> H <sub>5</sub> SO <sub>3</sub> ) <sub>2</sub> *6H <sub>2</sub> O	22,7596 /6,3312/7,0445		
Fe(C6H5SO3)2*2H2O22,586/16,3293			6,9989
Co(C6H5SO3)2*6H2O22,4279/6,3180			7,0068
Ni(C6H5SO3)2*6H2O22,3799/6,3016			6,9709
Zn(C6H5SO3)2*2H2O22,5006/6,3087			6,9890
Cu(C6H5SO3)2*6H2O22,5297/6,2980			7,0364
Compound	α [°]	β [°]	γ[°]
LiC <sub>6</sub> H <sub>5</sub> SO <sub>3</sub>	90	96,741	90
NaC <sub>6</sub> H <sub>5</sub> SO <sub>3</sub>	90	90	90
KC <sub>6</sub> H <sub>5</sub> SO <sub>3</sub>	90	90	90
$Mg(C_6H_5SO_3)_2$	90	97,884	90
$Ca(C_6H_5SO_3)_2$	90	90	90
$Sr(C_6H_5SO_3)_2$	90	90	90
Ba(C <sub>6</sub> H <sub>5</sub> SO <sub>3</sub> ) <sub>2</sub>	90	90	90
$Ag(C_6H_5SO_3)_2$	86,75	84,56	61,0
Mn(C <sub>6</sub> H <sub>5</sub> SO <sub>3</sub> ) <sub>2</sub> *6H <sub>2</sub> O90 93,564		93,564	90
Fe(C6H5SO3)2*2H2O90		93,608	90
Co(C6H5SO3)2*6H2O90		93,754	90
Ni(C6H5SO3)2*6H2O90		93,771	90
Zn(C6H5SO3)2*2H2O90		93,635	90
Cu(C6H5SO3)2*6H2O90		93,722	90

Tab. 1: Lattice parameters of Benzolsulfonates

[1] Gal, S., Meisel, T., Halmos, Z. und Erdey, L.: Mikrochim. Acta, (1966) 903.

#### m22.p11

### The Crystal Structure of Manganese modified Brownmillerite Solid Solutions

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Brownmillerite originally describes the chemical composition  $Ca_2AIFeO_5$  on the join  $Ca_2Fe_2O_5$ - $Ca_2Al_2O_5$  ( $Ca_2Fe_{2-x}Al_xO_5 x = 0,5$ ) in different cement types. The main interest was focused on the solid solution series  $Ca_2Fe_{2-x}Al_xO_5$  for their ability to react with water to hydration phases and form together with other hydration phases and quartz a 3-dimensional framework. With the addition of secondary manganese raw materials to cement its hydration properties were improved and  $Mn^{3+}$  is fixed in the Brownmillerite crystal structure.

The crystal structure of different solid solutions in the system  $Ca_2Fe_{2,x}Al_xO_5$  with  $0 \le x \le 1.33$  were recently investigated [1] and is best described as a 2-dimensional, oxygen deficient alternative of simple perovskite phases (e.g.  $CaMnO_{3-d}$ ) [2]. The structure comprises alternating layers of cations sited in corner sharing octahedra and corner sharing tetrahedra, perpendicular [010]. However, the orthorhombic phases are not isostructural, because the space group Pnma for iron rich members changes into s. g. I2mb at approximately x = 0.6 for alumina rich phases. Recently, metric parameters of solid solutions  $Ca_2(Fe_{1-y}Mn_y)_2AlO_5$  with  $0 \le y \le 0.5$  were determined [3]. However, no structural data of these phases are available. Samples with the composition  $Ca_{v}(Fe_{v}Mn_{v})_{2}$  $2xAl_{2x}O_5$  with y = 0.5; x = 0.042; 0.085; 0.25; 0.415; 0.585 were synthesized. Stochiometric oxide - concentrations were sintered at 1300°C for 96 h with several grinding steps in a tube furnace using Pt crucibles. The sinter products were quenched and analyzed by XRD. The Oxygen concentration was determined by Iodometric titration. The phase chemistry was determined by microprobe analysis. Structure refinements were performed at the BENSC Hahn-Meitner Institute Berlin. Neutron radiation was applied due to the fact, that Mn and Fe show strong electronic similarities: Mn (atomic number 25) and Fe (atomic number 26).

[1] Colville A. A. and Geller S., Crystal Structures Ca<sub>2</sub>Fe<sub>1.43</sub>Al<sub>0.57</sub>O<sub>5</sub> and Ca<sub>2</sub>Fe<sub>1.63</sub>Al<sub>0.57</sub>O<sub>5</sub> Acta Cryst 1972 B28: p. 3196-3200

[3] Puertas F., Blanco Varela M.T., and Dominguez R., Characterisation of Ca<sub>2</sub>AlMnO<sub>5</sub>. A Comperative Study between Ca<sub>2</sub>AlMnO<sub>5</sub> and Ca<sub>2</sub>AlFeO<sub>5</sub>. Cem. Concr. Res., 1990. 20: p. 429-438.

<sup>and Ca<sub>2</sub>Fe<sub>1.28</sub>Al<sub>0.72</sub>O<sub>5</sub>. Acta Cryst., 1972. B28: p. 3196-3200.
[2] Poeppelmeier K.R., Leonowicz M. E., and Longo J. M., CaMnO<sub>2.5</sub> and CaMnO<sub>3.5</sub>: New Oxigen Defect Perovskite-Type Oxides. J. Solid State Chemistry, 1982. 44: p. 89-98.</sup>